

APPROVED DRUG PRODUCTS

WITH

**THERAPEUTIC
EQUIVALENCE
EVALUATIONS**

39th EDITION

**THE PRODUCTS IN THIS LIST HAVE BEEN APPROVED UNDER
SECTION 505 OF THE FEDERAL FOOD, DRUG, AND COSMETIC ACT.**

**U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION
OFFICE OF MEDICAL PRODUCTS AND TOBACCO
CENTER FOR DRUG EVALUATION AND RESEARCH
OFFICE OF GENERIC DRUGS
OFFICE OF GENERIC DRUG POLICY**

2019

APPROVED DRUG PRODUCTS with THERAPEUTIC EQUIVALENCE EVALUATIONS

The products in this list have been approved under section 505 of the Federal Food, Drug, and Cosmetic Act. This volume is current through December 31, 2018.

39th EDITION



U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION
OFFICE OF MEDICAL PRODUCTS AND TOBACCO
CENTER FOR DRUG EVALUATION AND RESEARCH
OFFICE OF GENERIC DRUGS
OFFICE OF GENERIC DRUG POLICY

2019

**FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH**

**APPROVED DRUG PRODUCTS
With
Therapeutic Equivalence Evaluations**

CONTENTS

| | <i>PAGE</i> |
|----------------------------------------------------------------------------------------------------------------------------------------------|-------------|
| PREFACE TO THIRTY-NINTH EDITION..... | iv |
| 1.0 INTRODUCTION | vi |
| 1.1 Content and Exclusion | vi |
| 1.2 Therapeutic Equivalence-Related Terms | vii |
| 1.3 Further Guidance on Bioequivalence | ix |
| 1.4 Reference Listed Drug and Reference Standard..... | ix |
| 1.5 General Policies and Legal Status | x |
| 1.6 Practitioner/User Responsibilities | xi |
| 1.7 Therapeutic Equivalence Evaluations Codes | xiii |
| 1.8 Description of Certain Special Situations | xxi |
| 1.9 Therapeutic Equivalence Code Change for a Drug Entity | xxiii |
| 1.10 Change of the Therapeutic Equivalence Evaluation for a Single Product..... | xxiv |
| 1.11 Discontinued Section | xxv |
| 1.12 Changes to the Orange Book..... | xxv |
| 1.13 Availability of the Edition | xxvi |
| | |
| 2.0 HOW TO USE THE DRUG PRODUCTS LISTS | 2-1 |
| 2.1 Key Sections for Using the Drug Product Lists | 2-1 |
| 2.2 Drug Product Illustration | 2-3 |
| 2.3 Therapeutic Equivalence Evaluations Illustration | 2-4 |
| | |
| DRUG PRODUCT LISTS | |
| Prescription Drug Product List | 3-1 |
| OTC Drug Product List | 4-1 |
| Drug Products with Approval under Section 505 of the FD&C Act Administered by the Center for Biologics Evaluation and Research List | 5-1 |
| Discontinued Drug Product List | 6-1 |
| Orphan Products Designations and Approvals List | 7-1 |
| Drug Products Which Must Demonstrate in vivo Bioavailability Only if Product Fails to Achieve Adequate Dissolution | 8-1 |
| | |
| APPENDICES | |
| A. Product Name Index | A-1 |
| B. Product Name Index Listed by Applicant | B-1 |
| C. Uniform Terms | C-1 |
| | |
| PATENT AND EXCLUSIVITY INFORMATION ADDENDUM | AD1 |
| A. Patent and Exclusivity Lists | ADA1 |
| B. Patent and Exclusivity Terms | ADB1 |

**FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH**

**APPROVED DRUG PRODUCTS
With
Therapeutic Equivalence Evaluations**

PREFACE TO THIRTY-NINTH EDITION

The publication, *Approved Drug Products With Therapeutic Equivalence Evaluations* (the List, commonly known as the Orange Book), identifies drug products approved on the basis of safety and effectiveness by the Food and Drug Administration (FDA) under the Federal Food, Drug, and Cosmetic Act (the FD&C Act). The main criterion for the inclusion of any product is that the product is the subject of an application with an approval that has not been withdrawn for safety or efficacy reasons. Inclusion of products in the Orange Book is independent of any current regulatory action being taken administratively or judicially against a drug product. In addition, the Orange Book contains therapeutic equivalence evaluations for approved multisource prescription drug products. These evaluations have been prepared to serve as public information and advice to state health agencies, prescribers, and pharmacists to promote public education in the area of drug product selection and to foster containment of health care costs. Therapeutic equivalence evaluations in this publication are not official FDA actions affecting the legal status of products under the FD&C Act.

Background of the Publication. To contain drug costs, virtually every state has adopted laws and/or regulations that encourage the substitution of drug products. These state laws generally require either that substitution be limited to drugs on a specific list (the positive formulary approach) or that it be permitted for all drugs except those prohibited by a particular list (the negative formulary approach). Because of the number of requests in the late 1970s for FDA assistance in preparing both positive and negative formularies, it became apparent that FDA could not serve the needs of each state on an individual basis. The Agency also recognized that providing a single list based on common criteria would be preferable to evaluating drug products on the basis of differing definitions and criteria in various state laws. As a result, on May 31, 1978, the Commissioner of the Food and Drug Administration sent a letter to officials of each state announcing FDA's intent to provide a list of all prescription drug products that are approved by FDA for safety and effectiveness, along with therapeutic equivalence determinations for multisource prescription products.

The Orange Book was distributed as a proposal in January 1979. It included only currently marketed prescription drug products approved by FDA through new drug applications (NDAs) and abbreviated new drug applications (ANDAs) under the provisions of Section 505 of the FD&C Act and FDA regulations at that time.

The therapeutic equivalence evaluations in the Orange Book reflect FDA's application of specific criteria to the multisource prescription drug products listed in the Orange Book and approved under Section 505 of the FD&C Act. These evaluations are presented in the form of code letters that indicate the basis for the evaluation made. An explanation of the codes appears in the *Introduction*.

A complete discussion of the background and basis of FDA's therapeutic equivalence evaluation policy was published in the *Federal Register* on January 12, 1979 (44 FR 2932). The final rule, which includes FDA's responses to the public comments on the proposal, was published in the *Federal Register* on October 31, 1980 (45 FR 72582). The first publication of the Orange Book in October 1980, concurrent with finalization of the rule, incorporated appropriate corrections and additions. Each subsequent edition has included new approvals and made appropriate changes in data.

On September 24, 1984, the President signed into law the Drug Price Competition and Patent Term Restoration Act of 1984 (Hatch-Waxman Amendments). The Hatch-Waxman Amendments require that FDA, among other things, make publicly available a list of approved drug products with monthly supplements. The Orange Book and its monthly Cumulative Supplements satisfy this requirement. The *Addendum* to this publication identifies drugs that qualify under the FD&C Act for periods of exclusivity and provides patent information concerning the approved drug products in the Orange Book. The *Addendum* also provides additional information that may be helpful to those submitting an NDA or ANDA to the Agency.

The Agency intends to use this publication to further its objective of obtaining input and comment on the publication itself and related Agency procedures. Therefore, if you have comments on how the publication can be improved, please send them to the Director, Division of Legal and Regulatory Support, Office of Generic Drug Policy, Office of Generic Drugs, Center for Drug Evaluation and Research, 7620 Standish Place, Rockville, MD 20855-2773. Comments received are publicly available to the extent allowable under the Freedom of Information Act and FDA regulations.

1.0 INTRODUCTION

1.1 Content and Exclusion

The Orange Book is composed of four parts: (1) approved prescription drug products with therapeutic equivalence evaluations; (2) approved over-the-counter (OTC) drug products for those drugs that may not be marketed without NDAs or ANDAs because they are not covered under existing OTC monographs; (3) drug products with approval under Section 505 of the FD&C Act administered by the Center for Biologics Evaluation and Research; and (4) a cumulative list of approved products that have never been marketed, are for exportation, are for military use, have been discontinued from marketing and we have not determined that they were withdrawn from sale for safety or effectiveness reasons, or have had their approvals withdrawn for other than safety or efficacy reasons subsequent to being discontinued from marketing.¹ This publication also includes indices of prescription and OTC drug products by trade name (proprietary name) or established name (if no trade name exists) and by applicant name (holder of the approved application), which have been abbreviated for this publication. Established names for active ingredients generally conform to official compendial names or *United States Adopted Names* (USAN) as described in 21 CFR 299.4(e). A list of uniform terms is provided in Appendix C.

The *Addendum* contains patent and exclusivity information for the Prescription, OTC, Discontinued Drug Product Lists, and for the Drug Products with Approval under Section 505 of the FD&C Act Administered by the Center for Biologics Evaluation and Research. The publication may include additional information that the Agency deems appropriate to disseminate.

Prior to the 6th Edition, the publication had excluded OTC drug products and drug products with approval under Section 505 of the FD&C Act administered by the Center for Biologics Evaluation and Research. The Hatch-Waxman Amendments required the Agency to begin publishing an up-to-date list of all marketed drug products, OTC as well as prescription, that have been approved for safety and efficacy and for which new drug applications are required.

Under the FD&C Act, some drug products are given tentative approvals. The Agency will not include drug products with tentative approvals in the Orange Book because a drug product that is granted tentative approval is not an approved drug product. Tentative approval lists by month are available on FDA's website Drugs@FDA. When the tentative approval becomes a final approval through a subsequent action letter to the applicant, the Agency will list the drug product and the date of approval in the appropriate approved drug product list. In addition, we note that Section 505(x) of the FD&C Act affects the date of approval for certain drug products subject to scheduling under the Controlled Substances Act. The Agency will list the drug product in the Orange Book and the date of approval as determined under Section 505(x).

The Orange Book identifies the application holder of a drug product and does not identify distributors or repackagers.

¹ Generally, newly approved products are added to the Active Section of the Orange Book (i.e., the Prescription Drug Product List or the Over-the-Counter Drug Product List), depending on the dispensing requirements (prescription or OTC) or approval authority, unless the Orange Book staff is otherwise notified before publication. See Section 1.12.

1.2 Therapeutic Equivalence-Related Terms

Pharmaceutical Equivalents. Pharmaceutical equivalents are drug products in identical dosage forms and route(s) of administration that contain identical amounts of the identical active drug ingredient, i.e., the same salt or ester of the same therapeutic moiety, or, in the case of modified-release dosage forms that require a reservoir or overage or such forms as prefilled syringes where the residual volume may vary, that deliver identical amounts of the active drug ingredient over the identical dosing period; do not necessarily contain the same inactive ingredients; and meet the identical compendial or other applicable standard of identity, strength, quality, and purity, including potency and, where applicable, content uniformity, disintegration times, and/or dissolution rates.² They may differ in characteristics such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration date/time, and, within certain limits, labeling.

Pharmaceutical Alternatives. Pharmaceutical alternatives are drug products that contain the identical therapeutic moiety, or its precursor, but not necessarily in the same amount or dosage form, or the same salt or ester (e.g., tetracycline hydrochloride, 250mg capsules vs. tetracycline phosphate complex, 250mg capsules; quinidine sulfate, 200mg tablets vs. quinidine sulfate, 200mg capsules).³ Each such drug product individually meets either the identical or its own respective compendial or other applicable standard of identity, strength, quality, and purity, including potency and, where applicable, content uniformity, disintegration times, and/or dissolution rates.⁴ Different dosage forms and strengths within a product line by a single manufacturer are pharmaceutical alternatives, as are extended-release products when compared with immediate-release or standard-release formulations of the same active ingredient.

Therapeutic Equivalents. Approved drug products are considered to be therapeutic equivalents only if they are pharmaceutical equivalents for which bioequivalence has been demonstrated, and they can be expected to have the same clinical effect and safety profile when administered to patients under the conditions specified in the labeling.⁵

FDA classifies as therapeutically equivalent those drug products that meet the following general criteria: (1) they are approved as safe and effective; (2) they are pharmaceutical equivalents in that they (a) contain identical amounts of the identical active drug ingredient in the identical dosage form and route of administration, and (b) meet compendial or other applicable standards of strength, quality, purity, and identity; (3) they are bioequivalent in that (a) they do not present a known or potential bioequivalence problem, and they meet an acceptable *in vitro* standard, or (b) if they do present such a known or potential problem, they are shown to meet an appropriate bioequivalence standard; (4) they are adequately labeled; and (5) they are manufactured in compliance with Current Good Manufacturing Practice regulations. The concept of therapeutic equivalence applies only to drug products containing the identical active ingredient(s) and does not encompass a comparison of different therapeutic agents used for the same condition (e.g., meperidine hydrochloride vs. morphine sulfate for the treatment of pain). Any drug product in the Orange Book repackaged and/or distributed by other than the applicant is considered to be therapeutically

2 21 CFR 314.3(b).

3 See 21 CFR 314.3(b).

4 21 CFR 314.3(b).

5 21 CFR 314.3(b).

equivalent to the applicant's drug product even if the applicant's drug product is single source or coded as non-equivalent (e.g., **BN**). Distributors or repackagers of an applicant's drug product are not identified in the Orange Book.

FDA considers drug products to be therapeutically equivalent if they meet the criteria outlined above, even though they may differ in certain other characteristics such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration date/time and certain aspects of labeling (e.g., the presence of specific pharmacokinetic information), and storage conditions. When such differences are important in the care of a particular patient, it may be appropriate for the prescribing physician to require that a specific product be dispensed as a medical necessity. With this limitation, however, FDA believes that products classified as therapeutically equivalent can be substituted with the full expectation that the substituted product can be expected to have the same clinical effect and safety profile as the prescribed product when administered to patients under the conditions specified in the labeling.

Strength. Strength refers to the amount of drug substance contained in, delivered, or deliverable from a drug product, which includes: (1)(a) the total quantity of drug substance in mass or units of activity in a dosage unit or container closure (e.g., weight/unit dose, weight/volume or weight/weight in a container closure, or units/volume or units/weight in a container closure); and/or, as applicable, (b) the concentration of the drug substance in mass or units of activity per unit volume or mass (e.g., weight/weight, weight/volume, or units/volume); or (2) such other criteria the Agency establishes for determining the amount of drug substance contained in, delivered, or deliverable from a drug product if the weights and measures described in clause (1)(a) do not apply (e.g., certain drug-device combination products for which the amount of drug substance is emitted per use or unit time).⁶ Note that if the criteria the Agency establishes for determining and expressing the amount of drug substance in a product evolves over time, the Agency generally does not intend to revise the expressions of strength for drug products already included in the Orange Book, but rather intends to apply the criteria prospectively to drug products added to the Orange Book.

Although the strength of drug products in the Orange Book is generally expressed in terms of the amount of drug substance (active ingredient) in the drug product, it is sometimes expressed in terms of the amount of the active moiety. For example, certain drug products included in the Orange Book include a designation of "EQ" next to their expression of strength. This "EQ" designation generally is used in connection with salt drug products to indicate that the strength of such drug product is being expressed in terms of the equivalent strength of the active moiety (e.g., "EQ 200MG BASE"), rather than in terms of the strength of the active ingredient.

Bioavailability. Bioavailability is the rate and extent to which the active ingredient or active moiety is absorbed from a drug product and becomes available at the site of drug action. For drug products that are not intended to be absorbed into the bloodstream, bioavailability may be assessed by scientifically valid measurements intended to reflect the rate and extent to which the active ingredient or active moiety becomes available at the site of drug action.⁷

⁶ See 21 CFR 314.3(b).

⁷ 21 CFR 314.3(b).

Bioequivalence. Bioequivalence is the absence of a significant difference in the rate and extent to which the active ingredient or active moiety in pharmaceutical equivalents or pharmaceutical alternatives becomes available at the site of drug action when administered at the same molar dose under similar conditions in an appropriately designed study.⁸ Section 505(j)(8)(B) of the FD&C Act describes certain conditions under which a test drug and reference listed drug (see Section 1.4) shall be considered bioequivalent:

- (i) the rate and extent of absorption of the [test] drug do not show a significant difference from the rate and extent of absorption of the [reference] listed drug when administered at the same molar dose of the therapeutic ingredient under similar experimental conditions in either a single dose or multiple doses; or
- (ii) the extent of absorption of the [test] drug does not show a significant difference from the extent of absorption of the [reference] listed drug when administered at the same molar dose of the therapeutic ingredient under similar experimental conditions in either a single dose or multiple doses and the difference from the [reference] listed drug in the rate of absorption of the drug is intentional, is reflected in its proposed labeling, is not essential to the attainment of effective body drug concentrations on chronic use, and is considered medically insignificant for the drug.

Where these above methods are not applicable (e.g., for drug products that are not intended to be absorbed into the bloodstream), other scientifically valid *in vivo* or *in vitro* test methods to demonstrate bioequivalence may be appropriate.

For example, bioequivalence may sometimes be demonstrated using an *in vitro* bioequivalence standard, especially when such an *in vitro* test has been correlated with human *in vivo* bioavailability data. In other situations, bioequivalence may sometimes be demonstrated through comparative clinical trials or pharmacodynamic studies.⁹

1.3 Further Guidance on Bioequivalence

FDA's regulations and guidance documents provide additional information regarding bioequivalence and bioavailability, including methodologies and statistical criteria used to establish the bioequivalence of drug products.¹⁰

1.4 Reference Listed Drug and Reference Standard

A reference listed drug is the listed drug identified by FDA as the drug product upon which an applicant relies in seeking approval of its ANDA.¹¹ Generally, a reference listed drug is a drug product approved in a new drug application under Section 505(c) of the FD&C Act based on full reports of

⁸ 21 CFR 314.3(b).

⁹ 21 CFR 320.24

¹⁰ We note that prior editions of the Preface to the Orange Book included a section entitled "Statistical Criteria for Bioequivalence." Please see FDA's regulations and guidance documents for additional information regarding bioequivalence and bioavailability. See generally 21 CFR part 320. See FDA Drugs guidance Web page at <http://www.fda.gov/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/default.htm>; FDA Drugs guidance (Product-Specific Recommendations for Generic Drug Development) Web page at <http://www.fda.gov/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/ucm075207.htm>.

¹¹ 21 CFR 314.3(b).

investigations of safety and effectiveness. For an ANDA based on an approved suitability petition (a petitioned ANDA), the reference listed drug generally is the listed drug referenced in the approved suitability petition.¹²

A reference standard is the drug product selected by FDA that an applicant seeking approval of an ANDA must use in conducting an *in vivo* bioequivalence study required for approval.¹³ FDA generally selects a single reference standard that ANDA applicants must use in *in vivo* bioequivalence testing. Ordinarily, FDA will select the reference listed drug as the reference standard. However, in some instances, the reference listed drug and the reference standard may be different. For example, where the reference listed drug has been withdrawn from sale and FDA has determined it was not withdrawn for reasons of safety or effectiveness, FDA may select an ANDA as the reference standard.

FDA identifies reference listed drugs in the Prescription Drug Product, OTC Drug Product, and Discontinued Drug Product Lists. Listed drugs identified as reference listed drugs represent drug products upon which an applicant can rely in seeking approval of an ANDA. FDA intends to update periodically the reference listed drugs identified in the Prescription Drug Product, OTC Drug Product, and Discontinued Drug Product Lists, as appropriate.

In some instances when FDA has not designated a listed drug as a reference listed drug, such listed drug may be shielded from generic competition. If FDA has not designated a reference listed drug for a drug product the applicant intends to duplicate, the potential applicant may submit a controlled correspondence to the Office of Generic Drugs to ask FDA to designate a reference listed drug for that drug product. Section 1.7, *Therapeutic Equivalence Evaluations Codes (products meeting necessary bioequivalence requirements)* explains the character coding system (e.g., **AB**, **AB1**, **AB2**, **AB3**...) for multisource drug products listed under the same heading with two or more reference listed drugs.

FDA also identifies reference standards in the Prescription Drug Product and OTC Drug Product Lists. Listed drugs identified as reference standards represent FDA's best judgment at this time as to the appropriate comparator for purposes of conducting any *in vivo* bioequivalence studies required for approval.

A potential applicant should consult Agency guidance related to referencing approved drug products in ANDA submissions for information on submitting a request for selection of a reference standard. FDA may, on its own initiative, select a new reference standard when doing so will help to ensure that applications for generic drugs may be submitted and evaluated, e.g., in the event that the listed drug currently selected as the reference standard has been withdrawn from sale for other than safety and efficacy reasons.

If an applicant has a question related to the appropriate reference standard, it is recommended that an applicant planning to conduct an *in vivo* bioequivalence study submit a controlled correspondence to the Office of Generic Drugs.

1.5 General Policies and Legal Status

¹² 21 CFR 314.94(a)(3)(i).

¹³ 21 CFR 314.3(b).

The Orange Book contains public information and advice. It does not mandate the drug products that are purchased, prescribed, dispensed, or substituted for one another, nor does it, conversely, mandate the products that should be avoided. To the extent that the Orange Book sets forth FDA's evaluations of the therapeutic equivalence of drug products that have been approved, it contains FDA's advice to the public, to practitioners, and to the states regarding drug product selection. These evaluations do not constitute determinations that any product is in violation of the FD&C Act or that any product is preferable to any other. Therapeutic equivalence evaluations are a scientific judgment based upon evidence, while generic substitution may involve social and economic policy administered by the states, e.g., reducing the cost of drugs to consumers. To the extent that the Orange Book identifies drug products approved under Section 505 of the FD&C Act, it sets forth information that the Agency is required to publish and that the public is entitled to under the Freedom of Information Act. Exclusion of a drug product from the Orange Book does not necessarily mean that the drug product is in violation of Section 505 of the FD&C Act, that such a product is not safe or effective, or that such a product is not therapeutically equivalent to other drug products. Rather, the exclusion may be based on the fact that FDA has not evaluated the safety, effectiveness, and quality of the drug product.

1.6 Practitioner/User Responsibilities

Professional care and judgment should be exercised in using the Orange Book. Evaluations of therapeutic equivalence for prescription drugs are based on scientific and medical evaluations by FDA. Products evaluated as therapeutically equivalent can be expected, in the judgment of FDA, to have the same clinical effect and safety profile when administered to patients under the conditions specified in the labeling. However, these products may differ in other characteristics that are not required by statute or regulation to be the same, such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration date/time, and, in some instances, labeling. If products with such differences are substituted for each other, there is a potential for patient confusion, e.g., due to differences in color or shape of tablets, inability to provide a given dose using a partial tablet if the proper scoring configuration is not available, or decreased patient acceptance of certain products because of flavor. There may also be patient-specific allergic reactions in rare cases due to a coloring or a preservative ingredient.

FDA evaluation of therapeutic equivalence in no way relieves practitioners of their professional responsibilities in prescribing and dispensing such products with due care and with appropriate information to individual patients. In those circumstances where the characteristics of a specific product, other than its active ingredient, are important in the therapy of a particular patient, the practitioner's prescribing of that product may be appropriate. Pharmacists must also be familiar with the different characteristics of therapeutically equivalent products, e.g., expiration dates/times and labeling directions for storage of the different products (particularly for reconstituted products), so they can properly advise patients when one product is substituted for another.

Multisource and single-source drug products. In the Orange Book, FDA has evaluated for therapeutic equivalence only multisource prescription drug products approved under Section 505 of the FD&C Act, which in most instances means those pharmaceutical equivalents available from more than one manufacturer. For such products, a therapeutic equivalence code is included

and product information is highlighted in bold face and underlined. Those products with approved applications that are single-source (i.e., there is only one approved product available for that active ingredient, dosage form, route of administration, and strength) are also included in the Orange Book, but no therapeutic equivalence code is included with such products. Any drug product in the Orange Book repackaged and/or distributed by the applicant or some other person authorized by the applicant (e.g., an authorized generic) is considered to be therapeutically equivalent to the applicant's drug product even if the applicant's drug product is single source or coded as non-equivalent (e.g., **BN**). Distributors or repackagers of an applicant's drug product are not identified in the Orange Book. The details of therapeutic equivalence codes and the policies underlying them are discussed in Section 1.7, *Therapeutic Equivalence Evaluations Codes*.

Products in the Orange Book are identified by the names of the holders of approved applications (applicants) who may not necessarily be the manufacturer of the product. There are numerous entities other than the applicant that may be involved in the development, manufacturing, and/or marketing of a product. The applicant may or may not be the manufacturer and may simply be distributing the product for which it has obtained approval. In many instances, however, the manufacturer of the product is also the applicant. The name of the manufacturer is permitted by regulation to appear on the label, even when the manufacturer is not the applicant or marketer.

Although the products in the Orange Book are identified by the names of the applicants, circumstances, such as changing corporate ownership, have sometimes made identification of the applicant difficult. The Agency believes, based on continuing document review and communication with firms, that the applicant designations in the Orange Book are, in most cases, correct.

To relate firm name information on a product label to that in the Orange Book, the following should be noted: the applicant's name always appears in the Orange Book. This applies whether the applicant (firm name on the Form FDA 356h in the application) is the manufacturer or marketer (firm name in largest letters on the label) or not. However, the applicant's name may not always appear on the label of the product.

If the applicant is the marketer, its name appears in the Orange Book and on the label; if the applicant is not the marketer, and the Agency is aware of a corporate relationship (e.g., parent and subsidiary) between the applicant and the marketer, the name of the applicant appears in the Orange Book and both firm names may appear on the label. Firms with known corporate relationships are displayed in Appendix B. If there is no known corporate relationship between the applicant and the marketer, the applicant's name appears in the Orange Book; however, unless the applicant is the manufacturer, packager, or distributor, the applicant's name may not appear on the label. In this case, the practitioner, from labeling alone, will not be able to relate the marketed product to an applicant cited in the Orange Book, and hence to a specific approved drug product. In such cases, to assure that the product in question is the subject of an approved application, the firm named on the label should be contacted.

To relate trade name (proprietary name) information on a product label to that in the Orange Book, the following should be noted: if the applicant is the marketer, the applicant's name appears in the Orange Book and on the label; if the Agency is aware of a corporate relationship between the applicant and the marketer, the trade name (proprietary name) of the drug product (established name of the active ingredient, if no trade name exists) appears in the Orange Book. If a corporate relationship exists between an applicant and a marketer and both firms are distributing the drug product,

the FDA reserves the right to select the trade name of either the marketer or the applicant to appear in the Orange Book. If there is no known corporate relationship between the applicant and the marketer, the established drug name (i.e., non-proprietary name) appears in the Orange Book.

Every product in the Orange Book is subject at all times to regulatory action. From time to time, approved products may be found in violation of one or more provisions of the FD&C Act. In such circumstances, the Agency may commence appropriate enforcement action to correct the violation, if necessary, by securing removal of the product from the market by voluntary recall, seizure, or other enforcement actions. Such regulatory actions are, however, independent of the inclusion of a product in the Orange Book. The main criterion for inclusion of a product is that it has an application that has been approved and that has not been withdrawn for safety or efficacy reasons. FDA believes that retention of a violative product in the Orange Book will not have any significant adverse health consequences, because other legal mechanisms are available to the Agency to prevent the product's actual marketing. FDA may, however, change a product's therapeutic equivalence rating if the circumstances giving rise to the violation change or otherwise call into question the Agency's assessment of whether a product meets the criteria for therapeutic equivalence.

1.7 Therapeutic Equivalence Evaluations Codes

Generally, drug products that the Agency considers multisource have been assigned a therapeutic equivalence code. The coding system for therapeutic equivalence evaluations is designed to allow users to determine quickly whether the Agency has evaluated a particular approved product (e.g., a particular strength of an approved drug) as therapeutically equivalent to other pharmaceutically equivalent products (first letter) and to provide additional information on the basis of FDA's evaluations (second letter). With some exceptions (e.g., therapeutic equivalence evaluations for certain 505(b)(2) applications), the therapeutic equivalence evaluation date is the same as the approval date.

The two basic categories into which multisource drugs have been placed are indicated by the first letter of the relevant therapeutic equivalence code as follows:

A Drug products that FDA considers to be therapeutically equivalent to other pharmaceutically equivalent products, i.e., drug products for which:

- (1) there are no known or suspected bioequivalence problems. These are designated **AA, AN, AO, AP, or AT**, depending on the dosage form; or
- (2) actual or potential bioequivalence problems have been resolved with adequate *in vivo* and/or *in vitro* evidence supporting bioequivalence. These are designated **AB**.

B Drug products that FDA at this time, considers not to be therapeutically equivalent to other pharmaceutically equivalent products, i.e.,

drug products for which actual or potential bioequivalence problems have not been resolved by adequate evidence of bioequivalence. Often the problem is with specific dosage forms rather than with the active ingredients. These are designated **BC, BD, BE, BN, BP, BR, BS, BT, BX, or B***.

Individual drug products have been evaluated as therapeutically equivalent to the reference product in accordance with the definitions and policies outlined below:

"A" CODES

Drug products that are considered to be therapeutically equivalent to other pharmaceutically equivalent products.

"A" products are those for which there are no known or suspected bioequivalence problems or for which actual or potential bioequivalence problems have been resolved with adequate *in vivo* and/or *in vitro* evidence supporting bioequivalence. Drug products designated with an "A" code fall under one of two main policies:

- (1) for those active ingredients or dosage forms for which no *in vivo* bioequivalence issue is known or suspected, the information necessary to show bioequivalence between pharmaceutically equivalent products is either presumed and considered self-evident (based on other information in the application for some dosage forms (e.g., solutions)), or satisfied by a showing that an acceptable *in vitro* approach is met. A therapeutically equivalent rating is assigned such products so long as they are manufactured in accordance with Current Good Manufacturing Practice regulations and meet the other requirements of their approved applications (these are designated **AA**, **AN**, **AO**, **AP**, or **AT**, depending on the dosage form, as described below); or
- (2) for those Drug Efficacy Study Implementation (DESI) drug products containing active ingredients or dosage forms that have been identified by FDA as having actual or potential bioequivalence problems, and for post-1962 drug products presenting a potential bioequivalence problem, an evaluation of therapeutic equivalence is assigned to pharmaceutical equivalents only if the approved application contains adequate scientific evidence establishing through *in vivo* and/or *in vitro* studies the bioequivalence of the product to a selected reference product (these products are designated as **AB**).

There are some general principles that may affect the substitution of pharmaceutically equivalent products in specific cases. Prescribers and dispensers of drugs should be alert to these principles so as to deal appropriately with situations that require professional judgment and discretion.

There may be labeling differences among pharmaceutically equivalent products that require attention on the part of the health professional (e.g., pharmaceutically equivalent powders to be reconstituted for administration as oral or injectable liquids may vary with respect to their expiration time or storage conditions after reconstitution). FDA's determination that such products are therapeutically equivalent is applicable only when each product is reconstituted, stored, and used under the conditions specified in its labeling.

The Agency may use notes in this publication to point out special situations, such as potential differences between two drug products that have been evaluated as bioequivalent and otherwise therapeutically equivalent, when they should be brought to the attention of health professionals. These notes are contained in Section 1.8, *Description of Certain Special Situations*.

For example, in certain instances, there may be variations among therapeutically equivalent products in their use or in conditions of administration. Such differences may be due to patent or exclusivity rights associated with such use. When such variations may, in the Agency's opinion, affect prescribing or substitution decisions by health professionals, a note may be added to Section 1.8.

Also, occasionally a situation may arise in which changes in a listed drug product after its approval (for example, a change in dosing interval) may have an impact on the substitutability of already approved generic versions of that product that were rated by the Agency as therapeutically equivalent to the listed product. When such changes in the listed drug product are considered by the Agency to have a significant impact on therapeutic equivalence, the Agency will change the therapeutic equivalence ratings for other versions of the drug product unless the manufacturers of those other versions of the product provide additional information to assure equivalence under the changed conditions. Pending receipt of the additional data, the Agency may add a note to Section 1.8, or, in rare cases, may even change the therapeutic equivalence rating.

In some cases (e.g., Isolyte® S w/ Dextrose 5% in Plastic Container and Plasma-Lyte® 148 and Dextrose 5% in Plastic Container), closely related products are listed as containing the same active ingredients, but in somewhat different amounts. In determining which of these products are pharmaceutically equivalent, generally the Agency has considered products to be pharmaceutically equivalent with labeled strengths of an ingredient that do not vary by more than 1%.

Different salts, esters or other noncovalent derivatives (such as a complex, chelate, or clathrate) of the same active moiety are regarded as different active ingredients. For the purpose of this publication, products containing such different active ingredients are considered pharmaceutical alternatives and, thus, not therapeutically equivalent. Anhydrous and hydrated entities, as well as different polymorphs, are considered to be the same active ingredient and are expected to meet the same standards for identity to be considered pharmaceutical equivalents and therapeutic equivalents.

The codes in this book are not intended to preclude health care professionals from converting pharmaceutically different concentrations into pharmaceutical equivalents using accepted professional practice.

Where package size variations have therapeutic implications, products so packaged have not been considered pharmaceutically equivalent. For example, some oral contraceptives are supplied in 21-tablet and 28-tablet packets; the 28-tablet packets contain 7 placebo or iron tablets. These two packaging configurations are not regarded as pharmaceutically equivalent; thus, they are not designated as therapeutically equivalent.

Preservatives and other inactive ingredients may differ among some therapeutically equivalent drug products. These differences do not affect FDA's evaluation of therapeutic equivalence except in cases where these components may influence bioequivalence or routes of administration.

The specific sub-codes for those drugs evaluated as therapeutically equivalent and the policies underlying these sub-codes follow:

AA Products in conventional dosage forms not presenting bioequivalence problems

Multisource drug products coded as **AA** contain active ingredients and are in dosage forms that are not regarded as presenting either actual or

potential bioequivalence problems or drug quality or standards issues. However, all oral dosage forms must, nonetheless, meet an appropriate *in vitro* bioequivalence standard that is acceptable to the Agency in order to be approved.

AB, AB1, AB2, AB3... Products meeting necessary bioequivalence requirements

Multisource drug products listed under the same heading (i.e., identical active ingredients(s), dosage form, and route(s) of administration) and having the same strength (see Section 1.2, *Therapeutic Equivalence-Related Terms, Strength*) generally will be coded **AB** if data and information are submitted demonstrating bioequivalence.

In certain instances, a number is added to the end of the **AB** code to make a three character code (i.e., **AB1, AB2, AB3, etc.**). Three-character codes generally are assigned only in situations when more than one reference listed drug of the same strength has been designated under the same heading. If a study is submitted that demonstrates bioequivalence to a reference listed drug product, the generic product will be given the same three-character code as the reference listed drug it was compared against. For example, Adalat® CC and Procardia XL®, extended-release tablets, are listed under the active ingredient nifedipine. These drug products, listed under the same heading, are not bioequivalent to each other. Adalat® CC and Procardia XL® have been assigned ratings of **AB1** and **AB2**, respectively. Generic drug products deemed by FDA to be bioequivalent to Adalat® CC and Procardia XL® have been approved. As a result, the generic drug products bioequivalent to Adalat® CC have been assigned a rating of **AB1** and those bioequivalent to Procardia XL® have been assigned a rating of **AB2**. (The assignment of an **AB1** or **AB2** rating to a specific product does not imply product preference.) Even though drug products of distributors and/or repackagers are not included in the Orange Book, they are considered therapeutically equivalent to the applicant's drug product if the applicant's drug product is rated either with an **AB** or three-character code or is single source in the Orange Book. Drugs coded as **AB** under a heading are considered therapeutically equivalent only to other drugs coded as **AB** under that heading. Drugs coded with a three-character code under a heading are considered therapeutically equivalent only to other drugs coded with the same three-character code under that heading.

AN Solutions and powders for aerosolization

Uncertainty regarding the therapeutic equivalence of aerosolized products arises primarily because of differences in the drug delivery system. Solutions and powders intended for aerosolization that are marketed for use in general-use delivery systems are considered to be pharmaceutically and therapeutically equivalent and are coded **AN**. Those products that are compatible only with a specific delivery system or those products that are packaged in and with a specific delivery system are coded **BN**, unless they have met an appropriate bioequivalence standard and are otherwise determined to be therapeutically equivalent. Solutions or suspensions in a specific delivery system will be coded **AN** if the bioequivalence standard is based upon *in vitro* methodology, if bioequivalence needs to be demonstrated by *in vivo* methodology then the drug products will be coded **AB**.

AO Injectable oil solutions

The absorption of drugs in injectable (parenteral) oil solutions may vary substantially with the type of oil employed as a vehicle and the

concentration of the active ingredient. Injectable oil solutions are therefore considered to be pharmaceutically and therapeutically equivalent only when the active ingredient, its concentration, and the type of oil used as a vehicle are all identical.

AP Injectable aqueous solutions and, in certain instances, intravenous non-aqueous solutions

It should be noted that even though injectable (parenteral) products under a specific listing may be evaluated as therapeutically equivalent, there may be important differences among the products in the general category, Injectable; Injection. For example, historically some injectable products that are rated therapeutically equivalent are labeled for different routes of administration. In addition, some products evaluated as therapeutically equivalent may have different preservatives or no preservatives at all. Injectable products available as dry powders for reconstitution, concentrated sterile solutions for dilution, or sterile solutions ready for injection are pharmaceutical alternative drug products. They are not rated as therapeutically equivalent (AP) to each other even if these pharmaceutical alternative drug products are designed to produce the same concentration prior to injection and are similarly labeled. Consistent with accepted professional practice, it is the responsibility of the prescriber, dispenser, or individual administering the product to be familiar with a product's labeling to assure that it is given only by the route(s) of administration stated in the labeling.

Certain commonly used large volume intravenous products in glass containers are not included in the Orange Book (e.g., dextrose injection 5%, dextrose injection 10%, sodium chloride injection 0.9%) since these products are on the market without FDA approval and FDA has not published conditions for marketing such parenteral products under approved NDAs. When packaged in plastic containers, however, FDA regulations require approved applications prior to marketing. Approval then depends on, among other things, the extent of the available safety data involving the specific plastic component of the product. All large volume parenteral products are manufactured under similar standards, regardless of whether they are packaged in glass or plastic. Thus, FDA has no reason to believe that the packaging container of large volume parenteral drug products that are pharmaceutically equivalent would have any effect on their therapeutic equivalence.

Consistent with the definition of strength included in Section 1.2, *Therapeutic Equivalence-Related Terms*, the strength of parenteral drug products generally is identified by both the total drug content and the concentration of drug substance in a container approved by FDA.¹⁴ In the past, the strength of liquid parenteral drug products in the Orange Book has not been fully displayed. Rather, the strength of liquid parenteral drug products in the Orange Book has been displayed in terms of concentration, expressed as xmg/mL. Generally, the amount of dry powder or lyophilized powder in a container is identified as the strength, expressed as xmg/vial.

After the enactment of the Medicare Prescription Drug, Improvement, and Modernization Act of 2003 (MMA), which amended the FD&C Act, it became evident that the format of the Orange Book with respect to parenteral

¹⁴ The strengths of certain parenteral drug products, including contrast agents, may be expressed as a percentage.

solutions should be changed to reflect that each strength of a drug is considered to be a separate listed drug. The Orange Book now displays the strength of all new approvals of parenteral solutions. Previously, we would have displayed only the concentration of an approved parenteral solution, e.g. 50mg/mL. For example, if this application had a 20 mL and 60 mL container approved, we would now display two product strengths, listing both total drug content and concentration of drug substance in the relevant approved container, e.g. 1gm/20mL (50mg/mL) and 3gm/60mL (50mg/mL).

AT Topical products

There are a variety of topical dosage forms available for dermatologic, ophthalmic, otic, rectal, and vaginal administration, including creams, gels, lotions, oils, ointments, pastes, solutions, sprays, and suppositories. Even though different topical dosage forms may contain the same active ingredient and potency, these dosage forms are not considered pharmaceutically equivalent. Therefore, they are not considered therapeutically equivalent. All solutions and DESI drug products containing the same active ingredient in the same topical dosage form for which a waiver of *in vivo* bioequivalence has been granted, or the application contains adequate scientific evidence establishing through an *in vitro* approach the bioequivalence of the product to a selected reference product, and for which chemistry and manufacturing processes are adequate to demonstrate bioequivalence, are considered therapeutically equivalent and coded **AT**. Pharmaceutically equivalent topical products that raise questions of bioequivalence and for which a waiver of *in vivo* bioequivalence has not been granted, including all post-1962 non-solution topical drug products, are coded **AB** when supported by adequate *in vivo* bioequivalence data, and **BT** in the absence of such data.

"B" CODES

Drug products that FDA, at this time, considers not to be therapeutically equivalent to other pharmaceutically equivalent products.

"B" products, for which actual or potential bioequivalence problems have not been resolved by adequate evidence of bioequivalence, often have a problem with specific dosage forms rather than with the active ingredients. Drug products designated with a "B" code fall under one of three main policies:

- (1) the drug products contain active ingredients or are manufactured in dosage forms that have been identified by the Agency as having documented bioequivalence problems or a significant potential for such problems and for which no adequate studies demonstrating bioequivalence have been submitted to FDA; or
- (2) the quality standards are inadequate or FDA has an insufficient basis to determine therapeutic equivalence; or
- (3) the drug products are under regulatory review.

The specific coding definitions and policies for the "B" sub-codes are as follows:

B* Drug products requiring further FDA investigation and review to determine therapeutic equivalence

The code **B*** is assigned to products previously assigned an **A** or **B** code when FDA receives new information that raises a significant question regarding therapeutic equivalence that can be resolved only through further Agency investigation and/or review of data and information submitted by the applicant. The **B*** code signifies that the Agency will take no position regarding the therapeutic equivalence of the product until the Agency completes its investigation and review.

BC Extended-release dosage forms (capsules, injectables and tablets)

Extended-release tablets are formulated in such a manner as to make the contained drug substance available over an extended period of time following ingestion.

Although bioavailability studies have been conducted on these dosage forms, they may be subject to bioavailability differences, primarily because applicants developing extended-release products for the same active ingredient rarely employ the same formulation approach. FDA, therefore, does not consider different extended-release dosage forms containing the same active ingredient in equal strength to be therapeutically equivalent unless equivalence between individual products in both rate and extent has been specifically demonstrated through appropriate bioequivalence studies. Extended-release products for which such bioequivalence data have not been submitted are coded **BC**, while those for which such data are available have been coded **AB**.

BD Active ingredients and dosage forms with documented bioequivalence problems

The **BD** code denotes products containing active ingredients with known bioequivalence problems and for which adequate studies have not been submitted to FDA demonstrating bioequivalence. Where studies showing bioequivalence have been submitted, the product has been coded **AB**.

BE Delayed-release oral dosage forms

Where the drug may be destroyed or inactivated by the gastric juice or where it may irritate the gastric mucosa, the use of "enteric" coatings is indicated. Such coatings are intended to delay the release of the medication until the tablet has passed through the stomach. Drug products in delayed-release dosage forms containing the same active ingredients are subject to significant differences in absorption. Unless otherwise specifically noted, the Agency considers different delayed-release products containing the same active ingredients as presenting a potential bioequivalence problem and codes these products **BE** in the absence of *in vivo* studies showing bioequivalence. If adequate *in vivo* studies have demonstrated the bioequivalence of specific delayed-release products, such products are coded **AB**.

BN Products in aerosol-nebulizer drug delivery systems

This code applies to drug solutions or powders that are marketed only as a component of, or as compatible with, a specific drug delivery system. There may, for example, be significant differences in the dose of drug and particle size delivered by different products of this type. Therefore,

the Agency does not consider different metered aerosol dosage forms containing the same active ingredient(s) in equal strengths to be therapeutically equivalent unless the drug products meet an appropriate bioequivalence standard; such products are coded **AB**.

BP Active ingredients and dosage forms with potential bioequivalence problems

FDA's bioequivalence regulations (21 CFR 320.33) contain criteria and procedures for determining whether a specific active ingredient in a specific dosage form has a potential for causing a bioequivalence problem. It is FDA's policy to consider an ingredient meeting these criteria as having a potential bioequivalence problem even in the absence of positive data demonstrating inequivalence. Pharmaceutically equivalent products containing these ingredients in oral dosage forms are coded **BP** until adequate bioequivalence data are submitted, after which such products are coded **AB**. Injectable suspensions containing an active ingredient suspended in an aqueous or oleaginous vehicle have also been coded **BP**. Injectable suspensions are subject to bioequivalence problems because differences in particle size, polymorphic structure of the suspended active ingredient, or the suspension formulation can significantly affect the rate of release and absorption. FDA does not consider pharmaceutical equivalents of these products bioequivalent without adequate evidence of bioequivalence; such products would be coded **AB**.

BR Suppositories or enemas that deliver drugs for systemic absorption

The absorption of active ingredients from suppositories or enemas that are intended to have a systemic effect (as distinct from suppositories administered for local effect) can vary significantly from product to product. Therefore, FDA considers pharmaceutically equivalent systemic suppositories or enemas bioequivalent only if *in vivo* evidence of bioequivalence is available. In those cases where *in vivo* evidence is available, the products are coded **AB**. If such evidence is not available, the products are coded **BR**.

BS Products having drug standard deficiencies

If the drug standards for an active ingredient in a particular dosage form are found by FDA to be deficient so as to prevent an FDA evaluation of either pharmaceutical or therapeutic equivalence, all drug products containing that active ingredient in that dosage form are coded **BS**. For example, if the standards permit a wide variation in pharmacologically active components of the active ingredient such that pharmaceutical equivalence is in question, all products containing that active ingredient in that dosage form are coded **BS**.

BT Topical products with bioequivalence issues

This code applies mainly to post-1962 dermatologic, ophthalmic, otic, rectal, and vaginal products for topical administration, including creams, gels, lotions, oils, ointments, pastes, solutions, and sprays, as well as suppositories not intended for systemic drug absorption. Topical products evaluated as having acceptable clinical performance, but that are not bioequivalent to other pharmaceutically equivalent products or that lack sufficient evidence of bioequivalence, will be coded **BT**.

BX Drug products for which the data are insufficient to determine therapeutic equivalence

The code **BX** is assigned to specific drug products for which the data that have been reviewed by the Agency are insufficient to determine therapeutic equivalence under the policies stated in this document. In these situations, the drug products are presumed to be therapeutically inequivalent until the Agency has determined that there is adequate information to make a full evaluation of therapeutic equivalence.

1.8 Description of Certain Special Situations

Certain drugs listed in the Orange Book present special situations that merit further discussion. The following are descriptions of certain examples of those special situations:

Amino Acid and Protein Hydrolysate Injections. These products differ in the amount and kinds of amino acids they contain and, therefore, are not considered pharmaceutical equivalents. For this reason, these products are not considered therapeutically equivalent. At the same time, the Agency believes that it is appropriate to point out that where nitrogen balance is the sole therapeutic objective and individual amino acid content is not a consideration, pharmaceutical alternatives with the same total amount of nitrogen content may be expected to have the same clinical effect and safety profile when administered to patients under the conditions specified in the labeling.

Gaviscon®. Gaviscon® is an OTC product that has been marketed since September 1970. The active ingredients in this product, aluminum hydroxide and magnesium trisilicate, were reviewed by the Agency's OTC Antacid Panel and were considered to be safe and effective ingredients (Category I) by that Panel. However, the tablet failed to pass the antacid test that is required of all antacid products. The Agency, therefore, placed the tablet in Category III for lack of effectiveness. A full NDA with clinical studies was submitted by Marion Laboratories, Inc., and approved by FDA on December 9, 1983. Gaviscon®'s activity in treating reflux acidity is made possible by the physical-chemical properties of the inactive ingredients, sodium bicarbonate and alginic acid. Therefore, *all ANDAs that cite Gaviscon® tablets as the reference listed drug must contain the inactive ingredients sodium bicarbonate and alginic acid.* A full NDA will be required to support the effectiveness of the drug product if different inactive ingredients are to be substituted for sodium bicarbonate or alginic acid or if different proportions of these ingredients are to be used.

Levothyroxine Sodium. Because there are multiple reference listed drugs for levothyroxine sodium tablets and some reference listed drugs' sponsors have conducted studies to establish their drugs' therapeutic equivalence to other reference listed drugs, FDA has determined that its usual practice of assigning two or three character TE codes may be potentially confusing and inadequate for these drug products. Accordingly, FDA provides the following explanation and chart of therapeutic equivalence evaluations for levothyroxine sodium tablet drug products.

Levothyroxine Sodium (Mylan ANDA 076187), Levoxyl (King Pharms NDA 021301), Synthroid (Abbvie NDA 021402), and Levo-T (Cediprof Inc NDA 021342) tablets have been determined to be therapeutically equivalent to corresponding strengths of Unithroid (Jerome Stevens NDA 021210) tablets.

Levo-T (Cediprof Inc NDA 021342), Euthyrox (Provell Pharma LLC NDA 021292), Levothyroxine Sodium (Mylan ANDA 076187), and Unithroid (Jerome Stevens NDA 021210) tablets have been determined to be therapeutically equivalent to

corresponding strengths of Synthroid (Abbvie NDA 021402) tablets.

Levo-T (Cediprof Inc NDA 021342), Unithroid (Jerome Stevens NDA 021210), and Levothyroxine Sodium (Mylan ANDA 076187) tablets have been determined to be therapeutically equivalent to corresponding strengths of Levoxyl (King Pharms NDA 021301) tablets.

Levothyroxine Sodium (Mylan ANDA 076187) tablets have been determined to be therapeutically equivalent to corresponding strengths of Thyro-Tabs (Lloyd NDA 021116) tablets.¹⁵

The chart outlines TE codes for all 0.025 mg products in the active section of the Orange Book. Other product strengths may be similar. Therapeutic equivalence has been established between products that have the same AB+number TE code. More than one TE code may apply to some products. One common TE code indicates therapeutic equivalence between products.

| Trade Name | Applicant | Strength | TE Code | Appl No | Product No |
|----------------------|---------------|----------|-------------------|---------|------------|
| UNITHROID | STEVENS J | 0.025MG | AB1 | 021210 | 001 |
| LEVOTHYROXINE SODIUM | MYLAN | 0.025MG | AB1 | 076187 | 001 |
| LEVOXYL | KING PHARMS | 0.025MG | AB1 | 021301 | 001 |
| SYNTHROID | ABBVIE | 0.025MG | AB1 | 021402 | 001 |
| LEVO-T | CEDIPROF INC | 0.025MG | AB1 | 021342 | 001 |
| | | | | | |
| SYNTHROID | ABBVIE | 0.025MG | AB2 | 021402 | 001 |
| LEVOTHYROXINE SODIUM | MYLAN | 0.025MG | AB2 | 076187 | 001 |
| LEVO-T | CEDIPROF INC | 0.025MG | AB2 | 021342 | 001 |
| UNITHROID | STEVENS J | 0.025MG | AB2 | 021210 | 001 |
| EUTHYROX | PROVELL PHARM | 0.025MG | AB2 | 021292 | 001 |
| | | | | | |
| LEVOXYL | KING PHARMS | 0.025MG | AB3 | 021301 | 001 |
| LEVO-T | CEDIPROF INC | 0.025MG | AB3 | 021342 | 001 |
| UNITHROID | STEVENS J | 0.025MG | AB3 | 021210 | 001 |
| LEVOTHYROXINE SODIUM | MYLAN | 0.025MG | AB3 | 076187 | 001 |
| | | | | | |
| THYRO-TABS | LLOYD | 0.025MG | N/A ¹⁶ | 021116 | 001 |
| LEVOTHYROXINE SODIUM | MYLAN | 0.025MG | AB4 | 076187 | 001 |

15 Lloyd's Thyro-Tabs tablets (NDA 021116) (previously known as Levothroid) is currently listed in the Discontinued Drug Product List section of the Orange Book. It is the RLD for the AB4 category. Mylan's levothyroxine product (ANDA 076187) has been selected as the reference standard for ANDA applicants to use to establish bioequivalence to Thyro-Tabs. If an ANDA that uses Mylan's levothyroxine product as its reference standard is approved, the ANDA will receive an AB4 rating. The ANDA applicant also may obtain an AB rating for its product to the other reference listed drugs (i.e., Unithroid, Synthroid, and Levoxyl) by submitting supplements that demonstrate that the generic product is bioequivalent to these other reference listed drugs and satisfies all other therapeutic equivalence criteria with respect to these reference listed drugs. See Letter from Janet Woodcock, M.D., Director, Center for Drug Evaluation and Research, FDA to Teri Nataline, Principal Consultant, Lachman Consultant Services, Inc., Docket No. FDA-2015-P-0403 (May 27, 2016).

16 Id. Thyro-Tabs is in the Discontinued Drug Product List and therefore no longer is assigned a TE code.

Patent Certification(s) and Reference Standard for ANDAs Duplicating a Drug Product Approved in a Petitioned ANDA. To submit an ANDA for a generic drug that is not the same as its reference listed drug because it has one different active ingredient in a fixed-combination drug product, or has a different route of administration, dosage form, or strength than that of the reference listed drug, an applicant first must obtain permission from FDA through what is known as a suitability petition pursuant to Section 505(j)(2)(C) of the FD&C Act. A petitioned ANDA relies on the reference listed drug described in the suitability petition. An ANDA seeking approval of a drug that is the same as a drug product approved in a petitioned ANDA should use as its reference listed drug, the reference listed drug that served as the basis for the approved suitability petition, and use the drug product approved in the petitioned ANDA as its reference standard for conducting an *in vivo* bioequivalence study required for approval. However, the reference listed drug for any such ANDA is generally the listed drug referenced in the approved suitability petition. The ANDA must include appropriate patent certification(s) and an exclusivity statement with respect to the reference listed drug that served as the basis for the approved suitability petition.¹⁷ (This concept also generally applies to an ANDA applicant that utilizes a reference standard that is not a reference listed drug, as such an application must include appropriate patent certification(s) and an exclusivity statement with respect to the reference listed drug.)

Waived exclusivity. If an NDA submitted under Section 505(b) of the FD&C Act qualifies for exclusivity under the FD&C Act, the exclusivity is generally listed in the Patent and Exclusivity Section of the Orange Book. If a drug product has received this exclusivity, FDA will not accept for review and/or will not approve a 505(b)(2) application or an ANDA under Section 505(j) of the FD&C Act, as applicable, in accordance with the relevant exclusivity.¹⁸ If the listed drug is also protected by one or more patents, the approval date for the ANDA or 505(b)(2) application will be determined based on an analysis of the applicant's patent certification(s) or statement(s) for each relevant patent and the effect of relevant exclusivity listed in the Orange Book. However, the holder of the NDA may waive its exclusivity as to any or all ANDAs and 505(b)(2) applications that might otherwise be blocked by such exclusivity. If an NDA sponsor waives its exclusivity, qualified ANDAs or 505(b)(2) applications may be accepted for review and/or approved, as applicable. An NDA for which the holder has waived its exclusivity as to all ANDAs and 505(b)(2) applications will be coded with a "W" in the Patent and Exclusivity Section of the Orange Book. The applicant whose product might otherwise be blocked by this exclusivity should indicate in the exclusivity statement in its application that the holder of the listed drug has waived its exclusivity.

1.9 Therapeutic Equivalence Code Change for a Category of Multisource Drug Products

The Agency will use the following procedures when, in response to a petition or on its own initiative, it is considering a change in the therapeutic equivalence code for approved multisource drug products. Such changes will generally occur when the Agency becomes aware of new scientific information affecting the therapeutic equivalence of an entire category of

¹⁷ If after approval of a suitability petition and before approval of an ANDA submitted pursuant to the approved petition, a drug product is approved in an NDA for the change described in the petition, the suitability petition and the listed drug identified in the petition can no longer be the basis of submission for such ANDA. Under these circumstances, an applicant seeking approval for a drug product with the change approved in the suitability petition must submit a new ANDA that identifies the drug product approved under such NDA as the RLD and comply with applicable regulatory requirements. See 21 CFR 314.93(f)(2).

¹⁸ See Patent and Exclusivity Information Addendum in the Orange Book.

multisource drug products in the Orange Book (e.g., information concerning the active ingredient or the dosage form), rather than information concerning a single drug product within the category. These procedures will be used when a change in therapeutic equivalence code is under consideration for all drug products found in the Prescription Drug Product List under a specific active ingredient and dosage form. The change may be from the code signifying that the drug does not present a bioequivalence problem (e.g., **AA**) to a code signifying an actual or potential bioequivalence problem (e.g., **BP**), or vice versa. This procedure does not apply to a change of a particular product code (e.g., a change from **BP** to **AB** or from **AB** to **BX**).

Before making a change in a therapeutic equivalence code for an entire category of multisource drug products as described above, the Agency will announce in the *Introduction* to the Cumulative Supplement that it is considering the change and will invite comments. Comments, along with scientific data, may be sent to the Director, Office of Bioequivalence, Food and Drug Administration, Office of Generic Drugs, Central Document Room, 5901-B Ammendale Rd., Beltsville, MD 20705-1266.

The comment period will generally be 60 days in length, and the closing date for comments will be listed in the description of the proposed change for each drug entity.

The most useful type of scientific data submitted to support comments is an *in vivo* bioavailability/bioequivalence study conducted on batches of the subject drug products. Comments including scientific data from an *in vivo* bioavailability/bioequivalence study should present a full description of the analytical procedures and equipment used, a validation of the analytical methodology, including the standard curve, a description of the method of calculating results, and a description of the pharmacokinetic and statistical models used in analyzing the data. Anecdotal or testimonial information is the least useful to the Agency, and submission of comments based on such information is discouraged. However, when there is supporting published or unpublished scientific literature, copies should be submitted with comments.

1.10 Change of the Therapeutic Equivalence Evaluation for a Single Product

The procedure described in Section 1.9 does not apply to a change in a single drug product code. For example, a change in a single drug product's code from **BP** to **AB** as a result of the submission of an acceptable bioequivalence study ordinarily will not be the subject of notice and comment in the Cumulative Supplement. Likewise, a change in a single drug product's code from **AB** to **BX** (e.g., as a result of new information raising a significant question as to bioequivalence) does not require notice and comment. The Agency's responsibility to provide the public with the Agency's most current information related to therapeutic equivalence may require a change in a drug product's code prior to any formal notice and opportunity for the applicant to be heard. The publication in the *Federal Register* of a proposal to withdraw approval of a drug product will ordinarily result in a change in a product's code from **AB** to **BX** if this action has not already been taken.

We recognize that certain drug products approved in 505(b)(2) applications may not have therapeutic equivalence codes, and that FDA may undertake therapeutic equivalence evaluations with respect to such drug products. A person seeking to have a therapeutic equivalence rating for a drug product approved in a 505(b)(2) application may petition the Agency through the citizen petition procedure (see 21 CFR 10.25(a) and 21 CFR 10.30).

1.11 Discontinued Section

Those drug products in the discontinued section of the Orange Book (Discontinued Drug Product List) for which a determination has been made that the products were not withdrawn for safety or effectiveness reasons have been annotated with a footnote following the product strength: "***Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons***". The determinations listed in the Orange Book are only reflective of determinations made since 1995 and published in the Federal Register. The identification of these drug products in the Discontinued Drug Product List should avoid the submission of multiple citizen petitions requesting a determination for the same drug product.

Generally, approved products are added to the Discontinued Drug Product List when the applicant notifies the Orange Book staff of the products' not-marketed status. Products may also be added to the Discontinued Drug Product List if annual reports or other submissions to the Agency indicate the product is not being marketed or as a result of other Agency administrative actions.¹⁹ Changes to the Orange Book are not affected by the drug registration and listing requirements of Section 510 of the FD&C Act.

1.12 Changes to the Orange Book

Every effort is made to ensure the Annual Edition is current and accurate. Applicants are requested to inform the FDA Orange Book Staff of any changes or corrections, including any change in ownership or a product's marketing status that would result in the product being moved to the Discontinued Drug Product List. FDA notes that under Section 506I(a) of the FD&C Act, applicants must notify the Agency in writing 180 days prior to withdrawing a drug product from sale, or if 180 days is not practicable, not later than the date of withdrawal from sale. Furthermore, Section 506I(b) of the FD&C Act requires that applicants notify the Agency in writing within 180 days of approval of a drug product if such drug product will not be available for sale within 180 days of approval. A request to include a newly approved product in the Discontinued Drug Product List, rather than parts 1, 2 or 3 of the Orange Book (as discussed in Section 1.1), must be submitted to the Orange Book staff by the end of the month in which the product is approved to ensure that the product is not included in the "active" portions of the next published Orange Book update.

In addition, the FDA Orange Book Staff generally will act on requests to change a proprietary name for a listed drug only after approval of a supplement for the relevant change in proprietary name. To the extent that conventions for describing product identification information (i.e., active ingredients, dosage forms, routes of administration, product names, applicants, strengths) evolve over time, the Agency generally does not intend to revise such information for drug products already included in the Orange Book, but rather intends to apply the change prospectively to drug products added to the Orange Book.

You can contact the Orange Book Staff by email at orangebook@fda.hhs.gov.

¹⁹ See, e.g., Section 506I(d) of the FD&C Act.

1.13 Availability of the Edition

Commencing with the 25th edition, the Annual Edition and current monthly Cumulative Supplement are available in a Portable Document Format (PDF) at the [Orange Book](#) home page by clicking on Publications. The PDF annual format duplicates previous paper versions except for the Orphan Products Designations and Approvals List. An annual subscription of the PDF format may be obtained from the U.S. Government Publishing Office, <https://www.gpo.gov/>.

2.0 HOW TO USE THE DRUG PRODUCT LISTS

2.1 Key Sections for Using the Drug Product Lists

This publication contains illustrations, along with Drug Product Lists, indices, and lists of abbreviations and terms which facilitate their use.

Illustrations. The annotated *Drug Product Illustration*, see Section 2.2, and the *Therapeutic Equivalence Evaluations Illustration*, see Section 2.3, are offered to provide further clarification. These depict the format found in the Prescription Drug Product List (the only list in which therapeutic equivalence evaluation codes are displayed).

Drug Product Lists. The Prescription and OTC Drug Product Lists, arranged alphabetically by active ingredient(s), contain product identification information (active ingredients, dosage forms, routes of administration, product names, applicants, strengths) for single and multiple ingredient drug products. Also shown are the application number and drug product number (FDA internal computer data use only) and approval dates for those drug products approved on or after January 1, 1982. The application number preceded by "N" is a New Drug Application (NDA or commonly the innovator). The application number preceded by an "A" is an Abbreviated New Drug Application (ANDA or commonly the generic).

The Discontinued Drug Product List, arranged alphabetically by active ingredient(s), contains product identification information (dosage form, product name, strength, and application number).

If a prescription drug product is available from more than one source (multisource), a therapeutic equivalence code will appear in front of the applicant's name. If a product is therapeutically equivalent to one or more products or to an appropriate reference, it will be designated with a code beginning with "A" and the entry will be underlined and printed in bold font for emphasis.

Active ingredient headings for multiple ingredient (combination) drug products are arranged alphabetically. For purposes of this publication, this alphabetical sort takes precedence over United States Pharmacopeia official monograph order (i.e., Reserpine, Hydralazine Hydrochloride, Hydrochlorothiazide). For example, product information labeled as Reserpine, Hydrochlorothiazide and Hydralazine Hydrochloride appears under the active ingredient heading *Hydralazine Hydrochloride; Hydrochlorothiazide; Reserpine*. A cross-reference to the product information (for prescription and OTC products) appears for each additional active ingredient in the product. For combination drug products, the ingredient strengths are separated by semicolons and appear in the same relative sequence as the ingredients in the heading. Available strengths of the dosage form from an applicant appear on separate lines.

To use the Drug Product Lists, determine by alphabetical order the ingredient under which the product information is listed, using the Product Name Index, if necessary. Then, find the ingredient in the applicable Drug Product List. Proceed to the dosage form and route of administration and compare products within that ingredient heading only. Therapeutic equivalence or inequivalence for prescription products is determined on the basis of the therapeutic equivalence codes provided within that specific dosage form and route heading. The OTC Drug Product List, Discontinued Drug Product List, and

Drug Products with Approval under Section 505 of the Act Administered by the Center for Biologics Evaluation and Research List have their data arranged similarly.

The Discontinued Drug Product List contains approved products that have never been marketed, have been discontinued from marketing and we have not determined that they were withdrawn for safety or effectiveness reasons, are for military use, or have had their approvals withdrawn for other than safety or efficacy reasons subsequent to being discontinued from marketing. All products having a "@" in the December Cumulative Supplement of the previous Edition List have been added to the Discontinued Drug Product List appearing in this Edition. In addition, approved drug products that are not in the commercial distribution channel e.g., approved drug products in applications for export only are also listed in the Discontinued Drug Product List.

Product Name Index (*Prescription and OTC Drug Product Lists*). This is an index of drug products by trade name or established name of the active ingredient, if no trade name exists. The second term of each entry indicates the active ingredient name under which product information can be found in the appropriate Drug Product List. For those drug products with multiple active ingredients, only the first active ingredient (in alphabetical order) will appear. OTC products are so designated.

Product Name Index Listed by Applicant (*Prescription and OTC Drug Product Lists*). This is an index that cross-references applicants to drug products. The bolded and underlined entry represents the applicant name abbreviation used in this publication. Each complete applicant name that is represented by the abbreviated name is marked with an asterisk (*). Listed under each complete applicant name is the first alphabetically arranged ingredient under which product information can be found in the appropriate Drug Product List. OTC products are so designated. To use the Drug Product Lists, determine by alphabetical order the ingredient under which the product information is listed, using the Product Name Index, if appropriate.

Uniform Terms. To improve readability, uniform terms are used to designate dosage forms, routes of administration, and abbreviations used to express strengths. These terms are listed in Appendix C. In some cases, the terms used may differ from those used in product labels and other labeling.

2.2 DRUG PRODUCT ILLUSTRATION

SINGLE INGREDIENT

| | | |
|-----------------------------------------------------------|---|--------------------------------------------------------------------------------|
| ACTIVE INGREDIENT | → | <u>MEPERIDINE HYDROCHLORIDE</u> |
| DOSAGE FORM; ROUTE OF ADMINISTRATION | → | INJECTABLE; INJECTION |
| TRADE OR GENERIC NAMES | → | <u>HEXANON</u> |
| REFERENCE LISTED DRUG* (+) | → | <u>AP</u> +! PAGE PHARMA <u>25MG/ML</u> <u>N013111</u> <u>001</u> AUG 22, 1983 |
| REFERENCE STANDARD * (!) | → | <u>AP</u> +! <u>50MG/ML</u> <u>N013111</u> <u>002</u> AUG 22, 1983 |
| | | <u>AP</u> +! <u>75MG/ML</u> <u>N013111</u> <u>003</u> AUG 22, 1983 |
| | | <u>AP</u> +! <u>100MG/ML</u> <u>N013111</u> <u>004</u> JAN 04, 1989 |
| | → | <u>MEPERIDINE HCL</u> |
| THERAPEUTIC EQUIVALENCE (TE) | → | <u>AP</u> GREENBERG PHARM <u>25MG/ML</u> <u>A064890</u> 001 FEB 29, 1987 |
| CODE FOR MULTISOURCE PRODUCT | → | <u>AP</u> <u>50MG/ML</u> <u>A064890</u> 002 FEB 29, 1987 |
| | | <u>AP</u> <u>75MG/ML</u> <u>A064890</u> 003 FEB 29, 1987 |
| | | <u>AP</u> <u>100MG/ML</u> <u>A064890</u> 004 MAR 08, 1992 |
| SINGLE SOURCE PRODUCT (NO TE CODE) | | ! TIMOKIM LLC 10MG/ML A099225 001 DEC 12, 1995 |
| | | <u>AP</u> JOHNSON MED <u>25MG/ML</u> <u>A099226</u> <u>001</u> NOV 27, 1993 |
| | | ! KENDRA PHARM 150MG/ML A079444 001 OCT 31, 1999 |
| APPLICANT | → | ↑ |
| AVAILABLE STRENGTH(S) OF A PRODUCT | → | ↑ |
| APPLICATION NUMBER AND PRODUCT NUMBER | → | ↑ |
| PRODUCT NUMBER IS FOR FDA INTERNAL COMPUTER DATA USE ONLY | → | ↑ |
| APPROVAL DATE | → | ↑ |

*NOTE: REFERENCE LISTED DRUG AND REFERENCE STANDARD ARE DISCUSSED IN THE PREFACE SECTION 1.4

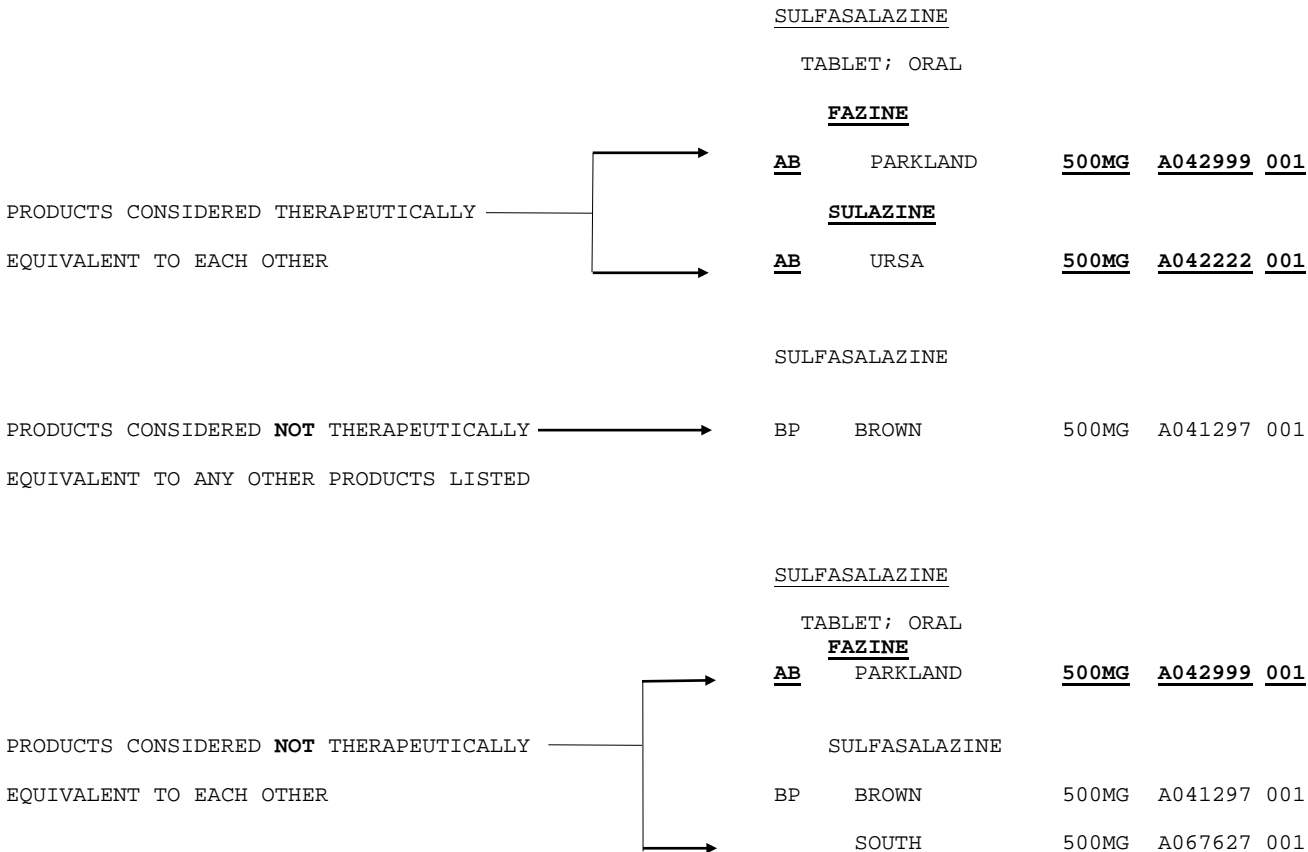
MULTIPLE INGREDIENTS WITH PRODUCT INFORMATION

| | | |
|--------------------------|---|------------------------------------------------------------------|
| ALPHABETICALLY SORTED BY | → | <u>HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE</u> |
| PRODUCT INFORMATION | → | TABLET; ORAL |
| | | HYDROCHLOROTHIAZIDE, RESERPINE AND HYDRALAZINE HCL |
| | | REINWALD LABS 25MG; 15MG; 0.1MG A069808 001 JAN 18, 1982 |

THIS EXAMPLE IS FOR PURPOSE OF ILLUSTRATION ONLY. IT DOES NOT REPRESENT ACTUAL PRODUCTS FROM THE PRESCRIPTION DRUG PRODUCT LIST.

2.3 THERAPEUTIC EQUIVALENCE EVALUATIONS ILLUSTRATION

DRUG PRODUCTS CODED **AB** (OR ANY CODE BEGINNING WITH AN "A") UNDER AN INGREDIENT AND DOSAGE FORM HEADING ARE CONSIDERED THERAPEUTICALLY EQUIVALENT ONLY TO OTHER PRODUCTS CODED **AB** (OR ANY CODE BEGINNING WITH AN "A") AND **NOT** TO THOSE CODED **BP** (OR ANY CODE BEGINNING WITH "B") AND ANY PRODUCTS NOT LISTED. DRUG PRODUCTS CODED **BP** (OR ANY CODE BEGINNING WITH A "B") ARE **NOT** CONSIDERED THERAPEUTICALLY EQUIVALENT TO ANY OTHER PRODUCT. FOR A COMPLETE EXPLANATION OF THE **TE** CODES REFER TO SECTION 1.7 OF THE *INTRODUCTION*.



NOTE: BOLD FONT AND UNDERLINING DENOTES MULTISOURCE PRODUCTS WHICH ARE CONSIDERED THERAPEUTICALLY EQUIVALENT.

THIS EXAMPLE IS FOR PURPOSES OF ILLUSTRATION ONLY. IT DOES NOT REPRESENT ACTUAL PRODUCTS FROM THE PRESCRIPTION DRUG PRODUCT LIST.

PRESCRIPTION DRUG PRODUCT LIST

ABACAVIR SULFATE

SOLUTION;ORAL

ABACAVIR SULFATE

| | | | | |
|-----------|----------------------|------------------------|--------------------|--------------|
| AA | AUROBINDO PHARMA LTD | EQ 20MG BASE/ML | A077950 001 | Mar 14, 2018 |
| AA | HETERO LABS LTD III | EQ 20MG BASE/ML | A201107 001 | Sep 26, 2016 |

ZIAGEN

| | | | | |
|-----------|-----------------|------------------------|--------------------|--------------|
| AA | +! VIIV HLHCARE | EQ 20MG BASE/ML | N020978 001 | Dec 17, 1998 |
|-----------|-----------------|------------------------|--------------------|--------------|

TABLET;ORAL

ABACAVIR SULFATE

| | | | | |
|-----------|----------------------|----------------------|--------------------|--------------|
| AB | APOTEX INC | EQ 300MG BASE | A201570 001 | Dec 17, 2012 |
| AB | AUROBINDO PHARMA LTD | EQ 300MG BASE | A077844 001 | Dec 17, 2012 |
| AB | CIPLA | EQ 300MG BASE | A078119 001 | Nov 21, 2017 |
| AB | HETERO LABS LTD III | EQ 300MG BASE | A091560 001 | Sep 13, 2013 |
| AB | MYLAN PHARMS INC | EQ 300MG BASE | A091294 001 | Jun 18, 2012 |
| AB | STRIDES PHARMA | EQ 300MG BASE | A091050 001 | Oct 28, 2016 |

ZIAGEN

| | | | | |
|-----------|-----------------|----------------------|--------------------|--------------|
| AB | +! VIIV HLHCARE | EQ 300MG BASE | N020977 001 | Dec 17, 1998 |
|-----------|-----------------|----------------------|--------------------|--------------|

ABACAVIR SULFATE; DOLUTEGRAVIR SODIUM; LAMIVUDINE

TABLET;ORAL

TRIUMEQ

| | | | | |
|----|--------------|-----------------------------------------|--------------------|--------------|
| +! | VIIV HLHCARE | EQ 600MG BASE;EQ 50MG BASE;300MG | N205551 001 | Aug 22, 2014 |
|----|--------------|-----------------------------------------|--------------------|--------------|

ABACAVIR SULFATE; LAMIVUDINE

TABLET;ORAL

ABACAVIR SULFATE AND LAMIVUDINE

| | | | | |
|-----------|----------------------|----------------------------|--------------------|--------------|
| AB | AUROBINDO PHARMA LTD | EQ 600MG BASE;300MG | A090159 001 | Nov 15, 2018 |
| AB | | EQ 600MG BASE;300MG | A206151 001 | Mar 28, 2017 |
| AB | CIPLA | EQ 600MG BASE;300MG | A091144 001 | Mar 28, 2017 |
| AB | LUPIN LTD | EQ 600MG BASE;300MG | A204990 001 | Mar 28, 2017 |
| AB | TEVA PHARMS USA | EQ 600MG BASE;300MG | A079246 001 | Sep 29, 2016 |
| AB | ZYDUS PHARMS USA INC | EQ 600MG BASE;300MG | A208990 001 | Nov 15, 2018 |

EPZICOM

| | | | | |
|-----------|-----------------|----------------------------|--------------------|--------------|
| AB | +! VIIV HLHCARE | EQ 600MG BASE;300MG | N021652 001 | Aug 02, 2004 |
|-----------|-----------------|----------------------------|--------------------|--------------|

ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE

TABLET;ORAL

ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE

| | | | | |
|-----------|-----------------|----------------------------------|--------------------|--------------|
| AB | LUPIN LTD | EQ 300MG BASE;150MG;300MG | A202912 001 | Dec 05, 2013 |
| AB | +! VIIV HLHCARE | EQ 300MG BASE;150MG;300MG | N021205 001 | Nov 14, 2000 |

TRIZIVIRABALOPARATIDE

SOLUTION;SUBCUTANEOUS

TYMLOS

| | | | | |
|----|-------------------|-------------------------------|--------------------|--------------|
| +! | RADIUS HEALTH INC | 3.12MG/1.56ML (2MG/ML) | N208743 001 | Apr 28, 2017 |
|----|-------------------|-------------------------------|--------------------|--------------|

ABEMACICLIB

TABLET;ORAL

VERZENIO

| | | | | |
|----|------------------|--------------|--------------------|--------------|
| + | ELI LILLY AND CO | 50MG | N208716 001 | Sep 28, 2017 |
| + | | 100MG | N208716 002 | Sep 28, 2017 |
| + | | 150MG | N208716 003 | Sep 28, 2017 |
| +! | | 200MG | N208716 004 | Sep 28, 2017 |

ABIRATERONE ACETATE

TABLET;ORAL

ABIRATERONE ACETATE

| | | | | |
|-----------|------------------|--------------|--------------------|--------------|
| AB | AMNEAL PHARMS | 250MG | A208327 001 | Jan 07, 2019 |
| AB | APOTEX INC | 250MG | A208453 001 | Oct 31, 2018 |
| AB | HIKMA PHARMS | 250MG | A208339 001 | Oct 31, 2018 |
| AB | MYLAN PHARMS INC | 250MG | A208446 001 | Oct 31, 2018 |
| AB | TEVA PHARMS USA | 250MG | A208432 001 | Oct 31, 2018 |

ZYTIGA

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| AB | + JANSSEN BIOTECH | 250MG | N202379 001 | Apr 28, 2011 |
| | YONSA | | | |
| | +! SUN PHARMA GLOBAL | 125MG | N210308 001 | May 22, 2018 |
| | ZYTIGA | | | |
| | +! JANSSEN BIOTECH | 500MG | N202379 002 | Apr 14, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

ACALABRUTINIB

CAPSULE;ORAL

CALQUENCE

+! ASTRAZENECA

100MG

N210259 001 Oct 31, 2017

ACAMPROSATE CALCIUM

TABLET, DELAYED RELEASE;ORAL

ACAMPROSATE CALCIUMAB BARR LABS DIV TEVA 333MGA200143 001 Nov 18, 2013AB ! GLENMARK GENERICS 333MGA202229 001 Jul 16, 2013AB MYLAN PHARMS INC 333MGA200142 001 Mar 11, 2014AB ZYDUS PHARMS USA 333MGA205995 001 May 26, 2017

INC

ACARBOSE

TABLET;ORAL

ACARBOSEAB EMCURE PHARMS LTD 25MGA202271 001 Feb 07, 2012AB 50MGA202271 002 Feb 07, 2012AB 100MGA202271 003 Feb 07, 2012AB IMPAX LABS 25MGA078441 001 May 14, 2009AB 50MGA078441 002 May 14, 2009AB 100MGA078441 003 May 14, 2009AB MYLAN 25MGA091053 001 Jan 06, 2011AB 50MGA091053 002 Jan 06, 2011AB 100MGA091053 003 Jan 06, 2011AB STRIDES PHARMA 25MGA090912 001 Jul 27, 2011AB 50MGA090912 002 Jul 27, 2011AB 100MGA090912 003 Jul 27, 2011AB VIRTUS PHARM 25MGA091343 001 Oct 17, 2013AB 50MGA091343 002 Oct 17, 2013AB 100MGA091343 003 Oct 17, 2013AB WATSON LABS 25MGA077532 001 May 07, 2008AB 50MGA077532 002 May 07, 2008AB 100MGA077532 003 May 07, 2008AB WEST-WARD PHARMS 25MGA078470 001 May 07, 2008

INT

AB 50MGA078470 002 May 07, 2008AB 100MGA078470 003 May 07, 2008PRECOSEAB +! BAYER HLTHCARE 25MGN020482 004 May 29, 1997AB + 50MGN020482 001 Sep 06, 1995AB + 100MGN020482 002 Sep 06, 1995ACEBUTOLOL HYDROCHLORIDE

CAPSULE;ORAL

ACEBUTOLOL HYDROCHLORIDEAB ! AMNEAL PHARM EQ 200MG BASEA075047 001 Dec 30, 1999AB ! EQ 400MG BASEA075047 002 Dec 30, 1999AB MYLAN EQ 200MG BASEA074288 001 Apr 24, 1995AB EQ 400MG BASEA074288 002 Apr 24, 1995ACETAMINOPHEN

SOLUTION;INTRAVENOUS

ACETAMINOPHENAP CUSTOPHARM INC 1GM/100ML (10MG/ML)A202605 001 Jun 13, 2016AP SANDOZ INC 1GM/100ML (10MG/ML)A204052 001 Mar 22, 2016OFIRMEVAP +! MALLINCKRODT HOSP 1GM/100ML (10MG/ML)N022450 001 Nov 02, 2010

ACETAMINOPHEN

FRESENIUS KABI USA

1GM/100ML (10MG/ML)

N204767 001 Oct 28, 2015

ACETAMINOPHEN; BENZHYDROCODONE HYDROCHLORIDE

TABLET;ORAL

APADAZ

+ KEMPHARM

325MG;EQ 6.12MG BASE

N208653 001 Feb 23, 2018

ACETAMINOPHEN; BUTALBITAL

CAPSULE;ORAL

BUTALBITAL AND ACETAMINOPHEN

! MAYNE PHARMA INC

300MG;50MG

A207313 001 Dec 27, 2017

TABLET;ORAL

BUTALBITAL AND ACETAMINOPHENAA CNTY LINE PHARMS 300MG;50MGA207635 001 Jun 05, 2017AA 325MG;50MGA205120 001 Oct 30, 2015AA LARKEN LABS INC 325MG;50MGA203484 002 Dec 04, 2015

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; BUTALBITAL

TABLET; ORAL

BUTALBITAL AND ACETAMINOPHEN

| | | | | |
|-----------|-----------------|--------------------|--------------------|--------------|
| AA | MIKART | 300MG; 50MG | A207386 001 | Nov 15, 2016 |
| AA | ! NEXGEN PHARMA | 300MG; 50MG | A090956 001 | Aug 23, 2011 |

BUTAPAP

| | | | | |
|-----------|-----------------|--------------------|--------------------|--------------|
| AA | ! MIKART | 325MG; 50MG | A089987 001 | Oct 26, 1992 |
| | ALLZITAL | | | |
| | LARKEN LABS INC | 325MG; 25MG | A203484 001 | Dec 04, 2015 |

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

| | | | | |
|-----------|---------------------|--------------------------|--------------------|--------------|
| AA | AUROLIFE PHARMA LLC | 325MG; 50MG; 40MG | A204733 001 | Sep 26, 2018 |
| AA | ! MAYNE PHARMA INC | 325MG; 50MG; 40MG | A089007 001 | Mar 17, 1986 |
| AA | ! NEXGEN PHARMA | 300MG; 50MG; 40MG | A040885 001 | Nov 16, 2009 |
| AA | NUVO PHARMS INC | 300MG; 50MG; 40MG | A207118 001 | Oct 28, 2016 |
| AA | WRASER PHARMS LLC | 300MG; 50MG; 40MG | A206615 001 | Aug 04, 2017 |

SOLUTION; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

| | | | | |
|---|--------|----------------------------------|-------------|--------------|
| ! | MIKART | 325MG/15ML; 50MG/15ML; 40MG/15ML | A040387 001 | Jan 31, 2003 |
|---|--------|----------------------------------|-------------|--------------|

TABLET; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

| | | | | |
|-----------|---------------------|--------------------------|--------------------|--------------|
| AA | ABHAI LLC | 325MG; 50MG; 40MG | A211106 001 | Sep 26, 2018 |
| AA | ACTAVIS LABS UT INC | 325MG; 50MG; 40MG | A088616 001 | Nov 09, 1984 |
| AA | CNTY LINE PHARMS | 325MG; 50MG; 40MG | A204984 001 | Jan 10, 2017 |
| AA | HIKMA PHARMS | 325MG; 50MG; 40MG | A089718 001 | Jun 12, 1995 |
| AA | LANNETT CO INC | 325MG; 50MG; 40MG | A200243 001 | Sep 13, 2012 |
| AA | MIKART | 325MG; 50MG; 40MG | A089175 001 | Jan 21, 1987 |
| AA | NEXGEN PHARMA INC | 325MG; 50MG; 40MG | A209587 001 | Oct 31, 2018 |
| AA | SPECGX LLC | 325MG; 50MG; 40MG | A087804 001 | Jan 24, 1985 |
| AA | ! VINTAGE PHARMS | 325MG; 50MG; 40MG | A040511 001 | Aug 27, 2003 |

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE

| | | | | |
|-----------|-------------------|--------------------------------|--------------------|--------------|
| AB | NEXGEN PHARMA INC | 325MG; 50MG; 40MG; 30MG | A076560 001 | Jun 10, 2004 |
| AB | VINTAGE PHARMS | 325MG; 50MG; 40MG; 30MG | A075929 001 | Apr 22, 2002 |

FIORICET W/ CODEINE

| | | | | |
|-----------|-----------------------------------------------------------|--------------------------------|--------------------|--------------|
| AB | +! ACTAVIS LABS UT INC | 325MG; 50MG; 40MG; 30MG | N020232 001 | Jul 30, 1992 |
| | BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE | | | |
| | NEXGEN PHARMA INC | 300MG; 50MG; 40MG; 30MG | A076560 002 | Jul 19, 2012 |

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

TREZIX

| | | | | |
|--|-------------------|---------------------|-------------|--------------|
| | WRASER PHARMS LLC | 320.5MG; 30MG; 16MG | A204785 001 | Nov 26, 2014 |
|--|-------------------|---------------------|-------------|--------------|

TABLET; ORAL

ACETAMINOPHEN, CAFFEINE AND DIHYDROCODEINE BITARTRATE

| | | | | |
|--|-----------------|-------------------|-------------|--------------|
| | LARKEN LABS INC | 325MG; 30MG; 16MG | A204209 001 | Sep 30, 2016 |
|--|-----------------|-------------------|-------------|--------------|

ACETAMINOPHEN; CODEINE PHOSPHATE

SOLUTION; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

| | | | | |
|-----------|------------------|----------------------------|--------------------|--------------|
| AA | HI TECH PHARMA | 120MG/5ML; 12MG/5ML | A040119 001 | Apr 26, 1996 |
| AA | LANNETT CO INC | 120MG/5ML; 12MG/5ML | A091238 001 | Nov 10, 2011 |
| AA | MIKART | 120MG/5ML; 12MG/5ML | A089450 001 | Oct 27, 1992 |
| AA | ! PHARM ASSOC | 120MG/5ML; 12MG/5ML | A087508 001 | |
| AA | WOCKHARDT BIO AG | 120MG/5ML; 12MG/5ML | A087006 001 | |

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

| | | | | |
|-----------|---------------------|--------------------|--------------------|--------------|
| AA | AMNEAL PHARMS NY | 300MG; 30MG | A040779 001 | May 29, 2008 |
| AA | AUROLIFE PHARMA LLC | 300MG; 15MG | A202800 001 | Apr 15, 2013 |
| AA | | 300MG; 30MG | A202800 002 | Apr 15, 2013 |
| AA | | 300MG; 60MG | A202800 003 | Apr 15, 2013 |
| AA | ! SPECGX LLC | 300MG; 15MG | A040419 001 | May 31, 2001 |
| AA | | 300MG; 30MG | A040419 002 | May 31, 2001 |
| AA | | 300MG; 60MG | A040419 003 | May 31, 2001 |
| AA | SUN PHARM INDS LTD | 300MG; 30MG | A085868 001 | |
| AA | | 300MG; 60MG | A087083 001 | |
| AA | TEVA | 300MG; 15MG | A088627 001 | Mar 06, 1985 |
| AA | | 300MG; 30MG | A088628 001 | Mar 06, 1985 |
| AA | ! | 300MG; 60MG | A088629 001 | Mar 06, 1985 |
| AA | VINTAGE | 300MG; 15MG | A089990 001 | Sep 30, 1988 |

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

| | | | | |
|-----------|---------------------------------|-------------------|--------------------|--------------|
| <u>AA</u> | | <u>300MG;30MG</u> | <u>A089805 001</u> | Sep 30, 1988 |
| <u>AA</u> | VINTAGE PHARMS | <u>300MG;60MG</u> | <u>A089828 001</u> | Sep 30, 1988 |
| | <u>TYLENOL W/ CODEINE NO. 3</u> | | | |
| <u>AA</u> | ! JANSSEN PHARMS | <u>300MG;30MG</u> | <u>A085055 003</u> | |
| | <u>TYLENOL W/ CODEINE NO. 4</u> | | | |
| <u>AA</u> | JANSSEN PHARMS | <u>300MG;60MG</u> | <u>A085055 004</u> | |

ACETAMINOPHEN; HYDROCODONE BITARTRATE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

| | | | | |
|-----------|--------------------|------------------------------|--------------------|--------------|
| <u>AA</u> | GENUS LIFESCIENCES | <u>325MG/15ML;7.5MG/15ML</u> | <u>A040894 001</u> | Jul 19, 2011 |
| <u>AA</u> | ! MIKART | <u>325MG/15ML;7.5MG/15ML</u> | <u>A040482 001</u> | Sep 25, 2003 |
| <u>AA</u> | PHARM ASSOC | <u>325MG/15ML;7.5MG/15ML</u> | <u>A040838 001</u> | May 10, 2013 |
| <u>AA</u> | VISTAPHARM | <u>325MG/15ML;7.5MG/15ML</u> | <u>A200343 001</u> | Jan 25, 2012 |
| | ! MIKART | 300MG/15ML;10MG/15ML | A040881 001 | Feb 25, 2010 |
| | ! PHARM ASSOC | 325MG/15ML;10MG/15ML | A040834 001 | Apr 18, 2008 |

TABLET; ORAL

ANEXSIA 5/325

| | | | | |
|-----------|------------|------------------|--------------------|--------------|
| <u>AA</u> | SPECGX LLC | <u>325MG;5MG</u> | <u>A040409 001</u> | Oct 20, 2000 |
|-----------|------------|------------------|--------------------|--------------|

ANEXSIA 7.5/325

| | | | | |
|-----------|------------|--------------------|--------------------|--------------|
| <u>AA</u> | SPECGX LLC | <u>325MG;7.5MG</u> | <u>A040405 001</u> | Sep 08, 2000 |
|-----------|------------|--------------------|--------------------|--------------|

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

| | | | | |
|-----------|---------------------|--------------------|--------------------|--------------|
| <u>AA</u> | ABHAI LLC | <u>300MG;5MG</u> | <u>A209036 001</u> | Jun 21, 2017 |
| <u>AA</u> | | <u>300MG;7.5MG</u> | <u>A209036 002</u> | Jun 21, 2017 |
| <u>AA</u> | | <u>300MG;10MG</u> | <u>A209036 003</u> | Jun 21, 2017 |
| <u>AA</u> | | <u>325MG;5MG</u> | <u>A209037 001</u> | Jun 21, 2017 |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A209037 002</u> | Jun 21, 2017 |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A209037 003</u> | Jun 21, 2017 |
| <u>AA</u> | ACTAVIS LABS FL INC | <u>300MG;5MG</u> | <u>A206470 001</u> | Jun 02, 2016 |
| <u>AA</u> | | <u>300MG;7.5MG</u> | <u>A206470 002</u> | Jun 02, 2016 |
| <u>AA</u> | | <u>300MG;10MG</u> | <u>A206470 003</u> | Jun 02, 2016 |
| <u>AA</u> | ALVOGEN PINE BROOK | <u>300MG;5MG</u> | <u>A208540 001</u> | Nov 08, 2018 |
| <u>AA</u> | | <u>300MG;7.5MG</u> | <u>A208540 002</u> | Nov 08, 2018 |
| <u>AA</u> | | <u>300MG;10MG</u> | <u>A208540 003</u> | Nov 08, 2018 |
| <u>AA</u> | | <u>325MG;2.5MG</u> | <u>A209958 001</u> | Oct 24, 2018 |
| <u>AA</u> | | <u>325MG;5MG</u> | <u>A209958 002</u> | Oct 24, 2018 |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A209958 003</u> | Oct 24, 2018 |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A209958 004</u> | Oct 24, 2018 |
| <u>AA</u> | AMNEAL PHARMS | <u>300MG;10MG</u> | <u>A207137 001</u> | Nov 29, 2016 |
| <u>AA</u> | AMNEAL PHARMS NY | <u>300MG;5MG</u> | <u>A206869 001</u> | Jun 23, 2017 |
| <u>AA</u> | | <u>325MG;5MG</u> | <u>A040736 001</u> | Aug 25, 2006 |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A040746 002</u> | May 10, 2016 |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A040746 001</u> | Aug 25, 2006 |
| <u>AA</u> | ASCENT PHARMS INC | <u>325MG;2.5MG</u> | <u>A211487 001</u> | Nov 07, 2018 |
| <u>AA</u> | | <u>325MG;5MG</u> | <u>A211487 002</u> | Nov 07, 2018 |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A211487 003</u> | Nov 07, 2018 |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A211487 004</u> | Nov 07, 2018 |
| <u>AA</u> | AUROLIFE PHARMA LLC | <u>300MG;5MG</u> | <u>A207709 001</u> | Sep 13, 2018 |
| <u>AA</u> | | <u>300MG;7.5MG</u> | <u>A207709 002</u> | Sep 13, 2018 |
| <u>AA</u> | | <u>300MG;10MG</u> | <u>A207709 003</u> | Sep 13, 2018 |
| <u>AA</u> | | <u>325MG;5MG</u> | <u>A201013 001</u> | Apr 11, 2012 |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A201013 002</u> | Apr 11, 2012 |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A201013 003</u> | Apr 11, 2012 |
| <u>AA</u> | ELITE LABS INC | <u>325MG;2.5MG</u> | <u>A209924 001</u> | Nov 16, 2018 |
| <u>AA</u> | | <u>325MG;5MG</u> | <u>A209924 002</u> | Nov 16, 2018 |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A209924 003</u> | Nov 16, 2018 |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A209924 004</u> | Nov 16, 2018 |
| <u>AA</u> | EPIC PHARMA LLC | <u>325MG;5MG</u> | <u>A203863 001</u> | Mar 30, 2018 |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A203863 002</u> | Mar 30, 2018 |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A203863 003</u> | Mar 30, 2018 |
| <u>AA</u> | LANNETT CO INC | <u>300MG;5MG</u> | <u>A207171 001</u> | Jun 20, 2017 |
| <u>AA</u> | | <u>300MG;7.5MG</u> | <u>A207171 002</u> | Jun 20, 2017 |
| <u>AA</u> | | <u>300MG;10MG</u> | <u>A207171 003</u> | Jun 20, 2017 |
| <u>AA</u> | | <u>325MG;5MG</u> | <u>A207172 001</u> | Jun 22, 2017 |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A207172 002</u> | Jun 22, 2017 |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A207172 003</u> | Jun 22, 2017 |
| <u>AA</u> | ! MIKART | <u>300MG;5MG</u> | <u>A040658 001</u> | Jan 19, 2006 |
| <u>AA</u> | ! | <u>300MG;7.5MG</u> | <u>A040658 002</u> | Mar 24, 2006 |
| <u>AA</u> | ! | <u>300MG;10MG</u> | <u>A040658 003</u> | Jun 23, 2004 |
| <u>AA</u> | ! | <u>325MG;2.5MG</u> | <u>A040846 001</u> | Jun 09, 2010 |

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

| | | | | |
|--------------|--------------------|---------------------|--------------------|--------------|
| <u>AA</u> | | <u>325MG; 7.5MG</u> | <u>A040432 001</u> | Jan 22, 2003 |
| <u>AA</u> | NOVEL LABS INC | <u>300MG; 5MG</u> | <u>A206142 001</u> | Nov 14, 2016 |
| <u>AA</u> | | <u>300MG; 7.5MG</u> | <u>A206142 002</u> | Nov 14, 2016 |
| <u>AA</u> | | <u>300MG; 10MG</u> | <u>A206142 003</u> | Nov 14, 2016 |
| <u>AA</u> | | <u>325MG; 5MG</u> | <u>A206245 001</u> | Dec 01, 2016 |
| <u>AA</u> | | <u>325MG; 7.5MG</u> | <u>A206245 002</u> | Dec 01, 2016 |
| <u>AA</u> | | <u>325MG; 10MG</u> | <u>A206245 003</u> | Dec 01, 2016 |
| <u>AA</u> | PAR PHARM | <u>300MG; 5MG</u> | <u>A205001 001</u> | Jul 05, 2016 |
| <u>AA</u> | | <u>300MG; 7.5MG</u> | <u>A205001 002</u> | Jul 05, 2016 |
| <u>AA</u> | | <u>300MG; 10MG</u> | <u>A205001 003</u> | Jul 05, 2016 |
| <u>AA</u> | | <u>325MG; 5MG</u> | <u>A202935 002</u> | Jun 15, 2016 |
| <u>AA</u> | | <u>325MG; 7.5MG</u> | <u>A202935 003</u> | Jun 15, 2016 |
| <u>AA</u> | | <u>325MG; 10MG</u> | <u>A202935 004</u> | Jun 15, 2016 |
| <u>AA</u> | RHODES PHARMS | <u>300MG; 5MG</u> | <u>A207808 001</u> | Mar 30, 2018 |
| <u>AA</u> | | <u>300MG; 7.5MG</u> | <u>A207808 002</u> | Mar 30, 2018 |
| <u>AA</u> | | <u>300MG; 10MG</u> | <u>A207808 003</u> | Mar 30, 2018 |
| <u>AA</u> | | <u>325MG; 5MG</u> | <u>A202991 001</u> | Apr 12, 2016 |
| <u>AA</u> | | <u>325MG; 7.5MG</u> | <u>A202991 002</u> | Apr 12, 2016 |
| <u>AA</u> | | <u>325MG; 10MG</u> | <u>A202991 003</u> | Apr 12, 2016 |
| <u>AA</u> | SPECGX LLC | <u>300MG; 5MG</u> | <u>A206718 001</u> | Mar 31, 2017 |
| <u>AA</u> | | <u>300MG; 7.5MG</u> | <u>A206718 002</u> | Mar 31, 2017 |
| <u>AA</u> | | <u>300MG; 10MG</u> | <u>A206718 003</u> | Mar 31, 2017 |
| <u>AA</u> | | <u>325MG; 10MG</u> | <u>A040400 001</u> | Jul 26, 2000 |
| <u>AA</u> | SUN PHARM INDS INC | <u>325MG; 5MG</u> | <u>A090118 001</u> | Dec 23, 2008 |
| <u>AA</u> | | <u>325MG; 7.5MG</u> | <u>A090118 002</u> | Dec 23, 2008 |
| <u>AA</u> | | <u>325MG; 10MG</u> | <u>A090118 003</u> | Dec 23, 2008 |
| <u>AA</u> | TRIS PHARMA INC | <u>300MG; 5MG</u> | <u>A202214 004</u> | Mar 15, 2016 |
| <u>AA</u> | | <u>300MG; 7.5MG</u> | <u>A202214 005</u> | Mar 15, 2016 |
| <u>AA</u> | | <u>300MG; 10MG</u> | <u>A202214 006</u> | Mar 15, 2016 |
| <u>AA</u> | | <u>325MG; 5MG</u> | <u>A202214 001</u> | Mar 27, 2013 |
| <u>AA</u> | | <u>325MG; 7.5MG</u> | <u>A202214 002</u> | Mar 27, 2013 |
| <u>AA</u> | | <u>325MG; 10MG</u> | <u>A202214 003</u> | Mar 27, 2013 |
| <u>AA</u> | UPSHER SMITH LABS | <u>325MG; 5MG</u> | <u>A206484 001</u> | Mar 24, 2017 |
| <u>AA</u> | | <u>325MG; 7.5MG</u> | <u>A206484 002</u> | Mar 24, 2017 |
| <u>AA</u> | | <u>325MG; 10MG</u> | <u>A206484 003</u> | Mar 24, 2017 |
| <u>AA</u> | VINTAGE PHARMS | <u>300MG; 5MG</u> | <u>A090415 001</u> | Jan 24, 2011 |
| <u>AA</u> | | <u>300MG; 7.5MG</u> | <u>A090415 002</u> | Jan 24, 2011 |
| <u>AA</u> | | <u>300MG; 10MG</u> | <u>A090415 003</u> | Jan 24, 2011 |
| <u>AA</u> | | <u>325MG; 5MG</u> | <u>A040655 001</u> | Jan 19, 2006 |
| <u>AA</u> | | <u>325MG; 7.5MG</u> | <u>A040656 001</u> | Jan 19, 2006 |
| <u>AA</u> | | <u>325MG; 10MG</u> | <u>A040355 001</u> | May 31, 2000 |
| <u>AA</u> | WES PHARMA INC | <u>300MG; 5MG</u> | <u>A207509 001</u> | Oct 29, 2018 |
| <u>AA</u> | | <u>300MG; 7.5MG</u> | <u>A207509 002</u> | Oct 29, 2018 |
| <u>AA</u> | | <u>300MG; 10MG</u> | <u>A207509 003</u> | Oct 29, 2018 |
| <u>AA</u> | | <u>325MG; 5MG</u> | <u>A210211 001</u> | Oct 30, 2017 |
| <u>AA</u> | | <u>325MG; 7.5MG</u> | <u>A210211 002</u> | Oct 30, 2017 |
| <u>AA</u> | | <u>325MG; 10MG</u> | <u>A210211 003</u> | Oct 30, 2017 |
| <u>NORCO</u> | | | | |
| <u>AA</u> | APIL | <u>325MG; 2.5MG</u> | <u>A040148 004</u> | Jul 07, 2014 |
| <u>AA</u> | ! | <u>325MG; 5MG</u> | <u>A040099 001</u> | Jun 25, 1997 |
| <u>AA</u> | | <u>325MG; 5MG</u> | <u>A040148 005</u> | Jul 07, 2014 |
| <u>AA</u> | ! | <u>325MG; 7.5MG</u> | <u>A040148 003</u> | Sep 12, 2000 |
| <u>AA</u> | ! | <u>325MG; 10MG</u> | <u>A040148 001</u> | Feb 14, 1997 |

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

SOLUTION; ORAL

OXYCODONE AND ACETAMINOPHEN

| | | | | |
|---|------------|---------------------|-------------|--------------|
| ! | ABHAI LLC | 325MG/5ML; 5MG/5ML | A211499 001 | Dec 31, 2018 |
| | MIKART INC | 300MG/5ML; 10MG/5ML | A202142 001 | Nov 27, 2018 |

TABLET; ORAL

OXYCET

| | | | | |
|-----------|------------|-------------------|--------------------|--------------|
| <u>AA</u> | SPECGX LLC | <u>325MG; 5MG</u> | <u>A087463 001</u> | Dec 07, 1983 |
|-----------|------------|-------------------|--------------------|--------------|

OXYCODONE AND ACETAMINOPHEN

| | | | | |
|-----------|-------------------|---------------------|--------------------|--------------|
| <u>AA</u> | ABHAI LLC | <u>325MG; 2.5MG</u> | <u>A210644 001</u> | Feb 09, 2018 |
| <u>AA</u> | | <u>325MG; 5MG</u> | <u>A210644 002</u> | Feb 09, 2018 |
| <u>AA</u> | | <u>325MG; 7.5MG</u> | <u>A210644 003</u> | Feb 09, 2018 |
| <u>AA</u> | | <u>325MG; 10MG</u> | <u>A210644 004</u> | Feb 09, 2018 |
| <u>AA</u> | ACTAVIS ELIZABETH | <u>325MG; 2.5MG</u> | <u>A201447 001</u> | Apr 12, 2013 |
| <u>AA</u> | | <u>325MG; 5MG</u> | <u>A201447 002</u> | Apr 12, 2013 |
| <u>AA</u> | | <u>325MG; 7.5MG</u> | <u>A201447 003</u> | Apr 12, 2013 |

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE AND ACETAMINOPHEN

| | | | | | |
|-----------------------------|---------------------|--------------------|--------------------|--------------------|--------------|
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A201447 004</u> | Apr 12, 2013 | |
| <u>AA</u> | ALVOGEN MALTA | <u>325MG;5MG</u> | <u>A202677 003</u> | Mar 08, 2016 | |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A202677 001</u> | Jul 26, 2012 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A202677 002</u> | Jul 26, 2012 | |
| <u>AA</u> | AMNEAL PHARMS | <u>325MG;5MG</u> | <u>A040777 001</u> | Nov 27, 2007 | |
| <u>AA</u> | AMNEAL PHARMS NY | <u>325MG;7.5MG</u> | <u>A040778 002</u> | Jun 27, 2014 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A040778 001</u> | Nov 27, 2007 | |
| <u>AA</u> | ASCENT PHARMS INC | <u>325MG;2.5MG</u> | <u>A207419 001</u> | Mar 22, 2017 | |
| <u>AA</u> | | <u>325MG;5MG</u> | <u>A207419 002</u> | Mar 22, 2017 | |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A207419 003</u> | Mar 22, 2017 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A207419 004</u> | Mar 22, 2017 | |
| <u>AA</u> | AUROLIFE PHARMA LLC | <u>325MG;2.5MG</u> | <u>A201972 001</u> | Jul 15, 2013 | |
| <u>AA</u> | | <u>325MG;5MG</u> | <u>A201972 002</u> | Jul 15, 2013 | |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A201972 003</u> | Jul 15, 2013 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A201972 004</u> | Jul 15, 2013 | |
| <u>AA</u> | CHEMO RESEARCH SL | <u>325MG;5MG</u> | <u>A207574 001</u> | Dec 13, 2016 | |
| <u>AA</u> | ELITE LABS INC | <u>325MG;5MG</u> | <u>A209385 001</u> | Jul 02, 2018 | |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A209385 002</u> | Jul 02, 2018 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A209385 003</u> | Jul 02, 2018 | |
| <u>AA</u> | EPIC PHARMA LLC | <u>325MG;5MG</u> | <u>A203864 001</u> | Jul 02, 2018 | |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A203864 002</u> | Jul 02, 2018 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A203864 003</u> | Jul 02, 2018 | |
| <u>AA</u> | LANNETT CO INC | <u>325MG;5MG</u> | <u>A207333 001</u> | Sep 25, 2017 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A207333 002</u> | Sep 25, 2017 | |
| <u>AA</u> | MAYNE PHARMA INC | <u>325MG;2.5MG</u> | <u>A090177 001</u> | Oct 20, 2008 | |
| <u>AA</u> | | <u>325MG;5MG</u> | <u>A090177 002</u> | Oct 20, 2008 | |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A090177 003</u> | Oct 20, 2008 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A090177 004</u> | Oct 20, 2008 | |
| <u>AA</u> | NESHER PHARMS | <u>325MG;2.5MG</u> | <u>A210079 001</u> | Dec 28, 2017 | |
| <u>AA</u> | | <u>325MG;5MG</u> | <u>A210079 002</u> | Dec 28, 2017 | |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A210079 003</u> | Dec 28, 2017 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A210079 004</u> | Dec 28, 2017 | |
| <u>AA</u> | NOVEL LABS INC | <u>325MG;2.5MG</u> | <u>A204407 001</u> | Feb 24, 2017 | |
| <u>AA</u> | | <u>325MG;5MG</u> | <u>A204407 002</u> | Feb 24, 2017 | |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A204407 003</u> | Feb 24, 2017 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A204407 004</u> | Feb 24, 2017 | |
| <u>AA</u> | RHODES PHARMS | <u>325MG;5MG</u> | <u>A201278 001</u> | Aug 28, 2014 | |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A201278 002</u> | Aug 28, 2014 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A201278 003</u> | Aug 28, 2014 | |
| <u>AA</u> | SPECGX LLC | <u>325MG;7.5MG</u> | <u>A040545 001</u> | Jun 30, 2004 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A040545 002</u> | Jun 30, 2004 | |
| <u>AA</u> | SUN PHARM INDS INC | <u>325MG;2.5MG</u> | <u>A090535 001</u> | Dec 26, 2013 | |
| <u>AA</u> | | <u>325MG;5MG</u> | <u>A090535 002</u> | Dec 26, 2013 | |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A090535 003</u> | Dec 26, 2013 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A090535 004</u> | Dec 26, 2013 | |
| <u>AA</u> | VINTAGE PHARMS | <u>325MG;2.5MG</u> | <u>A090733 001</u> | Jul 11, 2013 | |
| <u>AA</u> | | <u>325MG;5MG</u> | <u>A040105 001</u> | Jul 30, 1996 | |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A090734 001</u> | Jul 11, 2013 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A090734 002</u> | Jul 11, 2013 | |
| <u>AA</u> | WATSON LABS | <u>325MG;5MG</u> | <u>A040171 001</u> | Oct 30, 1997 | |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A040535 001</u> | Sep 05, 2003 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A040535 002</u> | Sep 05, 2003 | |
| <u>AA</u> | WES PHARMA INC | <u>325MG;5MG</u> | <u>A207510 001</u> | Mar 21, 2018 | |
| <u>AA</u> | | <u>325MG;7.5MG</u> | <u>A207510 002</u> | Mar 21, 2018 | |
| <u>AA</u> | | <u>325MG;10MG</u> | <u>A207510 003</u> | Mar 21, 2018 | |
| <u>PERCOCET</u> | | | | | |
| <u>AA</u> | ! | VINTAGE PHARMS LLC | <u>325MG;2.5MG</u> | <u>A040330 001</u> | Jun 25, 1999 |
| <u>AA</u> | ! | | <u>325MG;5MG</u> | <u>A040330 002</u> | Jun 25, 1999 |
| <u>AA</u> | ! | | <u>325MG;7.5MG</u> | <u>A040330 003</u> | Nov 23, 2001 |
| <u>AA</u> | ! | | <u>325MG;10MG</u> | <u>A040330 004</u> | Nov 23, 2001 |
| <u>ROXICET</u> | | | | | |
| <u>AA</u> | | WEST-WARD PHARMS | <u>325MG;5MG</u> | <u>A087003 001</u> | |
| OXYCODONE AND ACETAMINOPHEN | | | | | |
| | ! | MIKART | 300MG;2.5MG | A040608 001 | Dec 30, 2005 |
| | ! | | 300MG;5MG | A040608 002 | Dec 30, 2005 |
| | ! | | 300MG;7.5MG | A040608 003 | Dec 30, 2005 |
| | ! | | 300MG;10MG | A040608 004 | Dec 30, 2005 |

PRESCRIPTION DRUG PRODUCT LIST

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

| | | | | |
|-----------|----------------------|----------------------|--------------------|--------------|
| AB | ALKEM LABS LTD | <u>325MG; 37.5MG</u> | <u>A202076 001</u> | Mar 30, 2012 |
| AB | AMNEAL PHARMS | <u>325MG; 37.5MG</u> | <u>A090485 001</u> | Dec 09, 2009 |
| AB | APOTEX INC | <u>325MG; 37.5MG</u> | <u>A078778 001</u> | Apr 07, 2014 |
| AB | AUROBINDO PHARMA LTD | <u>325MG; 37.5MG</u> | <u>A207152 001</u> | Mar 22, 2017 |
| AB | MACLEODS PHARMS LTD | <u>325MG; 37.5MG</u> | <u>A206885 001</u> | May 02, 2017 |
| AB | MICRO LABS LTD INDIA | <u>325MG; 37.5MG</u> | <u>A201952 001</u> | Dec 14, 2012 |
| AB | MYLAN | <u>325MG; 37.5MG</u> | <u>A077858 001</u> | Sep 26, 2008 |
| AB | PAR PHARM | <u>325MG; 37.5MG</u> | <u>A076475 001</u> | Apr 21, 2005 |
| AB | SUN PHARM INDS INC | <u>325MG; 37.5MG</u> | <u>A077184 001</u> | Dec 16, 2005 |
| AB | ZYDUS PHARMS USA INC | <u>325MG; 37.5MG</u> | <u>A090460 001</u> | Sep 06, 2012 |

ULTRACET

| | | | | |
|-----------|---------------------------|----------------------|--------------------|--------------|
| AB | + ! JANSSEN PHARMS | <u>325MG; 37.5MG</u> | <u>N021123 001</u> | Aug 15, 2001 |
|-----------|---------------------------|----------------------|--------------------|--------------|

ACETAZOLAMIDE

CAPSULE, EXTENDED RELEASE; ORAL

ACETAZOLAMIDE

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| AB | ACCORD HLTHCARE | <u>500MG</u> | <u>A207659 001</u> | Oct 18, 2018 |
| AB | HERITAGE PHARMS INC | <u>500MG</u> | <u>A090779 001</u> | Jul 14, 2011 |
| AB | NOSTRUM LABS INC | <u>500MG</u> | <u>A204691 001</u> | Mar 29, 2016 |
| AB | NOVAST LABS | <u>500MG</u> | <u>A203434 001</u> | Sep 30, 2016 |
| AB | XYLOPIA | <u>500MG</u> | <u>A205301 001</u> | Jan 16, 2019 |
| AB | ZYDUS PHARMS USA INC | <u>500MG</u> | <u>A040904 001</u> | Dec 10, 2008 |

DIAMOX

| | | | | |
|-----------|-------------------------------|--------------|--------------------|--|
| AB | + ! TEVA BRANDED PHARM | <u>500MG</u> | <u>N012945 001</u> | |
|-----------|-------------------------------|--------------|--------------------|--|

TABLET; ORAL

ACETAZOLAMIDE

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| AB | HERITAGE PHARMA | <u>125MG</u> | <u>A205530 001</u> | Oct 27, 2016 |
| AB | | <u>250MG</u> | <u>A205530 002</u> | Oct 27, 2016 |
| AB | LANNETT | <u>250MG</u> | <u>A084840 001</u> | |
| AB | STRIDES PHARMA | <u>125MG</u> | <u>A209734 001</u> | Nov 20, 2017 |
| AB | | <u>250MG</u> | <u>A209734 002</u> | Nov 20, 2017 |
| AB | SUN PHARM INDUSTRIES | <u>125MG</u> | <u>A089753 002</u> | Jun 22, 1988 |
| AB | TARO | <u>125MG</u> | <u>A040195 001</u> | May 28, 1997 |
| AB | ! | <u>250MG</u> | <u>A040195 002</u> | May 28, 1997 |

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION

ACETAZOLAMIDE SODIUM

| | | | | |
|-----------|-----------------------|---------------------------|--------------------|--------------|
| AP | EMCURE PHARMS LTD | <u>EQ 500MG BASE/VIAL</u> | <u>A202693 001</u> | Dec 19, 2014 |
| AP | MYLAN ASI | <u>EQ 500MG BASE/VIAL</u> | <u>A200880 001</u> | May 09, 2012 |
| AP | PAR STERILE PRODUCTS | <u>EQ 500MG BASE/VIAL</u> | <u>A205358 001</u> | Jun 20, 2017 |
| AP | WEST-WARD PHARMS INT | <u>EQ 500MG BASE/VIAL</u> | <u>A040089 001</u> | Feb 28, 1995 |
| AP | ! X GEN PHARMS | <u>EQ 500MG BASE/VIAL</u> | <u>A040784 001</u> | Dec 10, 2008 |

ACETIC ACID, GLACIAL

SOLUTION; IRRIGATION, URETHRAL

ACETIC ACID 0.25% IN PLASTIC CONTAINER

| | | | | |
|-----------|----------------------------|--------------------|--------------------|--------------|
| AT | B BRAUN | <u>250MG/100ML</u> | <u>N018161 001</u> | |
| AT | BAXTER HLTHCARE | <u>250MG/100ML</u> | <u>N018523 001</u> | Feb 19, 1982 |
| AT | + ! ICU MEDICAL INC | <u>250MG/100ML</u> | <u>N017656 001</u> | |

SOLUTION/DROPS; OTIC

ACETIC ACID

| | | | | |
|-----------|---------------------------|-----------|--------------------|--------------|
| AT | LANNETT CO INC | <u>2%</u> | <u>A040607 001</u> | Feb 24, 2005 |
| AT | RISING PHARMS | <u>2%</u> | <u>A207280 001</u> | Mar 09, 2018 |
| AT | TARO | <u>2%</u> | <u>A088638 001</u> | Sep 06, 1984 |
| AT | ! WOCKHARDT BIO AG | <u>2%</u> | <u>A040166 001</u> | Jul 26, 1996 |

VOSOL

| | | | | |
|-----------|----------------|-----------|--------------------|--|
| AT | HI TECH PHARMA | <u>2%</u> | <u>N012179 001</u> | |
|-----------|----------------|-----------|--------------------|--|

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS; OTIC

ACETASOL HC

| | | | | |
|-----------|----------------------|---------------|--------------------|--------------|
| AT | ACTAVIS MID ATLANTIC | <u>2%; 1%</u> | <u>A087143 001</u> | Jan 13, 1982 |
|-----------|----------------------|---------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS;OTIC

HYDROCORTISONE AND ACETIC ACID

| | | | | |
|-----------|---------------------|--------------|--------------------|--------------|
| <u>AT</u> | TARO PHARM INDS LTD | <u>2%:1%</u> | <u>A088759 001</u> | Mar 04, 1985 |
| <u>AT</u> | VINTAGE | <u>2%:1%</u> | <u>A040609 001</u> | Feb 06, 2006 |

VOSOL HC

| | | | | |
|-----------|-------------------|--------------|--------------------|--|
| <u>AT</u> | +! HI TECH PHARMA | <u>2%:1%</u> | <u>N012770 001</u> | |
|-----------|-------------------|--------------|--------------------|--|

ACETOHYDROXAMIC ACID

TABLET;ORAL

LITHOSTAT

| | | | | |
|----|----------------|-------|-------------|--------------|
| +! | MISSION PHARMA | 250MG | N018749 001 | May 31, 1983 |
|----|----------------|-------|-------------|--------------|

ACETYLCHOLINE CHLORIDE

FOR SOLUTION;OPHTHALMIC

MIOCHOL-E

| | | | | |
|----|-----------------|-----------|-------------|--------------|
| +! | BAUSCH AND LOMB | 20MG/VIAL | N020213 001 | Sep 22, 1993 |
|----|-----------------|-----------|-------------|--------------|

ACETYLCYSTEINE

INJECTABLE;INTRAVENOUS

ACETADOTE

| | | | | |
|-----------|----------------------|----------------------------|--------------------|--------------|
| <u>AP</u> | +! CUMBERLAND PHARMS | <u>6GM/30ML (200MG/ML)</u> | <u>N021539 001</u> | Jan 23, 2004 |
|-----------|----------------------|----------------------------|--------------------|--------------|

ACETYLCYSTEINE

| | | | | |
|-----------|-----------|----------------------------|--------------------|--------------|
| <u>AP</u> | AKORN INC | <u>6GM/30ML (200MG/ML)</u> | <u>A203173 001</u> | Mar 24, 2015 |
|-----------|-----------|----------------------------|--------------------|--------------|

| | | | | |
|-----------|----------------------|----------------------------|--------------------|--------------|
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>6GM/30ML (200MG/ML)</u> | <u>A207358 001</u> | Feb 29, 2016 |
|-----------|----------------------|----------------------------|--------------------|--------------|

| | | | | |
|-----------|--------------------|----------------------------|--------------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>6GM/30ML (200MG/ML)</u> | <u>A200644 001</u> | Nov 07, 2012 |
|-----------|--------------------|----------------------------|--------------------|--------------|

| | | | | |
|-----------|---------------------|----------------------------|--------------------|--------------|
| <u>AP</u> | MYLAN INSTITUTIONAL | <u>6GM/30ML (200MG/ML)</u> | <u>A203624 001</u> | Jun 19, 2015 |
|-----------|---------------------|----------------------------|--------------------|--------------|

| | | | | |
|-----------|---------------|----------------------------|--------------------|--------------|
| <u>AP</u> | SAGENT PHARMS | <u>6GM/30ML (200MG/ML)</u> | <u>A091684 001</u> | Oct 31, 2017 |
|-----------|---------------|----------------------------|--------------------|--------------|

| | | | | |
|-----------|----------------------|----------------------------|--------------------|--------------|
| <u>AP</u> | ZYDUS PHARMS USA INC | <u>6GM/30ML (200MG/ML)</u> | <u>A208166 001</u> | Jul 20, 2018 |
|-----------|----------------------|----------------------------|--------------------|--------------|

SOLUTION;INHALATION, ORAL

ACETYLCYSTEINE

| | | | | |
|-----------|-------------|------------|--------------------|--------------|
| <u>AN</u> | ALVOGEN INC | <u>10%</u> | <u>A204674 001</u> | Feb 11, 2014 |
| <u>AN</u> | | <u>20%</u> | <u>A203853 001</u> | Jun 21, 2012 |

| | | | | |
|-----------|---------|------------|--------------------|--------------|
| <u>AN</u> | HOSPIRA | <u>10%</u> | <u>A073664 001</u> | Aug 30, 1994 |
|-----------|---------|------------|--------------------|--------------|

| | | | | |
|-----------|--|------------|--------------------|--------------|
| <u>AN</u> | | <u>20%</u> | <u>A074037 001</u> | Aug 30, 1994 |
|-----------|--|------------|--------------------|--------------|

| | | | | |
|-----------|------------|------------|--------------------|--------------|
| <u>AN</u> | ! LUITPOLD | <u>10%</u> | <u>A072489 001</u> | Jul 28, 1995 |
|-----------|------------|------------|--------------------|--------------|

| | | | | |
|-----------|---|------------|--------------------|--------------|
| <u>AN</u> | ! | <u>20%</u> | <u>A072547 001</u> | Jul 28, 1995 |
|-----------|---|------------|--------------------|--------------|

TABLET, EFFERVESCENT;ORAL

CETYLEV

| | | | | |
|---|------------------|-------|-------------|--------------|
| + | ARBOR PHARMS LLC | 500MG | N207916 001 | Jan 29, 2016 |
|---|------------------|-------|-------------|--------------|

| | | | | |
|----|--|-------|-------------|--------------|
| +! | | 2.5GM | N207916 002 | Jan 29, 2016 |
|----|--|-------|-------------|--------------|

ACITRETIN

CAPSULE;ORAL

ACITRETIN

| | | | | |
|-----------|---------------|-------------|--------------------|--------------|
| <u>AB</u> | BARR LABS INC | <u>10MG</u> | <u>A091455 001</u> | Apr 04, 2013 |
|-----------|---------------|-------------|--------------------|--------------|

| | | | | |
|-----------|--|-------------|--------------------|--------------|
| <u>AB</u> | | <u>25MG</u> | <u>A091455 002</u> | Apr 04, 2013 |
|-----------|--|-------------|--------------------|--------------|

| | | | | |
|-----------|----------------|-------------|--------------------|--------------|
| <u>AB</u> | IMPAX LABS INC | <u>10MG</u> | <u>A202552 001</u> | Dec 23, 2015 |
|-----------|----------------|-------------|--------------------|--------------|

| | | | | |
|-----------|--|---------------|--------------------|--------------|
| <u>AB</u> | | <u>17.5MG</u> | <u>A202552 002</u> | Dec 23, 2015 |
|-----------|--|---------------|--------------------|--------------|

| | | | | |
|-----------|--|---------------|--------------------|--------------|
| <u>AB</u> | | <u>22.5MG</u> | <u>A202552 003</u> | Dec 23, 2015 |
|-----------|--|---------------|--------------------|--------------|

| | | | | |
|-----------|--|-------------|--------------------|--------------|
| <u>AB</u> | | <u>25MG</u> | <u>A202552 004</u> | Dec 23, 2015 |
|-----------|--|-------------|--------------------|--------------|

| | | | | |
|-----------|------------------|-------------|--------------------|--------------|
| <u>AB</u> | MYLAN PHARMS INC | <u>10MG</u> | <u>A202148 001</u> | Sep 10, 2015 |
|-----------|------------------|-------------|--------------------|--------------|

| | | | | |
|-----------|--|-------------|--------------------|--------------|
| <u>AB</u> | | <u>25MG</u> | <u>A202148 002</u> | Sep 10, 2015 |
|-----------|--|-------------|--------------------|--------------|

| | | | | |
|-----------|---------------------|-------------|--------------------|--------------|
| <u>AB</u> | SIGMAPHARM LABS LLC | <u>10MG</u> | <u>A204633 001</u> | May 22, 2015 |
|-----------|---------------------|-------------|--------------------|--------------|

| | | | | |
|-----------|--|---------------|--------------------|--------------|
| <u>AB</u> | | <u>17.5MG</u> | <u>A204633 002</u> | May 22, 2015 |
|-----------|--|---------------|--------------------|--------------|

| | | | | |
|-----------|--|---------------|--------------------|--------------|
| <u>AB</u> | | <u>22.5MG</u> | <u>A204633 003</u> | May 22, 2015 |
|-----------|--|---------------|--------------------|--------------|

| | | | | |
|-----------|--|-------------|--------------------|--------------|
| <u>AB</u> | | <u>25MG</u> | <u>A204633 004</u> | May 22, 2015 |
|-----------|--|-------------|--------------------|--------------|

| | | | | |
|-----------|-----------------|---------------|--------------------|--------------|
| <u>AB</u> | TEVA PHARMS USA | <u>17.5MG</u> | <u>A202897 001</u> | Apr 04, 2013 |
|-----------|-----------------|---------------|--------------------|--------------|

| | | | | |
|-----------|--|---------------|--------------------|--------------|
| <u>AB</u> | | <u>22.5MG</u> | <u>A202897 002</u> | Apr 04, 2013 |
|-----------|--|---------------|--------------------|--------------|

SORIATANE

| | | | | |
|-----------|--------------------|-------------|--------------------|--------------|
| <u>AB</u> | + STIEFEL LABS INC | <u>10MG</u> | <u>N019821 001</u> | Oct 28, 1996 |
|-----------|--------------------|-------------|--------------------|--------------|

| | | | | |
|-----------|---|---------------|--------------------|--------------|
| <u>AB</u> | + | <u>17.5MG</u> | <u>N019821 003</u> | Aug 06, 2009 |
|-----------|---|---------------|--------------------|--------------|

| | | | | |
|-----------|---|---------------|--------------------|--------------|
| <u>AB</u> | + | <u>22.5MG</u> | <u>N019821 004</u> | Aug 06, 2009 |
|-----------|---|---------------|--------------------|--------------|

| | | | | |
|-----------|----|-------------|--------------------|--------------|
| <u>AB</u> | +! | <u>25MG</u> | <u>N019821 002</u> | Oct 28, 1996 |
|-----------|----|-------------|--------------------|--------------|

ACLDINIUM BROMIDE

POWDER, METERED;INHALATION

TUDORZA PRESSAIR

| | | | | |
|----|--------------------|-----------|-------------|--------------|
| +! | ASTRAZENECA PHARMS | 0.4MG/INH | N202450 001 | Jul 23, 2012 |
|----|--------------------|-----------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

ACRIVASTINE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE; ORAL

SEMPREX-D

+! AUXILIUM PHARMS LLC 8MG;60MG

N019806 001 Mar 25, 1994

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

| | | | | | | |
|-----------|---|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ! | APOTEX INC | <u>200MG</u> | <u>A075677</u> | <u>001</u> | Sep 28, 2005 |
| <u>AB</u> | | BOSCOGEN | <u>200MG</u> | <u>A075090</u> | <u>001</u> | Jan 26, 1999 |
| <u>AB</u> | | CADILA PHARMS LTD | <u>200MG</u> | <u>A201445</u> | <u>001</u> | Mar 06, 2014 |
| <u>AB</u> | | CARLSBAD TECHNOLOGY | <u>200MG</u> | <u>A206261</u> | <u>001</u> | Aug 16, 2017 |
| <u>AB</u> | | DAVA PHARMS INC | <u>200MG</u> | <u>A074833</u> | <u>001</u> | Apr 22, 1997 |
| <u>AB</u> | | HERITAGE PHARMS INC | <u>200MG</u> | <u>A074889</u> | <u>001</u> | Oct 31, 1997 |
| <u>AB</u> | | TEVA | <u>200MG</u> | <u>A074578</u> | <u>001</u> | Apr 22, 1997 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>200MG</u> | <u>A204313</u> | <u>001</u> | Mar 25, 2016 |

ZOVIRAX

| | | | | | | |
|-----------|---|------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | MYLAN PHARMS INC | <u>200MG</u> | <u>N018828</u> | <u>001</u> | Jan 25, 1985 |
|-----------|---|------------------|--------------|----------------|------------|--------------|

CREAM; TOPICAL

ZOVIRAX

+! VIB 5%

N021478 001 Dec 30, 2002

OINTMENT; TOPICAL

ACYCLOVIR

| | | | | | | |
|-----------|--|---------------------|-----------|----------------|------------|--------------|
| <u>AB</u> | | ALEMERIC PHARMS LTD | <u>5%</u> | <u>A209000</u> | <u>001</u> | Apr 06, 2018 |
| <u>AB</u> | | AMNEAL PHARMS | <u>5%</u> | <u>A204605</u> | <u>001</u> | Jun 18, 2014 |
| <u>AB</u> | | FOUGERA PHARMS INC | <u>5%</u> | <u>A206633</u> | <u>001</u> | May 11, 2016 |
| <u>AB</u> | | G AND W LABS INC | <u>5%</u> | <u>A205591</u> | <u>001</u> | Nov 13, 2017 |
| <u>AB</u> | | GLENMARK PHARMS LTD | <u>5%</u> | <u>A205510</u> | <u>001</u> | Jul 31, 2017 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>5%</u> | <u>A202459</u> | <u>001</u> | Apr 03, 2013 |
| <u>AB</u> | | TARO | <u>5%</u> | <u>A205469</u> | <u>001</u> | Dec 21, 2016 |
| <u>AB</u> | | TOLMAR | <u>5%</u> | <u>A206437</u> | <u>001</u> | Jul 31, 2017 |
| <u>AB</u> | | TORRENT PHARMS LTD | <u>5%</u> | <u>A209971</u> | <u>001</u> | Jan 11, 2019 |

ZOVIRAX

| | | | | | | |
|-----------|---|-----------------|-----------|----------------|------------|--------------|
| <u>AB</u> | ! | VALEANT BERMUDA | <u>5%</u> | <u>N018604</u> | <u>001</u> | Mar 29, 1982 |
|-----------|---|-----------------|-----------|----------------|------------|--------------|

SUSPENSION; ORAL

ACYCLOVIR

| | | | | | | |
|-----------|--|-------------------------|------------------|----------------|------------|--------------|
| <u>AB</u> | | ACTAVIS MID ATLANTIC | <u>200MG/5ML</u> | <u>A074738</u> | <u>001</u> | Apr 28, 1997 |
| <u>AB</u> | | HI TECH PHARMA | <u>200MG/5ML</u> | <u>A077026</u> | <u>001</u> | Jun 07, 2005 |

ZOVIRAX

| | | | | | | |
|-----------|---|------------------|------------------|----------------|------------|--------------|
| <u>AB</u> | ! | MYLAN PHARMS INC | <u>200MG/5ML</u> | <u>N019909</u> | <u>001</u> | Dec 22, 1989 |
|-----------|---|------------------|------------------|----------------|------------|--------------|

TABLET; BUCCAL

SITAVIG

+! EPI HLTH 50MG

N203791 001 Apr 12, 2013

TABLET; ORAL

ACYCLOVIR

| | | | | | | |
|-----------|--|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | APOTEX INC | <u>400MG</u> | <u>A077309</u> | <u>001</u> | Sep 29, 2005 |
| <u>AB</u> | | | <u>800MG</u> | <u>A077309</u> | <u>002</u> | Sep 29, 2005 |
| <u>AB</u> | | CADILA PHARMS LTD | <u>400MG</u> | <u>A202168</u> | <u>001</u> | Nov 15, 2013 |
| <u>AB</u> | | | <u>800MG</u> | <u>A202168</u> | <u>002</u> | Nov 15, 2013 |
| <u>AB</u> | | CARLSBAD | <u>400MG</u> | <u>A075382</u> | <u>001</u> | Apr 30, 1999 |
| <u>AB</u> | | | <u>800MG</u> | <u>A075382</u> | <u>002</u> | Apr 30, 1999 |
| <u>AB</u> | | DAVA PHARMS INC | <u>400MG</u> | <u>A074946</u> | <u>001</u> | Nov 19, 1997 |
| <u>AB</u> | | | <u>800MG</u> | <u>A074946</u> | <u>002</u> | Nov 19, 1997 |
| <u>AB</u> | | HERITAGE PHARMS INC | <u>400MG</u> | <u>A074891</u> | <u>001</u> | Oct 31, 1997 |
| <u>AB</u> | | | <u>800MG</u> | <u>A074891</u> | <u>002</u> | Oct 31, 1997 |
| <u>AB</u> | | HETERO LABS LTD V | <u>400MG</u> | <u>A203834</u> | <u>001</u> | Oct 29, 2013 |
| <u>AB</u> | | | <u>800MG</u> | <u>A203834</u> | <u>002</u> | Oct 29, 2013 |
| <u>AB</u> | | TEVA | <u>400MG</u> | <u>A074556</u> | <u>002</u> | Apr 22, 1997 |
| <u>AB</u> | | | <u>800MG</u> | <u>A074556</u> | <u>003</u> | Apr 22, 1997 |
| <u>AB</u> | | YILING PHARM LTD | <u>400MG</u> | <u>A210401</u> | <u>001</u> | Mar 07, 2018 |
| <u>AB</u> | | | <u>800MG</u> | <u>A210401</u> | <u>002</u> | Mar 07, 2018 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>400MG</u> | <u>A204314</u> | <u>001</u> | Aug 19, 2014 |
| <u>AB</u> | | | <u>800MG</u> | <u>A204314</u> | <u>002</u> | Aug 19, 2014 |

ZOVIRAX

| | | | | | | |
|-----------|---|------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | MYLAN PHARMS INC | <u>400MG</u> | <u>N020089</u> | <u>001</u> | Apr 30, 1991 |
|-----------|---|------------------|--------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|--|--------------|----------------|------------|--------------|
| <u>AB</u> | ! | | <u>800MG</u> | <u>N020089</u> | <u>002</u> | Apr 30, 1991 |
|-----------|---|--|--------------|----------------|------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR SODIUM

| | | | | |
|-----------|----------------------|------------------------|--------------------|--------------|
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>EQ 50MG BASE/ML</u> | <u>A203701 001</u> | Oct 11, 2013 |
| <u>AP</u> | ! FRESENIUS KABI USA | <u>EQ 50MG BASE/ML</u> | <u>A074930 001</u> | May 13, 1998 |
| | ZYDUS PHARMS USA INC | EQ 500MG BASE/VIAL | A206535 001 | Aug 31, 2018 |
| | | EQ 1GM BASE/VIAL | A206535 002 | Aug 31, 2018 |

ACYCLOVIR; HYDROCORTISONE

CREAM; TOPICAL

XERESE

| | | | | |
|---|-----------------|-------|-------------|--------------|
| + | VALEANT BERMUDA | 5%;1% | N022436 001 | Jul 31, 2009 |
|---|-----------------|-------|-------------|--------------|

ADAPALENE

CREAM; TOPICAL

ADAPALENE

| | | | | |
|-----------|----------------|-------------|--------------------|--------------|
| <u>AB</u> | FOUGERA PHARMS | <u>0.1%</u> | <u>A090824 001</u> | Jun 30, 2010 |
|-----------|----------------|-------------|--------------------|--------------|

DIFFERIN

| | | | | |
|-----------|--------------------|-------------|--------------------|--------------|
| <u>AB</u> | ! GALDERMA LABS LP | <u>0.1%</u> | <u>N020748 001</u> | May 26, 2000 |
|-----------|--------------------|-------------|--------------------|--------------|

GEL; TOPICAL

ADAPALENE

| | | | | |
|-----------|----------------------|-------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS MID ATLANTIC | <u>0.3%</u> | <u>A201000 001</u> | Oct 27, 2014 |
|-----------|----------------------|-------------|--------------------|--------------|

| | | | | |
|-----------|-------------------|-------------|--------------------|--------------|
| <u>AB</u> | GLENMARK GENERICS | <u>0.1%</u> | <u>A091314 001</u> | Jul 01, 2010 |
|-----------|-------------------|-------------|--------------------|--------------|

| | | | | |
|-----------|--------------------|-------------|--------------------|--------------|
| <u>AB</u> | PLIVA HRVATSKA DOO | <u>0.1%</u> | <u>A090962 001</u> | Jun 02, 2010 |
|-----------|--------------------|-------------|--------------------|--------------|

| | | | | |
|-----------|------|-------------|--------------------|--------------|
| <u>AB</u> | TARO | <u>0.3%</u> | <u>A208322 001</u> | Jun 23, 2016 |
|-----------|------|-------------|--------------------|--------------|

| | | | | |
|-----------|--------|-------------|--------------------|--------------|
| <u>AB</u> | TOLMAR | <u>0.3%</u> | <u>A200298 001</u> | Jun 14, 2012 |
|-----------|--------|-------------|--------------------|--------------|

DIFFERIN

| | | | | |
|-----------|--------------------|-------------|--------------------|--------------|
| <u>AB</u> | ! GALDERMA LABS LP | <u>0.3%</u> | <u>N021753 001</u> | Jun 19, 2007 |
|-----------|--------------------|-------------|--------------------|--------------|

LOTION; TOPICAL

DIFFERIN

| | | | | |
|---|------------------|------|-------------|--------------|
| + | GALDERMA LABS LP | 0.1% | N022502 001 | Mar 17, 2010 |
|---|------------------|------|-------------|--------------|

SOLUTION; TOPICAL

ADAPALENE

| | | | | |
|-----------|----------|-------------|--------------------|--------------|
| <u>AB</u> | CALL INC | <u>0.1%</u> | <u>A203981 001</u> | Sep 23, 2016 |
|-----------|----------|-------------|--------------------|--------------|

| | | | | |
|-----------|--|-------------|--------------------|--------------|
| <u>AB</u> | | <u>0.1%</u> | <u>A204593 001</u> | Jan 05, 2016 |
|-----------|--|-------------|--------------------|--------------|

ADAPALENE; BENZOYL PEROXIDE

GEL; TOPICAL

ADAPALENE AND BENZOYL PEROXIDE

| | | | | |
|-----------|----------------------|------------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS MID ATLANTIC | <u>0.1%;2.5%</u> | <u>A203790 001</u> | Sep 30, 2015 |
|-----------|----------------------|------------------|--------------------|--------------|

| | | | | |
|-----------|----------------|------------------|--------------------|--------------|
| <u>AB</u> | PERRIGO ISRAEL | <u>0.1%;2.5%</u> | <u>A205033 001</u> | Jan 23, 2018 |
|-----------|----------------|------------------|--------------------|--------------|

| | | | | |
|-----------|------|------------------|--------------------|--------------|
| <u>AB</u> | TARO | <u>0.1%;2.5%</u> | <u>A206959 001</u> | Jan 24, 2018 |
|-----------|------|------------------|--------------------|--------------|

| | | | | |
|-----------|--|------------------|--------------------|--------------|
| <u>AB</u> | | <u>0.3%;2.5%</u> | <u>A209148 001</u> | Oct 17, 2018 |
|-----------|--|------------------|--------------------|--------------|

| | | | | |
|-----------|--------|------------------|--------------------|--------------|
| <u>AB</u> | TOLMAR | <u>0.1%;2.5%</u> | <u>A206164 001</u> | May 23, 2018 |
|-----------|--------|------------------|--------------------|--------------|

EPIDUO

| | | | | |
|-----------|--------------------|------------------|--------------------|--------------|
| <u>AB</u> | ! GALDERMA LABS LP | <u>0.1%;2.5%</u> | <u>N022320 001</u> | Dec 08, 2008 |
|-----------|--------------------|------------------|--------------------|--------------|

EPIDUO FORTE

| | | | | |
|-----------|-----------------|------------------|--------------------|--------------|
| <u>AB</u> | ! GALDERMA LABS | <u>0.3%;2.5%</u> | <u>N207917 001</u> | Jul 15, 2015 |
|-----------|-----------------|------------------|--------------------|--------------|

ADEFOVIR DIPIVOXIL

TABLET; ORAL

ADEFOVIR DIPIVOXIL

| | | | | |
|-----------|------------|-------------|--------------------|--------------|
| <u>AB</u> | APOTEX INC | <u>10MG</u> | <u>A205459 001</u> | Jul 06, 2018 |
|-----------|------------|-------------|--------------------|--------------|

| | | | | |
|-----------|---------------------|-------------|--------------------|--------------|
| <u>AB</u> | SIGMAPHARM LABS LLC | <u>10MG</u> | <u>A202051 001</u> | Aug 29, 2013 |
|-----------|---------------------|-------------|--------------------|--------------|

HEPSERA

| | | | | |
|-----------|----------|-------------|--------------------|--------------|
| <u>AB</u> | ! GILEAD | <u>10MG</u> | <u>N021449 001</u> | Sep 20, 2002 |
|-----------|----------|-------------|--------------------|--------------|

ADENOSINE

INJECTABLE; INJECTION

ADENOSINE

| | | | | |
|-----------|---------|---------------|--------------------|--------------|
| <u>AP</u> | ! AKORN | <u>3MG/ML</u> | <u>A078076 001</u> | Oct 31, 2008 |
|-----------|---------|---------------|--------------------|--------------|

| | | | | |
|-----------|--------------------|---------------|--------------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>3MG/ML</u> | <u>A077133 001</u> | Apr 27, 2005 |
|-----------|--------------------|---------------|--------------------|--------------|

| | | | | |
|-----------|--|---------------|--------------------|--------------|
| <u>AP</u> | | <u>3MG/ML</u> | <u>A205568 001</u> | Apr 16, 2018 |
|-----------|--|---------------|--------------------|--------------|

| | | | | |
|-----------|------------------|---------------|--------------------|--------------|
| <u>AP</u> | GLAND PHARMA LTD | <u>3MG/ML</u> | <u>A077283 001</u> | Jun 14, 2007 |
|-----------|------------------|---------------|--------------------|--------------|

| | | | | |
|-----------|--|---------------|--------------------|--------------|
| <u>AP</u> | | <u>3MG/ML</u> | <u>A206778 001</u> | Feb 16, 2018 |
|-----------|--|---------------|--------------------|--------------|

| | | | | |
|-----------|----------|---------------|--------------------|--------------|
| <u>AP</u> | LUITPOLD | <u>3MG/ML</u> | <u>A090010 001</u> | Apr 28, 2009 |
|-----------|----------|---------------|--------------------|--------------|

| | | | | |
|-----------|----------------|---------------|--------------------|--------------|
| <u>AP</u> | MYLAN LABS LTD | <u>3MG/ML</u> | <u>A078640 001</u> | Mar 21, 2014 |
|-----------|----------------|---------------|--------------------|--------------|

| | | | | |
|-----------|--|---------------|--------------------|--------------|
| <u>AP</u> | | <u>3MG/ML</u> | <u>A078686 001</u> | May 13, 2009 |
|-----------|--|---------------|--------------------|--------------|

| | | | | |
|-----------|----------------------|---------------|--------------------|--------------|
| <u>AP</u> | WEST-WARD PHARMS INT | <u>3MG/ML</u> | <u>A076404 001</u> | Jun 16, 2004 |
|-----------|----------------------|---------------|--------------------|--------------|

| | | | | |
|-----------|--|---------------|--------------------|--------------|
| <u>AP</u> | | <u>3MG/ML</u> | <u>A076500 001</u> | Jun 16, 2004 |
|-----------|--|---------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

ADENOSINE

SOLUTION; INTRAVENOUS

ADENOSINE

| | | | | |
|-----------|----------------------|---------------------------|--------------------|--------------|
| AP | AKORN | 60MG/20ML (3MG/ML) | A090450 001 | Oct 02, 2014 |
| AP | | 90MG/30ML (3MG/ML) | A090450 002 | Oct 02, 2014 |
| AP | AUROBINDO PHARMA LTD | 60MG/20ML (3MG/ML) | A205331 001 | Nov 02, 2017 |
| AP | | 90MG/30ML (3MG/ML) | A205331 002 | Nov 02, 2017 |
| AP | EMCURE PHARMS LTD | 60MG/20ML (3MG/ML) | A202313 001 | Sep 15, 2014 |
| AP | | 90MG/30ML (3MG/ML) | A202313 002 | Sep 15, 2014 |
| AP | FRESENIUS KABI USA | 60MG/20ML (3MG/ML) | A077897 001 | Nov 27, 2017 |
| AP | | 90MG/30ML (3MG/ML) | A077897 002 | Nov 27, 2017 |
| AP | HOSPIRA INC | 60MG/20ML (3MG/ML) | A203883 001 | Mar 24, 2014 |
| AP | | 90MG/30ML (3MG/ML) | A203883 002 | Mar 24, 2014 |
| AP | MYLAN ASI | 60MG/20ML (3MG/ML) | A090212 001 | Mar 28, 2014 |
| AP | | 90MG/30ML (3MG/ML) | A090212 002 | Mar 28, 2014 |
| AP | ! TEVA PHARMS USA | 60MG/20ML (3MG/ML) | A077425 001 | Aug 29, 2013 |
| AP | ! | 90MG/30ML (3MG/ML) | A077425 002 | Aug 29, 2013 |

AFATINIB DIMALEATE

TABLET; ORAL

GILOTRIF

| | | | | |
|---|----------------------|--------------|-------------|--------------|
| + | BOEHRINGER INGELHEIM | EQ 20MG BASE | N201292 001 | Jul 12, 2013 |
| + | | EQ 30MG BASE | N201292 002 | Jul 12, 2013 |
| + | ! | EQ 40MG BASE | N201292 003 | Jul 12, 2013 |

ALBENDAZOLE

TABLET; ORAL

ALBENDAZOLE

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| AB | CIPLA LTD | 200MG | A210434 001 | Sep 21, 2018 |
| AB | STRIDES VIVIMED | 200MG | A210011 001 | Dec 07, 2018 |
| AB | ZYDUS PHARMS USA INC | 200MG | A208979 001 | Dec 14, 2018 |
| AB | ! IMPAX LABS INC | 200MG | N020666 001 | Jun 11, 1996 |

ALBUMIN HUMAN

INJECTABLE; INJECTION

OPTISON

| | | | | |
|---|---------------|---------|-------------|--------------|
| + | GE HEALTHCARE | 10MG/ML | N020899 001 | Dec 31, 1997 |
|---|---------------|---------|-------------|--------------|

ALBUMIN IODINATED I-125 SERUM

INJECTABLE; INJECTION

JEANATOPE

| | | | | |
|---|---------|------------------------|-------------|--------------|
| + | ISO TEX | 100uCi/10ML (10uCi/ML) | N017836 003 | Jun 08, 2004 |
| + | | 500uCi/0.5ML | N017836 001 | |
| + | ! | 1,000uCi/ML | N017836 002 | |

ALBUMIN IODINATED I-131 SERUM

INJECTABLE; INJECTION

MEGATOPE

| | | | | |
|---|---------|-------------|-------------|--|
| + | ISO TEX | 0.5mCi/VIAL | N017837 001 | |
| + | ! | 1mCi/VIAL | N017837 002 | |

ALBUTEROL SULFATE

AEROSOL, METERED; INHALATION

PROAIR HFA

| | | | | |
|----|-----------------------------|---------------------|-------------|--------------|
| BX | ! TEVA BRANDED PHARM | EQ 0.09MG BASE/INH | N021457 001 | Oct 29, 2004 |
| BX | ! 3M DRUG DELIVERY | EQ 0.09MG BASE/INH | N020503 001 | Aug 15, 1996 |
| BX | ! GLAXOSMITHKLINE | EQ 0.09MG BASE/INH | N020983 001 | Apr 19, 2001 |
| | POWDER, METERED; INHALATION | | | |
| | PROAIR RESPICLICK | | | |
| | ! TEVA BRANDED PHARM | EQ 0.090MG BASE/INH | N205636 001 | Mar 31, 2015 |

SOLUTION; INHALATION

ACCUNEB

| | | | | |
|-----------|-----------------------|-----------------------|--------------------|--------------|
| AN | ! MYLAN SPECIALITY LP | EQ 0.021% BASE | N020949 002 | Apr 30, 2001 |
| AN | ! | EQ 0.042% BASE | N020949 001 | Apr 30, 2001 |

ALBUTEROL SULFATE

| | | | | |
|-----------|----------------------|-----------------------|--------------------|--------------|
| AN | AUROBINDO PHARMA LTD | EQ 0.083% BASE | A206224 001 | Oct 17, 2017 |
| AN | ! BAUSCH AND LOMB | EQ 0.5% BASE | A075050 001 | Jun 18, 1998 |
| AN | HI TECH PHARMA | EQ 0.5% BASE | A074543 001 | Jan 15, 1998 |
| AN | NEPHRON | EQ 0.021% BASE | A076355 002 | Mar 31, 2010 |

PRESCRIPTION DRUG PRODUCT LIST

ALBUTEROL SULFATE

SOLUTION; INHALATION

ALBUTEROL SULFATE

| | | | | |
|-----------|-------------------|-----------------------|--------------------|--------------|
| <u>AN</u> | | <u>EQ 0.042% BASE</u> | <u>A076355 001</u> | Jun 28, 2004 |
| <u>AN</u> | ! | <u>EQ 0.083% BASE</u> | <u>A074880 001</u> | Sep 17, 1997 |
| <u>AN</u> | | <u>EQ 0.5% BASE</u> | <u>A075664 001</u> | Jun 26, 2001 |
| <u>AN</u> | RITEDOSE CORP | <u>EQ 0.083% BASE</u> | <u>A077839 001</u> | Dec 16, 2008 |
| <u>AN</u> | SUN PHARMA GLOBAL | <u>EQ 0.083% BASE</u> | <u>A207857 001</u> | Jul 21, 2017 |
| <u>AN</u> | WATSON LABS | <u>EQ 0.021% BASE</u> | <u>A077772 001</u> | Sep 25, 2007 |
| <u>AN</u> | | <u>EQ 0.042% BASE</u> | <u>A077772 002</u> | Sep 25, 2007 |

SYRUP; ORAL

ALBUTEROL SULFATE

| | | | | |
|-----------|------------------|------------------------|--------------------|--------------|
| <u>AA</u> | AMNEAL PHARMS | <u>EQ 2MG BASE/5ML</u> | <u>A079241 001</u> | May 12, 2010 |
| <u>AA</u> | G AND W LABS INC | <u>EQ 2MG BASE/5ML</u> | <u>A074454 001</u> | Sep 25, 1995 |
| <u>AA</u> | HI TECH PHARMA | <u>EQ 2MG BASE/5ML</u> | <u>A074749 001</u> | Jan 30, 1998 |
| <u>AA</u> | LANNETT CO INC | <u>EQ 2MG BASE/5ML</u> | <u>A078105 001</u> | Dec 27, 2006 |
| <u>AA</u> | ! | <u>EQ 2MG BASE/5ML</u> | <u>A073419 001</u> | Mar 30, 1992 |
| <u>AA</u> | VISTAPHARM | <u>EQ 2MG BASE/5ML</u> | <u>A077788 001</u> | Jun 26, 2007 |

TABLET; ORAL

ALBUTEROL SULFATE

| | | | | |
|-----------|----------------------|--------------------|--------------------|--------------|
| <u>AB</u> | AMNEAL PHARMS CO | <u>EQ 2MG BASE</u> | <u>A208804 001</u> | May 21, 2018 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A208804 002</u> | May 21, 2018 |
| <u>AB</u> | MYLAN | <u>EQ 2MG BASE</u> | <u>A072894 002</u> | Jan 17, 1991 |
| <u>AB</u> | ! | <u>EQ 4MG BASE</u> | <u>A072894 001</u> | Jan 17, 1991 |
| <u>AB</u> | RISING PHARMS | <u>EQ 2MG BASE</u> | <u>A207046 001</u> | Jun 29, 2018 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A207046 002</u> | Jun 29, 2018 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>EQ 2MG BASE</u> | <u>A072637 002</u> | Dec 05, 1989 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A072637 001</u> | Dec 05, 1989 |
| <u>AB</u> | VIRTUS PHARM | <u>EQ 2MG BASE</u> | <u>A211397 001</u> | Oct 26, 2018 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A211397 002</u> | Oct 26, 2018 |

TABLET, EXTENDED RELEASE; ORAL

ALBUTEROL SULFATE

| | | | | |
|-----------|-------------------|--------------------|--------------------|--------------------------|
| <u>AB</u> | MYLAN | <u>EQ 4MG BASE</u> | <u>A078092 002</u> | Jan 29, 2007 |
| | <u>VOSPIRE ER</u> | | | |
| <u>AB</u> | DAVA PHARMS INC | <u>EQ 4MG BASE</u> | <u>A076130 002</u> | Sep 26, 2002 |
| | ALBUTEROL SULFATE | | | |
| | ! | MYLAN | EQ 8MG BASE | A078092 001 Jan 29, 2007 |

ALBUTEROL SULFATE; IPRATROPIUM BROMIDE

SOLUTION; INHALATION

ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE

| | | | | |
|-----------|-------------------|-------------------------------|--------------------|--------------|
| <u>AN</u> | CIPLA | <u>EQ 0.083% BASE; 0.017%</u> | <u>A077559 001</u> | Dec 31, 2007 |
| <u>AN</u> | NEPHRON | <u>EQ 0.083% BASE; 0.017%</u> | <u>A076749 001</u> | Dec 31, 2007 |
| <u>AN</u> | RITEDOSE CORP | <u>EQ 0.083% BASE; 0.017%</u> | <u>A202496 001</u> | Oct 01, 2012 |
| <u>AN</u> | SUN PHARMA GLOBAL | <u>EQ 0.083% BASE; 0.017%</u> | <u>A207875 001</u> | Aug 07, 2017 |
| <u>AN</u> | WATSON LABS TEVA | <u>EQ 0.083% BASE; 0.017%</u> | <u>A077063 001</u> | Dec 31, 2007 |

SPRAY, METERED; INHALATION

COMBIVENT RESPIMAT

| | | | | |
|---|---|-------------------------|-------------------------------|--------------------------|
| + | ! | BOEHRINGER INGELHEIM | EQ 0.1MG BASE/INH; 0.02MG/INH | N021747 001 Oct 07, 2011 |
|---|---|-------------------------|-------------------------------|--------------------------|

ALCAFTADINE

SOLUTION/DROPS; OPHTHALMIC

LASTACFT

| | | | | |
|---|---|----------|-------|--------------------------|
| + | ! | ALLERGAN | 0.25% | N022134 001 Jul 28, 2010 |
|---|---|----------|-------|--------------------------|

ALCLOMETASONE DIPROPIONATE

CREAM; TOPICAL

ALCLOMETASONE DIPROPIONATE

| | | | | | |
|-----------|---|-------------------|--------------|--------------------|--------------|
| <u>AB</u> | ! | FOUGERA PHARMS | <u>0.05%</u> | <u>A076973 001</u> | Jul 12, 2005 |
| <u>AB</u> | | GLENMARK GENERICS | <u>0.05%</u> | <u>A079061 001</u> | Jun 23, 2009 |
| <u>AB</u> | | TARO | <u>0.05%</u> | <u>A076587 001</u> | Sep 15, 2005 |

OINTMENT; TOPICAL

ALCLOMETASONE DIPROPIONATE

| | | | | | |
|-----------|---|-------------------|--------------|--------------------|--------------|
| <u>AB</u> | ! | FOUGERA PHARMS | <u>0.05%</u> | <u>A076884 001</u> | Jul 18, 2005 |
| <u>AB</u> | | GLENMARK GENERICS | <u>0.05%</u> | <u>A079227 001</u> | Jul 30, 2009 |
| <u>AB</u> | | TARO | <u>0.05%</u> | <u>A076730 001</u> | Jul 29, 2004 |

PRESCRIPTION DRUG PRODUCT LIST

ALCOHOL

SOLUTION; INTRA-ARTERIAL

ABLYSINOL

| | | | | | |
|---|--------------------|-----------|---------|-----|--------------|
| + | BELCHER PHARMS LLC | 99% (1ML) | N207987 | 001 | Jun 21, 2018 |
| + | ! | 99% (5ML) | N207987 | 002 | Jun 21, 2018 |

ALECTINIB HYDROCHLORIDE

CAPSULE; ORAL

ALECENSA

| | | | | | | |
|---|---|-------------------|---------------|---------|-----|--------------|
| + | ! | HOFFMANN-LA ROCHE | EQ 150MG BASE | N208434 | 001 | Dec 11, 2015 |
|---|---|-------------------|---------------|---------|-----|--------------|

ALENDRONATE SODIUM

SOLUTION; ORAL

ALENDRONATE SODIUM

| | | | | | |
|---|-------------------------|-------------------|---------|-----|--------------|
| ! | WEST-WARD PHARMS INT | EQ 70MG BASE/75ML | A090520 | 001 | Feb 25, 2013 |
|---|-------------------------|-------------------|---------|-----|--------------|

TABLET; ORAL

ALENDRONATE SODIUM

| | | | | | |
|-----------|-------------------|---------------------|----------------|------------|--------------|
| <u>AB</u> | APOTEX | <u>EQ 5MG BASE</u> | <u>A077982</u> | <u>001</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A077982</u> | <u>002</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 35MG BASE</u> | <u>A077982</u> | <u>003</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 70MG BASE</u> | <u>A077982</u> | <u>004</u> | Aug 04, 2008 |
| <u>AB</u> | AUROBINDO PHARMA | <u>EQ 10MG BASE</u> | <u>A090124</u> | <u>001</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 35MG BASE</u> | <u>A090124</u> | <u>002</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 70MG BASE</u> | <u>A090124</u> | <u>003</u> | Aug 04, 2008 |
| <u>AB</u> | CIPLA | <u>EQ 5MG BASE</u> | <u>A076768</u> | <u>001</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A076768</u> | <u>002</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 35MG BASE</u> | <u>A076768</u> | <u>003</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A076768</u> | <u>004</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 70MG BASE</u> | <u>A076768</u> | <u>005</u> | Aug 04, 2008 |
| <u>AB</u> | HANGZHOU BINJIANG | <u>EQ 5MG BASE</u> | <u>A090258</u> | <u>001</u> | Sep 24, 2009 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A090258</u> | <u>002</u> | Sep 24, 2009 |
| <u>AB</u> | | <u>EQ 35MG BASE</u> | <u>A090258</u> | <u>003</u> | Sep 24, 2009 |
| <u>AB</u> | | <u>EQ 70MG BASE</u> | <u>A090258</u> | <u>004</u> | Sep 24, 2009 |
| <u>AB</u> | IMPAX LABS INC | <u>EQ 5MG BASE</u> | <u>A075710</u> | <u>001</u> | Feb 06, 2008 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A075710</u> | <u>002</u> | Feb 06, 2008 |
| <u>AB</u> | | <u>EQ 35MG BASE</u> | <u>A075710</u> | <u>003</u> | Feb 06, 2008 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A075710</u> | <u>004</u> | Feb 06, 2008 |
| <u>AB</u> | | <u>EQ 70MG BASE</u> | <u>A075710</u> | <u>005</u> | Feb 06, 2008 |
| <u>AB</u> | JUBILANT CADISTA | <u>EQ 5MG BASE</u> | <u>A090557</u> | <u>001</u> | Feb 18, 2010 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A090557</u> | <u>002</u> | Feb 18, 2010 |
| <u>AB</u> | | <u>EQ 35MG BASE</u> | <u>A090557</u> | <u>003</u> | Feb 18, 2010 |
| <u>AB</u> | | <u>EQ 70MG BASE</u> | <u>A090557</u> | <u>004</u> | Feb 18, 2010 |
| <u>AB</u> | MYLAN | <u>EQ 5MG BASE</u> | <u>A076584</u> | <u>001</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A076584</u> | <u>002</u> | Aug 04, 2008 |
| <u>AB</u> | NEOPHARMA | <u>EQ 5MG BASE</u> | <u>A079049</u> | <u>003</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A079049</u> | <u>004</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 35MG BASE</u> | <u>A079049</u> | <u>001</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 70MG BASE</u> | <u>A079049</u> | <u>002</u> | Aug 04, 2008 |
| <u>AB</u> | SUN PHARMA GLOBAL | <u>EQ 5MG BASE</u> | <u>A090022</u> | <u>001</u> | Sep 10, 2008 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A090022</u> | <u>002</u> | Sep 10, 2008 |
| <u>AB</u> | | <u>EQ 35MG BASE</u> | <u>A090022</u> | <u>003</u> | Sep 10, 2008 |
| <u>AB</u> | | <u>EQ 70MG BASE</u> | <u>A090022</u> | <u>004</u> | Sep 10, 2008 |
| <u>AB</u> | WATSON LABS | <u>EQ 35MG BASE</u> | <u>A076984</u> | <u>001</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A076984</u> | <u>002</u> | Aug 04, 2008 |
| <u>AB</u> | | <u>EQ 70MG BASE</u> | <u>A076984</u> | <u>003</u> | Aug 04, 2008 |

FOSAMAX

| | | | | | | | |
|----------------------------|---|----------------|------------------|---------------------|----------------|--------------|--------------|
| <u>AB</u> | + | ! | MERCK AND CO INC | <u>EQ 70MG BASE</u> | <u>N020560</u> | <u>005</u> | Oct 20, 2000 |
| TABLET, EFFERVESCENT; ORAL | | | | | | | |
| BINOSTO | | | | | | | |
| + | ! | MISSION PHARMA | EQ 70MG BASE | N202344 | 001 | Mar 12, 2012 | |

ALENDRONATE SODIUM; CHOLECALCIFEROL

TABLET; ORAL

FOSAMAX PLUS D

| | | | | | |
|---|-------|------------------------|---------|-----|--------------|
| + | MERCK | EQ 70MG BASE; 2,800 IU | N021762 | 001 | Apr 07, 2005 |
| + | ! | EQ 70MG BASE; 5,600 IU | N021762 | 002 | Apr 26, 2007 |

ALFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

ALFENTA

| | | | | | | | |
|-----------|---|---|-------|-------------------------|----------------|------------|--------------|
| <u>AP</u> | + | ! | AKORN | <u>EQ 0.5MG BASE/ML</u> | <u>N019353</u> | <u>001</u> | Dec 29, 1986 |
|-----------|---|---|-------|-------------------------|----------------|------------|--------------|

ALFENTANIL

| | | | | | | | |
|-----------|--|--|---------|-------------------------|----------------|------------|--------------|
| <u>AP</u> | | | HOSPIRA | <u>EQ 0.5MG BASE/ML</u> | <u>A075221</u> | <u>001</u> | Oct 28, 1999 |
|-----------|--|--|---------|-------------------------|----------------|------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

ALFUZOSIN HYDROCHLORIDE

| | | | | | |
|-----------|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | APOTEX INC | <u>10MG</u> | <u>A079013</u> | <u>001</u> | Jul 18, 2011 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>10MG</u> | <u>A079060</u> | <u>001</u> | Aug 30, 2012 |
| <u>AB</u> | INVAGEN PHARMS | <u>10MG</u> | <u>A090284</u> | <u>001</u> | Jan 17, 2012 |
| <u>AB</u> | MYLAN | <u>10MG</u> | <u>A079014</u> | <u>001</u> | Jul 18, 2011 |
| <u>AB</u> | SUN PHARMA GLOBAL | <u>10MG</u> | <u>A079057</u> | <u>001</u> | Jul 18, 2011 |
| <u>AB</u> | TEVA PHARMS | <u>10MG</u> | <u>A079056</u> | <u>001</u> | Jul 18, 2011 |
| <u>AB</u> | TORRENT PHARMS | <u>10MG</u> | <u>A079054</u> | <u>001</u> | Jul 18, 2011 |
| <u>AB</u> | UNICHEM LABS LTD | <u>10MG</u> | <u>A203192</u> | <u>001</u> | Jan 28, 2016 |

UROXATRAL

| | | | | | |
|-----------|----------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | +! CONCORDIA PHARMS INC | <u>10MG</u> | <u>N021287</u> | <u>001</u> | Jun 12, 2003 |
|-----------|----------------------------|-------------|----------------|------------|--------------|

ALISKIREN HEMIFUMARATE

TABLET;ORAL

TEKTURNA

| | | | | | |
|----|--------------|---------------|---------|-----|--------------|
| + | NODEN PHARMA | EQ 150MG BASE | N021985 | 001 | Mar 05, 2007 |
| +! | | EQ 300MG BASE | N021985 | 002 | Mar 05, 2007 |

ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

TEKTURNA HCT

| | | | | | |
|----|--------------|----------------------|---------|-----|--------------|
| + | NODEN PHARMA | EQ 150MG BASE;12.5MG | N022107 | 001 | Jan 18, 2008 |
| + | | EQ 150MG BASE;25MG | N022107 | 002 | Jan 18, 2008 |
| +! | | EQ 300MG BASE;12.5MG | N022107 | 003 | Jan 18, 2008 |
| +! | | EQ 300MG BASE;25MG | N022107 | 004 | Jan 18, 2008 |

ALITRETINOIN

GEL;TOPICAL

PANRETIN

| | | | | | |
|---|-----------|--------------|---------|-----|--------------|
| + | EISAI INC | EQ 0.1% BASE | N020886 | 001 | Feb 02, 1999 |
|---|-----------|--------------|---------|-----|--------------|

ALLOPURINOL

TABLET;ORAL

ALLOPURINOL

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>100MG</u> | <u>A203154</u> | <u>001</u> | May 06, 2013 |
| <u>AB</u> | | <u>300MG</u> | <u>A203154</u> | <u>002</u> | May 06, 2013 |
| <u>AB</u> | APOTEX INC | <u>100MG</u> | <u>A077353</u> | <u>001</u> | Sep 08, 2005 |
| <u>AB</u> | | <u>300MG</u> | <u>A077353</u> | <u>002</u> | Sep 08, 2005 |
| <u>AB</u> | INDOCO REMEDIES | <u>100MG</u> | <u>A204467</u> | <u>001</u> | Jul 28, 2016 |
| <u>AB</u> | | <u>300MG</u> | <u>A204467</u> | <u>002</u> | Jul 28, 2016 |
| <u>AB</u> | IPCA LABS LTD | <u>100MG</u> | <u>A090637</u> | <u>001</u> | Mar 16, 2011 |
| <u>AB</u> | | <u>300MG</u> | <u>A090637</u> | <u>002</u> | Mar 16, 2011 |
| <u>AB</u> | MYLAN | <u>100MG</u> | <u>A018659</u> | <u>001</u> | Oct 24, 1986 |
| <u>AB</u> | | <u>300MG</u> | <u>A018659</u> | <u>002</u> | Oct 24, 1986 |
| <u>AB</u> | NORTHSTAR HLTHCARE | <u>100MG</u> | <u>A078253</u> | <u>001</u> | Sep 11, 2007 |
| <u>AB</u> | | <u>300MG</u> | <u>A078253</u> | <u>002</u> | Sep 11, 2007 |
| <u>AB</u> | SUN PHARM INDS INC | <u>100MG</u> | <u>A078390</u> | <u>001</u> | Aug 30, 2007 |
| <u>AB</u> | | <u>300MG</u> | <u>A078390</u> | <u>002</u> | Aug 30, 2007 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>100MG</u> | <u>A071450</u> | <u>002</u> | Jan 09, 1987 |
| <u>AB</u> | | <u>300MG</u> | <u>A071450</u> | <u>001</u> | Jan 09, 1987 |
| <u>AB</u> | VINTAGE PHARMS | <u>100MG</u> | <u>A075798</u> | <u>001</u> | Jun 27, 2003 |
| <u>AB</u> | | <u>300MG</u> | <u>A075798</u> | <u>002</u> | Jun 27, 2003 |
| <u>AB</u> | WATSON LABS | <u>100MG</u> | <u>N018832</u> | <u>002</u> | Sep 28, 1984 |
| <u>AB</u> | | <u>300MG</u> | <u>N018877</u> | <u>001</u> | Sep 28, 1984 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>100MG</u> | <u>A210117</u> | <u>001</u> | Oct 12, 2017 |
| <u>AB</u> | | <u>300MG</u> | <u>A210117</u> | <u>002</u> | Oct 12, 2017 |

LOPURIN

| | | | | | |
|-----------|--------------|--------------|----------------|------------|--------------|
| <u>AB</u> | DR REDDYS LA | <u>100MG</u> | <u>A071586</u> | <u>001</u> | Apr 02, 1987 |
| <u>AB</u> | | <u>300MG</u> | <u>A071587</u> | <u>001</u> | Apr 02, 1987 |

ZYLOPRIM

| | | | | | |
|-----------|---------------------|--------------|----------------|------------|--|
| <u>AB</u> | + CASPER PHARMA LLC | <u>100MG</u> | <u>N016084</u> | <u>001</u> | |
| <u>AB</u> | +! | <u>300MG</u> | <u>N016084</u> | <u>002</u> | |

ALLOPURINOL SODIUM

INJECTABLE;INJECTION

ALLOPURINOL SODIUM

| | | | | | |
|-----------|-------------------------|---------------------------|----------------|------------|--------------|
| <u>AP</u> | WEST-WARD PHARMS INT | <u>EQ 500MG BASE/VIAL</u> | <u>A076870</u> | <u>001</u> | Aug 26, 2004 |
|-----------|-------------------------|---------------------------|----------------|------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

ALLOPURINOL SODIUM

INJECTABLE; INJECTION

ALOPRIM

| | | | | | | |
|-----------|----------|---------------------|---------------------------|----------------|------------|--------------|
| AP | + | MYLAN INSTITUTIONAL | EQ 500MG BASE/VIAL | N020298 | 001 | May 17, 1996 |
|-----------|----------|---------------------|---------------------------|----------------|------------|--------------|

ALLOPURINOL; LESINURAD

TABLET; ORAL

DUZALLO

| | | | | | |
|---|---------------------|--------------|---------|-----|--------------|
| + | IRONWOOD PHARMS INC | 200MG; 200MG | N209203 | 001 | Aug 18, 2017 |
|---|---------------------|--------------|---------|-----|--------------|

| | | | | | |
|---|--|--------------|---------|-----|--------------|
| + | | 300MG; 200MG | N209203 | 002 | Aug 18, 2017 |
|---|--|--------------|---------|-----|--------------|

ALMOTRIPTAN MALATE

TABLET; ORAL

ALMOTRIPTAN MALATE

| | | | | | | |
|-----------|--|-------------------|-----------------------|----------------|------------|--------------|
| AB | | AJANTA PHARMA LTD | EQ 6.25MG BASE | A205523 | 001 | Mar 03, 2016 |
|-----------|--|-------------------|-----------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-----------------------|----------------|------------|--------------|
| AB | | | EQ 12.5MG BASE | A205523 | 002 | Mar 03, 2016 |
|-----------|--|--|-----------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|------------------|-----------------------|----------------|------------|--------------|
| AB | | MYLAN PHARMS INC | EQ 6.25MG BASE | A205171 | 001 | Nov 09, 2015 |
|-----------|--|------------------|-----------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-----------------------|----------------|------------|--------------|
| AB | | | EQ 12.5MG BASE | A205171 | 002 | Nov 09, 2015 |
|-----------|--|--|-----------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-----------------|-----------------------|----------------|------------|--------------|
| AB | | TEVA PHARMS USA | EQ 6.25MG BASE | A078027 | 001 | Jul 07, 2015 |
|-----------|--|-----------------|-----------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-----------------------|----------------|------------|--------------|
| AB | | | EQ 12.5MG BASE | A078027 | 002 | Jul 07, 2015 |
|-----------|--|--|-----------------------|----------------|------------|--------------|

AXERT

| | | | | | | |
|-----------|----------|----------------|-----------------------|----------------|------------|--------------|
| AB | + | JANSSEN PHARMS | EQ 6.25MG BASE | N021001 | 001 | May 07, 2001 |
|-----------|----------|----------------|-----------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|----------|--|-----------------------|----------------|------------|--------------|
| AB | + | | EQ 12.5MG BASE | N021001 | 002 | May 07, 2001 |
|-----------|----------|--|-----------------------|----------------|------------|--------------|

ALOGLIPTIN BENZOATE

TABLET; ORAL

NESINA

| | | | | | |
|---|-------------------|----------------|---------|-----|--------------|
| + | TAKEDA PHARMS USA | EQ 6.25MG BASE | N022271 | 001 | Jan 25, 2013 |
|---|-------------------|----------------|---------|-----|--------------|

| | | | | | |
|---|--|----------------|---------|-----|--------------|
| + | | EQ 12.5MG BASE | N022271 | 002 | Jan 25, 2013 |
|---|--|----------------|---------|-----|--------------|

| | | | | | |
|---|--|--------------|---------|-----|--------------|
| + | | EQ 25MG BASE | N022271 | 003 | Jan 25, 2013 |
|---|--|--------------|---------|-----|--------------|

ALOGLIPTIN BENZOATE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

KAZANO

| | | | | | |
|---|-------------------|-----------------------|---------|-----|--------------|
| + | TAKEDA PHARMS USA | EQ 12.5MG BASE; 500MG | N203414 | 001 | Jan 25, 2013 |
|---|-------------------|-----------------------|---------|-----|--------------|

| | | | | | |
|---|--|---------------------|---------|-----|--------------|
| + | | EQ 12.5MG BASE; 1GM | N203414 | 002 | Jan 25, 2013 |
|---|--|---------------------|---------|-----|--------------|

ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

OSEN

| | | | | | |
|---|-------------------|------------------------------|---------|-----|--------------|
| + | TAKEDA PHARMS USA | EQ 12.5MG BASE; EQ 15MG BASE | N022426 | 004 | Jan 25, 2013 |
|---|-------------------|------------------------------|---------|-----|--------------|

| | | | | | |
|---|--|------------------------------|---------|-----|--------------|
| + | | EQ 12.5MG BASE; EQ 30MG BASE | N022426 | 005 | Jan 25, 2013 |
|---|--|------------------------------|---------|-----|--------------|

| | | | | | |
|---|--|------------------------------|---------|-----|--------------|
| + | | EQ 12.5MG BASE; EQ 45MG BASE | N022426 | 006 | Jan 25, 2013 |
|---|--|------------------------------|---------|-----|--------------|

| | | | | | |
|---|--|----------------------------|---------|-----|--------------|
| + | | EQ 25MG BASE; EQ 15MG BASE | N022426 | 001 | Jan 25, 2013 |
|---|--|----------------------------|---------|-----|--------------|

| | | | | | |
|---|--|----------------------------|---------|-----|--------------|
| + | | EQ 25MG BASE; EQ 30MG BASE | N022426 | 002 | Jan 25, 2013 |
|---|--|----------------------------|---------|-----|--------------|

| | | | | | |
|---|--|----------------------------|---------|-----|--------------|
| + | | EQ 25MG BASE; EQ 45MG BASE | N022426 | 003 | Jan 25, 2013 |
|---|--|----------------------------|---------|-----|--------------|

ALOSETRON HYDROCHLORIDE

TABLET; ORAL

ALOSETRON HYDROCHLORIDE

| | | | | | | |
|-----------|--|---------------|----------------------|----------------|------------|--------------|
| AB | | AMNEAL PHARMS | EQ 0.5MG BASE | A206647 | 001 | Dec 22, 2016 |
|-----------|--|---------------|----------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|--------------------|----------------|------------|--------------|
| AB | | | EQ 1MG BASE | A206647 | 002 | Dec 22, 2016 |
|-----------|--|--|--------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-------|----------------------|----------------|------------|--------------|
| AB | | CIPLA | EQ 0.5MG BASE | A209180 | 001 | Jan 14, 2019 |
|-----------|--|-------|----------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|--------------------|----------------|------------|--------------|
| AB | | | EQ 1MG BASE | A209180 | 002 | Jan 14, 2019 |
|-----------|--|--|--------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|---------------|----------------------|----------------|------------|--------------|
| AB | | PAR PHARM INC | EQ 0.5MG BASE | A206113 | 001 | Feb 23, 2018 |
|-----------|--|---------------|----------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|--------------------|----------------|------------|--------------|
| AB | | | EQ 1MG BASE | A206113 | 002 | Feb 23, 2018 |
|-----------|--|--|--------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|------------------|----------------------|----------------|------------|--------------|
| AB | | WEST-WARD PHARMS | EQ 0.5MG BASE | A200652 | 001 | May 04, 2015 |
|-----------|--|------------------|----------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-----|--|--|--|--|
| AB | | INT | | | | |
|-----------|--|-----|--|--|--|--|

| | | | | | | |
|-----------|--|--|--------------------|----------------|------------|--------------|
| AB | | | EQ 1MG BASE | A200652 | 002 | May 04, 2015 |
|-----------|--|--|--------------------|----------------|------------|--------------|

LOTRONEX

| | | | | | | |
|-----------|----------|--------------------|----------------------|----------------|------------|--------------|
| AB | + | SEBELA IRELAND LTD | EQ 0.5MG BASE | N021107 | 002 | Dec 23, 2003 |
|-----------|----------|--------------------|----------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|----------|--|--------------------|----------------|------------|--------------|
| AB | + | | EQ 1MG BASE | N021107 | 001 | Feb 09, 2000 |
|-----------|----------|--|--------------------|----------------|------------|--------------|

ALPHA-TOCOPHEROL ACETATE; ASCORBIC ACID; BIOTIN; CHOLECALCIFEROL; CYANOCOBALAMIN; DEXPANTHENOL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A PALMITATE; VITAMIN K

SOLUTION; INTRAVENOUS

INFUVITE ADULT

| | | | | | |
|---|------------|--------------------------------|---------|-----|--------------|
| + | SANDOZ INC | 2 IU/ML; 40MG/ML; 12MCG/ML; 40 | N021163 | 001 | May 18, 2000 |
|---|------------|--------------------------------|---------|-----|--------------|

| | | | | | |
|--|--|----------------------------------------------|--|--|--|
| | | IU/ML; 1MCG/ML; 3MG/ML; 120MCG/ML; 8MG/ML; 1 | | | |
|--|--|----------------------------------------------|--|--|--|

| | | | | | |
|--|--|-----------------------------------|--|--|--|
| | | .2MG/ML; 0.72MG/ML; 1.2MG/ML; 660 | | | |
|--|--|-----------------------------------|--|--|--|

| | | | | | |
|--|--|------------------|--|--|--|
| | | IU/ML; 0.03MG/ML | | | |
|--|--|------------------|--|--|--|

| | | | | | |
|---|--|--------------------------------|---------|-----|--------------|
| + | | 2 IU/ML; 40MG/ML; 12MCG/ML; 40 | N021559 | 001 | Jun 16, 2003 |
|---|--|--------------------------------|---------|-----|--------------|

| | | | | | |
|--|--|----------------------------------------------|--|--|--|
| | | IU/ML; 1MCG/ML; 3MG/ML; 120MCG/ML; 8MG/ML; 1 | | | |
|--|--|----------------------------------------------|--|--|--|

| | | | | | |
|--|--|-----------------------------------|--|--|--|
| | | .2MG/ML; 0.72MG/ML; 1.2MG/ML; 660 | | | |
|--|--|-----------------------------------|--|--|--|

| | | | | | |
|--|--|-----------------|--|--|--|
| | | IU/ML; 30MCG/ML | | | |
|--|--|-----------------|--|--|--|

PRESCRIPTION DRUG PRODUCT LIST

ALPRAZOLAM

CONCENTRATE; ORAL

ALPRAZOLAM! WEST-WARD PHARMS
INT

1MG/ML

A074312 001 Oct 31, 1993

TABLET; ORAL

ALPRAZOLAM

| | | | | |
|--------------------------------|---------------------------|---------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS ELIZABETH | <u>0.25MG</u> | <u>A074342 001</u> | Oct 31, 1993 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A074342 002</u> | Oct 31, 1993 |
| <u>AB</u> | | <u>1MG</u> | <u>A074342 003</u> | Oct 31, 1993 |
| <u>AB</u> | | <u>2MG</u> | <u>A074342 004</u> | Oct 31, 1993 |
| <u>AB</u> | APOTEX INC | <u>0.25MG</u> | <u>A077741 001</u> | Jan 19, 2007 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A077741 002</u> | Jan 19, 2007 |
| <u>AB</u> | | <u>1MG</u> | <u>A077741 003</u> | Jan 19, 2007 |
| <u>AB</u> | | <u>2MG</u> | <u>A077741 004</u> | Jan 19, 2007 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>0.25MG</u> | <u>A203346 001</u> | Jul 31, 2015 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A203346 002</u> | Jul 31, 2015 |
| <u>AB</u> | | <u>1MG</u> | <u>A203346 003</u> | Jul 31, 2015 |
| <u>AB</u> | | <u>2MG</u> | <u>A203346 004</u> | Jul 31, 2015 |
| <u>AB</u> | BRECKENRIDGE PHARM | <u>0.25MG</u> | <u>A207507 001</u> | Jul 09, 2018 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A207507 002</u> | Jul 09, 2018 |
| <u>AB</u> | | <u>1MG</u> | <u>A207507 003</u> | Jul 09, 2018 |
| <u>AB</u> | | <u>2MG</u> | <u>A207507 004</u> | Jul 09, 2018 |
| <u>AB</u> | DAVA INTL INC | <u>0.25MG</u> | <u>A074174 001</u> | Oct 19, 1993 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A074174 002</u> | Oct 19, 1993 |
| <u>AB</u> | | <u>1MG</u> | <u>A074174 003</u> | Oct 19, 1993 |
| <u>AB</u> | | <u>2MG</u> | <u>A074174 004</u> | Oct 19, 1993 |
| <u>AB</u> | MYLAN | <u>0.25MG</u> | <u>A074215 001</u> | Jan 27, 1994 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A074215 002</u> | Jan 27, 1994 |
| <u>AB</u> | | <u>1MG</u> | <u>A074215 003</u> | Jan 27, 1994 |
| <u>AB</u> | | <u>2MG</u> | <u>A074215 004</u> | Jan 27, 1994 |
| <u>AB</u> | NATCO PHARMA LTD | <u>0.25MG</u> | <u>A200739 001</u> | Apr 15, 2015 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A200739 002</u> | Apr 15, 2015 |
| <u>AB</u> | | <u>1MG</u> | <u>A200739 003</u> | Apr 15, 2015 |
| <u>AB</u> | | <u>2MG</u> | <u>A200739 004</u> | Apr 15, 2015 |
| <u>AB</u> | OXFORD PHARMS | <u>0.25MG</u> | <u>A078491 001</u> | Sep 25, 2008 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A078491 002</u> | Sep 25, 2008 |
| <u>AB</u> | | <u>1MG</u> | <u>A078491 003</u> | Sep 25, 2008 |
| <u>AB</u> | | <u>2MG</u> | <u>A078491 004</u> | Dec 12, 2008 |
| <u>AB</u> | SANDOZ | <u>0.25MG</u> | <u>A074112 001</u> | Dec 29, 1995 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A074112 002</u> | Dec 29, 1995 |
| <u>AB</u> | | <u>1MG</u> | <u>A074112 003</u> | Dec 29, 1995 |
| <u>AB</u> | | <u>2MG</u> | <u>A074909 001</u> | Mar 25, 1998 |
| <u>AB</u> | SUN PHARMA GLOBAL | <u>0.25MG</u> | <u>A090082 001</u> | Jun 17, 2010 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A090082 002</u> | Jun 17, 2010 |
| <u>AB</u> | | <u>1MG</u> | <u>A090082 003</u> | Jun 17, 2010 |
| <u>AB</u> | | <u>2MG</u> | <u>A090082 004</u> | Jun 17, 2010 |
| <u>AB</u> | VINTAGE PHARMS | <u>0.25MG</u> | <u>A090248 001</u> | Sep 17, 2010 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A090248 002</u> | Sep 17, 2010 |
| <u>AB</u> | | <u>1MG</u> | <u>A090248 003</u> | Sep 17, 2010 |
| <u>AB</u> | | <u>2MG</u> | <u>A090248 004</u> | Sep 17, 2010 |
| <u>XANAX</u> | | | | |
| <u>AB</u> | + PHARMACIA AND UPJOHN | <u>0.25MG</u> | <u>N018276 001</u> | |
| <u>AB</u> | + | <u>0.5MG</u> | <u>N018276 002</u> | |
| <u>AB</u> | +! | <u>1MG</u> | <u>N018276 003</u> | |
| <u>AB</u> | + | <u>2MG</u> | <u>N018276 004</u> | Nov 27, 1985 |
| TABLET, EXTENDED RELEASE; ORAL | | | | |
| <u>ALPRAZOLAM</u> | | | | |
| <u>AB</u> | ACTAVIS ELIZABETH | <u>0.5MG</u> | <u>A078056 001</u> | Feb 13, 2007 |
| <u>AB</u> | | <u>1MG</u> | <u>A078056 002</u> | Feb 13, 2007 |
| <u>AB</u> | | <u>2MG</u> | <u>A078056 003</u> | Feb 13, 2007 |
| <u>AB</u> | | <u>3MG</u> | <u>A078056 004</u> | Feb 13, 2007 |
| <u>AB</u> | AMNEAL PHARMS NY | <u>0.5MG</u> | <u>A078387 001</u> | May 30, 2008 |
| <u>AB</u> | | <u>1MG</u> | <u>A078387 002</u> | May 30, 2008 |
| <u>AB</u> | | <u>2MG</u> | <u>A078387 003</u> | May 30, 2008 |
| <u>AB</u> | | <u>3MG</u> | <u>A078387 004</u> | May 30, 2008 |
| <u>AB</u> | ANCHEN PHARMS | <u>0.5MG</u> | <u>A078469 001</u> | Sep 29, 2011 |
| <u>AB</u> | | <u>1MG</u> | <u>A078469 002</u> | Sep 29, 2011 |
| <u>AB</u> | | <u>2MG</u> | <u>A078469 003</u> | Sep 29, 2011 |
| <u>AB</u> | | <u>3MG</u> | <u>A078469 004</u> | Sep 29, 2011 |
| <u>AB</u> | ANI PHARMS INC | <u>0.5MG</u> | <u>A077725 001</u> | Jul 31, 2006 |

PRESCRIPTION DRUG PRODUCT LIST

ALPRAZOLAM

TABLET, EXTENDED RELEASE;ORAL

ALPRAZOLAM

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| <u>AB</u> | | <u>1MG</u> | <u>A077725 002</u> | Jul 31, 2006 |
| <u>AB</u> | | <u>2MG</u> | <u>A077725 004</u> | Jul 31, 2006 |
| <u>AB</u> | | <u>3MG</u> | <u>A077725 003</u> | Jul 31, 2006 |
| <u>AB</u> | APOTEX INC | <u>0.5MG</u> | <u>A078449 001</u> | Nov 12, 2008 |
| <u>AB</u> | | <u>1MG</u> | <u>A078449 004</u> | Dec 23, 2015 |
| <u>AB</u> | | <u>2MG</u> | <u>A078449 002</u> | Nov 12, 2008 |
| <u>AB</u> | | <u>3MG</u> | <u>A078449 003</u> | Nov 12, 2008 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>0.5MG</u> | <u>A090871 001</u> | Jun 07, 2011 |
| <u>AB</u> | | <u>1MG</u> | <u>A090871 002</u> | Jun 07, 2011 |
| <u>AB</u> | | <u>2MG</u> | <u>A090871 003</u> | Jun 07, 2011 |
| <u>AB</u> | | <u>3MG</u> | <u>A090871 004</u> | Jun 07, 2011 |
| <u>AB</u> | HERITAGE PHARMS INC | <u>0.5MG</u> | <u>A078489 001</u> | Oct 17, 2008 |
| <u>AB</u> | | <u>1MG</u> | <u>A078489 002</u> | Oct 17, 2008 |
| <u>AB</u> | | <u>2MG</u> | <u>A078489 003</u> | Oct 17, 2008 |
| <u>AB</u> | | <u>3MG</u> | <u>A078489 004</u> | Oct 17, 2008 |

XANAX XR

| | | | | | |
|-----------|---|----------------------|--------------|--------------------|--------------|
| <u>AB</u> | + | PHARMACIA AND UPJOHN | <u>0.5MG</u> | <u>N021434 001</u> | Jan 17, 2003 |
| <u>AB</u> | + | | <u>1MG</u> | <u>N021434 002</u> | Jan 17, 2003 |
| <u>AB</u> | + | | <u>2MG</u> | <u>N021434 003</u> | Jan 17, 2003 |
| <u>AB</u> | + | | <u>3MG</u> | <u>N021434 004</u> | Jan 17, 2003 |

TABLET, ORALLY DISINTEGRATING;ORAL

ALPRAZOLAM

| | | | | | |
|-----------|---|-------------------|---------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS ELIZABETH | <u>0.25MG</u> | <u>A078561 001</u> | Mar 16, 2010 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A078561 002</u> | Mar 16, 2010 |
| <u>AB</u> | | | <u>1MG</u> | <u>A078561 003</u> | Mar 16, 2010 |
| <u>AB</u> | | | <u>2MG</u> | <u>A078561 004</u> | Mar 16, 2010 |
| <u>AB</u> | | PAR PHARM | <u>0.25MG</u> | <u>A078088 001</u> | Jan 09, 2009 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A078088 002</u> | Jan 09, 2009 |
| <u>AB</u> | ! | | <u>1MG</u> | <u>A078088 003</u> | Jan 09, 2009 |
| <u>AB</u> | | | <u>2MG</u> | <u>A078088 004</u> | Jan 09, 2009 |

ALPROSTADIL

INJECTABLE; INJECTION

ALPROSTADIL

| | | | | | |
|-----------|--|----------------------|-----------------|--------------------|--------------|
| <u>AP</u> | | TEVA PHARMS USA | <u>0.5MG/ML</u> | <u>A075196 001</u> | Apr 30, 1999 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>0.5MG/ML</u> | <u>A074815 001</u> | Jan 20, 1998 |

CAVERJECT

| | | | | | |
|-----------|---|----------------------|--------------------|--------------------|--------------|
| <u>AP</u> | + | PHARMACIA AND UPJOHN | <u>0.01MG/VIAL</u> | <u>N020379 001</u> | Jul 06, 1995 |
| <u>AP</u> | + | | <u>0.02MG/VIAL</u> | <u>N020379 002</u> | Jul 06, 1995 |
| <u>AP</u> | + | | <u>0.04MG/VIAL</u> | <u>N020379 004</u> | May 19, 1997 |

EDEX

| | | | | | |
|-----------|---|---------------------|--------------------|--------------------|--------------|
| <u>AP</u> | + | AUXILIUM PHARMS LLC | <u>0.01MG/VIAL</u> | <u>N020649 002</u> | Jun 12, 1997 |
| <u>AP</u> | + | | <u>0.02MG/VIAL</u> | <u>N020649 003</u> | Jun 12, 1997 |
| <u>AP</u> | + | | <u>0.04MG/VIAL</u> | <u>N020649 004</u> | Jun 12, 1997 |

PROSTIN VR PEDIATRIC

| | | | | | |
|-----------|---|----------------------|-----------------|--------------------|--------------|
| <u>AP</u> | + | PHARMACIA AND UPJOHN | <u>0.5MG/ML</u> | <u>N018484 001</u> | |
| | | CAVERJECT IMPULSE | 0.01MG/VIAL | N021212 001 | Jun 11, 2002 |
| | | PHARMACIA AND UPJOHN | 0.02MG/VIAL | N021212 002 | Jun 11, 2002 |
| | | EDEX | | | |
| | + | AUXILIUM PHARMS LLC | 0.01MG/VIAL | N020649 005 | Jul 30, 1998 |
| | + | | 0.02MG/VIAL | N020649 006 | Jul 30, 1998 |
| | + | | 0.04MG/VIAL | N020649 007 | Jul 30, 1998 |

SUPPOSITORY; URETHRAL

MUSE

| | | | | | |
|--|---|---------------------|---------|-------------|--------------|
| | + | MYLAN SPECIALITY LP | 0.125MG | N020700 001 | Nov 19, 1996 |
| | + | | 0.25MG | N020700 002 | Nov 19, 1996 |
| | + | | 0.5MG | N020700 003 | Nov 19, 1996 |
| | + | | 1MG | N020700 004 | Nov 19, 1996 |

PRESCRIPTION DRUG PRODUCT LIST

ALTRETAMINE

CAPSULE; ORAL

HEXALEN

+! EISAI INC

50MG

N019926 001 Dec 26, 1990

ALVIMOPAN

CAPSULE; ORAL

ENTEREG

+! CUBIST PHARMS

12MG

N021775 001 May 20, 2008

AMANTADINE HYDROCHLORIDE

CAPSULE; ORAL

AMANTADINE HYDROCHLORIDE**AB** ALEMBIC PHARMS LTD **100MG****A208966 001** Jun 21, 2017**AB** BIONPHARMA INC **100MG****A078720 001** May 29, 2008**AB** GRAVITI PHARMS **100MG****A207570 001** Sep 30, 2016**AB** HERITAGE PHARMA **100MG****A209171 001** Jun 12, 2017**AB** ! SANDOZ **100MG****A071293 001** Feb 18, 1987**AB** STRIDES PHARMA **100MG****A209047 001** Jun 07, 2017**AB** USL PHARMA **100MG****A070589 001** Aug 05, 1986**AB** WATSON LABS INC **100MG****A208107 001** Dec 06, 2016**AB** ZYDUS PHARMS USA **100MG****A208278 001** May 31, 2016

INC

CAPSULE, EXTENDED RELEASE; ORAL

GOCOVRI

+ ADAMAS PHARMA

EQ 68.5MG BASE

N208944 001 Aug 24, 2017

+!

EQ 137MG BASE

N208944 002 Aug 24, 2017

SYRUP; ORAL

AMANTADINE HYDROCHLORIDE**AA** ! CMP PHARMA INC **50MG/5ML****A075819 001** Sep 11, 2002**AA** ! HI TECH PHARMA **50MG/5ML****A074170 001** Oct 28, 1994**AA** ! MIKART **50MG/5ML****A074028 001** Jun 28, 1993**AA** ! PHARM ASSOC **50MG/5ML****A074509 001** Jul 17, 1995**AA** ! WOCKHARDT BIO AG **50MG/5ML****A075060 001** Dec 24, 1998

TABLET; ORAL

AMANTADINE HYDROCHLORIDE**AB** GRAVITI PHARMS **100MG****A207571 001** Jan 31, 2017**AB** JUBILANT GENERICS **100MG****A210403 001** Feb 07, 2018**AB** STRIDES PHARMA **100MG****A209035 001** Jun 09, 2017**AB** ! USL PHARMA **100MG****A076186 001** Dec 16, 2002**AB** WATSON LABS INC **100MG****A208096 001** Dec 15, 2016

TABLET, EXTENDED RELEASE; ORAL

OSMOLEX ER

+ OSMOTICA PHARM

EQ 129MG BASE

N209410 001 Feb 16, 2018

+

EQ 193MG BASE

N209410 002 Feb 16, 2018

+!

EQ 258MG BASE

N209410 003 Feb 16, 2018

AMBRISENTAN

TABLET; ORAL

LETAIRIS

+ GILEAD

5MG

N022081 001 Jun 15, 2007

+!

10MG

N022081 002 Jun 15, 2007

AMCINONIDE

CREAM; TOPICAL

AMCINONIDE**AB** ! FOUGERA PHARMS **0.1%****A076065 001** May 15, 2003**AB** TARO PHARM INDS **0.1%****A076229 001** May 31, 2002

LOTION; TOPICAL

AMCINONIDE

! FOUGERA PHARMS

0.1%

A076329 001 Nov 06, 2002

OINTMENT; TOPICAL

AMCINONIDE**AB** ! FOUGERA PHARMS **0.1%****A076096 001** Nov 19, 2002**AB** TARO PHARM INDS **0.1%****A076367 001** Mar 19, 2003AMIFAMPRIDINE PHOSPHATE

TABLET; ORAL

FIRDAPSE

+! CATALYST PHARMS

EQ 10MG BASE

N208078 001 Nov 28, 2018

PRESCRIPTION DRUG PRODUCT LIST

AMIFOSTINE

INJECTABLE; INJECTION

AMIFOSTINE

| | | | | |
|-----------|-------------------|-------------------|--------------------|--------------|
| <u>AP</u> | MYLAN LABS LTD | <u>500MG/VIAL</u> | <u>A204363 001</u> | Jul 17, 2017 |
| <u>AP</u> | SUN PHARMA GLOBAL | <u>500MG/VIAL</u> | <u>A077126 001</u> | Mar 14, 2008 |

ETHYOL

| | | | | |
|-----------|----------------------|-------------------|--------------------|--------------|
| <u>AP</u> | +! CLINIGEN HLTHCARE | <u>500MG/VIAL</u> | <u>N020221 001</u> | Dec 08, 1995 |
|-----------|----------------------|-------------------|--------------------|--------------|

AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKACIN SULFATE

| | | | | |
|-----------|---------------------------|-------------------------|--------------------|--------------|
| <u>AP</u> | ! EMCURE PHARMS LTD | <u>EQ 250MG BASE/ML</u> | <u>A204040 001</u> | Dec 12, 2013 |
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 50MG BASE/ML</u> | <u>A205605 001</u> | Dec 09, 2015 |
| <u>AP</u> | | <u>EQ 250MG BASE/ML</u> | <u>A205604 001</u> | Dec 09, 2015 |
| <u>AP</u> | SAGENT PHARMS | <u>EQ 250MG BASE/ML</u> | <u>A203323 001</u> | May 12, 2016 |
| <u>AP</u> | TEVA PHARMS USA | <u>EQ 250MG BASE/ML</u> | <u>A064045 002</u> | Sep 28, 1993 |
| <u>AP</u> | ! WEST-WARD PHARMS INT | <u>EQ 50MG BASE/ML</u> | <u>A063313 001</u> | Apr 11, 1994 |
| <u>AP</u> | | <u>EQ 250MG BASE/ML</u> | <u>A063315 001</u> | Apr 11, 1994 |

SUSPENSION, LIPOSOMAL; INHALATION

ARIKAYCE KIT

| | | | | |
|----|------------|---------------------|-------------|--------------|
| +! | INSMED INC | EQ 590MG BASE/8.4ML | N207356 001 | Sep 28, 2018 |
|----|------------|---------------------|-------------|--------------|

AMILORIDE HYDROCHLORIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE

| | | | | |
|-----------|---------------------|------------|--------------------|--------------|
| <u>AB</u> | ! PAR PHARM | <u>5MG</u> | <u>A070346 001</u> | Jan 22, 1986 |
| <u>AB</u> | SIGMAPHARM LABS LLC | <u>5MG</u> | <u>A079133 001</u> | Jan 30, 2009 |
| <u>AB</u> | USPHARMA WINDLAS | <u>5MG</u> | <u>A204180 001</u> | Aug 07, 2015 |

MIDAMOR

| | | | | |
|-----------|---------------|------------|--------------------|--|
| <u>AB</u> | + PADDOCK LLC | <u>5MG</u> | <u>N018200 001</u> | |
|-----------|---------------|------------|--------------------|--|

AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

| | | | | |
|-----------|---------|-------------------------------|--------------------|--------------|
| <u>AB</u> | BARR | <u>EQ 5MG ANHYDROUS; 50MG</u> | <u>A071111 001</u> | May 10, 1988 |
| <u>AB</u> | ! MYLAN | <u>EQ 5MG ANHYDROUS; 50MG</u> | <u>A073209 001</u> | Oct 31, 1991 |

AMINO ACIDS

INJECTABLE; INJECTION

AMINO ACIDS

| | | | |
|------------------------------------------------|--------------------|-------------|--------------|
| B BRAUN | 15% (150GM/1000ML) | A091112 001 | Apr 13, 2012 |
| | 15% (300GM/2000ML) | A091112 002 | Apr 13, 2012 |
| AMINOSYN 10% | | | |
| ICU MEDICAL INC | 10% (10GM/100ML) | N017673 003 | |
| AMINOSYN 8.5% | | | |
| ICU MEDICAL INC | 8.5% (8.5GM/100ML) | N017673 004 | |
| AMINOSYN II 10% IN PLASTIC CONTAINER | | | |
| ICU MEDICAL INC | 10% (10GM/100ML) | N020015 001 | Dec 19, 1991 |
| AMINOSYN II 15% IN PLASTIC CONTAINER | | | |
| ICU MEDICAL INC | 15% (15GM/100ML) | N020041 001 | Dec 19, 1991 |
| AMINOSYN-PF 10% | | | |
| ICU MEDICAL INC | 10% (10GM/100ML) | N019492 002 | Oct 17, 1986 |
| AMINOSYN-PF 7% | | | |
| ICU MEDICAL INC | 7% (7GM/100ML) | N019398 001 | Sep 06, 1985 |
| CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 15% (15GM/100ML) | A020512 001 | Aug 30, 1996 |
| FREAMINE HBC 6.9% | | | |
| B BRAUN | 6.9% (6.9GM/100ML) | N016822 006 | May 17, 1983 |
| FREAMINE III 10% | | | |
| B BRAUN | 10% (10GM/100ML) | N016822 005 | |
| FREAMINE III 8.5% | | | |
| B BRAUN | 8.5% (8.5GM/100ML) | N016822 004 | |
| HEPATAMINE 8% | | | |
| B BRAUN | 8% (8GM/100ML) | N018676 001 | Aug 03, 1982 |
| NEPHRAMINE 5.4% | | | |
| B BRAUN | 5.4% (5.4GM/100ML) | N017766 001 | |
| PREMASOL 10% IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 10% (10GM/100ML) | A075880 002 | Jun 19, 2003 |
| PREMASOL 6% IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 6% (6GM/100ML) | A075880 001 | Jun 19, 2003 |
| PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER | | | |
| +! BAXTER HLTHCARE | 20% (20GM/100ML) | N020849 001 | Aug 26, 1998 |

PRESCRIPTION DRUG PRODUCT LIST

AMINO ACIDS

INJECTABLE; INJECTION

| | | | |
|------------------------------------|--------------------|-------------|--------------|
| TRAVASOL 10% IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 10% (10GM/100ML) | N018931 003 | Aug 23, 1984 |
| TRAVASOL 5.5% IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 5.5% (5.5GM/100ML) | N018931 001 | Aug 23, 1984 |
| TRAVASOL 8.5% IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 8.5% (8.5GM/100ML) | N018931 002 | Aug 23, 1984 |
| TROPHAMINE | | | |
| +! B BRAUN | 6% (6GM/100ML) | N019018 001 | Jul 20, 1984 |
| TROPHAMINE 10% | | | |
| +! B BRAUN | 10% (10GM/100ML) | N019018 003 | Sep 07, 1988 |

AMINO ACIDS; CALCIUM ACETATE; GLYCERIN; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | | |
|-------------|------------------------------------------------------------------------------------------|-------------|--------------|
| PROCALAMINE | | | |
| B BRAUN | 3%; 26MG/100ML; 3GM/100ML; 54MG/100ML; 41MG/100ML; 150MG/100ML; 200MG/100ML; 120MG/100ML | N018582 001 | May 08, 1982 |

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | | |
|------------------------------------------------------------------------------------------|----------------------------------------------------------------------------------|-------------|--------------|
| CLINIMIX E 2.75/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER | | | |
| +! BAXTER HLTHCARE | 2.75%; 33MG/100ML; 10GM/100ML; 51MG/100ML; 261MG/100ML; 217MG/100ML; 112MG/100ML | N020678 002 | Mar 26, 1997 |
| CLINIMIX E 2.75/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER | | | |
| +! BAXTER HLTHCARE | 2.75%; 33MG/100ML; 25GM/100ML; 51MG/100ML; 261MG/100ML; 217MG/100ML; 112MG/100ML | N020678 005 | Mar 26, 1997 |
| CLINIMIX E 2.75/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER | | | |
| +! BAXTER HLTHCARE | 2.75%; 33MG/100ML; 5GM/100ML; 51MG/100ML; 261MG/100ML; 217MG/100ML; 112MG/100ML | N020678 001 | Mar 26, 1997 |
| CLINIMIX E 4.25/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER | | | |
| +! BAXTER HLTHCARE | 4.25%; 33MG/100ML; 10GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML | N020678 009 | Mar 26, 1997 |
| CLINIMIX E 4.25/20 SULFITE FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER | | | |
| +! BAXTER HLTHCARE | 4.25%; 33MG/100ML; 20GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML | N020678 011 | Mar 26, 1997 |
| CLINIMIX E 4.25/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER | | | |
| +! BAXTER HLTHCARE | 4.25%; 33MG/100ML; 25GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML | N020678 012 | Mar 26, 1997 |
| CLINIMIX E 4.25/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER | | | |
| +! BAXTER HLTHCARE | 4.25%; 33MG/100ML; 5GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML | N020678 008 | Mar 26, 1997 |
| CLINIMIX E 5/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER | | | |
| +! BAXTER HLTHCARE | 5%; 33MG/100ML; 10GM/100ML; 51MG/100ML; 261MG/100ML; 340MG/100ML; 59MG/100ML | N020678 016 | Mar 26, 1997 |
| CLINIMIX E 5/15 SULFITE FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER | | | |
| +! BAXTER HLTHCARE | 5%; 33MG/100ML; 15GM/100ML; 51MG/100ML; 261MG/100ML; 340MG/100ML; 59MG/100ML | N020678 017 | Mar 26, 1997 |
| CLINIMIX E 5/20 SULFITE FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER | | | |
| +! BAXTER HLTHCARE | 5%; 33MG/100ML; 20GM/100ML; 51MG/100ML; 261MG/100ML; 340MG/100ML; 59MG/100ML | N020678 018 | Mar 26, 1997 |
| CLINIMIX E 5/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER | | | |
| +! BAXTER HLTHCARE | 5%; 33MG/100ML; 25GM/100ML; 51MG/100ML; 261MG/100ML; 340MG/100ML; 59MG/100ML | N020678 019 | Mar 26, 1997 |
| CLINIMIX E 5/35 SULFITE FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER | | | |
| +! BAXTER HLTHCARE | 5%; 33MG/100ML; 35GM/100ML; 51MG/100ML; 261MG/100ML; 340MG/100ML; 59MG/100ML | N020678 021 | Mar 26, 1997 |

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM GLYCEROPHOSPHATE; SOYBEAN OIL

EMULSION; INTRAVENOUS

| | | | |
|------------------------------|--------------------------------------------------------------------------------------------------------|-------------|--------------|
| KABIVEN IN PLASTIC CONTAINER | | | |
| + FRESENIUS KABI USA | 3.3%; 29MG/100ML; 9.8GM/100ML; 96MG/100ML; 174MG/100ML; 239MG/100ML; 147MG/100ML; 3.9GM/100ML (1026ML) | N200656 004 | Aug 25, 2014 |
| + | 3.3%; 29MG/100ML; 9.8GM/100ML; 96MG/100ML; 174MG/100ML; 239MG/100ML; 147MG/100ML; 3.9GM/100ML (1540ML) | N200656 005 | Aug 25, 2014 |
| + | 3.3%; 29MG/100ML; 9.8GM/100ML; 96MG/100ML; 174MG/100ML; 239MG/100ML; 147MG/100ML; 3.9GM/100ML (2053ML) | N200656 006 | Aug 25, 2014 |
| +! | 3.3%; 29MG/100ML; 9.8GM/100ML; 96MG/100ML; 174MG/100ML; 239MG/100ML; 147MG/100ML; 3.9GM/100ML (2566ML) | N200656 007 | Aug 25, 2014 |

PRESCRIPTION DRUG PRODUCT LISTAMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM GLYCEROPHOSPHATE; SOYBEAN OIL

EMULSION; INTRAVENOUS

PERIKABIVEN IN PLASTIC CONTAINER

| | | | | |
|---|--------------------|--------------------------------------------------------------------------------------------------------|-------------|--------------|
| + | FRESENIUS KABI USA | 2.4%; 20MG/100ML; 6.8GM/100ML; 68MG/100ML; 124MG/100ML; 170MG/100ML; 105MG/100ML; 3.5GM/100ML (1440ML) | N200656 001 | Aug 25, 2014 |
| + | | 2.4%; 20MG/100ML; 6.8GM/100ML; 68MG/100ML; 124MG/100ML; 170MG/100ML; 105MG/100ML; 3.5GM/100ML (1920ML) | N200656 002 | Aug 25, 2014 |
| + | | 2.4%; 20MG/100ML; 6.8GM/100ML; 68MG/100ML; 124MG/100ML; 170MG/100ML; 105MG/100ML (2400ML) | N200656 003 | Aug 25, 2014 |

AMINO ACIDS; DEXTROSE

INJECTABLE; INJECTION

| | | | | |
|--|--------------------------------------------------------------------|-------------------|-------------|--------------|
| | CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | 2.75%; 10GM/100ML | N020734 002 | Sep 29, 1997 |
| | CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | 2.75%; 25GM/100ML | N020734 005 | Sep 29, 1997 |
| | CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | 2.75%; 5GM/100ML | N020734 001 | Sep 29, 1997 |
| | CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | 4.25%; 10GM/100ML | N020734 008 | Sep 29, 1997 |
| | CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | 4.25%; 20GM/100ML | N020734 010 | Sep 29, 1997 |
| | CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | 4.25%; 25GM/100ML | N020734 011 | Sep 29, 1997 |
| | CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | 4.25%; 5GM/100ML | N020734 007 | Sep 29, 1997 |
| | CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | 5%; 10GM/100ML | N020734 014 | Sep 29, 1997 |
| | CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | 5%; 15GM/100ML | N020734 015 | Sep 29, 1997 |
| | CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | 5%; 20GM/100ML | N020734 016 | Sep 29, 1997 |
| | CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | 5%; 25GM/100ML | N020734 017 | Sep 29, 1997 |
| | CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | 5%; 35GM/100ML | N020734 018 | Sep 29, 1997 |

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM ACETATE; POTASSIUM CHLORIDE; SODIUM ACETATE

INJECTABLE; INJECTION

FREAMINE III 8.5% W/ ELECTROLYTES

| | | | | |
|--|---------|----------------------------------------------------------------------|-------------|--------------|
| | B BRAUN | 8.5%; 110MG/100ML; 230MG/100ML; 10MG/100ML; 440MG/100ML; 690MG/100ML | N016822 007 | Jul 01, 1988 |
|--|---------|----------------------------------------------------------------------|-------------|--------------|

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 3.5% M

| | | | | |
|--|-----------------|--------------------------------------------------------|-------------|--|
| | ICU MEDICAL INC | 3.5%; 21MG/100ML; 40MG/100ML; 128MG/100ML; 234MG/100ML | N017789 003 | |
|--|-----------------|--------------------------------------------------------|-------------|--|

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

FREAMINE III 3% W/ ELECTROLYTES

| | | | | |
|--|---------|-------------------------------------------------------------------|-------------|--|
| | B BRAUN | 3%; 54MG/100ML; 40MG/100ML; 150MG/100ML; 200MG/100ML; 120MG/100ML | N016822 003 | |
|--|---------|-------------------------------------------------------------------|-------------|--|

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC

INJECTABLE; INJECTION

AMINOSYN 8.5% W/ELECTROLYTES

| | | | | |
|--|-----------------|---------------------------------------------------------|-------------|--------------|
| | ICU MEDICAL INC | 8.5%; 102MG/100ML; 487MG/100ML; 28MG/100ML; 425MG/100ML | N017673 009 | Oct 25, 2002 |
|--|-----------------|---------------------------------------------------------|-------------|--------------|

AMINOCAPROIC ACID

INJECTABLE; INJECTION

AMINOCAPROIC ACID

| | | | | |
|-----------|----------|-----------------|--------------------|--------------|
| AP | LUITPOLD | 250MG/ML | A071192 001 | Dec 01, 1987 |
|-----------|----------|-----------------|--------------------|--------------|

AMINOCAPROIC ACID IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------|-----------------|--------------------|--------------|
| AP | ! HOSPIRA | 250MG/ML | A070010 001 | Mar 09, 1987 |
|-----------|-----------|-----------------|--------------------|--------------|

SYRUP; ORAL

AMICAR

| | | | | |
|---|---------------|------------|-------------|--|
| + | CLOVER PHARMS | 1.25GM/5ML | N015230 002 | |
|---|---------------|------------|-------------|--|

PRESCRIPTION DRUG PRODUCT LIST

AMINOCAPROIC ACID

TABLET; ORAL

AMICAR

| | | | | | | |
|-----------|---|---------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | CLOVER PHARMS | <u>500MG</u> | <u>N015197</u> | <u>001</u> | |
| <u>AB</u> | + | ! | <u>1GM</u> | <u>N015197</u> | <u>002</u> | Jun 24, 2004 |

AMINOCAPROIC ACID

| | | | | | | |
|-----------|--|---------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | SUNNY PHARMTECH INC | <u>500MG</u> | <u>A209060</u> | <u>001</u> | Nov 27, 2018 |
| <u>AB</u> | | | <u>1GM</u> | <u>A209060</u> | <u>002</u> | Nov 27, 2018 |

AMINOLEVULINIC ACID HYDROCHLORIDE

FOR SOLUTION; ORAL

GLEOLAN

| | | | | | | |
|---|---|------|------------|---------|-----|--------------|
| + | ! | NXDC | 1.5GM/VIAL | N208630 | 001 | Jun 06, 2017 |
|---|---|------|------------|---------|-----|--------------|

GEL; TOPICAL

AMELUZ

| | | | | | | |
|---|---|-------------|-----|---------|-----|--------------|
| + | ! | BIOFRONTERA | 10% | N208081 | 001 | May 10, 2016 |
|---|---|-------------|-----|---------|-----|--------------|

SOLUTION; TOPICAL

LEVULAN

| | | | | | | |
|---|---|------|-----|---------|-----|--------------|
| + | ! | DUSA | 20% | N020965 | 001 | Dec 03, 1999 |
|---|---|------|-----|---------|-----|--------------|

AMINOPHYLLINE

INJECTABLE; INJECTION

AMINOPHYLLINE

| | | | | | | |
|-----------|---|----------|----------------|----------------|------------|--------------|
| <u>AP</u> | ! | HOSPIRA | <u>25MG/ML</u> | <u>A087242</u> | <u>001</u> | Oct 26, 1983 |
| <u>AP</u> | | LUITPOLD | <u>25MG/ML</u> | <u>A087600</u> | <u>001</u> | |

AMINOSALICYLIC ACID

GRANULE, DELAYED RELEASE; ORAL

PASER

| | | | | | | |
|---|--|---------|------------|---------|-----|--------------|
| ! | | JACOBUS | 4GM/PACKET | A074346 | 001 | Jun 30, 1994 |
|---|--|---------|------------|---------|-----|--------------|

AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION

AMIODARONE HYDROCHLORIDE

| | | | | | | |
|-----------|---|----------------------|----------------|----------------|------------|--------------|
| <u>AP</u> | | AUROBINDO PHARMA LTD | <u>50MG/ML</u> | <u>A204550</u> | <u>001</u> | Oct 25, 2017 |
| <u>AP</u> | ! | FRESENIUS KABI USA | <u>50MG/ML</u> | <u>A075761</u> | <u>001</u> | Oct 15, 2002 |
| <u>AP</u> | ! | GLAND PHARMA LTD | <u>50MG/ML</u> | <u>A077161</u> | <u>001</u> | Apr 20, 2005 |
| <u>AP</u> | | HIKMA FARMACEUTICA | <u>50MG/ML</u> | <u>A077234</u> | <u>001</u> | Feb 25, 2008 |
| <u>AP</u> | ! | HOSPIRA | <u>50MG/ML</u> | <u>A075955</u> | <u>001</u> | Oct 18, 2002 |
| <u>AP</u> | | HOSPIRA INC | <u>50MG/ML</u> | <u>A203884</u> | <u>001</u> | Nov 25, 2013 |
| <u>AP</u> | | | <u>50MG/ML</u> | <u>A203885</u> | <u>001</u> | Nov 25, 2013 |
| <u>AP</u> | ! | MYLAN INSTITUTIONAL | <u>50MG/ML</u> | <u>A076217</u> | <u>001</u> | Oct 15, 2002 |
| <u>AP</u> | | WOCKHARDT | <u>50MG/ML</u> | <u>A077610</u> | <u>001</u> | Oct 30, 2008 |
| <u>AP</u> | | | <u>50MG/ML</u> | <u>A077834</u> | <u>001</u> | Oct 30, 2008 |

NEXTERONE

| | | | | | | |
|---|---|-----------------|------------------------|---------|-----|--------------|
| + | ! | BAXTER HLTHCARE | 150MG/100ML (1.5MG/ML) | N022325 | 002 | Nov 16, 2010 |
|---|---|-----------------|------------------------|---------|-----|--------------|

| | | | | | | |
|---|---|--|------------------------|---------|-----|--------------|
| + | ! | | 360MG/200ML (1.8MG/ML) | N022325 | 003 | Nov 16, 2010 |
|---|---|--|------------------------|---------|-----|--------------|

TABLET; ORAL

AMIODARONE HYDROCHLORIDE

| | | | | | | |
|-----------|---|----------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | APOTEX INC | <u>200MG</u> | <u>A078578</u> | <u>001</u> | Nov 06, 2008 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>200MG</u> | <u>A204742</u> | <u>001</u> | Jun 03, 2016 |
| <u>AB</u> | | MAYNE PHARMA INC | <u>100MG</u> | <u>A075389</u> | <u>002</u> | Dec 28, 2017 |
| <u>AB</u> | | | <u>200MG</u> | <u>A075389</u> | <u>001</u> | Jan 25, 2001 |
| <u>AB</u> | | | <u>400MG</u> | <u>A075389</u> | <u>003</u> | Dec 28, 2017 |
| <u>AB</u> | | MURTY PHARMS | <u>100MG</u> | <u>A077069</u> | <u>003</u> | Oct 04, 2016 |
| <u>AB</u> | | | <u>200MG</u> | <u>A077069</u> | <u>001</u> | Apr 08, 2005 |
| <u>AB</u> | | | <u>400MG</u> | <u>A077069</u> | <u>002</u> | Apr 08, 2005 |
| <u>AB</u> | ! | SANDOZ | <u>200MG</u> | <u>A075315</u> | <u>001</u> | Dec 23, 1998 |
| <u>AB</u> | | | <u>400MG</u> | <u>A075315</u> | <u>002</u> | Jun 30, 2000 |
| <u>AB</u> | | TARO PHARM | <u>100MG</u> | <u>A075424</u> | <u>002</u> | Dec 18, 2002 |
| <u>AB</u> | | | <u>200MG</u> | <u>A075424</u> | <u>001</u> | Mar 30, 2001 |
| <u>AB</u> | | | <u>400MG</u> | <u>A076362</u> | <u>001</u> | Nov 29, 2002 |
| <u>AB</u> | | TEVA PHARMS | <u>200MG</u> | <u>A074739</u> | <u>001</u> | Nov 30, 1998 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>200MG</u> | <u>A079029</u> | <u>001</u> | Sep 16, 2008 |

PACERONE

| | | | | | | |
|-----------|--|-------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | UPSHER SMITH LABS | <u>100MG</u> | <u>A075135</u> | <u>002</u> | Apr 12, 2005 |
| <u>AB</u> | | | <u>200MG</u> | <u>A075135</u> | <u>001</u> | Apr 30, 1998 |

AMIODARONE HYDROCHLORIDE

| | | | | | | |
|--|--|------------|-------|---------|-----|--------------|
| | | TARO PHARM | 300MG | A076362 | 002 | Dec 02, 2003 |
|--|--|------------|-------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

AMITRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

AMITRIPTYLINE HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>10MG</u> | <u>A202446 001</u> | Jun 04, 2014 |
| <u>AB</u> | | <u>25MG</u> | <u>A202446 002</u> | Jun 04, 2014 |
| <u>AB</u> | | <u>50MG</u> | <u>A202446 003</u> | Jun 04, 2014 |
| <u>AB</u> | | <u>75MG</u> | <u>A202446 004</u> | Jun 04, 2014 |
| <u>AB</u> | | <u>100MG</u> | <u>A202446 005</u> | Jun 04, 2014 |
| <u>AB</u> | | <u>150MG</u> | <u>A202446 006</u> | Jun 04, 2014 |
| <u>AB</u> | MYLAN | <u>10MG</u> | <u>A086009 002</u> | |
| <u>AB</u> | | <u>25MG</u> | <u>A086009 003</u> | |
| <u>AB</u> | | <u>50MG</u> | <u>A086009 001</u> | |
| <u>AB</u> | | <u>75MG</u> | <u>A086009 004</u> | |
| <u>AB</u> | | <u>100MG</u> | <u>A086009 005</u> | |
| <u>AB</u> | | <u>150MG</u> | <u>A086009 006</u> | |
| <u>AB</u> | SANDOZ | <u>10MG</u> | <u>A085969 001</u> | |
| <u>AB</u> | ! | <u>25MG</u> | <u>A085966 001</u> | |
| <u>AB</u> | | <u>50MG</u> | <u>A085968 001</u> | |
| <u>AB</u> | | <u>75MG</u> | <u>A085971 001</u> | |
| <u>AB</u> | | <u>100MG</u> | <u>A085967 001</u> | |
| <u>AB</u> | | <u>150MG</u> | <u>A085970 001</u> | |
| <u>AB</u> | SUN PHARM INDS INC | <u>10MG</u> | <u>A089399 002</u> | Jul 14, 1987 |
| <u>AB</u> | | <u>25MG</u> | <u>A089399 001</u> | Jul 14, 1987 |
| <u>AB</u> | | <u>50MG</u> | <u>A089399 003</u> | Jul 14, 1987 |
| <u>AB</u> | | <u>75MG</u> | <u>A089399 004</u> | Jul 14, 1987 |
| <u>AB</u> | | <u>100MG</u> | <u>A089399 005</u> | Jul 14, 1987 |
| <u>AB</u> | | <u>150MG</u> | <u>A089399 006</u> | Jul 14, 1987 |
| <u>AB</u> | VINTAGE PHARMS | <u>10MG</u> | <u>A040218 001</u> | Sep 11, 1997 |
| <u>AB</u> | | <u>25MG</u> | <u>A040218 002</u> | Sep 11, 1997 |
| <u>AB</u> | | <u>50MG</u> | <u>A040218 003</u> | Sep 11, 1997 |
| <u>AB</u> | | <u>75MG</u> | <u>A040218 004</u> | Sep 11, 1997 |
| <u>AB</u> | | <u>100MG</u> | <u>A040218 005</u> | Sep 11, 1997 |
| <u>AB</u> | | <u>150MG</u> | <u>A040218 006</u> | Sep 11, 1997 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>10MG</u> | <u>A210086 001</u> | Oct 06, 2017 |
| <u>AB</u> | | <u>25MG</u> | <u>A210086 002</u> | Oct 06, 2017 |
| <u>AB</u> | | <u>50MG</u> | <u>A210086 003</u> | Oct 06, 2017 |
| <u>AB</u> | | <u>75MG</u> | <u>A210086 004</u> | Oct 06, 2017 |
| <u>AB</u> | | <u>100MG</u> | <u>A210086 005</u> | Oct 06, 2017 |
| <u>AB</u> | | <u>150MG</u> | <u>A210086 006</u> | Oct 06, 2017 |

AMITRIPTYLINE HYDROCHLORIDE; CHLORDIAZEPOXIDE

TABLET;ORAL

CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE

| | | | | |
|--|------------------|--------------------|-------------|--------------|
| | MYLAN PHARMS INC | EQ 12.5MG BASE;5MG | A071297 002 | Dec 10, 1986 |
| | ! | EQ 25MG BASE;10MG | A071297 001 | Dec 10, 1986 |

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET;ORAL

PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE

| | | | | |
|--|-------|----------|-------------|--------------|
| | MYLAN | 10MG;2MG | A071443 002 | Nov 10, 1988 |
| | | 10MG;4MG | A071443 003 | Nov 10, 1988 |
| | ! | 25MG;2MG | A071443 004 | Nov 10, 1988 |
| | ! | 25MG;4MG | A071443 005 | Nov 10, 1988 |
| | ! | 50MG;4MG | A071443 001 | Nov 10, 1988 |

AMLODIPINE BESYLATE

TABLET;ORAL

AMLODIPINE BESYLATE

| | | | | |
|-----------|------------------|----------------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>EQ 2.5MG BASE</u> | <u>A202553 001</u> | Apr 29, 2013 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A202553 002</u> | Apr 29, 2013 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A202553 003</u> | Apr 29, 2013 |
| <u>AB</u> | ALKEM | <u>EQ 2.5MG BASE</u> | <u>A078925 001</u> | May 04, 2009 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A078925 002</u> | May 04, 2009 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A078925 003</u> | May 04, 2009 |
| <u>AB</u> | APOTEX | <u>EQ 2.5MG BASE</u> | <u>A076719 001</u> | May 23, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A076719 002</u> | May 23, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A076719 003</u> | May 23, 2007 |
| <u>AB</u> | AUROBINDO PHARMA | <u>EQ 2.5MG BASE</u> | <u>A078021 001</u> | Jul 17, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A078021 002</u> | Jul 17, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A078021 003</u> | Jul 17, 2007 |
| <u>AB</u> | CHINA RESOURCES | <u>EQ 2.5MG BASE</u> | <u>A090752 003</u> | May 16, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A090752 001</u> | Apr 15, 2011 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A090752 002</u> | Apr 15, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE

TABLET; ORAL

AMLODIPINE BESYLATE

| | | | | |
|-----------|-------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | CIPLA | <u>EQ 2.5MG BASE</u> | <u>A077073 001</u> | Sep 26, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A077073 002</u> | Sep 26, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A077073 003</u> | Sep 26, 2007 |
| <u>AB</u> | EPIC PHARMA LLC | <u>EQ 2.5MG BASE</u> | <u>A078552 001</u> | Apr 08, 2009 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A078552 002</u> | Apr 08, 2009 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A078552 003</u> | Apr 08, 2009 |
| <u>AB</u> | HEBEI CHANGSHAN | <u>EQ 2.5MG BASE</u> | <u>A076692 001</u> | Jul 20, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A076692 002</u> | Jul 20, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A076692 003</u> | Jul 20, 2007 |
| <u>AB</u> | HIKMA PHARMS | <u>EQ 2.5MG BASE</u> | <u>A077771 001</u> | Apr 12, 2011 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A077771 002</u> | Apr 12, 2011 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A077771 003</u> | Apr 12, 2011 |
| <u>AB</u> | INVAGEN PHARMS | <u>EQ 2.5MG BASE</u> | <u>A077955 001</u> | Aug 28, 2007 |
| <u>AB</u> | | <u>EQ 2.5MG BASE</u> | <u>A206367 001</u> | Dec 10, 2015 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A077955 002</u> | Aug 28, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A206367 002</u> | Dec 10, 2015 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A077955 003</u> | Aug 28, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A206367 003</u> | Dec 10, 2015 |
| <u>AB</u> | LUPIN | <u>EQ 2.5MG BASE</u> | <u>A078043 001</u> | Jul 12, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A078043 002</u> | Jul 12, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A078043 003</u> | Jul 12, 2007 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>EQ 5MG BASE</u> | <u>A201380 001</u> | Apr 13, 2012 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A201380 002</u> | Apr 13, 2012 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 2.5MG BASE</u> | <u>A076418 001</u> | Oct 03, 2005 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A076418 002</u> | Oct 03, 2005 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A076418 003</u> | Oct 03, 2005 |
| <u>AB</u> | ORCHID HLTHCARE | <u>EQ 2.5MG BASE</u> | <u>A078453 001</u> | Jul 02, 2009 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A078453 002</u> | Jul 02, 2009 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A078453 003</u> | Jul 02, 2009 |
| <u>AB</u> | OXFORD PHARMS | <u>EQ 2.5MG BASE</u> | <u>A078414 001</u> | Apr 07, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A078414 002</u> | Apr 07, 2010 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A078414 003</u> | Apr 07, 2010 |
| <u>AB</u> | POLYGEN PHARMS | <u>EQ 2.5MG BASE</u> | <u>A207821 001</u> | Jul 11, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A207821 002</u> | Jul 11, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A207821 003</u> | Jul 11, 2016 |
| <u>AB</u> | STRIDES VIVIMED | <u>EQ 2.5MG BASE</u> | <u>A077516 001</u> | Jul 11, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A077516 002</u> | Jul 11, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A077516 003</u> | Jul 11, 2007 |
| <u>AB</u> | SUN PHARM INDS INC | <u>EQ 2.5MG BASE</u> | <u>A078231 001</u> | Nov 30, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A078231 002</u> | Nov 30, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A078231 003</u> | Nov 30, 2007 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>EQ 2.5MG BASE</u> | <u>A077974 001</u> | Jul 09, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A077974 002</u> | Jul 09, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A077974 003</u> | Jul 09, 2007 |
| <u>AB</u> | TEVA | <u>EQ 2.5MG BASE</u> | <u>A076846 001</u> | Jun 28, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A076846 002</u> | Jun 28, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A076846 003</u> | Jun 28, 2007 |
| <u>AB</u> | TORRENT PHARMS | <u>EQ 2.5MG BASE</u> | <u>A078573 001</u> | Sep 22, 2008 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A078573 002</u> | Sep 22, 2008 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A078573 003</u> | Sep 22, 2008 |
| <u>AB</u> | UNICHEM LABS LTD | <u>EQ 2.5MG BASE</u> | <u>A203245 001</u> | Oct 21, 2013 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A203245 002</u> | Oct 21, 2013 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A203245 003</u> | Oct 21, 2013 |
| <u>AB</u> | UPSHER SMITH LABS | <u>EQ 2.5MG BASE</u> | <u>A077759 001</u> | Jul 09, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A077759 002</u> | Jul 09, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A077759 003</u> | Jul 09, 2007 |
| <u>AB</u> | WATSON LABS | <u>EQ 2.5MG BASE</u> | <u>A077671 001</u> | Jul 19, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A077671 002</u> | Jul 19, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A077671 003</u> | Jul 19, 2007 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>EQ 2.5MG BASE</u> | <u>A077262 001</u> | Jul 09, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A077262 002</u> | Jul 09, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A077262 003</u> | Jul 09, 2007 |
| <u>AB</u> | WOCKHARDT | <u>EQ 2.5MG BASE</u> | <u>A078500 001</u> | Sep 06, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A078500 002</u> | Sep 06, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A078500 003</u> | Sep 06, 2007 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>EQ 2.5MG BASE</u> | <u>A078226 001</u> | Jul 09, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A078226 002</u> | Jul 09, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A078226 003</u> | Jul 09, 2007 |

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE

TABLET; ORAL

NORVASC

| | | | | | |
|-----------|---|--------|----------------------|--------------------|--------------|
| <u>AB</u> | + | PFIZER | <u>EQ 2.5MG BASE</u> | <u>N019787 001</u> | Jul 31, 1992 |
| <u>AB</u> | + | | <u>EQ 5MG BASE</u> | <u>N019787 002</u> | Jul 31, 1992 |
| <u>AB</u> | + | | <u>EQ 10MG BASE</u> | <u>N019787 003</u> | Jul 31, 1992 |

AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM

TABLET; ORAL

AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM

| | | | | | |
|---------------|---|-------------------------|-----------------------------------|--------------------|--------------|
| <u>AB</u> | | DR REDDYS LABS LTD | <u>EQ 2.5MG BASE;EQ 10MG BASE</u> | <u>A203874 001</u> | Mar 07, 2014 |
| <u>AB</u> | | | <u>EQ 2.5MG BASE;EQ 20MG BASE</u> | <u>A203874 002</u> | Mar 07, 2014 |
| <u>AB</u> | | | <u>EQ 2.5MG BASE;EQ 40MG BASE</u> | <u>A203874 003</u> | Mar 07, 2014 |
| <u>AB</u> | | | <u>EQ 5MG BASE;EQ 10MG BASE</u> | <u>A203874 004</u> | Mar 07, 2014 |
| <u>AB</u> | | | <u>EQ 5MG BASE;EQ 20MG BASE</u> | <u>A203874 005</u> | Mar 07, 2014 |
| <u>AB</u> | | | <u>EQ 5MG BASE;EQ 40MG BASE</u> | <u>A203874 006</u> | Mar 07, 2014 |
| <u>AB</u> | | | <u>EQ 5MG BASE;EQ 80MG BASE</u> | <u>A203874 007</u> | Mar 07, 2014 |
| <u>AB</u> | | | <u>EQ 10MG BASE;EQ 10MG BASE</u> | <u>A203874 008</u> | Mar 07, 2014 |
| <u>AB</u> | | | <u>EQ 10MG BASE;EQ 20MG BASE</u> | <u>A203874 009</u> | Mar 07, 2014 |
| <u>AB</u> | | | <u>EQ 10MG BASE;EQ 40MG BASE</u> | <u>A203874 010</u> | Mar 07, 2014 |
| <u>AB</u> | | | <u>EQ 10MG BASE;EQ 80MG BASE</u> | <u>A203874 011</u> | Mar 07, 2014 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>EQ 2.5MG BASE;EQ 10MG BASE</u> | <u>A200465 001</u> | Nov 29, 2013 |
| <u>AB</u> | | | <u>EQ 2.5MG BASE;EQ 20MG BASE</u> | <u>A200465 002</u> | Nov 29, 2013 |
| <u>AB</u> | | | <u>EQ 2.5MG BASE;EQ 40MG BASE</u> | <u>A200465 003</u> | Nov 29, 2013 |
| <u>AB</u> | | | <u>EQ 5MG BASE;EQ 10MG BASE</u> | <u>A200465 004</u> | Nov 29, 2013 |
| <u>AB</u> | | | <u>EQ 5MG BASE;EQ 20MG BASE</u> | <u>A200465 005</u> | Nov 29, 2013 |
| <u>AB</u> | | | <u>EQ 5MG BASE;EQ 40MG BASE</u> | <u>A200465 006</u> | Nov 29, 2013 |
| <u>AB</u> | | | <u>EQ 5MG BASE;EQ 80MG BASE</u> | <u>A200465 007</u> | Nov 29, 2013 |
| <u>AB</u> | | | <u>EQ 10MG BASE;EQ 10MG BASE</u> | <u>A200465 008</u> | Nov 29, 2013 |
| <u>AB</u> | | | <u>EQ 10MG BASE;EQ 20MG BASE</u> | <u>A200465 009</u> | Nov 29, 2013 |
| <u>AB</u> | | | <u>EQ 10MG BASE;EQ 40MG BASE</u> | <u>A200465 010</u> | Nov 29, 2013 |
| <u>AB</u> | | | <u>EQ 10MG BASE;EQ 80MG BASE</u> | <u>A200465 011</u> | Nov 29, 2013 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>EQ 2.5MG BASE;EQ 10MG BASE</u> | <u>A207762 001</u> | Jan 11, 2019 |
| <u>AB</u> | | | <u>EQ 2.5MG BASE;EQ 20MG BASE</u> | <u>A207762 002</u> | Jan 11, 2019 |
| <u>AB</u> | | | <u>EQ 2.5MG BASE;EQ 40MG BASE</u> | <u>A207762 003</u> | Jan 11, 2019 |
| <u>AB</u> | | | <u>EQ 5MG BASE;EQ 10MG BASE</u> | <u>A207762 004</u> | Jan 11, 2019 |
| <u>AB</u> | | | <u>EQ 5MG BASE;EQ 20MG BASE</u> | <u>A207762 005</u> | Jan 11, 2019 |
| <u>AB</u> | | | <u>EQ 5MG BASE;EQ 40MG BASE</u> | <u>A207762 006</u> | Jan 11, 2019 |
| <u>AB</u> | | | <u>EQ 5MG BASE;EQ 80MG BASE</u> | <u>A207762 007</u> | Jan 11, 2019 |
| <u>AB</u> | | | <u>EQ 10MG BASE;EQ 10MG BASE</u> | <u>A207762 008</u> | Jan 11, 2019 |
| <u>AB</u> | | | <u>EQ 10MG BASE;EQ 20MG BASE</u> | <u>A207762 009</u> | Jan 11, 2019 |
| <u>AB</u> | | | <u>EQ 10MG BASE;EQ 40MG BASE</u> | <u>A207762 010</u> | Jan 11, 2019 |
| <u>AB</u> | | | <u>EQ 10MG BASE;EQ 80MG BASE</u> | <u>A207762 011</u> | Jan 11, 2019 |
| <u>CADUET</u> | | | | | |
| <u>AB</u> | + | PFIZER | <u>EQ 2.5MG BASE;EQ 10MG BASE</u> | <u>N021540 009</u> | Jul 29, 2004 |
| <u>AB</u> | + | | <u>EQ 2.5MG BASE;EQ 20MG BASE</u> | <u>N021540 010</u> | Jul 29, 2004 |
| <u>AB</u> | + | | <u>EQ 2.5MG BASE;EQ 40MG BASE</u> | <u>N021540 011</u> | Jul 29, 2004 |
| <u>AB</u> | + | | <u>EQ 5MG BASE;EQ 10MG BASE</u> | <u>N021540 001</u> | Jan 30, 2004 |
| <u>AB</u> | + | | <u>EQ 5MG BASE;EQ 20MG BASE</u> | <u>N021540 002</u> | Jan 30, 2004 |
| <u>AB</u> | + | | <u>EQ 5MG BASE;EQ 40MG BASE</u> | <u>N021540 003</u> | Jan 30, 2004 |
| <u>AB</u> | + | | <u>EQ 5MG BASE;EQ 80MG BASE</u> | <u>N021540 004</u> | Jan 30, 2004 |
| <u>AB</u> | + | | <u>EQ 10MG BASE;EQ 10MG BASE</u> | <u>N021540 005</u> | Jan 30, 2004 |
| <u>AB</u> | + | | <u>EQ 10MG BASE;EQ 20MG BASE</u> | <u>N021540 006</u> | Jan 30, 2004 |
| <u>AB</u> | + | | <u>EQ 10MG BASE;EQ 40MG BASE</u> | <u>N021540 007</u> | Jan 30, 2004 |
| <u>AB</u> | + | | <u>EQ 10MG BASE;EQ 80MG BASE</u> | <u>N021540 008</u> | Jan 30, 2004 |

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE; ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

| | | | | | |
|-----------|--|-------------------------|---------------------------|--------------------|--------------|
| <u>AB</u> | | APOTEX INC | <u>EQ 2.5MG BASE;10MG</u> | <u>A091431 001</u> | Dec 30, 2013 |
| <u>AB</u> | | | <u>EQ 5MG BASE;10MG</u> | <u>A091431 002</u> | Dec 30, 2013 |
| <u>AB</u> | | | <u>EQ 5MG BASE;20MG</u> | <u>A091431 003</u> | Dec 30, 2013 |
| <u>AB</u> | | | <u>EQ 5MG BASE;40MG</u> | <u>A091431 004</u> | Dec 30, 2013 |
| <u>AB</u> | | | <u>EQ 10MG BASE;20MG</u> | <u>A091431 005</u> | Dec 30, 2013 |
| <u>AB</u> | | | <u>EQ 10MG BASE;40MG</u> | <u>A091431 006</u> | Dec 30, 2013 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>EQ 2.5MG BASE;10MG</u> | <u>A202239 001</u> | Sep 05, 2012 |
| <u>AB</u> | | | <u>EQ 5MG BASE;10MG</u> | <u>A202239 002</u> | Sep 05, 2012 |
| <u>AB</u> | | | <u>EQ 5MG BASE;20MG</u> | <u>A202239 003</u> | Sep 05, 2012 |
| <u>AB</u> | | | <u>EQ 5MG BASE;40MG</u> | <u>A202239 004</u> | Sep 05, 2012 |
| <u>AB</u> | | | <u>EQ 10MG BASE;20MG</u> | <u>A202239 005</u> | Sep 05, 2012 |
| <u>AB</u> | | | <u>EQ 10MG BASE;40MG</u> | <u>A202239 006</u> | Sep 05, 2012 |
| <u>AB</u> | | CIPLA | <u>EQ 2.5MG BASE;10MG</u> | <u>A077215 001</u> | Dec 07, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE;ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

| | | | | |
|---------------|--------------------|---------------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 5MG BASE;10MG</u> | <u>A077215 002</u> | Dec 07, 2018 |
| <u>AB</u> | | <u>EQ 5MG BASE;20MG</u> | <u>A077215 003</u> | Dec 07, 2018 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A077215 004</u> | Dec 07, 2018 |
| <u>AB</u> | DR REDDYS LABS INC | <u>EQ 2.5MG BASE;10MG</u> | <u>A077183 001</u> | Apr 15, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE;10MG</u> | <u>A077183 002</u> | Apr 15, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE;20MG</u> | <u>A077183 003</u> | Apr 15, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A090149 001</u> | Jul 05, 2011 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A077183 004</u> | Apr 15, 2010 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A090149 002</u> | Jul 05, 2011 |
| <u>AB</u> | LUPIN PHARMS | <u>EQ 2.5MG BASE;10MG</u> | <u>A078466 001</u> | Feb 05, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE;10MG</u> | <u>A078466 002</u> | Feb 05, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE;20MG</u> | <u>A078466 003</u> | Feb 05, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A078466 005</u> | Jul 05, 2011 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A078466 004</u> | Feb 05, 2010 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A078466 006</u> | Jul 05, 2011 |
| <u>AB</u> | MYLAN | <u>EQ 2.5MG BASE;10MG</u> | <u>A077375 001</u> | May 21, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE;10MG</u> | <u>A077375 002</u> | May 21, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE;20MG</u> | <u>A077375 003</u> | May 21, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A079047 001</u> | Jul 05, 2011 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A077375 004</u> | May 21, 2010 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A079047 002</u> | Jul 05, 2011 |
| <u>AB</u> | PAR PHARM | <u>EQ 2.5MG BASE;10MG</u> | <u>A078381 001</u> | Jul 29, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE;10MG</u> | <u>A078381 002</u> | Jul 29, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE;20MG</u> | <u>A078381 003</u> | Jul 29, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A078381 005</u> | Jul 29, 2010 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A078381 004</u> | Jul 29, 2010 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A078381 006</u> | Jul 29, 2010 |
| <u>AB</u> | TEVA PHARMS | <u>EQ 2.5MG BASE;10MG</u> | <u>A077179 001</u> | May 18, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE;10MG</u> | <u>A077179 002</u> | May 18, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE;20MG</u> | <u>A077179 003</u> | May 18, 2007 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A077179 005</u> | Jul 05, 2011 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A077179 004</u> | May 18, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A077179 006</u> | Jul 05, 2011 |
| <u>AB</u> | WATSON LABS | <u>EQ 2.5MG BASE;10MG</u> | <u>A077890 001</u> | Oct 14, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE;10MG</u> | <u>A077890 002</u> | Oct 14, 2010 |
| <u>AB</u> | | <u>EQ 5MG BASE;20MG</u> | <u>A077890 003</u> | Oct 14, 2010 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A077890 004</u> | Oct 14, 2010 |
| <u>AB</u> | WATSON LABS INC | <u>EQ 5MG BASE;40MG</u> | <u>A090364 001</u> | Jul 05, 2011 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A090364 002</u> | Jul 05, 2011 |
| <u>LOTREL</u> | | | | |
| <u>AB</u> | + | <u>EQ 2.5MG BASE;10MG</u> | <u>N020364 002</u> | Mar 03, 1995 |
| <u>AB</u> | + | <u>EQ 5MG BASE;10MG</u> | <u>N020364 003</u> | Mar 03, 1995 |
| <u>AB</u> | + | <u>EQ 5MG BASE;20MG</u> | <u>N020364 004</u> | Mar 03, 1995 |
| <u>AB</u> | + | <u>EQ 5MG BASE;40MG</u> | <u>N020364 007</u> | Apr 11, 2006 |
| <u>AB</u> | + | <u>EQ 10MG BASE;20MG</u> | <u>N020364 005</u> | Jun 20, 2002 |
| <u>AB</u> | + | <u>EQ 10MG BASE;40MG</u> | <u>N020364 006</u> | Apr 11, 2006 |

AMLODIPINE BESYLATE; CELECOXIB

TABLET;ORAL

CONSENSI

| | | | | |
|---|------------------|---------------------|-------------|--------------|
| + | KITOV PHARMS LTD | EQ 2.5MG BASE;200MG | N210045 001 | May 31, 2018 |
| + | | EQ 5MG BASE;200MG | N210045 002 | May 31, 2018 |
| + | | EQ 10MG BASE;200MG | N210045 003 | May 31, 2018 |

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET;ORAL

OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE

| | | | | |
|-----------|--------------------|---------------------------------|--------------------|--------------|
| <u>AB</u> | PAR PHARM INC | <u>EQ 5MG BASE;12.5MG;20MG</u> | <u>A206137 001</u> | Oct 26, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE;12.5MG;40MG</u> | <u>A206137 002</u> | Oct 26, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE;25MG;40MG</u> | <u>A206137 003</u> | Oct 26, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;12.5MG;40MG</u> | <u>A206137 004</u> | Oct 26, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;25MG;40MG</u> | <u>A206137 005</u> | Oct 26, 2016 |
| <u>AB</u> | TEVA PHARMS USA | <u>EQ 5MG BASE;12.5MG;20MG</u> | <u>A202491 001</u> | Nov 03, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE;12.5MG;40MG</u> | <u>A202491 002</u> | Nov 03, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE;25MG;40MG</u> | <u>A202491 003</u> | Nov 03, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;12.5MG;40MG</u> | <u>A202491 004</u> | Nov 03, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;25MG;40MG</u> | <u>A202491 005</u> | Nov 03, 2016 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>EQ 5MG BASE;12.5MG;20MG</u> | <u>A203580 001</u> | Oct 26, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE;12.5MG;40MG</u> | <u>A203580 002</u> | Oct 26, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE;25MG;40MG</u> | <u>A203580 003</u> | Oct 26, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE

| | | | | |
|------------------|---|---------------------------------|---------------------------------|---------------------------------|
| <u>AB</u> | | <u>EQ 10MG BASE;12.5MG;40MG</u> | <u>A203580 004</u> | Oct 26, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;25MG;40MG</u> | <u>A203580 005</u> | Oct 26, 2016 |
| <u>TRIBENZOR</u> | | | | |
| <u>AB</u> | + | DAIICHI SANKYO | <u>EQ 5MG BASE;12.5MG;20MG</u> | <u>N200175 001</u> Jul 23, 2010 |
| <u>AB</u> | + | | <u>EQ 5MG BASE;12.5MG;40MG</u> | <u>N200175 002</u> Jul 23, 2010 |
| <u>AB</u> | + | | <u>EQ 5MG BASE;25MG;40MG</u> | <u>N200175 003</u> Jul 23, 2010 |
| <u>AB</u> | + | | <u>EQ 10MG BASE;12.5MG;40MG</u> | <u>N200175 004</u> Jul 23, 2010 |
| <u>AB</u> | + | | <u>EQ 10MG BASE;25MG;40MG</u> | <u>N200175 005</u> Jul 23, 2010 |

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE

| | | | | |
|--------------------|----------------------|--------------------------|--------------------------|---------------------------------|
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>5MG;12.5MG;160MG</u> | <u>A206180 001</u> | Dec 19, 2017 |
| <u>AB</u> | | <u>5MG;25MG;160MG</u> | <u>A206180 002</u> | Dec 19, 2017 |
| <u>AB</u> | | <u>10MG;12.5MG;160MG</u> | <u>A206180 003</u> | Dec 19, 2017 |
| <u>AB</u> | | <u>10MG;25MG;160MG</u> | <u>A206180 004</u> | Dec 19, 2017 |
| <u>AB</u> | | <u>10MG;25MG;320MG</u> | <u>A206180 005</u> | Dec 19, 2017 |
| <u>AB</u> | LUPIN LTD | <u>5MG;12.5MG;160MG</u> | <u>A200797 001</u> | Jun 03, 2015 |
| <u>AB</u> | | <u>5MG;25MG;160MG</u> | <u>A200797 002</u> | Jun 03, 2015 |
| <u>AB</u> | | <u>10MG;12.5MG;160MG</u> | <u>A200797 003</u> | Jun 03, 2015 |
| <u>AB</u> | | <u>10MG;25MG;160MG</u> | <u>A200797 004</u> | Jun 03, 2015 |
| <u>AB</u> | | <u>10MG;25MG;320MG</u> | <u>A200797 005</u> | Jun 03, 2015 |
| <u>AB</u> | PAR PHARM | <u>5MG;12.5MG;160MG</u> | <u>A201087 001</u> | Jun 01, 2015 |
| <u>AB</u> | | <u>5MG;25MG;160MG</u> | <u>A201087 002</u> | Jun 01, 2015 |
| <u>AB</u> | | <u>10MG;12.5MG;160MG</u> | <u>A201087 003</u> | Jun 01, 2015 |
| <u>AB</u> | | <u>10MG;25MG;160MG</u> | <u>A201087 004</u> | Jun 01, 2015 |
| <u>AB</u> | | <u>10MG;25MG;320MG</u> | <u>A201087 005</u> | Jun 01, 2015 |
| <u>AB</u> | TEVA PHARMS | <u>5MG;12.5MG;160MG</u> | <u>A200435 001</u> | Sep 25, 2012 |
| <u>AB</u> | | <u>5MG;25MG;160MG</u> | <u>A200435 002</u> | Sep 25, 2012 |
| <u>AB</u> | | <u>10MG;12.5MG;160MG</u> | <u>A200435 005</u> | Sep 25, 2012 |
| <u>AB</u> | | <u>10MG;25MG;160MG</u> | <u>A200435 003</u> | Sep 25, 2012 |
| <u>AB</u> | | <u>10MG;25MG;320MG</u> | <u>A200435 004</u> | Sep 25, 2012 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>5MG;12.5MG;160MG</u> | <u>A201593 001</u> | Jun 03, 2015 |
| <u>AB</u> | | <u>5MG;25MG;160MG</u> | <u>A201593 002</u> | Jun 03, 2015 |
| <u>AB</u> | | <u>10MG;12.5MG;160MG</u> | <u>A201593 003</u> | Jun 03, 2015 |
| <u>AB</u> | | <u>10MG;25MG;160MG</u> | <u>A201593 004</u> | Jun 03, 2015 |
| <u>AB</u> | | <u>10MG;25MG;320MG</u> | <u>A201593 005</u> | Jun 03, 2015 |
| <u>EXFORGE HCT</u> | | | | |
| <u>AB</u> | + | NOVARTIS | <u>5MG;12.5MG;160MG</u> | <u>N022314 001</u> Apr 30, 2009 |
| <u>AB</u> | + | | <u>5MG;25MG;160MG</u> | <u>N022314 002</u> Apr 30, 2009 |
| <u>AB</u> | + | | <u>10MG;12.5MG;160MG</u> | <u>N022314 003</u> Apr 30, 2009 |
| <u>AB</u> | + | | <u>10MG;25MG;160MG</u> | <u>N022314 004</u> Apr 30, 2009 |
| <u>AB</u> | + | | <u>10MG;25MG;320MG</u> | <u>N022314 005</u> Apr 30, 2009 |

AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

AMLODIPINE AND OLMESARTAN MEDOXOMIL

| | | | | |
|-----------|----------------------|--------------------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE INC | <u>EQ 5MG BASE;20MG</u> | <u>A209600 001</u> | Aug 30, 2018 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A209600 003</u> | Aug 30, 2018 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A209600 002</u> | Aug 30, 2018 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A209600 004</u> | Aug 30, 2018 |
| <u>AB</u> | AJANTA PHARMA LTD | <u>EQ 5MG BASE;20MG</u> | <u>A207216 001</u> | Oct 28, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A207216 002</u> | Oct 28, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A207216 003</u> | Oct 28, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A207216 004</u> | Oct 28, 2016 |
| <u>AB</u> | ALEMBIC PHARMS LTD | <u>EQ 5MG BASE;20MG</u> | <u>A207073 001</u> | Jul 17, 2017 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A207073 002</u> | Jul 17, 2017 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A207073 003</u> | Jul 17, 2017 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A207073 004</u> | Jul 17, 2017 |
| <u>AB</u> | ALKEM LABS LTD | <u>EQ 5MG BASE;20MG</u> | <u>A209042 001</u> | Aug 14, 2017 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A209042 002</u> | Aug 14, 2017 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A209042 003</u> | Aug 14, 2017 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A209042 004</u> | Aug 14, 2017 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 5MG BASE;20MG</u> | <u>A206906 001</u> | May 15, 2017 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A206906 002</u> | May 15, 2017 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A206906 003</u> | May 15, 2017 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A206906 004</u> | May 15, 2017 |
| <u>AB</u> | GLENMARK PHARMS LTD | <u>EQ 5MG BASE;20MG</u> | <u>A207807 001</u> | Jul 05, 2017 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A207807 002</u> | Jul 05, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

AMLODIPINE AND OLMESARTAN MEDOXOMIL

| | | | | |
|-------------|-------------------------|--------------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A207807 003</u> | Jul 05, 2017 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A207807 004</u> | Jul 05, 2017 |
| <u>AB</u> | JUBILANT GENERICS | <u>EQ 5MG BASE;20MG</u> | <u>A207450 001</u> | May 15, 2017 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A207450 002</u> | May 15, 2017 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A207450 003</u> | May 15, 2017 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A207450 004</u> | May 15, 2017 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>EQ 5MG BASE;20MG</u> | <u>A206884 001</u> | Oct 26, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A206884 003</u> | Oct 26, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A206884 002</u> | Oct 26, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A206884 004</u> | Oct 26, 2016 |
| <u>AB</u> | MICRO LABS | <u>EQ 5MG BASE;20MG</u> | <u>A207435 001</u> | Nov 02, 2017 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A207435 002</u> | Nov 02, 2017 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A207435 003</u> | Nov 02, 2017 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A207435 004</u> | Nov 02, 2017 |
| <u>AB</u> | SCIEGEN PHARMS INC | <u>EQ 5MG BASE;20MG</u> | <u>A209010 001</u> | Dec 03, 2018 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A209010 002</u> | Dec 03, 2018 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A209010 003</u> | Dec 03, 2018 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A209010 004</u> | Dec 03, 2018 |
| <u>AB</u> | TEVA PHARMS USA | <u>EQ 5MG BASE;20MG</u> | <u>A091154 001</u> | Oct 26, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A091154 002</u> | Oct 26, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A091154 003</u> | Oct 26, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A091154 004</u> | Oct 26, 2016 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>EQ 5MG BASE;20MG</u> | <u>A202933 001</u> | Nov 25, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A202933 002</u> | Nov 25, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A202933 003</u> | Nov 25, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A202933 004</u> | Nov 25, 2016 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 5MG BASE;20MG</u> | <u>A207771 001</u> | Sep 22, 2017 |
| <u>AB</u> | | <u>EQ 5MG BASE;40MG</u> | <u>A207771 002</u> | Sep 22, 2017 |
| <u>AB</u> | | <u>EQ 10MG BASE;20MG</u> | <u>A207771 003</u> | Sep 22, 2017 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A207771 004</u> | Sep 22, 2017 |
| <u>AZOR</u> | | | | |
| <u>AB</u> | + DAIICHI SANKYO | <u>EQ 5MG BASE;20MG</u> | <u>N022100 001</u> | Sep 26, 2007 |
| <u>AB</u> | + | <u>EQ 5MG BASE;40MG</u> | <u>N022100 002</u> | Sep 26, 2007 |
| <u>AB</u> | + | <u>EQ 10MG BASE;20MG</u> | <u>N022100 003</u> | Sep 26, 2007 |
| <u>AB</u> | +! | <u>EQ 10MG BASE;40MG</u> | <u>N022100 004</u> | Sep 26, 2007 |

AMLODIPINE BESYLATE; PERINDOPRIL ARGININE

TABLET; ORAL

PRESTALIA

| | | | | |
|----|----------------|---------------------|-------------|--------------|
| + | MARINA BIOTECH | EQ 2.5MG BASE;3.5MG | N205003 001 | Jan 21, 2015 |
| + | | EQ 5MG BASE;7MG | N205003 002 | Jan 21, 2015 |
| +! | | EQ 10MG BASE;14MG | N205003 003 | Jan 21, 2015 |

AMLODIPINE BESYLATE; TELMISARTAN

TABLET; ORAL

TELMISARTAN AND AMLODIPINE

| | | | | |
|----------------|---------------------------|--------------------------|--------------------|--------------|
| <u>AB</u> | ALEMbic PHARMS LTD | <u>EQ 5MG BASE;40MG</u> | <u>A205234 001</u> | Nov 17, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE;80MG</u> | <u>A205234 003</u> | Nov 17, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A205234 002</u> | Nov 17, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE;80MG</u> | <u>A205234 004</u> | Nov 17, 2016 |
| <u>AB</u> | LUPIN LTD | <u>EQ 5MG BASE;40MG</u> | <u>A201586 001</u> | Jan 08, 2014 |
| <u>AB</u> | | <u>EQ 5MG BASE;80MG</u> | <u>A201586 003</u> | Jan 08, 2014 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A201586 002</u> | Jan 08, 2014 |
| <u>AB</u> | | <u>EQ 10MG BASE;80MG</u> | <u>A201586 004</u> | Jan 08, 2014 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 5MG BASE;40MG</u> | <u>A202516 001</u> | Aug 26, 2014 |
| <u>AB</u> | | <u>EQ 5MG BASE;80MG</u> | <u>A202516 003</u> | Aug 26, 2014 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A202516 002</u> | Aug 26, 2014 |
| <u>AB</u> | | <u>EQ 10MG BASE;80MG</u> | <u>A202516 004</u> | Aug 26, 2014 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>EQ 5MG BASE;40MG</u> | <u>A202517 001</u> | Jan 08, 2014 |
| <u>AB</u> | | <u>EQ 5MG BASE;80MG</u> | <u>A202517 003</u> | Jan 08, 2014 |
| <u>AB</u> | | <u>EQ 10MG BASE;40MG</u> | <u>A202517 002</u> | Jan 08, 2014 |
| <u>AB</u> | | <u>EQ 10MG BASE;80MG</u> | <u>A202517 004</u> | Jan 08, 2014 |
| <u>TWYNSTA</u> | | | | |
| <u>AB</u> | + BOEHRINGER INGELHEIM | <u>EQ 5MG BASE;40MG</u> | <u>N022401 001</u> | Oct 16, 2009 |
| <u>AB</u> | + | <u>EQ 5MG BASE;80MG</u> | <u>N022401 003</u> | Oct 16, 2009 |
| <u>AB</u> | + | <u>EQ 10MG BASE;40MG</u> | <u>N022401 002</u> | Oct 16, 2009 |
| <u>AB</u> | +! | <u>EQ 10MG BASE;80MG</u> | <u>N022401 004</u> | Oct 16, 2009 |

PRESCRIPTION DRUG PRODUCT LIST

AMLODIPINE BESYLATE; VALSARTAN

TABLET; ORAL

AMLODIPINE BESYLATE AND VALSARTAN

| | | | | |
|----------------|----------------------|---------------------------|--------------------|--------------|
| <u>AB</u> | ALEMIC PHARMS LTD | <u>EQ 5MG BASE:160MG</u> | <u>A202713 001</u> | Apr 03, 2015 |
| <u>AB</u> | | <u>EQ 5MG BASE:320MG</u> | <u>A202713 003</u> | Apr 03, 2015 |
| <u>AB</u> | | <u>EQ 10MG BASE:160MG</u> | <u>A202713 002</u> | Apr 03, 2015 |
| <u>AB</u> | | <u>EQ 10MG BASE:320MG</u> | <u>A202713 004</u> | Apr 03, 2015 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 5MG BASE:160MG</u> | <u>A206512 001</u> | Apr 22, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE:320MG</u> | <u>A206512 002</u> | Apr 22, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE:160MG</u> | <u>A206512 003</u> | Apr 22, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE:320MG</u> | <u>A206512 004</u> | Apr 22, 2016 |
| <u>AB</u> | INVAGEN PHARMS | <u>EQ 5MG BASE:160MG</u> | <u>A205137 001</u> | Sep 16, 2016 |
| <u>AB</u> | | <u>EQ 5MG BASE:320MG</u> | <u>A205137 003</u> | Sep 16, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE:160MG</u> | <u>A205137 002</u> | Sep 16, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE:320MG</u> | <u>A205137 004</u> | Sep 16, 2016 |
| <u>AB</u> | LUPIN | <u>EQ 5MG BASE:160MG</u> | <u>A090245 001</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 5MG BASE:320MG</u> | <u>A090245 003</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 10MG BASE:160MG</u> | <u>A090245 002</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 10MG BASE:320MG</u> | <u>A090245 004</u> | Mar 30, 2015 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 5MG BASE:160MG</u> | <u>A090483 001</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 5MG BASE:320MG</u> | <u>A090483 003</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 10MG BASE:160MG</u> | <u>A090483 002</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 10MG BASE:320MG</u> | <u>A090483 004</u> | Mar 30, 2015 |
| <u>AB</u> | NOVEL LABS INC | <u>EQ 5MG BASE:160MG</u> | <u>A202829 001</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 5MG BASE:320MG</u> | <u>A202829 003</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 10MG BASE:160MG</u> | <u>A202829 002</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 10MG BASE:320MG</u> | <u>A202829 004</u> | Mar 30, 2015 |
| <u>AB</u> | PAR PHARM INC | <u>EQ 5MG BASE:160MG</u> | <u>A090011 001</u> | Mar 28, 2013 |
| <u>AB</u> | | <u>EQ 5MG BASE:320MG</u> | <u>A090011 003</u> | Mar 28, 2013 |
| <u>AB</u> | | <u>EQ 10MG BASE:160MG</u> | <u>A090011 002</u> | Mar 28, 2013 |
| <u>AB</u> | | <u>EQ 10MG BASE:320MG</u> | <u>A090011 004</u> | Mar 28, 2013 |
| <u>AB</u> | TEVA PHARMS USA | <u>EQ 5MG BASE:160MG</u> | <u>A091235 001</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 5MG BASE:320MG</u> | <u>A091235 003</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 10MG BASE:160MG</u> | <u>A091235 002</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 10MG BASE:320MG</u> | <u>A091235 004</u> | Mar 30, 2015 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>EQ 5MG BASE:160MG</u> | <u>A202377 001</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 5MG BASE:320MG</u> | <u>A202377 002</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 10MG BASE:160MG</u> | <u>A202377 003</u> | Mar 30, 2015 |
| <u>AB</u> | | <u>EQ 10MG BASE:320MG</u> | <u>A202377 004</u> | Mar 30, 2015 |
| <u>EXFORGE</u> | | | | |
| <u>AB</u> | + NOVARTIS | <u>EQ 5MG BASE:160MG</u> | <u>N021990 002</u> | Jun 20, 2007 |
| <u>AB</u> | + | <u>EQ 5MG BASE:320MG</u> | <u>N021990 004</u> | Jun 20, 2007 |
| <u>AB</u> | +! | <u>EQ 10MG BASE:160MG</u> | <u>N021990 003</u> | Jun 20, 2007 |
| <u>AB</u> | +! | <u>EQ 10MG BASE:320MG</u> | <u>N021990 005</u> | Jun 20, 2007 |

AMMONIA N-13

INJECTABLE; INTRAVENOUS

AMMONIA N 13

| | | | | |
|-----------|---------------------|-------------------------------------------------|--------------------|--------------|
| <u>AP</u> | 3D IMAGING DRUG | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>A203779 001</u> | Oct 19, 2015 |
| <u>AP</u> | BIOMEDCL RES FDN | <u>48.75mCi-487.5mCi/13ML (3.75-37.5mCi/ML)</u> | <u>A204352 001</u> | May 01, 2015 |
| <u>AP</u> | BRIGHAM WOMENS HOSP | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>A203783 001</u> | Oct 30, 2014 |
| <u>AP</u> | CARDINAL HEALTH 414 | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>A203700 001</u> | Feb 25, 2013 |
| <u>AP</u> | +! FEINSTEIN | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>N022119 001</u> | Aug 23, 2007 |
| <u>AP</u> | GEN HOSP | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>A207025 001</u> | Feb 03, 2016 |
| <u>AP</u> | GLOBAL ISOTOPEs LLC | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>A204465 001</u> | Oct 23, 2014 |
| <u>AP</u> | IONETIX | <u>22.5mCi-225mCi/6ML (3.75-37.5mCi/ML)</u> | <u>A210524 001</u> | Dec 21, 2018 |
| <u>AP</u> | JOHNS HOPKINS UNIV | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>A204514 001</u> | Aug 19, 2014 |
| <u>AP</u> | KREITCHMAN PET CTR | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>A203938 001</u> | Dec 09, 2013 |
| <u>AP</u> | MCPRF | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>A203321 001</u> | Feb 25, 2013 |
| <u>AP</u> | MIDWEST MEDCL | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>A204457 001</u> | Nov 18, 2015 |
| <u>AP</u> | MIPS CRF | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>A204535 001</u> | Nov 20, 2014 |
| <u>AP</u> | PETNET | <u>30mCi-300mCi (3.75-37.5mCi/ML)</u> | <u>A204510 001</u> | Nov 02, 2015 |
| <u>AP</u> | SOFIE | <u>18.8mCi-188mCi/5ML (3.75-37.5mCi/ML)</u> | <u>A204667 001</u> | Apr 22, 2015 |
| <u>AP</u> | SPECTRON MRC LLC | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>A204455 001</u> | Apr 23, 2015 |
| <u>AP</u> | UCLA BIOMEDICAL | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>A203812 001</u> | Jun 27, 2013 |
| <u>AP</u> | UCSF RODIOPHARM | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>A204496 001</u> | Mar 28, 2014 |
| <u>AP</u> | WA UNIV SCH MED | <u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u> | <u>A204506 001</u> | Feb 07, 2014 |
| | ESSENTIAL ISOTOPEs | 3.75-260mCi/ML | A205687 001 | Dec 17, 2015 |
| | HOUSTON CYCLOTROn | 3.75-260mCi/ML | A203543 001 | Dec 14, 2012 |
| | NCM USA BRONX LLC | 3.75-260mCi/mL | A204515 001 | Feb 04, 2015 |
| | PRECISION NUCLEAR | 3.75-260mCi/ML | A204547 001 | Aug 14, 2015 |

PRESCRIPTION DRUG PRODUCT LIST

AMMONIA N-13

INJECTABLE; INTRAVENOUS

AMMONIA N 13

| | | | | |
|--------------------|----------------|---------|-----|--------------|
| SHERTECH LABS LLC | 3.75-260mCi/ML | A204366 | 001 | Sep 19, 2014 |
| WI MEDCL CYCLOTRON | 3.75-260mCi/ML | A204356 | 001 | Dec 18, 2014 |

AMMONIUM CHLORIDE

INJECTABLE; INJECTION

AMMONIUM CHLORIDE IN PLASTIC CONTAINER

| | | | | |
|-----------|---------|---------|-----|--------------|
| ! HOSPIRA | 5MEQ/ML | A088366 | 001 | Jun 13, 1984 |
|-----------|---------|---------|-----|--------------|

AMMONIUM LACTATE

CREAM; TOPICAL

AMMONIUM LACTATE

| | | | | |
|------------------------------|--------------------|----------------|------------|--------------|
| <u>AB</u> ! PERRIGO NEW YORK | <u>EQ 12% BASE</u> | <u>A075774</u> | <u>001</u> | May 01, 2002 |
| <u>AB</u> TARO | <u>EQ 12% BASE</u> | <u>A075883</u> | <u>001</u> | Apr 10, 2003 |
| <u>AB</u> WATSON LABS INC | <u>EQ 12% BASE</u> | <u>A076829</u> | <u>001</u> | Feb 07, 2006 |

LOTION; TOPICAL

AMMONIUM LACTATE

| | | | | |
|------------------------------|--------------------|----------------|------------|--------------|
| <u>AB</u> ! PERRIGO NEW YORK | <u>EQ 12% BASE</u> | <u>A075570</u> | <u>001</u> | Jun 23, 2004 |
| <u>AB</u> TARO | <u>EQ 12% BASE</u> | <u>A076216</u> | <u>001</u> | May 28, 2004 |
| <u>AB</u> WATSON LABS INC | <u>EQ 12% BASE</u> | <u>A075575</u> | <u>001</u> | Jun 11, 2002 |

AMOXAPINE

TABLET; ORAL

AMOXAPINE

| | | | | |
|-------------|-------|---------|-----|--------------|
| WATSON LABS | 25MG | A072691 | 002 | Aug 28, 1992 |
| | 50MG | A072691 | 003 | Aug 28, 1992 |
| | 100MG | A072691 | 004 | Aug 28, 1992 |
| ! | 150MG | A072691 | 001 | Aug 28, 1992 |

AMOXICILLIN

CAPSULE; ORAL

AMOXICILLIN

| | | | | |
|---------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> AM ANTIBIOTICS | <u>250MG</u> | <u>A062058</u> | <u>001</u> | |
| <u>AB</u> | <u>500MG</u> | <u>A062058</u> | <u>002</u> | |
| <u>AB</u> AUROBINDO | <u>250MG</u> | <u>A065271</u> | <u>001</u> | Nov 09, 2005 |
| <u>AB</u> | <u>500MG</u> | <u>A065271</u> | <u>002</u> | Nov 09, 2005 |
| <u>AB</u> DAVA PHARMS INC | <u>250MG</u> | <u>A062884</u> | <u>001</u> | Feb 25, 1988 |
| <u>AB</u> | <u>500MG</u> | <u>A062881</u> | <u>001</u> | Feb 25, 1988 |
| <u>AB</u> HIKMA PHARMS | <u>250MG</u> | <u>A065291</u> | <u>001</u> | Feb 05, 2007 |
| <u>AB</u> | <u>500MG</u> | <u>A065291</u> | <u>002</u> | Feb 05, 2007 |
| <u>AB</u> SANDOZ | <u>250MG</u> | <u>A064076</u> | <u>001</u> | Sep 30, 1994 |
| <u>AB</u> | <u>500MG</u> | <u>A064076</u> | <u>002</u> | Sep 30, 1994 |
| <u>AB</u> TEVA | <u>250MG</u> | <u>A061926</u> | <u>001</u> | |
| <u>AB</u> ! | <u>500MG</u> | <u>A061926</u> | <u>003</u> | |

AMOXIL

| | | | | |
|---------------------|--------------|----------------|------------|--|
| <u>AB</u> NEOPHARMA | <u>250MG</u> | <u>A062216</u> | <u>001</u> | |
| <u>AB</u> | <u>500MG</u> | <u>A062216</u> | <u>004</u> | |

FOR SUSPENSION; ORAL

AMOXICILLIN

| | | | | |
|--------------------------------|------------------|----------------|------------|--------------|
| <u>AB</u> AUROBINDO | <u>200MG/5ML</u> | <u>A065334</u> | <u>001</u> | Dec 28, 2006 |
| <u>AB</u> | <u>400MG/5ML</u> | <u>A065334</u> | <u>002</u> | Dec 28, 2006 |
| <u>AB</u> AUROBINDO PHARMA LTD | <u>125MG/5ML</u> | <u>A204030</u> | <u>001</u> | Sep 15, 2014 |
| <u>AB</u> | <u>250MG/5ML</u> | <u>A204030</u> | <u>002</u> | Sep 15, 2014 |
| <u>AB</u> DAVA PHARMS INC | <u>125MG/5ML</u> | <u>A062927</u> | <u>001</u> | Nov 25, 1988 |
| <u>AB</u> | <u>250MG/5ML</u> | <u>A062927</u> | <u>002</u> | Nov 25, 1988 |
| <u>AB</u> HIKMA | <u>125MG/5ML</u> | <u>A065322</u> | <u>002</u> | Jun 19, 2006 |
| <u>AB</u> | <u>200MG/5ML</u> | <u>A065325</u> | <u>002</u> | Jun 19, 2006 |
| <u>AB</u> | <u>250MG/5ML</u> | <u>A065322</u> | <u>001</u> | Jun 19, 2006 |
| <u>AB</u> | <u>400MG/5ML</u> | <u>A065325</u> | <u>001</u> | Jun 19, 2006 |
| <u>AB</u> SANDOZ | <u>125MG/5ML</u> | <u>A065387</u> | <u>001</u> | Mar 26, 2007 |
| <u>AB</u> | <u>200MG/5ML</u> | <u>A065378</u> | <u>001</u> | Mar 26, 2007 |
| <u>AB</u> | <u>250MG/5ML</u> | <u>A065387</u> | <u>002</u> | Mar 26, 2007 |
| <u>AB</u> | <u>400MG/5ML</u> | <u>A065378</u> | <u>002</u> | Mar 26, 2007 |
| <u>AB</u> TEVA | <u>125MG/5ML</u> | <u>A061931</u> | <u>001</u> | |
| <u>AB</u> | <u>200MG/5ML</u> | <u>A065119</u> | <u>001</u> | Dec 04, 2002 |
| <u>AB</u> ! | <u>250MG/5ML</u> | <u>A061931</u> | <u>002</u> | |
| <u>AB</u> ! | <u>400MG/5ML</u> | <u>A065119</u> | <u>002</u> | Dec 04, 2002 |
| <u>AB</u> WOCKHARDT BIO AG | <u>400MG/5ML</u> | <u>A065319</u> | <u>002</u> | Jun 18, 2007 |

AMOXICILLIN PEDIATRIC

| | | | | |
|----------------|----------------|----------------|------------|--------------|
| <u>AB</u> TEVA | <u>50MG/ML</u> | <u>A061931</u> | <u>003</u> | Dec 01, 1982 |
|----------------|----------------|----------------|------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

AMOXICILLIN

FOR SUSPENSION;ORAL

AMOXIL

| | | | | | |
|-----------|-----------|------------------|----------------|------------|--|
| <u>AB</u> | NEOPHARMA | <u>50MG/ML</u> | <u>A062226</u> | <u>005</u> | |
| <u>AB</u> | | <u>125MG/5ML</u> | <u>A062226</u> | <u>001</u> | |
| <u>AB</u> | | <u>250MG/5ML</u> | <u>A062226</u> | <u>002</u> | |

LAROTID

| | | | | | |
|-----------|-----------|------------------|----------------|------------|--|
| <u>AB</u> | NEOPHARMA | <u>125MG/5ML</u> | <u>A062226</u> | <u>003</u> | |
| <u>AB</u> | | <u>250MG/5ML</u> | <u>A062226</u> | <u>004</u> | |

TABLET;ORAL

AMOXICILLIN

| | | | | | |
|-----------|-----------|--------------|----------------|------------|--------------|
| <u>AB</u> | AUROBINDO | <u>500MG</u> | <u>A065256</u> | <u>001</u> | Nov 09, 2005 |
| <u>AB</u> | | <u>875MG</u> | <u>A065256</u> | <u>002</u> | Nov 09, 2005 |
| <u>AB</u> | HIKMA | <u>875MG</u> | <u>A065255</u> | <u>001</u> | Mar 29, 2006 |
| <u>AB</u> | SANDOZ | <u>500MG</u> | <u>A065228</u> | <u>001</u> | Jul 13, 2005 |
| <u>AB</u> | | <u>875MG</u> | <u>A065228</u> | <u>002</u> | Jul 13, 2005 |
| <u>AB</u> | TEVA | <u>500MG</u> | <u>A065056</u> | <u>001</u> | Sep 18, 2000 |
| <u>AB</u> | ! | <u>875MG</u> | <u>A065056</u> | <u>002</u> | Sep 18, 2000 |

TABLET, CHEWABLE;ORAL

AMOXICILLIN

| | | | | | |
|--|------|-------|---------|-----|--------------|
| | TEVA | 125MG | A064013 | 002 | Sep 11, 1995 |
| | ! | 250MG | A064013 | 001 | Dec 22, 1992 |

AMOXICILLIN; CLARITHROMYCIN; LANSOPRAZOLE

CAPSULE, CAPSULE, DELAYED REL PELLETS, TABLET;ORAL

LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN

| | | | | | | |
|-----------|---|---------------|-------------------------------------------------|----------------|------------|--------------|
| <u>AB</u> | ! | RISING PHARMS | <u>500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,30MG</u> | <u>A206006</u> | <u>001</u> | Oct 07, 2016 |
| <u>AB</u> | | SANDOZ INC | <u>500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,30MG</u> | <u>A202588</u> | <u>001</u> | Mar 04, 2014 |

AMOXICILLIN; CLARITHROMYCIN; OMEPRAZOLE

CAPSULE, TABLET, CAPSULE, DELAYED RELEASE;ORAL

OMEPRazole AND CLARITHROMYCIN AND AMOXICILLIN

| | | | | | | |
|--|---|-------------------|------------------------------------------|---------|-----|--------------|
| | + | CUMBERLAND PHARMS | 500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,20MG | N050824 | 001 | Feb 08, 2011 |
|--|---|-------------------|------------------------------------------|---------|-----|--------------|

AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

| | | | | | |
|-----------|----------------------|-------------------------------------|----------------|------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>200MG/5ML;EQ 28.5MG BASE/5ML</u> | <u>A201090</u> | <u>001</u> | Dec 20, 2011 |
| <u>AB</u> | | <u>400MG/5ML;EQ 57MG BASE/5ML</u> | <u>A201090</u> | <u>002</u> | Dec 20, 2011 |
| <u>AB</u> | | <u>600MG/5ML;EQ 42.9MG BASE/5ML</u> | <u>A201091</u> | <u>001</u> | Dec 20, 2011 |
| <u>AB</u> | HIKMA PHARMS | <u>200MG/5ML;EQ 28.5MG BASE/5ML</u> | <u>A065191</u> | <u>002</u> | Jan 25, 2005 |
| <u>AB</u> | | <u>400MG/5ML;EQ 57MG BASE/5ML</u> | <u>A065191</u> | <u>001</u> | Jan 25, 2005 |
| <u>AB</u> | | <u>600MG/5ML;EQ 42.9MG BASE/5ML</u> | <u>A065373</u> | <u>001</u> | Nov 09, 2007 |
| <u>AB</u> | SANDOZ | <u>200MG/5ML;EQ 28.5MG BASE/5ML</u> | <u>A065066</u> | <u>001</u> | Jun 05, 2002 |
| <u>AB</u> | | <u>400MG/5ML;EQ 57MG BASE/5ML</u> | <u>A065066</u> | <u>002</u> | Jun 05, 2002 |
| <u>AB</u> | SANDOZ INC | <u>200MG/5ML;EQ 28.5MG BASE/5ML</u> | <u>A065098</u> | <u>001</u> | Dec 16, 2002 |
| <u>AB</u> | | <u>400MG/5ML;EQ 57MG BASE/5ML</u> | <u>A065098</u> | <u>002</u> | Dec 16, 2002 |
| <u>AB</u> | | <u>600MG/5ML;EQ 42.9MG BASE/5ML</u> | <u>A065358</u> | <u>001</u> | Aug 13, 2007 |
| <u>AB</u> | TEVA | <u>200MG/5ML;EQ 28.5MG BASE/5ML</u> | <u>A065089</u> | <u>001</u> | May 25, 2004 |
| <u>AB</u> | ! | <u>400MG/5ML;EQ 57MG BASE/5ML</u> | <u>A065089</u> | <u>002</u> | May 25, 2004 |
| <u>AB</u> | ! | <u>600MG/5ML;EQ 42.9MG BASE/5ML</u> | <u>A065162</u> | <u>001</u> | Mar 12, 2004 |
| <u>AB</u> | WOCKHARDT BIO AG | <u>250MG/5ML;EQ 62.5MG BASE/5ML</u> | <u>A065431</u> | <u>001</u> | Nov 25, 2008 |
| <u>AB</u> | | <u>600MG/5ML;EQ 42.9MG BASE/5ML</u> | <u>A065420</u> | <u>001</u> | Dec 02, 2013 |

AUGMENTIN '250'

| | | | | | | |
|-----------|---|-----------------|-------------------------------------|----------------|------------|--------------|
| <u>AB</u> | + | NEOPHARMA | <u>250MG/5ML;EQ 62.5MG BASE/5ML</u> | <u>N050575</u> | <u>002</u> | Aug 06, 1984 |
| | | AUGMENTIN '125' | | | | |
| | + | NEOPHARMA | 125MG/5ML;EQ 31.25MG BASE/5ML | N050575 | 001 | Aug 06, 1984 |

TABLET;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

| | | | | | | |
|-----------|----------------------|----------------------------|----------------------------|----------------|--------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>250MG;EQ 125MG BASE</u> | <u>A091569</u> | <u>001</u> | Jan 20, 2012 | |
| <u>AB</u> | | <u>500MG;EQ 125MG BASE</u> | <u>A091569</u> | <u>002</u> | Jan 20, 2012 | |
| <u>AB</u> | | <u>875MG;EQ 125MG BASE</u> | <u>A091568</u> | <u>001</u> | Jan 20, 2012 | |
| <u>AB</u> | HIKMA PHARMS | <u>875MG;EQ 125MG BASE</u> | <u>A203824</u> | <u>001</u> | Aug 23, 2016 | |
| <u>AB</u> | MICRO LABS LTD INDIA | <u>250MG;EQ 125MG BASE</u> | <u>A205707</u> | <u>001</u> | Dec 30, 2016 | |
| <u>AB</u> | | <u>500MG;EQ 125MG BASE</u> | <u>A205707</u> | <u>002</u> | Dec 30, 2016 | |
| <u>AB</u> | | <u>875MG;EQ 125MG BASE</u> | <u>A204755</u> | <u>003</u> | Dec 30, 2016 | |
| <u>AB</u> | ! | SANDOZ | <u>250MG;EQ 125MG BASE</u> | <u>A065189</u> | <u>001</u> | Aug 23, 2005 |
| <u>AB</u> | | <u>500MG;EQ 125MG BASE</u> | <u>A065064</u> | <u>001</u> | Mar 15, 2002 | |
| <u>AB</u> | ! | <u>875MG;EQ 125MG BASE</u> | <u>A065063</u> | <u>001</u> | Mar 14, 2002 | |

PRESCRIPTION DRUG PRODUCT LIST

AMOXICILLIN; CLAVULANATE POTASSIUM

TABLET; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

| | | | | | |
|-----------|----------|-----------------|----------------------------|--------------------|--------------|
| AB | ! | SANDOZ INC | <u>500MG;EQ 125MG BASE</u> | <u>A065117 001</u> | Nov 27, 2002 |
| AB | | | <u>875MG;EQ 125MG BASE</u> | <u>A065093 001</u> | Nov 21, 2002 |
| AB | | TEVA | <u>500MG;EQ 125MG BASE</u> | <u>A065101 001</u> | Oct 30, 2002 |
| AB | | TEVA PHARMS USA | <u>875MG;EQ 125MG BASE</u> | <u>A065096 001</u> | Oct 29, 2002 |

AUGMENTIN '875'

| | | | | | |
|-----------|----------|--------------------|----------------------------|--------------------|--------------|
| AB | + | DR REDDYS LABS INC | <u>875MG;EQ 125MG BASE</u> | <u>N050720 001</u> | Feb 13, 1996 |
|-----------|----------|--------------------|----------------------------|--------------------|--------------|

TABLET, CHEWABLE; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

| | | | | | |
|--|--|----------|----------------------|-------------|--------------|
| | | TEVA | 200MG;EQ 28.5MG BASE | A065205 001 | Feb 09, 2005 |
| | | ! | 400MG;EQ 57MG BASE | A065205 002 | Feb 09, 2005 |

TABLET, EXTENDED RELEASE; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

| | | | | | |
|-----------|--|--------|---------------------------|--------------------|--------------|
| AB | | SANDOZ | <u>1GM;EQ 62.5MG BASE</u> | <u>A090227 001</u> | Apr 21, 2010 |
|-----------|--|--------|---------------------------|--------------------|--------------|

AUGMENTIN XR

| | | | | | |
|-----------|----------|-----------|---------------------------|--------------------|--------------|
| AB | + | NEOPHARMA | <u>1GM;EQ 62.5MG BASE</u> | <u>N050785 001</u> | Sep 25, 2002 |
|-----------|----------|-----------|---------------------------|--------------------|--------------|

AMPHETAMINE

SUSPENSION, EXTENDED RELEASE; ORAL

ADZENYS ER

| | | | | | |
|--|----------|------------------|-------------------|-------------|--------------|
| | + | NEOS THERAPS INC | EQ 1.25MG BASE/ML | N204325 001 | Sep 15, 2017 |
|--|----------|------------------|-------------------|-------------|--------------|

DYANAVEL XR

| | | | | | |
|--|----------|-----------------|------------------|-------------|--------------|
| | + | TRIS PHARMA INC | EQ 2.5MG BASE/ML | N208147 001 | Oct 19, 2015 |
|--|----------|-----------------|------------------|-------------|--------------|

TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE; ORAL

ADZENYS XR-ODT

| | | | | | |
|--|----------|--------------|----------------|-------------|--------------|
| | + | NEOS THERAPS | EQ 3.1MG BASE | N204326 001 | Jan 27, 2016 |
| | + | | EQ 6.3MG BASE | N204326 002 | Jan 27, 2016 |
| | + | | EQ 9.4MG BASE | N204326 003 | Jan 27, 2016 |
| | + | | EQ 12.5MG BASE | N204326 004 | Jan 27, 2016 |
| | + | | EQ 15.7MG BASE | N204326 005 | Jan 27, 2016 |
| | + | | EQ 18.8MG BASE | N204326 006 | Jan 27, 2016 |

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

ADDERALL XR 10

| | | | | | |
|-----------|----------|-------|--------------------------------|--------------------|--------------|
| AB | + | SHIRE | <u>2.5MG;2.5MG;2.5MG;2.5MG</u> | <u>N021303 001</u> | Oct 11, 2001 |
|-----------|----------|-------|--------------------------------|--------------------|--------------|

ADDERALL XR 15

| | | | | | |
|-----------|----------|-------|------------------------------------|--------------------|--------------|
| AB | + | SHIRE | <u>3.75MG;3.75MG;3.75MG;3.75MG</u> | <u>N021303 006</u> | May 22, 2002 |
|-----------|----------|-------|------------------------------------|--------------------|--------------|

ADDERALL XR 20

| | | | | | |
|-----------|----------|-------|------------------------|--------------------|--------------|
| AB | + | SHIRE | <u>5MG;5MG;5MG;5MG</u> | <u>N021303 002</u> | Oct 11, 2001 |
|-----------|----------|-------|------------------------|--------------------|--------------|

ADDERALL XR 25

| | | | | | |
|-----------|----------|-------|------------------------------------|--------------------|--------------|
| AB | + | SHIRE | <u>6.25MG;6.25MG;6.25MG;6.25MG</u> | <u>N021303 004</u> | May 22, 2002 |
|-----------|----------|-------|------------------------------------|--------------------|--------------|

ADDERALL XR 30

| | | | | | |
|-----------|----------|-------|--------------------------------|--------------------|--------------|
| AB | + | SHIRE | <u>7.5MG;7.5MG;7.5MG;7.5MG</u> | <u>N021303 003</u> | Oct 11, 2001 |
|-----------|----------|-------|--------------------------------|--------------------|--------------|

ADDERALL XR 5

| | | | | | |
|-----------|----------|-------|------------------------------------|--------------------|--------------|
| AB | + | SHIRE | <u>1.25MG;1.25MG;1.25MG;1.25MG</u> | <u>N021303 005</u> | May 22, 2002 |
|-----------|----------|-------|------------------------------------|--------------------|--------------|

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

| | | | | | |
|-----------|--|-------------------|------------------------------------|--------------------|--------------|
| AB | | ACTAVIS ELIZABETH | <u>1.25MG;1.25MG;1.25MG;1.25MG</u> | <u>A077302 001</u> | Jun 22, 2012 |
| AB | | | <u>2.5MG;2.5MG;2.5MG;2.5MG</u> | <u>A077302 002</u> | Jun 22, 2012 |
| AB | | | <u>3.75MG;3.75MG;3.75MG;3.75MG</u> | <u>A077302 003</u> | Jun 22, 2012 |
| AB | | | <u>5MG;5MG;5MG;5MG</u> | <u>A077302 004</u> | Jun 22, 2012 |
| AB | | | <u>6.25MG;6.25MG;6.25MG;6.25MG</u> | <u>A077302 005</u> | Jun 22, 2012 |
| AB | | | <u>7.5MG;7.5MG;7.5MG;7.5MG</u> | <u>A077302 006</u> | Jun 22, 2012 |
| AB | | IMPAX LABS | <u>1.25MG;1.25MG;1.25MG;1.25MG</u> | <u>A076852 001</u> | Feb 16, 2016 |
| AB | | | <u>2.5MG;2.5MG;2.5MG;2.5MG</u> | <u>A076852 002</u> | Feb 16, 2016 |
| AB | | | <u>3.75MG;3.75MG;3.75MG;3.75MG</u> | <u>A076852 003</u> | Feb 16, 2016 |
| AB | | | <u>5MG;5MG;5MG;5MG</u> | <u>A076852 004</u> | Feb 16, 2016 |
| AB | | | <u>6.25MG;6.25MG;6.25MG;6.25MG</u> | <u>A076852 005</u> | Feb 16, 2016 |
| AB | | | <u>7.5MG;7.5MG;7.5MG;7.5MG</u> | <u>A076852 006</u> | Feb 16, 2016 |
| AB | | TEVA | <u>1.25MG;1.25MG;1.25MG;1.25MG</u> | <u>A077488 001</u> | Apr 29, 2013 |
| AB | | | <u>2.5MG;2.5MG;2.5MG;2.5MG</u> | <u>A077488 002</u> | Apr 29, 2013 |
| AB | | | <u>3.75MG;3.75MG;3.75MG;3.75MG</u> | <u>A077488 003</u> | Apr 29, 2013 |
| AB | | | <u>5MG;5MG;5MG;5MG</u> | <u>A077488 004</u> | Apr 29, 2013 |
| AB | | | <u>6.25MG;6.25MG;6.25MG;6.25MG</u> | <u>A077488 005</u> | Apr 29, 2013 |
| AB | | | <u>7.5MG;7.5MG;7.5MG;7.5MG</u> | <u>A077488 006</u> | Apr 29, 2013 |

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

| | | | | | |
|-----------|--|---------------|------------------------------------|--------------------|--------------|
| AB | | BARR LABS INC | <u>1.25MG;1.25MG;1.25MG;1.25MG</u> | <u>A076536 001</u> | Feb 12, 2013 |
| AB | | | <u>2.5MG;2.5MG;2.5MG;2.5MG</u> | <u>A076536 002</u> | Feb 12, 2013 |
| AB | | | <u>3.75MG;3.75MG;3.75MG;3.75MG</u> | <u>A076536 003</u> | Feb 12, 2013 |
| AB | | | <u>5MG;5MG;5MG;5MG</u> | <u>A076536 004</u> | Feb 12, 2013 |
| AB | | | <u>6.25MG;6.25MG;6.25MG;6.25MG</u> | <u>A076536 005</u> | Feb 12, 2013 |

PRESCRIPTION DRUG PRODUCT LIST

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

| | | | | |
|-----------|---------|--------------------------------|---------------------------------|--------------------------|
| AB | | 7.5MG;7.5MG;7.5MG;7.5MG | A076536 006 | Feb 12, 2013 |
| | MYDAYIS | | | |
| | + | SHIRE DEV LLC | 3.125MG;3.125MG;3.125MG;3.125MG | N022063 001 Jun 20, 2017 |
| | + | | 6.25MG;6.25MG;6.25MG;6.25MG | N022063 002 Jun 20, 2017 |
| | + | | 9.375MG;9.375MG;9.375MG;9.375MG | N022063 003 Jun 20, 2017 |
| | + | ! | 12.5MG;12.5MG;12.5MG;12.5MG | N022063 004 Jun 20, 2017 |

TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

| | | | | |
|-----------|---------------------|----------------------------------------|--------------------|--------------|
| AB | ACTAVIS ELIZABETH | 1.25MG;1.25MG;1.25MG;1.25MG | A206340 001 | Feb 05, 2016 |
| AB | | 1.875MG;1.875MG;1.875MG;1.875MG | A206340 002 | Feb 05, 2016 |
| AB | | 2.5MG;2.5MG;2.5MG;2.5MG | A206340 003 | Feb 05, 2016 |
| AB | | 3.125MG;3.125MG;3.125MG;3.125MG | A206340 004 | Feb 05, 2016 |
| AB | | 3.75MG;3.75MG;3.75MG;3.75MG | A206340 005 | Feb 05, 2016 |
| AB | | 5MG;5MG;5MG;5MG | A206340 006 | Feb 05, 2016 |
| AB | | 7.5MG;7.5MG;7.5MG;7.5MG | A206340 007 | Feb 05, 2016 |
| AB | ALVOGEN MALTA | 1.25MG;1.25MG;1.25MG;1.25MG | A207388 001 | Jul 28, 2017 |
| AB | | 1.875MG;1.875MG;1.875MG;1.875MG | A207388 002 | Jul 28, 2017 |
| AB | | 2.5MG;2.5MG;2.5MG;2.5MG | A207388 003 | Jul 28, 2017 |
| AB | | 3.125MG;3.125MG;3.125MG;3.125MG | A207388 004 | Jul 28, 2017 |
| AB | | 3.75MG;3.75MG;3.75MG;3.75MG | A207388 005 | Jul 28, 2017 |
| AB | | 5MG;5MG;5MG;5MG | A207388 006 | Jul 28, 2017 |
| AB | | 7.5MG;7.5MG;7.5MG;7.5MG | A207388 007 | Jul 28, 2017 |
| AB | AUROLIFE PHARMA LLC | 1.25MG;1.25MG;1.25MG;1.25MG | A202424 001 | Nov 27, 2013 |
| AB | | 1.875MG;1.875MG;1.875MG;1.875MG | A202424 002 | Nov 27, 2013 |
| AB | | 2.5MG;2.5MG;2.5MG;2.5MG | A202424 003 | Nov 27, 2013 |
| AB | | 3.125MG;3.125MG;3.125MG;3.125MG | A202424 004 | Nov 27, 2013 |
| AB | | 3.75MG;3.75MG;3.75MG;3.75MG | A202424 005 | Nov 27, 2013 |
| AB | | 5MG;5MG;5MG;5MG | A202424 006 | Nov 27, 2013 |
| AB | | 7.5MG;7.5MG;7.5MG;7.5MG | A202424 007 | Nov 27, 2013 |
| AB | BARR | 1.25MG;1.25MG;1.25MG;1.25MG | A040422 001 | Feb 11, 2002 |
| AB | | 1.875MG;1.875MG;1.875MG;1.875MG | A040422 005 | Mar 19, 2003 |
| AB | | 2.5MG;2.5MG;2.5MG;2.5MG | A040422 002 | Feb 11, 2002 |
| AB | | 3.125MG;3.125MG;3.125MG;3.125MG | A040422 006 | Mar 19, 2003 |
| AB | | 3.75MG;3.75MG;3.75MG;3.75MG | A040422 007 | Mar 19, 2003 |
| AB | | 5MG;5MG;5MG;5MG | A040422 003 | Feb 11, 2002 |
| AB | ! | 7.5MG;7.5MG;7.5MG;7.5MG | A040422 004 | Feb 11, 2002 |
| AB | EPIC PHARMA LLC | 1.25MG;1.25MG;1.25MG;1.25MG | A040444 001 | Jun 19, 2002 |
| AB | | 1.875MG;1.875MG;1.875MG;1.875MG | A040444 005 | Nov 03, 2014 |
| AB | | 2.5MG;2.5MG;2.5MG;2.5MG | A040444 002 | Jun 19, 2002 |
| AB | | 3.125MG;3.125MG;3.125MG;3.125MG | A040444 006 | Nov 03, 2014 |
| AB | | 3.75MG;3.75MG;3.75MG;3.75MG | A040444 007 | Nov 03, 2014 |
| AB | | 5MG;5MG;5MG;5MG | A040444 003 | Jun 19, 2002 |
| AB | | 7.5MG;7.5MG;7.5MG;7.5MG | A040444 004 | Jun 19, 2002 |
| AB | MYLAN PHARMS INC | 1.25MG;1.25MG;1.25MG;1.25MG | A206721 001 | Nov 10, 2015 |
| AB | | 1.875MG;1.875MG;1.875MG;1.875MG | A206721 002 | Nov 10, 2015 |
| AB | | 2.5MG;2.5MG;2.5MG;2.5MG | A206721 003 | Nov 10, 2015 |
| AB | | 3.125MG;3.125MG;3.125MG;3.125MG | A206721 004 | Nov 10, 2015 |
| AB | | 3.75MG;3.75MG;3.75MG;3.75MG | A206721 005 | Nov 10, 2015 |
| AB | | 5MG;5MG;5MG;5MG | A206721 006 | Nov 10, 2015 |
| AB | | 7.5MG;7.5MG;7.5MG;7.5MG | A206721 007 | Nov 10, 2015 |
| AB | NESHER PHARMS | 1.25MG;1.25MG;1.25MG;1.25MG | A207340 001 | Oct 31, 2017 |
| AB | | 1.875MG;1.875MG;1.875MG;1.875MG | A207340 002 | Oct 31, 2017 |
| AB | | 2.5MG;2.5MG;2.5MG;2.5MG | A207340 003 | Oct 31, 2017 |
| AB | | 3.125MG;3.125MG;3.125MG;3.125MG | A207340 004 | Oct 31, 2017 |
| AB | | 3.75MG;3.75MG;3.75MG;3.75MG | A207340 005 | Oct 31, 2017 |
| AB | | 5MG;5MG;5MG;5MG | A207340 006 | Oct 31, 2017 |
| AB | | 7.5MG;7.5MG;7.5MG;7.5MG | A207340 007 | Oct 31, 2017 |
| AB | NUVO PHARM | 1.25MG;1.25MG;1.25MG;1.25MG | A209799 001 | Dec 28, 2017 |
| AB | | 1.875MG;1.875MG;1.875MG;1.875MG | A209799 002 | Dec 28, 2017 |
| AB | | 2.5MG;2.5MG;2.5MG;2.5MG | A209799 003 | Dec 28, 2017 |
| AB | | 3.125MG;3.125MG;3.125MG;3.125MG | A209799 004 | Dec 28, 2017 |
| AB | | 3.75MG;3.75MG;3.75MG;3.75MG | A209799 005 | Dec 28, 2017 |
| AB | | 5MG;5MG;5MG;5MG | A209799 006 | Dec 28, 2017 |
| AB | | 7.5MG;7.5MG;7.5MG;7.5MG | A209799 007 | Dec 28, 2017 |
| AB | SANDOZ | 1.25MG;1.25MG;1.25MG;1.25MG | A040439 004 | Sep 27, 2002 |
| AB | | 2.5MG;2.5MG;2.5MG;2.5MG | A040439 001 | Jun 14, 2002 |
| AB | | 5MG;5MG;5MG;5MG | A040439 002 | Jun 14, 2002 |
| AB | | 7.5MG;7.5MG;7.5MG;7.5MG | A040439 003 | Jun 14, 2002 |
| AB | SPECGX LLC | 1.25MG;1.25MG;1.25MG;1.25MG | A040440 001 | Oct 07, 2003 |

PRESCRIPTION DRUG PRODUCT LIST

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

| | | | | |
|-----------|-------------------------|----------------------------------------|--------------------|--------------|
| <u>AB</u> | | <u>1.875MG;1.875MG;1.875MG;1.875MG</u> | <u>A040440 002</u> | Oct 07, 2003 |
| <u>AB</u> | | <u>2.5MG;2.5MG;2.5MG;2.5MG</u> | <u>A040440 003</u> | Oct 07, 2003 |
| <u>AB</u> | | <u>3.125MG;3.125MG;3.125MG;3.125MG</u> | <u>A040440 004</u> | Oct 07, 2003 |
| <u>AB</u> | | <u>3.75MG;3.75MG;3.75MG;3.75MG</u> | <u>A040440 005</u> | Oct 07, 2003 |
| <u>AB</u> | | <u>5MG;5MG;5MG;5MG</u> | <u>A040440 006</u> | Oct 07, 2003 |
| <u>AB</u> | | <u>7.5MG;7.5MG;7.5MG;7.5MG</u> | <u>A040440 007</u> | Oct 07, 2003 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>1.25MG;1.25MG;1.25MG;1.25MG</u> | <u>A040480 001</u> | Sep 09, 2003 |
| <u>AB</u> | | <u>1.875MG;1.875MG;1.875MG;1.875MG</u> | <u>A040480 002</u> | Sep 09, 2003 |
| <u>AB</u> | | <u>2.5MG;2.5MG;2.5MG;2.5MG</u> | <u>A040480 003</u> | Sep 09, 2003 |
| <u>AB</u> | | <u>3.125MG;3.125MG;3.125MG;3.125MG</u> | <u>A040480 004</u> | Sep 09, 2003 |
| <u>AB</u> | | <u>3.75MG;3.75MG;3.75MG;3.75MG</u> | <u>A040480 005</u> | Sep 09, 2003 |
| <u>AB</u> | | <u>5MG;5MG;5MG;5MG</u> | <u>A040480 006</u> | Sep 09, 2003 |
| <u>AB</u> | | <u>7.5MG;7.5MG;7.5MG;7.5MG</u> | <u>A040480 007</u> | Sep 09, 2003 |
| <u>AB</u> | SUNGEN PHARMA | <u>1.25MG;1.25MG;1.25MG;1.25MG</u> | <u>A211352 001</u> | Dec 07, 2018 |
| <u>AB</u> | | <u>1.875MG;1.875MG;1.875MG;1.875MG</u> | <u>A211352 002</u> | Dec 07, 2018 |
| <u>AB</u> | | <u>2.5MG;2.5MG;2.5MG;2.5MG</u> | <u>A211352 003</u> | Dec 07, 2018 |
| <u>AB</u> | | <u>3.125MG;3.125MG;3.125MG;3.125MG</u> | <u>A211352 004</u> | Dec 07, 2018 |
| <u>AB</u> | | <u>3.75MG;3.75MG;3.75MG;3.75MG</u> | <u>A211352 005</u> | Dec 07, 2018 |
| <u>AB</u> | | <u>5MG;5MG;5MG;5MG</u> | <u>A211352 006</u> | Dec 07, 2018 |
| <u>AB</u> | | <u>7.5MG;7.5MG;7.5MG;7.5MG</u> | <u>A211352 007</u> | Dec 07, 2018 |

AMPHETAMINE SULFATE

TABLET; ORAL

AMPHETAMINE SULFATE

| | | | | |
|-----------|------------------|-------------|--------------------|--------------|
| <u>AA</u> | AMNEAL PHARMS | <u>5MG</u> | <u>A211139 001</u> | Sep 26, 2018 |
| <u>AA</u> | | <u>10MG</u> | <u>A211139 002</u> | Sep 26, 2018 |
| | <u>EVEKEO</u> | | | |
| <u>AA</u> | ARBOR PHARMS LLC | <u>5MG</u> | <u>A200166 001</u> | Aug 09, 2012 |
| <u>AA</u> | ! | <u>10MG</u> | <u>A200166 002</u> | Aug 09, 2012 |

AMPHOTERICIN B

INJECTABLE; INJECTION

AMPHOTERICIN B

| | | | | |
|---|---------------------------------------------|-----------|-------------|--------------|
| ! | X GEN PHARMS | 50MG/VIAL | A063206 001 | Apr 29, 1992 |
| | <u>INJECTABLE, LIPID COMPLEX; INJECTION</u> | | | |
| | <u>ABELCET</u> | | | |
| + | LEADIANT BIOSCI INC | 5MG/ML | N050724 001 | Nov 20, 1995 |
| | <u>INJECTABLE, LIPOSOMAL; INJECTION</u> | | | |
| | <u>AMBISOME</u> | | | |
| + | ASTELLAS | 50MG/VIAL | N050740 001 | Aug 11, 1997 |

AMPICILLIN SODIUM

INJECTABLE; INJECTION

AMPICILLIN SODIUM

| | | | | |
|-----------|-------------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | ACS DOBFAR SPA | <u>EQ 10GM BASE/VIAL</u> | <u>A090889 001</u> | Apr 03, 2013 |
| <u>AP</u> | ANTIBIOTICE | <u>EQ 250MG BASE/VIAL</u> | <u>A090354 001</u> | Dec 28, 2009 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A090354 002</u> | Dec 28, 2009 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A090354 003</u> | Dec 28, 2009 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A090354 004</u> | Dec 28, 2009 |
| <u>AP</u> | AUROBINDO PHARMA | <u>EQ 250MG BASE/VIAL</u> | <u>A065499 002</u> | Aug 17, 2010 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A065499 003</u> | Aug 17, 2010 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A065499 004</u> | Aug 17, 2010 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A065499 005</u> | Aug 17, 2010 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A065493 001</u> | Aug 17, 2010 |
| <u>AP</u> | HANFORD GC | <u>EQ 250MG BASE/VIAL</u> | <u>A062772 006</u> | Apr 15, 1993 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A062772 007</u> | Apr 15, 1993 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A062772 001</u> | Apr 15, 1993 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A062772 003</u> | Apr 15, 1993 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A063142 001</u> | Apr 15, 1993 |
| <u>AP</u> | HOSPIRA INC | <u>EQ 250MG BASE/VIAL</u> | <u>A202864 001</u> | Sep 04, 2015 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A202864 002</u> | Sep 04, 2015 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A202864 003</u> | Sep 04, 2015 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A202864 004</u> | Sep 04, 2015 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A202865 001</u> | Sep 04, 2015 |
| <u>AP</u> | ISTITUTO BIO ITA SPA | <u>EQ 10GM BASE/VIAL</u> | <u>A201404 001</u> | Dec 20, 2013 |
| <u>AP</u> | | <u>EQ 250MG BASE/VIAL</u> | <u>A062719 001</u> | May 12, 1987 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A062719 003</u> | May 12, 1987 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A062719 002</u> | May 12, 1987 |

PRESCRIPTION DRUG PRODUCT LIST

AMPICILLIN SODIUM

INJECTABLE; INJECTION

AMPICILLIN SODIUM

| | | | | |
|-----------|----------------|---------------------------|--------------------|--------------|
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A062797 002</u> | Jul 12, 1993 |
| <u>AP</u> | MYLAN LABS LTD | <u>EQ 250MG BASE/VIAL</u> | <u>A201025 001</u> | Apr 09, 2014 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A201025 002</u> | Apr 09, 2014 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A201025 003</u> | Apr 09, 2014 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A201025 004</u> | Apr 09, 2014 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A202198 001</u> | Apr 07, 2014 |
| <u>AP</u> | SAGENT PHARMS | <u>EQ 125MG BASE/VIAL</u> | <u>A090583 001</u> | Nov 27, 2015 |
| <u>AP</u> | | <u>EQ 250MG BASE/VIAL</u> | <u>A090583 002</u> | Nov 27, 2015 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A090583 003</u> | Nov 27, 2015 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A090583 004</u> | Nov 27, 2015 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A090583 005</u> | Nov 27, 2015 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A090581 001</u> | Oct 20, 2015 |
| <u>AP</u> | ! | <u>EQ 125MG BASE/VIAL</u> | <u>A061395 001</u> | |
| <u>AP</u> | ! | <u>EQ 250MG BASE/VIAL</u> | <u>A061395 002</u> | |
| <u>AP</u> | ! | <u>EQ 500MG BASE/VIAL</u> | <u>A061395 003</u> | |
| <u>AP</u> | ! | <u>EQ 1GM BASE/VIAL</u> | <u>A061395 004</u> | |
| <u>AP</u> | ! | <u>EQ 2GM BASE/VIAL</u> | <u>A061395 005</u> | |
| <u>AP</u> | ! | <u>EQ 10GM BASE/VIAL</u> | <u>A061395 006</u> | |

POWDER; INTRAVENOUS

AMPICILLIN SODIUM

| | | | | |
|-----------|---|-------------------------|--------------------|--------------|
| <u>AP</u> | ! | <u>EQ 1GM BASE/VIAL</u> | <u>A062738 001</u> | Feb 19, 1987 |
| <u>AP</u> | ! | <u>EQ 2GM BASE/VIAL</u> | <u>A062738 002</u> | Feb 19, 1987 |

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

| | | | | |
|-----------|----------------------|--------------------------------------------|--------------------|--------------|
| <u>AP</u> | ACS DOBFAR | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A065406 001</u> | Dec 22, 2009 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A065406 002</u> | Dec 22, 2009 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u> | <u>A065403 001</u> | Dec 23, 2009 |
| <u>AP</u> | ANTIBIOTICE | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A201406 001</u> | Dec 07, 2015 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A201406 002</u> | Dec 07, 2015 |
| <u>AP</u> | ASTRAL | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A090579 001</u> | Jan 08, 2016 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A090579 002</u> | Jan 08, 2016 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u> | <u>A090578 001</u> | Jan 11, 2016 |
| <u>AP</u> | AUROBINDO PHARMA | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A090340 001</u> | Sep 20, 2010 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A090349 001</u> | Sep 20, 2010 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A090340 002</u> | Sep 20, 2010 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A090349 002</u> | Sep 20, 2010 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u> | <u>A090339 001</u> | Sep 20, 2010 |
| <u>AP</u> | HANFORD GC | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A065176 001</u> | Nov 30, 2005 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A065176 002</u> | Nov 30, 2005 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u> | <u>A065188 001</u> | Nov 25, 2005 |
| <u>AP</u> | HOSPIRA INC | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A090375 001</u> | Dec 21, 2011 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A090653 001</u> | Dec 21, 2011 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A090375 002</u> | Dec 21, 2011 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A090653 002</u> | Dec 21, 2011 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u> | <u>A090646 001</u> | Dec 21, 2011 |
| <u>AP</u> | ISTITUTO BIO ITA SPA | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A065222 001</u> | Nov 29, 2005 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A065222 002</u> | Nov 29, 2005 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u> | <u>A065314 001</u> | Nov 27, 2006 |
| <u>AP</u> | MUSTAFA NEVZAT ILAC | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A065316 001</u> | Jun 29, 2007 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A065316 002</u> | Jun 29, 2007 |
| <u>AP</u> | MYLAN LABS LTD | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A201024 001</u> | Apr 07, 2014 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A201024 002</u> | Apr 07, 2014 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u> | <u>A202197 001</u> | Apr 07, 2014 |
| <u>AP</u> | SANDOZ | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A065241 001</u> | Jul 25, 2006 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A065310 001</u> | Jul 25, 2006 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A065241 002</u> | Jul 25, 2006 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A065310 002</u> | Jul 25, 2006 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u> | <u>A065240 001</u> | Jul 25, 2006 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A065074 001</u> | Mar 19, 2002 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A065074 002</u> | Mar 19, 2002 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u> | <u>A065076 001</u> | Mar 19, 2002 |

UNASYN

| | | | | |
|-----------|---|--------------------------------------------|--------------------|--------------|
| <u>AP</u> | ! | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>A062901 002</u> | Feb 27, 1992 |
| <u>AP</u> | ! | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A062901 001</u> | Nov 23, 1988 |
| <u>AP</u> | + | <u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>N050608 002</u> | Dec 31, 1986 |
| <u>AP</u> | + | <u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u> | <u>N050608 001</u> | Dec 31, 1986 |

PRESCRIPTION DRUG PRODUCT LIST

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

UNASYN

| | | | | |
|-----------|------------|-------------------------------------------|--------------------|--------------|
| AP | + ! | EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL | N050608 005 | Dec 10, 1993 |
|-----------|------------|-------------------------------------------|--------------------|--------------|

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMPICILLIN TRIHYDRATE

| | | | | |
|-----------|-----------------|----------------------|--------------------|--------------|
| AB | DAVA PHARMS INC | EQ 250MG BASE | A062883 001 | Feb 25, 1988 |
| AB | ! | EQ 500MG BASE | A062882 001 | Feb 25, 1988 |
| AB | SANDOZ | EQ 250MG BASE | A064082 001 | Aug 29, 1995 |
| AB | | EQ 500MG BASE | A064082 002 | Aug 29, 1995 |

FOR SUSPENSION; ORAL

AMPICILLIN TRIHYDRATE

DAVA PHARMS INC

EQ 125MG BASE/5ML

A062982 001 Feb 10, 1989

!

EQ 250MG BASE/5ML

A062982 002 Feb 10, 1989

ANAGRELIDE HYDROCHLORIDE

CAPSULE; ORAL

AGRYLIN

| | | | | |
|-----------|-----------|----------------------|--------------------|--------------|
| AB | SHIRE LLC | EQ 0.5MG BASE | N020333 001 | Mar 14, 1997 |
|-----------|-----------|----------------------|--------------------|--------------|

ANAGRELIDE HYDROCHLORIDE

| | | | | |
|-----------|----------------------|----------------------|--------------------|--------------|
| AB | BARR | EQ 0.5MG BASE | A076530 001 | Apr 18, 2005 |
| AB | | EQ 1MG BASE | A076530 002 | Apr 18, 2005 |
| AB | IMPAX LABS | EQ 0.5MG BASE | A076910 001 | Apr 18, 2005 |
| AB | | EQ 1MG BASE | A076910 002 | Apr 18, 2005 |
| AB | IVAX SUB TEVA PHARMS | EQ 0.5MG BASE | A076468 001 | Apr 18, 2005 |
| AB | ! | EQ 1MG BASE | A076468 002 | Apr 18, 2005 |
| AB | TORRENT PHARMS LTD | EQ 0.5MG BASE | A209151 001 | Jun 30, 2017 |
| AB | | EQ 1MG BASE | A209151 002 | Jun 30, 2017 |

ANASTROZOLE

TABLET; ORAL

ANASTROZOLE

| | | | | | |
|-----------|----------------------|-----------------------|--------------------|--------------------|--------------|
| AB | ACCORD HLTHCARE | 1MG | A090568 001 | Jun 28, 2010 | |
| AB | APOTEX INC | 1MG | A200654 001 | May 11, 2012 | |
| AB | BEIJING YILING | 1MG | A206037 001 | Nov 09, 2018 | |
| AB | BOSCOGEN | 1MG | A078944 001 | Jun 28, 2010 | |
| AB | CIPLA | 1MG | A091164 001 | Jun 28, 2010 | |
| AB | FRESENIUS KABI ONCOL | 1MG | A090088 001 | Jun 28, 2010 | |
| AB | MYLAN | 1MG | A091051 001 | Jun 28, 2010 | |
| AB | NATCO PHARMA LTD | 1MG | A079220 001 | Jun 28, 2010 | |
| AB | NEOPHARMA | 1MG | A090732 001 | Jun 28, 2010 | |
| AB | TEVA PHARMS | 1MG | A078058 001 | Jun 28, 2010 | |
| AB | WEST-WARD PHARMS INT | 1MG | A078485 001 | Jun 28, 2010 | |
| AB | ZYDUS PHARMS USA INC | 1MG | A078921 001 | Jun 28, 2010 | |
| AB | + ! | ANI PHARMS INC | 1MG | N020541 001 | Dec 27, 1995 |

ANGIOTENSIN II ACETATE

SOLUTION; INTRAVENOUS

GIAPREZA

+! LA JOLLA PHARMA

EQ 2.5MG BASE/ML (EQ 2.5MG BASE/ML)

N209360 001 Dec 21, 2017

ANIDULAFUNGIN

POWDER; INTRAVENOUS

ERAXIS

+! VICURON

50MG/VIAL

N021632 001 Feb 17, 2006

+!

100MG/VIAL

N021632 002 Nov 14, 2006

APALUTAMIDE

TABLET; ORAL

ERLEADA

+! JANSSEN BIOTECH

60MG

N210951 001 Feb 14, 2018

APIXABAN

TABLET; ORAL

ELIQUIS

+ BRISTOL MYERS

2.5MG

N202155 001 Dec 28, 2012

SQUIBB

+!

5MG

N202155 002 Dec 28, 2012

PRESCRIPTION DRUG PRODUCT LIST

APOMORPHINE HYDROCHLORIDE

INJECTABLE; SUBCUTANEOUS

APOKYN

+! US WORLDMEDS 30MG/3ML (10MG/ML) N021264 002 Apr 20, 2004

APRACLONIDINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

APRACLONIDINE HYDROCHLORIDE**AT** AKORN INC **EQ 0.5% BASE** **A077764 001** Mar 12, 2009IOPIDINE**AT** +! NOVARTIS PHARMS **EQ 0.5% BASE** **N020258 001** Jul 30, 1993

CORP

+! EQ 1% BASE N019779 001 Dec 31, 1987

APREMILAST

TABLET; ORAL

OTEZLA

+ CELGENE CORP 10MG N205437 001 Mar 21, 2014

+ 20MG N205437 002 Mar 21, 2014

+! 30MG N205437 003 Mar 21, 2014

APREPITANT

CAPSULE; ORAL

APREPITANT**AB** GLENMARK PHARMS SA **40MG** **A207777 001** Oct 12, 2017**AB** **80MG** **A207777 002** Oct 12, 2017**AB** **125MG** **A207777 003** Oct 12, 2017**AB** SANDOZ **40MG** **A090999 001** Sep 24, 2012**AB** **80MG** **A090999 002** Sep 24, 2012**AB** **125MG** **A090999 003** Sep 24, 2012EMEND**AB** + MERCK **40MG** **N021549 003** Jun 30, 2006**AB** + **80MG** **N021549 001** Mar 26, 2003**AB** +! **125MG** **N021549 002** Mar 26, 2003

EMULSION; INTRAVENOUS

CINVANTI

+! HERON THERAPS INC 130MG/18ML (7.2MG/ML) N209296 001 Nov 09, 2017

FOR SUSPENSION; ORAL

EMEND

+! MSD MERCK CO 125MG/KIT N207865 001 Dec 17, 2015

ARFORMOTEROL TARTRATE

SOLUTION; INHALATION

BROVANA

+! SUNOVION EQ 0.015MG BASE/2ML N021912 001 Oct 06, 2006

ARGATROBAN

INJECTABLE; INJECTION

ARGATROBAN**AP** AMNEAL PHARMS CO **250MG/2.5ML (100MG/ML)** **A206698 001** Jan 26, 2018**AP** FRESENIUS KABI USA **250MG/2.5ML (100MG/ML)** **N201811 001** Mar 23, 2015**AP** HIKMA PHARM CO LTD **250MG/2.5ML (100MG/ML)** **N203049 001** Jan 05, 2012**AP** HOSPIRA INC **250MG/2.5ML (100MG/ML)** **A204120 001** Sep 21, 2016**AP** MYLAN INSTITUTIONAL **250MG/2.5ML (100MG/ML)** **A202626 001** Jun 30, 2014**AP** +! NOVARTIS PHARMS **250MG/2.5ML (100MG/ML)** **N020883 001** Jun 30, 2000

CORP

AP PAR STERILE **250MG/2.5ML (100MG/ML)** **A091665 001** Jun 30, 2014

PRODUCTS

+! HIKMA PHARM CO LTD 50MG/50ML (1MG/ML) N203049 002 Sep 30, 2016

INJECTABLE; INTRAVENOUS

ARGATROBAN IN SODIUM CHLORIDE**AP** GLAND PHARMA LTD **125MG/125ML (1MG/ML)** **A205570 001** May 22, 2017**AP** +! SANDOZ **125MG/125ML (1MG/ML)** **N022485 001** May 09, 2011

ARGATROBAN IN 0.9% SODIUM CHLORIDE

TEVA PHARMS USA 250MG/250ML (1MG/ML) N206769 001 Dec 15, 2014

ARGATROBAN IN SODIUM CHLORIDE

+! EAGLE PHARMS 50MG/50ML (1MG/ML) N022434 001 Jun 29, 2011

SOLUTION; INTRAVENOUS

ARGATROBAN IN SODIUM CHLORIDE

AUROBINDO PHARMA 50MG/50ML (1MG/ML) N209552 001 Nov 27, 2018

LTD

PRESCRIPTION DRUG PRODUCT LIST

ARGININE HYDROCHLORIDE

INJECTABLE; INJECTION

R-GENE 10

+! PHARMACIA AND UPJOHN 10GM/100ML

N016931 001

ARIPIRAZOLE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

ABILIFY MAINTENA KIT

+ OTSUKA PHARM CO LTD 300MG/VIAL
+ 300MG
+! 400MG/VIAL
+ 400MGN202971 001 Feb 28, 2013
N202971 003 Sep 29, 2014
N202971 002 Feb 28, 2013
N202971 004 Sep 29, 2014

SOLUTION; ORAL

ARIPIRAZOLEAA ! AMNEAL PHARMS 1MG/ML A203906 001 Aug 14, 2015AA APOTEX INC 1MG/ML A204094 001 Sep 30, 2015AA LANNETT CO INC 1MG/ML A204171 001 Aug 14, 2015

TABLET; ORAL

ABILIFYAB + OTSUKA 2MG N021436 006 Nov 15, 2002AB +! 5MG N021436 005 Nov 15, 2002AB +! 10MG N021436 001 Nov 15, 2002AB + 15MG N021436 002 Nov 15, 2002AB + 20MG N021436 003 Nov 15, 2002AB + 30MG N021436 004 Nov 15, 2002ARIPIRAZOLEAB ACCORD HLTHCARE 2MG A206251 001 Dec 07, 2016AB 5MG A206251 002 Dec 07, 2016AB 10MG A206251 003 Dec 07, 2016AB 15MG A206251 004 Dec 07, 2016AB 20MG A206251 005 Dec 07, 2016AB 30MG A206251 006 Dec 07, 2016AB AJANTA PHARMA LTD 2MG A206174 001 Sep 12, 2016AB 5MG A206174 002 Sep 12, 2016AB 10MG A206174 003 Sep 12, 2016AB 15MG A206174 004 Sep 12, 2016AB 20MG A206174 005 Sep 12, 2016AB 30MG A206174 006 Sep 12, 2016AB ALEMBIC PHARMS LTD 2MG A202101 001 Apr 28, 2015AB 5MG A202101 002 Apr 28, 2015AB 10MG A202101 003 Apr 28, 2015AB 15MG A202101 004 Apr 28, 2015AB 20MG A202101 005 Apr 28, 2015AB 30MG A202101 006 Apr 28, 2015AB AMNEAL PHARMS 2MG A204838 001 Jun 17, 2016AB 5MG A204838 002 Jun 17, 2016AB 10MG A204838 003 Jun 17, 2016AB 15MG A204838 004 Jun 17, 2016AB 20MG A204838 005 Jun 17, 2016AB 30MG A204838 006 Jun 17, 2016AB APOTEX INC 2MG A078583 001 Jul 24, 2015AB 5MG A078583 002 Jul 24, 2015AB 10MG A078583 003 Jul 24, 2015AB 15MG A078583 004 Jul 24, 2015AB 20MG A078583 005 Jul 24, 2015AB 30MG A078583 006 Jul 24, 2015AB AUROBINDO PHARMA LTD 2MG A203908 001 Oct 08, 2015AB 5MG A203908 002 Oct 08, 2015AB 10MG A203908 003 Oct 08, 2015AB 15MG A203908 004 Oct 08, 2015AB 20MG A203908 005 Oct 08, 2015AB 30MG A203908 006 Oct 08, 2015AB BOSCOGEN 2MG A091279 001 Jan 09, 2017AB 5MG A091279 002 Jan 09, 2017AB 10MG A091279 003 Jan 09, 2017AB 15MG A091279 004 Jan 09, 2017AB 20MG A091279 005 Jan 09, 2017AB 30MG A091279 006 Jan 09, 2017AB HETERO LABS LTD V 2MG A205064 001 Apr 28, 2015AB 5MG A205064 002 Apr 28, 2015AB 10MG A205064 003 Apr 28, 2015

PRESCRIPTION DRUG PRODUCT LIST

ARIPIPIRAZOLE

TABLET; ORAL

ARIPIPIRAZOLE

| | | | | | |
|-----------|---------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | <u>15MG</u> | <u>A205064</u> | <u>004</u> | Apr 28, 2015 |
| <u>AB</u> | | <u>20MG</u> | <u>A205064</u> | <u>005</u> | Apr 28, 2015 |
| <u>AB</u> | | <u>30MG</u> | <u>A205064</u> | <u>006</u> | Apr 28, 2015 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>2MG</u> | <u>A204111</u> | <u>001</u> | Oct 07, 2016 |
| <u>AB</u> | | <u>5MG</u> | <u>A204111</u> | <u>002</u> | Oct 07, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A204111</u> | <u>003</u> | Oct 07, 2016 |
| <u>AB</u> | | <u>15MG</u> | <u>A204111</u> | <u>004</u> | Oct 07, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A204111</u> | <u>005</u> | Oct 07, 2016 |
| <u>AB</u> | | <u>30MG</u> | <u>A204111</u> | <u>006</u> | Oct 07, 2016 |
| <u>AB</u> | ORCHID HLTHCARE | <u>2MG</u> | <u>A202683</u> | <u>001</u> | May 23, 2017 |
| <u>AB</u> | | <u>5MG</u> | <u>A202683</u> | <u>002</u> | May 23, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A202683</u> | <u>003</u> | May 23, 2017 |
| <u>AB</u> | | <u>15MG</u> | <u>A202683</u> | <u>004</u> | May 23, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A202683</u> | <u>005</u> | May 23, 2017 |
| <u>AB</u> | | <u>30MG</u> | <u>A202683</u> | <u>006</u> | May 23, 2017 |
| <u>AB</u> | PRINSTON INC | <u>2MG</u> | <u>A205363</u> | <u>001</u> | Dec 04, 2017 |
| <u>AB</u> | | <u>5MG</u> | <u>A205363</u> | <u>002</u> | Dec 04, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A205363</u> | <u>003</u> | Dec 04, 2017 |
| <u>AB</u> | | <u>15MG</u> | <u>A205363</u> | <u>004</u> | Dec 04, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A205363</u> | <u>005</u> | Dec 04, 2017 |
| <u>AB</u> | | <u>30MG</u> | <u>A205363</u> | <u>006</u> | Dec 04, 2017 |
| <u>AB</u> | SCIEGEN PHARMS INC | <u>2MG</u> | <u>A206383</u> | <u>001</u> | Sep 29, 2016 |
| <u>AB</u> | | <u>5MG</u> | <u>A206383</u> | <u>002</u> | Sep 29, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A206383</u> | <u>003</u> | Sep 29, 2016 |
| <u>AB</u> | | <u>15MG</u> | <u>A206383</u> | <u>004</u> | Sep 29, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A206383</u> | <u>005</u> | Sep 29, 2016 |
| <u>AB</u> | | <u>30MG</u> | <u>A206383</u> | <u>006</u> | Sep 29, 2016 |
| <u>AB</u> | TEVA PHARMS USA | <u>2MG</u> | <u>A078607</u> | <u>001</u> | Apr 28, 2015 |
| <u>AB</u> | | <u>5MG</u> | <u>A078607</u> | <u>002</u> | Apr 28, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A078608</u> | <u>001</u> | Apr 28, 2015 |
| <u>AB</u> | | <u>15MG</u> | <u>A078708</u> | <u>001</u> | Apr 28, 2015 |
| <u>AB</u> | | <u>20MG</u> | <u>A078708</u> | <u>002</u> | Apr 28, 2015 |
| <u>AB</u> | | <u>30MG</u> | <u>A078708</u> | <u>003</u> | Apr 28, 2015 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>2MG</u> | <u>A201519</u> | <u>001</u> | Apr 28, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A201519</u> | <u>003</u> | Apr 28, 2015 |
| <u>AB</u> | | <u>5MG</u> | <u>A201519</u> | <u>002</u> | Apr 28, 2015 |
| <u>AB</u> | | <u>15MG</u> | <u>A201519</u> | <u>004</u> | Apr 28, 2015 |
| <u>AB</u> | | <u>20MG</u> | <u>A201519</u> | <u>005</u> | Apr 28, 2015 |
| <u>AB</u> | | <u>30MG</u> | <u>A201519</u> | <u>006</u> | Apr 28, 2015 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>2MG</u> | <u>A090472</u> | <u>001</u> | Jan 07, 2019 |
| <u>AB</u> | INC | <u>5MG</u> | <u>A090472</u> | <u>002</u> | Jan 07, 2019 |
| <u>AB</u> | | <u>10MG</u> | <u>A090472</u> | <u>003</u> | Jan 07, 2019 |
| <u>AB</u> | | <u>15MG</u> | <u>A090472</u> | <u>004</u> | Jan 07, 2019 |
| <u>AB</u> | | <u>20MG</u> | <u>A090472</u> | <u>005</u> | Jan 07, 2019 |
| <u>AB</u> | | <u>30MG</u> | <u>A090472</u> | <u>006</u> | Jan 07, 2019 |

ABILIFY MYCITE KIT

| | | | | | |
|---|---------------------|------|---------|-----|--------------|
| + | OTSUKA PHARM CO LTD | 2MG | N207202 | 001 | Nov 13, 2017 |
| + | ! | 5MG | N207202 | 002 | Nov 13, 2017 |
| + | | 10MG | N207202 | 003 | Nov 13, 2017 |
| + | | 15MG | N207202 | 004 | Nov 13, 2017 |
| + | | 20MG | N207202 | 005 | Nov 13, 2017 |
| + | | 30MG | N207202 | 006 | Nov 13, 2017 |

TABLET, ORALLY DISINTEGRATING; ORAL

ARIPIPIRAZOLE

| | | | | | | |
|-----------|---|--------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | ! | ALEMBIC PHARMS LTD | <u>10MG</u> | <u>A202102</u> | <u>001</u> | Apr 28, 2015 |
| <u>AB</u> | | | <u>15MG</u> | <u>A202102</u> | <u>002</u> | Apr 28, 2015 |
| <u>AB</u> | | ORCHID HLTHCARE | <u>10MG</u> | <u>A202547</u> | <u>001</u> | Dec 11, 2017 |
| <u>AB</u> | | | <u>15MG</u> | <u>A202547</u> | <u>002</u> | Dec 11, 2017 |
| <u>AB</u> | | SCIEGEN PHARMS INC | <u>10MG</u> | <u>A207240</u> | <u>001</u> | Apr 18, 2018 |
| <u>AB</u> | | | <u>15MG</u> | <u>A207240</u> | <u>002</u> | Apr 18, 2018 |
| <u>AB</u> | | ZYDUS PHARMS USA | <u>10MG</u> | <u>A090165</u> | <u>001</u> | Aug 28, 2018 |
| <u>AB</u> | | INC | <u>15MG</u> | <u>A090165</u> | <u>002</u> | Aug 28, 2018 |
| | | | 20MG | A090165 | 003 | Aug 28, 2018 |
| | | | 30MG | A090165 | 004 | Aug 28, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

ARIPRAZOLE LAUROXIL

SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

ARISTADA

| | | | | | |
|---------------------|--------------|----------------------------|---------|-----|--------------|
| + | ALKERMES INC | 441MG/1.6ML (275.63MG/ML) | N207533 | 001 | Oct 05, 2015 |
| + | | 662MG/2.4ML (275.83MG/ML) | N207533 | 002 | Oct 05, 2015 |
| + | ! | 882MG/3.2ML (275.63MG/ML) | N207533 | 003 | Oct 05, 2015 |
| + | | 1064MG/3.9ML (272.82MG/ML) | N207533 | 004 | Jun 05, 2017 |
| ARISTADA INITIO KIT | | | | | |
| + | ALKERMES INC | 675MG/2.4ML | N209830 | 001 | Jun 29, 2018 |

ARMODAFINIL

TABLET; ORAL

ARMODAFINIL

| | | | | | |
|----------------|----------------------|--------------|----------------|------------|--------------|
| AB | AUROBINDO PHARMA LTD | 50MG | A206069 | 001 | Mar 06, 2018 |
| AB | | 150MG | A206069 | 002 | Mar 06, 2018 |
| AB | | 250MG | A206069 | 003 | Mar 06, 2018 |
| AB | LUPIN LTD | 50MG | A200751 | 001 | Nov 28, 2016 |
| AB | | 150MG | A200751 | 003 | Nov 28, 2016 |
| AB | | 200MG | A200751 | 004 | Nov 28, 2016 |
| AB | | 250MG | A200751 | 005 | Nov 28, 2016 |
| AB | MYLAN PHARMS INC | 50MG | A200043 | 001 | Jun 01, 2012 |
| AB | | 150MG | A200043 | 002 | Jun 01, 2012 |
| AB | | 250MG | A200043 | 003 | Jun 01, 2012 |
| AB | NATCO PHARMA LTD | 50MG | A202768 | 001 | Nov 28, 2016 |
| AB | | 150MG | A202768 | 002 | Nov 28, 2016 |
| AB | | 200MG | A202768 | 005 | Sep 28, 2017 |
| AB | | 250MG | A202768 | 003 | Nov 28, 2016 |
| <u>NUVIGIL</u> | | | | | |
| AB | + CEPHALON | 50MG | N021875 | 001 | Jun 15, 2007 |
| AB | + | 150MG | N021875 | 003 | Jun 15, 2007 |
| AB | + | 200MG | N021875 | 005 | Mar 26, 2009 |
| AB | + | 250MG | N021875 | 004 | Jun 15, 2007 |
| ARMODAFINIL | | | | | |
| | NATCO PHARMA LTD | 100MG | A202768 | 004 | Sep 28, 2017 |

ARSENIC TRIOXIDE

INJECTABLE; INJECTION

ARSENIC TRIOXIDE

| | | | | | |
|-----------|----------------------|---------------|----------------|------------|--------------|
| AP | AMRING PHARMS | 1MG/ML | A210802 | 001 | Nov 13, 2018 |
| AP | FRESENIUS KABI USA | 1MG/ML | A208231 | 001 | Aug 31, 2018 |
| AP | INGENUS PHARMS LLC | 1MG/ML | A209315 | 001 | Nov 15, 2018 |
| AP | NEXUS PHARMS | 1MG/ML | A209780 | 001 | Nov 15, 2018 |
| AP | ZYDUS PHARMS USA INC | 1MG/ML | A206228 | 001 | Nov 13, 2018 |
| TRISENOX | | | | | |
| + | CEPHALON | 2MG/ML | N021248 | 002 | Oct 13, 2017 |

ARTEMETHER; LUMEFANTRINE

TABLET; ORAL

COARTEM

| | | | | | |
|---|----------|-------------|---------|-----|--------------|
| + | NOVARTIS | 20MG; 120MG | N022268 | 001 | Apr 07, 2009 |
|---|----------|-------------|---------|-----|--------------|

ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

ARTICAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE

| | | | | | |
|----------------------|--------------|----------------------------------------------------------|----------------|------------|--------------|
| AP | HOSPIRA | 4%;EQ 0.017MG BASE/1.7ML (4%;EQ 0.01MG BASE/ML) | A079138 | 001 | Jun 18, 2010 |
| <u>SEPTOCAINE</u> | | | | | |
| AP | +! DEPROCO | 4%;EQ 0.0085MG BASE/1.7ML (4%;EQ 0.005MG BASE/ML) | N022010 | 001 | Mar 30, 2006 |
| AP | + | 4%;EQ 0.017MG BASE/1.7ML (4%;EQ 0.01MG BASE/ML) | N020971 | 001 | Apr 03, 2000 |
| <u>ULTACAN</u> | | | | | |
| AP | HANSAMED INC | 4%;EQ 0.0085MG BASE/1.7ML (4%;EQ 0.005MG BASE/ML) | A201751 | 001 | Jul 11, 2017 |
| <u>ULTACAN FORTE</u> | | | | | |
| AP | HANSAMED INC | 4%;EQ 0.017MG BASE/1.7ML (4%;EQ 0.01MG BASE/ML) | A201750 | 001 | Jul 11, 2017 |
| ORABLOC | | | | | |
| + | PIERREL | 4%;EQ 0.009MG BASE/1.8ML (EQ 0.005MG BASE/ML) | N022466 | 001 | Feb 26, 2010 |
| + | ! | 4%;EQ 0.018MG BASE/1.8ML (EQ 0.01MG BASE/ML) | N022466 | 002 | Feb 26, 2010 |

PRESCRIPTION DRUG PRODUCT LIST

ASCORBIC ACID

SOLUTION; INTRAVENOUS

ASCOR

+! MCGUFF 25,000MG/50ML (500MG/ML) N209112 001 Oct 02, 2017

ASCORBIC ACID; BIOTIN; CHOLECALCIFEROL; CYANOCOBALAMIN; DEXPANTHENOL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE; RIBOFLAVIN; THIAMINE; TOCOPHEROL ACETATE; VITAMIN A; VITAMIN K

INJECTABLE; INTRAVENOUS

INFUVITE PEDIATRIC

+! SANDOZ INC 80MG/VIAL; 0.02MG/VIAL; 400 IU/VIAL; 0.001MG/VIAL; 5MG/VIAL; 0.14MG/VIAL; 17MG/VIAL; 1MG/VIAL; 1.4MG/VIAL; 1.2MG/VIAL; 7 IU/VIAL; 2,300 IU/VIAL; 0.2MG/VIAL N021265 001 Feb 21, 2001

INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE)

+! SANDOZ INC 80MG/VIAL; 0.02MG/VIAL; 400 IU/VIAL; 0.001MG/VIAL; 5MG/VIAL; 0.14MG/VIAL; 17MG/VIAL; 1MG/VIAL; 1.4MG/VIAL; 1.2MG/VIAL; 7 IU/VIAL; 2,300 IU/VIAL; 0.2MG/VIAL N021646 001 Jan 29, 2004

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PHYTONADIONE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

FOR SOLUTION; INTRAVENOUS

M.V.I. PEDIATRIC

+! HOSPIRA 80MG/VIAL; 0.02MG/VIAL; 0.001MG/VIAL; 5MG/VIAL; 0.01MG/VIAL; 0.14MG/VIAL; 17MG/VIAL; 0.2MG/VIAL; 1MG/VIAL; 1.4MG/VIAL; EQ 1.2MG BASE/VIAL; 0.7MG/VIAL; 7MG/VIAL N018920 001 Sep 21, 2000

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E; VITAMIN K

INJECTABLE; INTRAVENOUS

M.V.I. ADULT

+! HOSPIRA 200MG/VIAL; 0.06MG/VIAL; 0.005MG/VIAL; 15MG/VIAL; 0.005MG/VIAL; 0.6MG/VIAL; 40MG/VIAL; 6MG/VIAL; 3.6MG/VIAL; 6MG/VIAL; 1MG/VIAL; 10MG/VIAL; 0.15MG/VIAL N021625 001 Jan 30, 2004

M.V.I. ADULT (PHARMACY BULK PACKAGE)

+! HOSPIRA 200MG/5ML; 0.06MG/5ML; 0.005MG/5ML; 15MG/5ML; 0.005MG/5ML; 0.6MG/5ML; 40MG/5ML; 6MG/5ML; 3.6MG/5ML; 6MG/5ML; 1MG/5ML; 10MG/5ML; 0.15MG/5ML N021643 001 Feb 18, 2004

ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE

FOR SOLUTION; ORAL

MOVIPREP

+! SALIX PHARMS 4.7GM; 100GM; 1.015GM; 5.9GM; 2.691GM; 7.5GM N021881 001 Aug 02, 2006

PLENVU

+! SALIX PHARMS INC 7.54GM; 140GM; 2.2GM; 48.11GM; 5.2GM; 9GM N209381 001 May 04, 2018

ASENAPINE MALEATE

TABLET; SUBLINGUAL

ASENAPINE MALEATEAB SIGMAPHARM LABS LLC EQ 5MG BASE A206107 001 Jul 17, 2018AB EQ 10MG BASE A206107 002 Jul 17, 2018SAPHRISAB + FOREST LABS LLC EQ 5MG BASE N022117 001 Aug 13, 2009AB +! EQ 10MG BASE N022117 002 Aug 13, 2009

+ EQ 2.5 BASE N022117 003 Mar 12, 2015

ASPIRIN

CAPSULE, EXTENDED RELEASE; ORAL

DURLAZA

+! ESPERO 162.5MG N200671 001 Sep 04, 2015

ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

FIORINALAA +! ALLERGAN SALES LLC 325MG; 50MG; 40MG N017534 005 Apr 16, 1986LANORINALAA LANNETT 325MG; 50MG; 40MG A086996 002 Oct 11, 1985

TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINEAA ! HIKMA INTL PHARMS 325MG; 50MG; 40MG A086162 002 Feb 16, 1984AA PII 325MG; 50MG; 40MG A204195 001 Sep 22, 2016

PRESCRIPTION DRUG PRODUCT LIST

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE;ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

| | | | | | |
|-----------|-------------------|-----------------------------|----------------|------------|--------------|
| AB | MAYNE PHARMA INC | <u>325MG;50MG;40MG;30MG</u> | <u>A203335</u> | <u>001</u> | Oct 30, 2015 |
| AB | NEXGEN PHARMA INC | <u>325MG;50MG;40MG;30MG</u> | <u>A075231</u> | <u>001</u> | Nov 30, 2001 |
| AB | STEVENS J | <u>325MG;50MG;40MG;30MG</u> | <u>A074951</u> | <u>001</u> | Aug 31, 1998 |

FIORINAL W/CODEINE

| | | | | | |
|-----------|-------------------------------|-----------------------------|----------------|------------|--------------|
| AB | + ! ALLERGAN SALES LLC | <u>325MG;50MG;40MG;30MG</u> | <u>N019429</u> | <u>003</u> | Oct 26, 1990 |
|-----------|-------------------------------|-----------------------------|----------------|------------|--------------|

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET;ORAL

ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE

| | | | | | |
|--|--------|-----------------|---------|-----|--------------|
| | SANDOZ | 385MG;30MG;25MG | A074654 | 001 | Dec 31, 1996 |
| | ! | 770MG;60MG;50MG | A074654 | 002 | Dec 31, 1996 |

ASPIRIN; CARISOPRODOL

TABLET;ORAL

CARISOPRODOL AND ASPIRIN

| | | | | | |
|-----------|------------------------------|--------------------|----------------|------------|--------------|
| AB | ! HERITAGE PHARMS INC | <u>325MG;200MG</u> | <u>A089594</u> | <u>001</u> | Mar 31, 1989 |
| AB | NOVAST LABS | <u>325MG;200MG</u> | <u>A040832</u> | <u>001</u> | Jan 07, 2010 |
| AB | SANDOZ | <u>325MG;200MG</u> | <u>A040116</u> | <u>001</u> | Apr 25, 1996 |

ASPIRIN; CARISOPRODOL; CODEINE PHOSPHATE

TABLET;ORAL

CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE

| | | | | | |
|-----------|-------------------|-------------------------|----------------|------------|--------------|
| AB | INGENUS PHARMS NJ | <u>325MG;200MG;16MG</u> | <u>A040860</u> | <u>001</u> | Jan 07, 2010 |
| AB | ! SANDOZ | <u>325MG;200MG;16MG</u> | <u>A040118</u> | <u>001</u> | Apr 16, 1996 |

ASPIRIN; DIPYRIDAMOLE

CAPSULE, EXTENDED RELEASE;ORAL

AGGRENOX

| | | | | | |
|-----------|------------------------------------|-------------------|----------------|------------|--------------|
| AB | + ! BOEHRINGER INGELHEIM | <u>25MG;200MG</u> | <u>N020884</u> | <u>001</u> | Nov 22, 1999 |
|-----------|------------------------------------|-------------------|----------------|------------|--------------|

ASPIRIN AND DIPYRIDAMOLE

| | | | | | |
|-----------|-------------------------|-------------------|----------------|------------|--------------|
| AB | AMNEAL PHARMS | <u>25MG;200MG</u> | <u>A206392</u> | <u>001</u> | Mar 08, 2016 |
| AB | ANI PHARMS INC | <u>25MG;200MG</u> | <u>A206964</u> | <u>001</u> | Jan 18, 2017 |
| AB | BARR | <u>25MG;200MG</u> | <u>A078804</u> | <u>001</u> | Aug 14, 2009 |
| AB | DR REDDYS LABS LTD | <u>25MG;200MG</u> | <u>A209048</u> | <u>001</u> | Oct 10, 2018 |
| AB | PAR PHARM INC | <u>25MG;200MG</u> | <u>A207944</u> | <u>001</u> | Jan 18, 2017 |
| AB | SANDOZ INC | <u>25MG;200MG</u> | <u>A206739</u> | <u>001</u> | Jan 18, 2017 |
| AB | SUN PHARMA GLOBAL | <u>25MG;200MG</u> | <u>A208572</u> | <u>001</u> | Aug 21, 2018 |
| AB | ZYDUS PHARMS USA INC | <u>25MG;200MG</u> | <u>A206753</u> | <u>001</u> | Aug 29, 2017 |

ASPIRIN; METHOCARBAMOL

TABLET;ORAL

METHOCARBAMOL AND ASPIRIN

| | | | | | | |
|--|---|-----------|-------------|---------|-----|--------------|
| | ! | STEVENS J | 325MG;400MG | A081145 | 001 | Jan 31, 1995 |
|--|---|-----------|-------------|---------|-----|--------------|

ASPIRIN; OMEPRAZOLE

TABLET, DELAYED RELEASE;ORAL

YOSPRALA

| | | | | | | |
|--|------------|--------------------|------------|---------|-----|--------------|
| | + | GENUS LIFESCIENCES | 81MG;40MG | N205103 | 001 | Sep 14, 2016 |
| | + ! | | 325MG;40MG | N205103 | 002 | Sep 14, 2016 |

ASPIRIN; OXYCODONE HYDROCHLORIDE

TABLET;ORAL

OXYCODONE AND ASPIRIN

| | | | | | |
|-----------|------------------------|-----------------------|----------------|------------|--------------|
| AA | ACTAVIS LABS FL INC | <u>325MG;4.8355MG</u> | <u>A090084</u> | <u>001</u> | Mar 22, 2011 |
| AA | MAYNE PHARMA INC | <u>325MG;4.8355MG</u> | <u>A091670</u> | <u>001</u> | Mar 16, 2011 |
| AA | + ! ENDO PHARMS | <u>325MG;4.8355MG</u> | <u>N007337</u> | <u>007</u> | Aug 05, 2005 |

ATAZANAVIR SULFATE

CAPSULE;ORAL

ATAZANAVIR SULFATE

| | | | | | |
|-----------|-------------------------|----------------------|----------------|------------|--------------|
| AB | AUROBINDO PHARMA LTD | <u>EQ 100MG BASE</u> | <u>A204806</u> | <u>001</u> | Jun 25, 2018 |
| AB | | <u>EQ 150MG BASE</u> | <u>A204806</u> | <u>002</u> | Jun 25, 2018 |
| AB | | <u>EQ 200MG BASE</u> | <u>A204806</u> | <u>003</u> | Jun 25, 2018 |
| AB | | <u>EQ 300MG BASE</u> | <u>A204806</u> | <u>004</u> | Jun 25, 2018 |
| AB | CIPLA | <u>EQ 100MG BASE</u> | <u>A200626</u> | <u>001</u> | Aug 09, 2018 |
| AB | | <u>EQ 150MG BASE</u> | <u>A200626</u> | <u>002</u> | Aug 09, 2018 |
| AB | | <u>EQ 200MG BASE</u> | <u>A200626</u> | <u>003</u> | Aug 09, 2018 |
| AB | | <u>EQ 300MG BASE</u> | <u>A200626</u> | <u>004</u> | Aug 09, 2018 |
| AB | MYLAN PHARMS INC | <u>EQ 150MG BASE</u> | <u>A208177</u> | <u>001</u> | Sep 24, 2018 |
| AB | | <u>EQ 200MG BASE</u> | <u>A208177</u> | <u>002</u> | Sep 24, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

ATAZANAVIR SULFATE

CAPSULE; ORAL

ATAZANAVIR SULFATE

| | | | | |
|-----------|-----------------|----------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A208177 003</u> | Sep 24, 2018 |
| <u>AB</u> | TEVA PHARMS USA | <u>EQ 100MG BASE</u> | <u>A091673 001</u> | Apr 22, 2014 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A091673 002</u> | Apr 22, 2014 |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A091673 003</u> | Apr 22, 2014 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A091673 004</u> | Apr 22, 2014 |

REYATAZ

| | | | | |
|-----------|---------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | + BRISTOL MYERS SQUIBB | <u>EQ 150MG BASE</u> | <u>N021567 002</u> | Jun 20, 2003 |
| <u>AB</u> | + | <u>EQ 200MG BASE</u> | <u>N021567 003</u> | Jun 20, 2003 |
| <u>AB</u> | +! | <u>EQ 300MG BASE</u> | <u>N021567 004</u> | Oct 16, 2006 |

POWDER; ORAL

REYATAZ

| | | | | |
|--|----------------------------|---------------------|-------------|--------------|
| | +! BRISTOL MYERS SQUIBB | EQ 50MG BASE/PACKET | N206352 001 | Jun 02, 2014 |
|--|----------------------------|---------------------|-------------|--------------|

ATAZANAVIR SULFATE; COBICISTAT

TABLET; ORAL

EVOTAZ

| | | | | |
|--|----------------------------|----------------------|-------------|--------------|
| | +! BRISTOL-MYERS SQUIBB | EQ 300MG BASE; 150MG | N206353 001 | Jan 29, 2015 |
|--|----------------------------|----------------------|-------------|--------------|

ATENOLOL

TABLET; ORAL

ATENOLOL

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | ALVOGEN MALTA | <u>25MG</u> | <u>A072304 002</u> | Jul 31, 1992 |
| <u>AB</u> | | <u>50MG</u> | <u>A072304 003</u> | Jul 18, 1988 |
| <u>AB</u> | | <u>100MG</u> | <u>A072304 001</u> | Jul 15, 1988 |
| <u>AB</u> | AUROBINDO PHARMA | <u>25MG</u> | <u>A078512 001</u> | Oct 31, 2007 |
| <u>AB</u> | | <u>50MG</u> | <u>A078512 002</u> | Oct 31, 2007 |
| <u>AB</u> | | <u>100MG</u> | <u>A078512 003</u> | Oct 31, 2007 |
| <u>AB</u> | DAVA PHARMS INC | <u>50MG</u> | <u>A073542 001</u> | Dec 19, 1991 |
| <u>AB</u> | | <u>100MG</u> | <u>A073543 001</u> | Dec 19, 1991 |
| <u>AB</u> | IPCA LABS LTD | <u>25MG</u> | <u>A077877 001</u> | Dec 27, 2006 |
| <u>AB</u> | | <u>50MG</u> | <u>A077877 002</u> | Dec 27, 2006 |
| <u>AB</u> | | <u>100MG</u> | <u>A077877 003</u> | Dec 27, 2006 |
| <u>AB</u> | MYLAN | <u>25MG</u> | <u>A073457 002</u> | Apr 26, 1999 |
| <u>AB</u> | | <u>50MG</u> | <u>A073457 003</u> | Jan 24, 1992 |
| <u>AB</u> | | <u>100MG</u> | <u>A073457 001</u> | Jan 24, 1992 |
| <u>AB</u> | SANDOZ | <u>25MG</u> | <u>A074052 001</u> | May 01, 1992 |
| <u>AB</u> | | <u>50MG</u> | <u>A073025 001</u> | Sep 17, 1991 |
| <u>AB</u> | | <u>100MG</u> | <u>A073026 001</u> | Sep 17, 1991 |
| <u>AB</u> | SUN PHARM INDS INC | <u>25MG</u> | <u>A078210 001</u> | Jul 10, 2007 |
| <u>AB</u> | | <u>50MG</u> | <u>A078210 002</u> | Jul 10, 2007 |
| <u>AB</u> | | <u>100MG</u> | <u>A078210 003</u> | Jul 10, 2007 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>25MG</u> | <u>A074499 001</u> | Jul 30, 1997 |
| <u>AB</u> | | <u>50MG</u> | <u>A073475 001</u> | Mar 30, 1993 |
| <u>AB</u> | | <u>100MG</u> | <u>A073476 001</u> | Mar 30, 1993 |
| <u>AB</u> | TEVA | <u>25MG</u> | <u>A074056 003</u> | Jul 19, 2004 |
| <u>AB</u> | | <u>50MG</u> | <u>A074056 001</u> | Jan 18, 1995 |
| <u>AB</u> | | <u>100MG</u> | <u>A074056 002</u> | Jan 18, 1995 |
| <u>AB</u> | UNIQUE PHARM LABS | <u>25MG</u> | <u>A077443 001</u> | Sep 13, 2006 |
| <u>AB</u> | | <u>50MG</u> | <u>A077443 002</u> | Sep 13, 2006 |
| <u>AB</u> | | <u>100MG</u> | <u>A077443 003</u> | Sep 13, 2006 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>25MG</u> | <u>A076900 001</u> | Jan 28, 2005 |
| <u>AB</u> | | <u>50MG</u> | <u>A076900 002</u> | Jan 28, 2005 |
| <u>AB</u> | | <u>100MG</u> | <u>A076900 003</u> | Jan 28, 2005 |

TENORMIN

| | | | | |
|-----------|-----------------|--------------|--------------------|--------------|
| <u>AB</u> | + ALVOGEN MALTA | <u>25MG</u> | <u>N018240 004</u> | Apr 09, 1990 |
| <u>AB</u> | + | <u>50MG</u> | <u>N018240 001</u> | |
| <u>AB</u> | +! | <u>100MG</u> | <u>N018240 002</u> | |

ATENOLOL; CHLORTHALIDONE

TABLET; ORAL

ATENOLOL AND CHLORTHALIDONE

| | | | | |
|-----------|-------------------------|--------------------|--------------------|--------------|
| <u>AB</u> | ALVOGEN MALTA | <u>50MG; 25MG</u> | <u>A072302 002</u> | May 31, 1990 |
| <u>AB</u> | | <u>100MG; 25MG</u> | <u>A072302 001</u> | May 31, 1990 |
| <u>AB</u> | MYLAN | <u>50MG; 25MG</u> | <u>A074203 001</u> | Oct 31, 1993 |
| <u>AB</u> | | <u>100MG; 25MG</u> | <u>A074203 002</u> | Oct 31, 1993 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>50MG; 25MG</u> | <u>A073582 002</u> | Apr 29, 1993 |
| <u>AB</u> | | <u>100MG; 25MG</u> | <u>A073582 001</u> | Apr 29, 1993 |

PRESCRIPTION DRUG PRODUCT LIST

ATENOLOL; CHLORTHALIDONE

TABLET; ORAL

ATENOLOL AND CHLORTHALIDONE

| | | | | |
|----------------------|------------------|-------------------|--------------------|--------------|
| AB | WATSON LABS | 50MG;25MG | A073665 001 | Jul 02, 1992 |
| AB | | 100MG;25MG | A073665 002 | Jul 02, 1992 |
| TENORETIC 100 | | | | |
| AB | +! ALVOGEN MALTA | 100MG;25MG | N018760 001 | Jun 08, 1984 |
| TENORETIC 50 | | | | |
| AB | + ALVOGEN MALTA | 50MG;25MG | N018760 002 | Jun 08, 1984 |

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

ATOMOXETINE HYDROCHLORIDE

| | | | | |
|------------------|----------------------|--------------|--------------------|--------------|
| AB | APOTEX INC | 10MG | A078983 001 | May 30, 2017 |
| AB | | 18MG | A078983 002 | May 30, 2017 |
| AB | | 25MG | A078983 003 | May 30, 2017 |
| AB | | 40MG | A078983 004 | May 30, 2017 |
| AB | | 60MG | A078983 005 | May 30, 2017 |
| AB | | 80MG | A078983 006 | May 30, 2017 |
| AB | | 100MG | A078983 007 | May 30, 2017 |
| AB | AUROBINDO PHARMA LTD | 10MG | A079016 001 | May 30, 2017 |
| AB | | 18MG | A079016 002 | May 30, 2017 |
| AB | | 25MG | A079016 003 | May 30, 2017 |
| AB | | 40MG | A079016 004 | May 30, 2017 |
| AB | | 60MG | A079016 005 | May 30, 2017 |
| AB | | 80MG | A079016 006 | May 30, 2017 |
| AB | | 100MG | A079016 007 | May 30, 2017 |
| AB | DR REDDYS LABS LTD | 10MG | A090609 001 | Feb 23, 2018 |
| AB | | 18MG | A090609 002 | Feb 23, 2018 |
| AB | | 25MG | A090609 003 | Feb 23, 2018 |
| AB | | 40MG | A090609 004 | Feb 23, 2018 |
| AB | | 60MG | A090609 005 | Feb 23, 2018 |
| AB | | 80MG | A090609 006 | Feb 23, 2018 |
| AB | | 100MG | A090609 007 | Feb 23, 2018 |
| AB | GLENMARK PHARMS LTD | 10MG | A079019 001 | May 30, 2017 |
| AB | | 18MG | A079019 002 | May 30, 2017 |
| AB | | 25MG | A079019 003 | May 30, 2017 |
| AB | | 40MG | A079019 004 | May 30, 2017 |
| AB | | 60MG | A079019 005 | May 30, 2017 |
| AB | | 80MG | A079019 006 | May 30, 2017 |
| AB | | 100MG | A079019 007 | May 30, 2017 |
| AB | TEVA PHARMS USA | 10MG | A079022 001 | May 30, 2017 |
| AB | | 18MG | A079022 002 | May 30, 2017 |
| AB | | 25MG | A079022 003 | May 30, 2017 |
| AB | | 40MG | A079022 004 | May 30, 2017 |
| AB | | 60MG | A079022 005 | May 30, 2017 |
| AB | | 80MG | A079022 006 | May 30, 2017 |
| AB | | 100MG | A079022 007 | May 30, 2017 |
| STRATTERA | | | | |
| AB | + LILLY | 10MG | N021411 002 | Nov 26, 2002 |
| AB | | 18MG | N021411 003 | Nov 26, 2002 |
| AB | | 25MG | N021411 004 | Nov 26, 2002 |
| AB | | 40MG | N021411 005 | Nov 26, 2002 |
| AB | +! | 60MG | N021411 006 | Nov 26, 2002 |
| AB | | 80MG | N021411 007 | Feb 14, 2005 |
| AB | | 100MG | N021411 008 | Feb 14, 2005 |

ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

| | | | | |
|-----------|--------------------|---------------------|--------------------|--------------|
| AB | ACCORD HLTHCARE | EQ 10MG BASE | A207687 001 | Mar 30, 2018 |
| AB | | EQ 20MG BASE | A207687 002 | Mar 30, 2018 |
| AB | | EQ 40MG BASE | A207687 003 | Mar 30, 2018 |
| AB | | EQ 80MG BASE | A207687 004 | Mar 30, 2018 |
| AB | APOTEX INC | EQ 10MG BASE | A090548 001 | May 29, 2012 |
| AB | | EQ 20MG BASE | A090548 002 | May 29, 2012 |
| AB | | EQ 40MG BASE | A090548 003 | May 29, 2012 |
| AB | | EQ 80MG BASE | A090548 004 | May 29, 2012 |
| AB | DR REDDYS LABS LTD | EQ 10MG BASE | A091650 001 | Jul 17, 2012 |
| AB | | EQ 20MG BASE | A091650 002 | Jul 17, 2012 |
| AB | | EQ 40MG BASE | A091650 003 | Jul 17, 2012 |
| AB | | EQ 80MG BASE | A202357 001 | Jul 17, 2012 |
| AB | GRAVITI PHARMS | EQ 10MG BASE | A209912 001 | Jun 18, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

| | | | | |
|----------------|----------------------|---------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A209912 002</u> | Jun 18, 2018 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A209912 003</u> | Jun 18, 2018 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A209912 004</u> | Jun 18, 2018 |
| <u>AB</u> | INVAGEN PHARMS | <u>EQ 10MG BASE</u> | <u>A204846 001</u> | Jan 09, 2017 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A204846 002</u> | Jan 09, 2017 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A204846 003</u> | Jan 09, 2017 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A204846 004</u> | Jan 09, 2017 |
| <u>AB</u> | LANNETT CO INC | <u>EQ 10MG BASE</u> | <u>A091624 001</u> | Apr 05, 2013 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A091624 002</u> | Apr 05, 2013 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A091624 003</u> | Apr 05, 2013 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A091624 004</u> | Apr 05, 2013 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 10MG BASE</u> | <u>A091226 001</u> | May 29, 2012 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A091226 002</u> | May 29, 2012 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A091226 003</u> | May 29, 2012 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A091226 004</u> | May 29, 2012 |
| <u>AB</u> | SANDOZ INC | <u>EQ 10MG BASE</u> | <u>A077575 001</u> | May 29, 2012 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077575 002</u> | May 29, 2012 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077575 003</u> | May 29, 2012 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A077575 004</u> | May 29, 2012 |
| <u>AB</u> | SCIEGEN PHARMS INC | <u>EQ 10MG BASE</u> | <u>A205519 001</u> | May 19, 2016 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A205519 002</u> | May 19, 2016 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A205519 003</u> | May 19, 2016 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A205519 004</u> | May 19, 2016 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>EQ 10MG BASE</u> | <u>A076477 001</u> | Nov 30, 2011 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A076477 002</u> | Nov 30, 2011 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A076477 003</u> | Nov 30, 2011 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A076477 004</u> | Nov 30, 2011 |
| <u>AB</u> | TEVA PHARMS USA | <u>EQ 10MG BASE</u> | <u>A205300 001</u> | Mar 27, 2017 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A205300 002</u> | Mar 27, 2017 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A205300 003</u> | Mar 27, 2017 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A205300 004</u> | Mar 27, 2017 |
| <u>AB</u> | THEPHARMANETWORK LLC | <u>EQ 10MG BASE</u> | <u>A209288 001</u> | Dec 21, 2018 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A209288 002</u> | Dec 21, 2018 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A209288 003</u> | Dec 21, 2018 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A209288 004</u> | Dec 21, 2018 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 10MG BASE</u> | <u>A206536 001</u> | Nov 20, 2018 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A206536 002</u> | Nov 20, 2018 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A206536 003</u> | Nov 20, 2018 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A206536 004</u> | Nov 20, 2018 |
| <u>LIPITOR</u> | | | | |
| <u>AB</u> | + | <u>EQ 10MG BASE</u> | <u>N020702 001</u> | Dec 17, 1996 |
| <u>AB</u> | + | <u>EQ 20MG BASE</u> | <u>N020702 002</u> | Dec 17, 1996 |
| <u>AB</u> | + | <u>EQ 40MG BASE</u> | <u>N020702 003</u> | Dec 17, 1996 |
| <u>AB</u> | + | <u>EQ 80MG BASE</u> | <u>N020702 004</u> | Apr 07, 2000 |

ATORVASTATIN CALCIUM; EZETIMIBE

TABLET; ORAL

EZETIMIBE AND ATORVASTATIN CALCIUM

| | | | | |
|--|------------------|--------------------------|--------------------|--------------|
| | WATSON LABS TEVA | <u>EQ 10MG BASE;10MG</u> | <u>A206084 001</u> | Apr 26, 2017 |
| | | <u>EQ 20MG BASE;10MG</u> | <u>A206084 002</u> | Apr 26, 2017 |
| | | <u>EQ 40MG BASE;10MG</u> | <u>A206084 003</u> | Apr 26, 2017 |
| | ! | <u>EQ 80MG BASE;10MG</u> | <u>A206084 004</u> | Apr 26, 2017 |

ATOVAQUONE

SUSPENSION; ORAL

ATOVAQUONE

| | | | | | |
|---------------|---------------------|----------------------------|--------------------|--------------------|--------------|
| <u>AB</u> | AMNEAL PHARMS | <u>750MG/5ML</u> | <u>A202960 001</u> | Mar 18, 2014 | |
| <u>AB</u> | APOTEX INC | <u>750MG/5ML</u> | <u>A209750 001</u> | Oct 11, 2017 | |
| <u>AB</u> | GLENMARK PHARMS | <u>750MG/5ML</u> | <u>A209685 001</u> | Nov 21, 2018 | |
| <u>AB</u> | HETERO LABS LTD III | <u>750MG/5ML</u> | <u>A210692 001</u> | Oct 11, 2018 | |
| <u>AB</u> | LUPIN LTD | <u>750MG/5ML</u> | <u>A209105 001</u> | Sep 11, 2018 | |
| <u>AB</u> | PADDOCK LLC | <u>750MG/5ML</u> | <u>A207833 001</u> | Apr 28, 2017 | |
| <u>MEPRON</u> | | | | | |
| <u>AB</u> | + | <u>GLAXOSMITHKLINE LLC</u> | <u>750MG/5ML</u> | <u>N020500 001</u> | Feb 08, 1995 |

PRESCRIPTION DRUG PRODUCT LIST

ATOVAQUONE; PROGUANIL HYDROCHLORIDE

TABLET; ORAL

ATOVAQUONE AND PROGUANIL HYDROCHLORIDE

| | | | | |
|---------------------------|----------------------------|--------------------|--------------------|--------------|
| AB | GLENMARK GENERICS | 62.5MG;25MG | A091211 002 | Apr 06, 2015 |
| AB | | 250MG;100MG | A091211 001 | Jan 12, 2011 |
| AB | MYLAN PHARMS INC | 62.5MG;25MG | A202362 001 | May 27, 2014 |
| AB | | 250MG;100MG | A202362 002 | May 27, 2014 |
| MALARONE | | | | |
| AB | + ! GLAXOSMITHKLINE | 250MG;100MG | N021078 001 | Jul 14, 2000 |
| MALARONE PEDIATRIC | | | | |
| AB | + GLAXOSMITHKLINE | 62.5MG;25MG | N021078 002 | Jul 14, 2000 |

ATACURIUM BESYLATE

INJECTABLE; INJECTION

ATACURIUM BESYLATE

| | | | | |
|---------------------------------------------|-------------------------------|----------------|--------------------|--------------|
| AP | AUROBINDO PHARMA LTD | 10MG/ML | A206011 001 | Apr 08, 2015 |
| AP | HOSPIRA INC | 10MG/ML | A090761 001 | Oct 18, 2012 |
| AP | MYLAN LABS LTD | 10MG/ML | A206096 001 | Jun 22, 2017 |
| AP | NANJING KING-FRIEND | 10MG/ML | A091489 001 | Feb 17, 2012 |
| AP | ! WEST-WARD PHARMS INT | 10MG/ML | A074901 001 | Jul 18, 1997 |
| ATACURIUM BESYLATE PRESERVATIVE FREE | | | | |
| AP | AUROBINDO PHARMA LTD | 10MG/ML | A206010 001 | Apr 08, 2015 |
| AP | HOSPIRA INC | 10MG/ML | A090782 001 | Oct 18, 2012 |
| AP | MYLAN LABS LTD | 10MG/ML | A206001 001 | Apr 07, 2017 |
| AP | NANJING KING-FRIEND | 10MG/ML | A091488 001 | Feb 17, 2012 |
| AP | ! WEST-WARD PHARMS INT | 10MG/ML | A074900 001 | Jul 18, 1997 |

ATROPINE SULFATE

SOLUTION; INTRAMUSCULAR

ATROPEN

| | | | | |
|------------|----------------------|--------------|-------------|--------------|
| + ! | MERIDIAN MEDCL TECHN | 0.25MG/0.3ML | N017106 004 | Sep 17, 2004 |
| + ! | | 0.5MG/0.7ML | N017106 003 | Jun 19, 2003 |
| + ! | | 1MG/0.7ML | N017106 002 | Jun 19, 2003 |
| + ! | | 2MG/0.7ML | N017106 001 | |

SOLUTION; INTRAVENOUS

ATROPINE SULFATE LIFESHIELD ABBOJECT SYRINGE

| | | | | |
|------------|---------|----------------------|-------------|--------------|
| + ! | HOSPIRA | 0.5MG/5ML (0.1MG/ML) | N021146 004 | Aug 17, 2017 |
| + ! | | 1MG/10ML (0.1MG/ML) | N021146 005 | Aug 17, 2017 |

SOLUTION; INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS, ENDOTRACHEAL

ATROPINE SULFATE ANSYR PLASTIC SYRINGE

| | | | | |
|------------|---------|------------------------|-------------|--------------|
| + ! | HOSPIRA | 0.25MG/5ML (0.05MG/ML) | N021146 002 | Jul 09, 2001 |
| + ! | | 1MG/10ML (0.1MG/ML) | N021146 003 | Jul 09, 2001 |

SOLUTION; IV (INFUSION), INTRAMUSCULAR, SUBCUTANEOUS, INTRAOSSEOUS, ENDOTRACHEAL

ATROPINE SULFATE

| | | | | |
|------------|--------------------|---------------------|-------------|--------------|
| + ! | FRESENIUS KABI USA | 8MG/20ML (0.4MG/ML) | N209260 001 | Jan 26, 2018 |
|------------|--------------------|---------------------|-------------|--------------|

SOLUTION/DROPS; OPHTHALMIC

ATROPINE SULFATE

| | | | | |
|------------|-------|----|-------------|--------------|
| + ! | AKORN | 1% | N206289 001 | Jul 18, 2014 |
|------------|-------|----|-------------|--------------|

ISOPTO ATROPINE

| | | | | |
|--|----------------|----|-------------|--------------|
| | ALCON LABS INC | 1% | N208151 001 | Dec 01, 2016 |
|--|----------------|----|-------------|--------------|

ATROPINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET; ORAL

MOTOFEN

| | | | | |
|------------|--------------------|-------------|-------------|--|
| + ! | SEBELA IRELAND LTD | 0.025MG;1MG | N017744 002 | |
|------------|--------------------|-------------|-------------|--|

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

SOLUTION; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

| | | | | |
|----------|----------------------|------------------------|-------------|--------------|
| ! | WEST-WARD PHARMS INT | 0.025MG/5ML; 2.5MG/5ML | A087708 001 | May 03, 1982 |
|----------|----------------------|------------------------|-------------|--------------|

TABLET; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

| | | | | |
|-----------|---------------------|----------------------|--------------------|--------------|
| AA | ANI PHARMS INC | 0.025MG;2.5MG | A086727 001 | |
| AA | BAYSHORE PHARMS LLC | 0.025MG;2.5MG | A210819 001 | Nov 13, 2018 |
| AA | LANNETT | 0.025MG;2.5MG | A085372 001 | |
| AA | MYLAN | 0.025MG;2.5MG | A085762 001 | |
| AA | PAR PHARM | 0.025MG;2.5MG | A040357 001 | May 02, 2000 |
| AA | UPSHER SMITH LABS | 0.025MG;2.5MG | A210571 001 | Aug 31, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

TABLET; ORAL

LOMOTIL

| | | | | | | |
|-----------|-----------|---------------|-----------------------|----------------|------------|--|
| AA | +! | GD SEARLE LLC | 0.025MG; 2.5MG | N012462 | 001 | |
|-----------|-----------|---------------|-----------------------|----------------|------------|--|

ATROPINE; PRALIDOXIME CHLORIDE

INJECTABLE; INTRAMUSCULAR

DUODOTE

| | | | | | |
|-----------|----------------|------------------------|---------|-----|--------------|
| +! | MERIDIAN MEDCL | 2.1MG/0.7ML; 600MG/2ML | N021983 | 001 | Sep 28, 2006 |
|-----------|----------------|------------------------|---------|-----|--------------|

AURANOFIN

CAPSULE; ORAL

RIDAURA

| | | | | | |
|-----------|--------------------|-----|---------|-----|--------------|
| +! | SEBELA IRELAND LTD | 3MG | N018689 | 001 | May 24, 1985 |
|-----------|--------------------|-----|---------|-----|--------------|

AVANAFIL

TABLET; ORAL

STENDRA

| | | | | | |
|----------|-----------------|------|---------|-----|--------------|
| + | METUCHEN PHARMS | 50MG | N202276 | 001 | Apr 27, 2012 |
|----------|-----------------|------|---------|-----|--------------|

| | | | | | |
|----------|--|-------|---------|-----|--------------|
| + | | 100MG | N202276 | 002 | Apr 27, 2012 |
|----------|--|-------|---------|-----|--------------|

| | | | | | |
|-----------|--|-------|---------|-----|--------------|
| +! | | 200MG | N202276 | 003 | Apr 27, 2012 |
|-----------|--|-------|---------|-----|--------------|

AVATROMBOPAG MALEATE

TABLET; ORAL

DOPTELET

| | | | | | |
|-----------|-----------|--------------|---------|-----|--------------|
| +! | AKARX INC | EQ 20MG BASE | N210238 | 001 | May 21, 2018 |
|-----------|-----------|--------------|---------|-----|--------------|

AVIBACTAM SODIUM; CEFTAZIDIME

POWDER; IV (INFUSION)

AVYCAZ

| | | | | | |
|-----------|--------------------|-------------------------|---------|-----|--------------|
| +! | ALLERGAN SALES LLC | EQ 0.5GM BASE; 2GM/VIAL | N206494 | 001 | Feb 25, 2015 |
|-----------|--------------------|-------------------------|---------|-----|--------------|

AXITINIB

TABLET; ORAL

INLYTA

| | | | | | |
|----------|-------------|-----|---------|-----|--------------|
| + | PF PRISM CV | 1MG | N202324 | 001 | Jan 27, 2012 |
|----------|-------------|-----|---------|-----|--------------|

| | | | | | |
|-----------|--|-----|---------|-----|--------------|
| +! | | 5MG | N202324 | 002 | Jan 27, 2012 |
|-----------|--|-----|---------|-----|--------------|

AZACITIDINE

POWDER; INTRAVENOUS, SUBCUTANEOUS

AZACITIDINE

| | | | | | |
|-----------|-----------------|-------------------|----------------|------------|--------------|
| AP | ACCORD HLTHCARE | 100MG/VIAL | A207475 | 001 | Jul 02, 2018 |
|-----------|-----------------|-------------------|----------------|------------|--------------|

| | | | | | |
|-----------|-------------|-------------------|----------------|------------|--------------|
| AP | ACTAVIS LLC | 100MG/VIAL | N208216 | 001 | Apr 29, 2016 |
|-----------|-------------|-------------------|----------------|------------|--------------|

| | | | | | |
|-----------|-------|-------------------|----------------|------------|--------------|
| AP | CIPLA | 100MG/VIAL | A209540 | 001 | May 04, 2018 |
|-----------|-------|-------------------|----------------|------------|--------------|

| | | | | | |
|-----------|--------------------|-------------------|----------------|------------|--------------|
| AP | DR REDDYS LABS LTD | 100MG/VIAL | A201537 | 001 | Sep 16, 2013 |
|-----------|--------------------|-------------------|----------------|------------|--------------|

| | | | | | |
|-----------|---------------------|-------------------|----------------|------------|--------------|
| AP | MYLAN INSTITUTIONAL | 100MG/VIAL | A204949 | 001 | Apr 28, 2016 |
|-----------|---------------------|-------------------|----------------|------------|--------------|

| | | | | | |
|-----------|------------------|-------------------|----------------|------------|--------------|
| AP | NATCO PHARMA LTD | 100MG/VIAL | A207234 | 001 | Jun 23, 2017 |
|-----------|------------------|-------------------|----------------|------------|--------------|

| | | | | | |
|-----------|-----------------|-------------------|----------------|------------|--------------|
| AP | SHILPA MEDICARE | 100MG/VIAL | A207518 | 001 | Sep 29, 2016 |
|-----------|-----------------|-------------------|----------------|------------|--------------|

VIDAZA

| | | | | | | |
|-----------|-----------|---------|-------------------|----------------|------------|--------------|
| AP | +! | CELGENE | 100MG/VIAL | N050794 | 001 | May 19, 2004 |
|-----------|-----------|---------|-------------------|----------------|------------|--------------|

AZATHIOPRINE

TABLET; ORAL

AZASAN

| | | | | | |
|-----------|---------------|-------------|----------------|------------|--------------|
| AB | AAIPHARMA LLC | 25MG | A075252 | 002 | Feb 03, 2003 |
|-----------|---------------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-------------|----------------|------------|--------------|
| AB | | 50MG | A075252 | 001 | Jun 07, 1999 |
|-----------|--|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-------------|----------------|------------|--------------|
| AB | | 75MG | A075252 | 003 | Feb 03, 2003 |
|-----------|--|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|--------------|----------------|------------|--------------|
| AB | | 100MG | A075252 | 004 | Feb 03, 2003 |
|-----------|--|--------------|----------------|------------|--------------|

AZATHIOPRINE

| | | | | | |
|-----------|-------------------|-------------|----------------|------------|--------------|
| AB | AMNEAL PHARMS LLC | 50MG | A074069 | 001 | Feb 16, 1996 |
|-----------|-------------------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|-------|-------------|----------------|------------|--------------|
| AB | MYLAN | 50MG | A075568 | 001 | Dec 13, 1999 |
|-----------|-------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|------------------|-------------|----------------|------------|--------------|
| AB | ZYDUS PHARMS USA | 25MG | A077621 | 002 | Sep 05, 2008 |
|-----------|------------------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-------------|----------------|------------|--------------|
| AB | | 50MG | A077621 | 001 | Mar 15, 2007 |
|-----------|--|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-------------|----------------|------------|--------------|
| AB | | 75MG | A077621 | 003 | Sep 05, 2008 |
|-----------|--|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|--------------|----------------|------------|--------------|
| AB | | 100MG | A077621 | 004 | Sep 05, 2008 |
|-----------|--|--------------|----------------|------------|--------------|

TMURAN

| | | | | | | |
|-----------|-----------|--------------------|-------------|----------------|------------|--|
| AB | +! | SEBELA IRELAND LTD | 50MG | N016324 | 001 | |
|-----------|-----------|--------------------|-------------|----------------|------------|--|

AZATHIOPRINE SODIUM

INJECTABLE; INJECTION

AZATHIOPRINE SODIUM

| | | | | | |
|----------|------------------|--------------------|---------|-----|--------------|
| ! | WEST-WARD PHARMS | EQ 100MG BASE/VIAL | A074419 | 001 | Mar 31, 1995 |
|----------|------------------|--------------------|---------|-----|--------------|

INT

PRESCRIPTION DRUG PRODUCT LIST

AZELAIC ACID

AEROSOL, FOAM;TOPICAL

FINACEA

+! LEO PHARMA AS 15% N207071 001 Jul 29, 2015

CREAM;TOPICAL

AZELEX

+! AQUA PHARMS LLC 20% N020428 001 Sep 13, 1995

GEL;TOPICAL

AZELAIC ACID**AB** ACTAVIS LABS UT INC **15%** **A208011 001** Nov 19, 2018**AB** GLENMARK PHARMS **15%** **A204637 001** Nov 19, 2018**AB** TOLMAR **15%** **A208724 001** Nov 19, 2018FINACEA**AB** +! LEO PHARMA AS **15%** **N021470 001** Dec 24, 2002AZELASTINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

AZELASTINE HYDROCHLORIDE**AT** AKORN **0.05%** **A203660 001** Nov 08, 2016**AT** APOTEX INC **0.05%** **A078621 001** Aug 03, 2009**AT** ! SANDOZ INC **0.05%** **A202305 001** May 31, 2012**AT** SUN PHARMA GLOBAL **0.05%** **A078738 001** Jun 21, 2010

SPRAY, METERED;NASAL

ASTELIN**AB** +! MYLAN SPECIALITY LP **EQ 0.125MG BASE/SPRAY** **N020114 001** Nov 01, 1996ASTEPRO**AB** +! MYLAN SPECIALITY LP **EQ 0.1876MG BASE/SPRAY** **N022203 002** Aug 31, 2009AZELASTINE HYDROCHLORIDE**AB** ALKEM LABS LTD **EQ 0.125MG BASE/SPRAY** **A208156 001** Aug 18, 2017**AB** AMNEAL PHARMS LLC **EQ 0.125MG BASE/SPRAY** **A204660 001** Aug 28, 2017**AB** **EQ 0.1876MG BASE/SPRAY** **A208199 001** Dec 15, 2017**AB** APOTEX INC **EQ 0.125MG BASE/SPRAY** **A077954 001** Apr 30, 2009**AB** **EQ 0.1876MG BASE/SPRAY** **A201846 001** Aug 31, 2012**AB** BRECKENRIDGE PHARM **EQ 0.125MG BASE/SPRAY** **A090176 001** Jul 28, 2015**AB** PERRIGO ISRAEL **EQ 0.1876MG BASE/SPRAY** **A202743 001** May 08, 2014**AB** SUN PHARMA GLOBAL **EQ 0.125MG BASE/SPRAY** **A090423 001** May 23, 2012**AB** UPSHER SMITH LABS **EQ 0.125MG BASE/SPRAY** **A202609 001** Mar 17, 2017**AB** WEST-WARD PHARMS **EQ 0.125MG BASE/SPRAY** **A091444 001** Oct 24, 2014**AB** **EQ 0.1876MG BASE/SPRAY** **A207243 001** Sep 22, 2017**AB** ZYDUS PHARMS USA **EQ 0.125MG BASE/SPRAY** **A091409 001** Aug 14, 2017**AB** INCAZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE

SPRAY, METERED;NASAL

DYMISTA

+! MYLAN SPECIALITY LP EQ 0.125MG BASE/SPRAY;0.05MG/SPRAY N202236 001 May 01, 2012

AZILSARTAN KAMEDOXOMIL

TABLET;ORAL

EDARBI

+ ARBOR PHARMS LLC EQ 40MG MEDOXOMIL N200796 001 Feb 25, 2011

+! EQ 80MG MEDOXOMIL N200796 002 Feb 25, 2011

AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE

TABLET;ORAL

EDARBYCLOR

+ ARBOR PHARMS LLC EQ 40MG MEDOXOMIL;12.5MG N202331 001 Dec 20, 2011

+! EQ 40MG MEDOXOMIL;25MG N202331 002 Dec 20, 2011

AZITHROMYCIN

FOR SUSPENSION;ORAL

AZITHROMYCIN**AB** AMNEAL PHARMS LLC **EQ 100MG BASE/5ML** **A205666 001** Jul 19, 2018**AB** **EQ 200MG BASE/5ML** **A205666 002** Jul 19, 2018**AB** AUROBINDO PHARMA **EQ 100MG BASE/5ML** **A209201 001** Oct 09, 2018**AB** LTD **EQ 200MG BASE/5ML** **A209201 002** Oct 09, 2018**AB** EPIC PHARMA LLC **EQ 100MG BASE/5ML** **A207531 001** Apr 09, 2018**AB** **EQ 200MG BASE/5ML** **A207531 002** Apr 09, 2018**AB** LUPIN LTD **EQ 100MG BASE/5ML** **A065488 001** May 15, 2015**AB** **EQ 200MG BASE/5ML** **A065488 002** May 15, 2015**AB** PLIVA **EQ 100MG BASE/5ML** **A065246 002** Jul 05, 2006**AB** **EQ 200MG BASE/5ML** **A065246 001** Jul 05, 2006**AB** TEVA PHARMS **EQ 100MG BASE/5ML** **A065419 001** Jun 24, 2008

PRESCRIPTION DRUG PRODUCT LIST

AZITHROMYCIN

FOR SUSPENSION;ORAL

AZITHROMYCIN

| | | | | |
|-----------|-----------------|--------------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 200MG BASE/5ML</u> | <u>A065419 002</u> | Jun 24, 2008 |
| <u>AB</u> | ZYDUS WORLDWIDE | <u>EQ 100MG BASE/5ML</u> | <u>A211147 001</u> | Jul 31, 2018 |
| <u>AB</u> | | <u>EQ 200MG BASE/5ML</u> | <u>A211147 002</u> | Jul 31, 2018 |

ZITHROMAX

| | | | | | |
|-----------|---|--------|--------------------------|--------------------|--------------|
| <u>AB</u> | + | PFIZER | <u>EQ 100MG BASE/5ML</u> | <u>N050710 001</u> | Oct 19, 1995 |
| <u>AB</u> | + | ! | <u>EQ 200MG BASE/5ML</u> | <u>N050710 002</u> | Oct 19, 1995 |
| | + | | EQ 1GM BASE/PACKET | N050693 001 | Sep 28, 1994 |

INJECTABLE; INJECTION

AZITHROMYCIN

| | | | | |
|-----------|----------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>EQ 500MG BASE/VIAL</u> | <u>A203294 001</u> | Jun 19, 2015 |
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 500MG BASE/VIAL</u> | <u>A065179 001</u> | Dec 13, 2005 |
| <u>AP</u> | GLAND PHARMA LTD | <u>EQ 500MG BASE/VIAL</u> | <u>A065501 001</u> | Nov 09, 2009 |
| <u>AP</u> | HAINAN POLY PHARM | <u>EQ 500MG BASE/VIAL</u> | <u>A203412 001</u> | Oct 09, 2018 |
| <u>AP</u> | HOSPIRA | <u>EQ 500MG BASE/VIAL</u> | <u>A065500 001</u> | Jun 26, 2009 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A065511 001</u> | Jun 26, 2009 |
| <u>AP</u> | MYLAN ASI | <u>EQ 500MG BASE/VIAL</u> | <u>A065506 001</u> | Mar 24, 2009 |
| <u>AP</u> | MYLAN LABS LTD | <u>EQ 500MG BASE/VIAL</u> | <u>A204732 001</u> | Jan 26, 2017 |
| <u>AP</u> | SUN PHARM INDS LTD | <u>EQ 500MG BASE/VIAL</u> | <u>A090923 001</u> | Apr 02, 2013 |

ZITHROMAX

| | | | | | |
|-----------|---|--------|---------------------------|--------------------|--------------|
| <u>AP</u> | + | PFIZER | <u>EQ 500MG BASE/VIAL</u> | <u>N050733 001</u> | Jan 30, 1997 |
|-----------|---|--------|---------------------------|--------------------|--------------|

SOLUTION/DROPS;OPHTHALMIC

AZASITE

| | | | | | |
|--|---|----------------|----|-------------|--------------|
| | + | OAK PHARMS INC | 1% | N050810 001 | Apr 27, 2007 |
|--|---|----------------|----|-------------|--------------|

TABLET;ORAL

AZITHROMYCIN

| | | | | |
|-----------|----------------------|----------------------|--------------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 250MG BASE</u> | <u>A207370 001</u> | Jul 05, 2018 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A207398 001</u> | Jul 05, 2018 |
| <u>AB</u> | BIONPHARMA INC | <u>EQ 600MG BASE</u> | <u>A209999 001</u> | Dec 26, 2018 |
| <u>AB</u> | CSPC OUYI PHARM CO | <u>EQ 500MG BASE</u> | <u>A208249 001</u> | Oct 25, 2018 |
| <u>AB</u> | | <u>EQ 600MG BASE</u> | <u>A207566 001</u> | Sep 24, 2018 |
| <u>AB</u> | LUPIN LTD | <u>EQ 250MG BASE</u> | <u>A065398 001</u> | May 15, 2015 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065399 001</u> | May 15, 2015 |
| <u>AB</u> | | <u>EQ 600MG BASE</u> | <u>A065400 001</u> | May 15, 2015 |
| <u>AB</u> | MYLAN | <u>EQ 600MG BASE</u> | <u>A065360 001</u> | Jan 08, 2007 |
| <u>AB</u> | PLIVA | <u>EQ 250MG BASE</u> | <u>A065225 001</u> | Nov 14, 2005 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065223 001</u> | Nov 14, 2005 |
| <u>AB</u> | | <u>EQ 600MG BASE</u> | <u>A065218 001</u> | Nov 14, 2005 |
| <u>AB</u> | SANDOZ | <u>EQ 250MG BASE</u> | <u>A065211 001</u> | Nov 14, 2005 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065212 001</u> | Nov 14, 2005 |
| <u>AB</u> | | <u>EQ 600MG BASE</u> | <u>A065209 001</u> | Nov 14, 2005 |
| <u>AB</u> | SUNSHINE LAKE | <u>EQ 250MG BASE</u> | <u>A209045 001</u> | Dec 07, 2018 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A209044 001</u> | Dec 07, 2018 |
| <u>AB</u> | | <u>EQ 600MG BASE</u> | <u>A209043 001</u> | Dec 06, 2018 |
| <u>AB</u> | TEVA | <u>EQ 250MG BASE</u> | <u>A065153 001</u> | Nov 14, 2005 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065193 001</u> | Nov 14, 2005 |
| <u>AB</u> | | <u>EQ 600MG BASE</u> | <u>A065150 001</u> | Nov 14, 2005 |
| <u>AB</u> | WOCKHARDT | <u>EQ 250MG BASE</u> | <u>A065404 001</u> | Feb 11, 2008 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065405 001</u> | Feb 11, 2008 |
| <u>AB</u> | | <u>EQ 600MG BASE</u> | <u>A065302 003</u> | Feb 11, 2008 |

ZITHROMAX

| | | | | | |
|-----------|---|--------|----------------------|--------------------|--------------|
| <u>AB</u> | + | PFIZER | <u>EQ 250MG BASE</u> | <u>N050711 001</u> | Jul 18, 1996 |
| <u>AB</u> | + | | <u>EQ 500MG BASE</u> | <u>N050784 001</u> | May 24, 2002 |
| <u>AB</u> | + | ! | <u>EQ 600MG BASE</u> | <u>N050730 001</u> | Jun 12, 1996 |

AZTREONAM

FOR SOLUTION;INHALATION

CAYSTON

| | | | | | |
|--|---|--------|-----------|-------------|--------------|
| | + | GILEAD | 75MG/VIAL | N050814 001 | Feb 22, 2010 |
|--|---|--------|-----------|-------------|--------------|

INJECTABLE; INJECTION

AZACTAM

| | | | | | |
|-----------|---|----------------------|-----------------|--------------------|--------------|
| <u>AP</u> | + | BRISTOL MYERS SQUIBB | <u>1GM/VIAL</u> | <u>N050580 002</u> | Dec 31, 1986 |
| <u>AP</u> | + | | <u>2GM/VIAL</u> | <u>N050580 003</u> | Dec 31, 1986 |

AZTREONAM

| | | | | |
|-----------|--------------------|-----------------|--------------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>1GM/VIAL</u> | <u>A065439 002</u> | Jun 18, 2010 |
| <u>AP</u> | | <u>2GM/VIAL</u> | <u>A065439 003</u> | Jun 18, 2010 |

AZACTAM IN PLASTIC CONTAINER

| | | | | | |
|--|---|----------------------|---------|-------------|--------------|
| | + | BRISTOL MYERS SQUIBB | 20MG/ML | N050632 002 | May 24, 1989 |
|--|---|----------------------|---------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

AZTREONAM

INJECTABLE; INJECTION

AZACTAM IN PLASTIC CONTAINER

+!

40MG/ML

N050632 001 May 24, 1989

AZTREONAM

FRESENIUS KABI USA 500MG/VIAL

A065439 001 Jun 18, 2010

BACITRACIN

INJECTABLE; INJECTION

BACIIM**AP** X GEN PHARMS**50,000 UNITS/VIAL****A064153 001** May 09, 1997BACITRACIN**AP** AKORN**50,000 UNITS/VIAL****A206719 001** Oct 20, 2017**AP** FRESENIUS KABI USA**50,000 UNITS/VIAL****A065116 001** Dec 03, 2002**AP** ! PHARMACIA AND**50,000 UNITS/VIAL****A060733 002**

UPJOHN

AP XELLIA PHARMS APS**50,000 UNITS/VIAL****A203177 001** Aug 25, 2014

OINTMENT; OPHTHALMIC

BACITRACIN

!

PERRIGO CO

500 UNITS/GM

A061212 001

TENNESSEE

BACITRACIN ZINC; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE

!

PERRIGO CO

400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM

A062166 002

TENNESSEE

BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE**AT** AKORN**400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM****A065213 001** Jul 25, 2012**AT** ! BAUSCH AND LOMB**400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM****A064068 001** Oct 30, 1995

OINTMENT; TOPICAL

CORTISPORIN

+!

MONARCH PHARMS

400 UNITS/GM;1%;EQ 3.5MG BASE/GM;5,000 UNITS/GM

N050168 002 May 04, 1984

BACITRACIN ZINC; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC**AT** AKORN**400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM****A065088 001** Feb 06, 2004**AT** ! BAUSCH AND LOMB**400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM****A064064 001** Oct 30, 1995**AT** PERRIGO CO**400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM****A060764 002**

TENNESSEE

NEOSPORIN**AT** + CASPER PHARMA LLC**400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM****N050417 001**BACITRACIN ZINC; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

BACITRACIN ZINC AND POLYMYXIN B SULFATE**AT** AKORN**500 UNITS/GM;10,000 UNITS/GM****A064028 001** Jan 30, 1995**AT** ! BAUSCH AND LOMB**500 UNITS/GM;10,000 UNITS/GM****A064046 001** Jan 26, 1995**AT** PERRIGO CO**500 UNITS/GM;10,000 UNITS/GM****A065022 001** Feb 27, 2002

TENNESSEE

BACLOFEN

INJECTABLE; INTRATHECAL

BACLOFEN**AP** EMERALD INTL LTD**0.05MG/ML****A091193 001** May 03, 2016**AP****0.5MG/ML****A091193 002** May 03, 2016**AP****2MG/ML****A091193 003** May 03, 2016**AP** MYLAN LABS LTD**0.5MG/ML****A209592 001** Mar 21, 2018**AP****1MG/ML****A209594 001** Mar 06, 2018**AP****2MG/ML****A209592 002** Mar 21, 2018GABLOFEN**AP** PIRAMAL CRITICAL**0.05MG/ML****N022462 001** Nov 19, 2010**AP****0.5MG/ML****N022462 002** Nov 19, 2010**AP** +!**1MG/ML****N022462 004** Jun 22, 2012**AP****2MG/ML****N022462 003** Nov 19, 2010LIORESAL**AP** +! SAOL THERAPS RES**0.05MG/ML****N020075 003** Nov 07, 1996

LTD

PRESCRIPTION DRUG PRODUCT LIST

BACLOFEN

INJECTABLE; INTRATHECAL

LIORESAL

| | | | | | |
|-----------|------------|-----------------|----------------|------------|--------------|
| <u>AB</u> | <u>+</u> ! | <u>0.5MG/ML</u> | <u>N020075</u> | <u>001</u> | Jun 17, 1992 |
| <u>AB</u> | <u>+</u> ! | <u>2MG/ML</u> | <u>N020075</u> | <u>002</u> | Jun 17, 1992 |

TABLET; ORAL

BACLOFEN

| | | | | | | |
|-----------|----------|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | IMPAX LABS | <u>10MG</u> | <u>A077971</u> | <u>001</u> | Oct 26, 2007 |
| <u>AB</u> | | | <u>20MG</u> | <u>A077971</u> | <u>002</u> | Oct 26, 2007 |
| <u>AB</u> | | IVAX SUB TEVA PHARMS | <u>10MG</u> | <u>A072234</u> | <u>001</u> | Jul 21, 1988 |
| <u>AB</u> | <u>!</u> | | <u>20MG</u> | <u>A072235</u> | <u>001</u> | Jul 21, 1988 |
| <u>AB</u> | | LANNETT CO INC | <u>10MG</u> | <u>A077241</u> | <u>002</u> | Jul 06, 2007 |
| <u>AB</u> | | | <u>20MG</u> | <u>A077241</u> | <u>001</u> | Dec 20, 2005 |
| <u>AB</u> | | MYLAN | <u>20MG</u> | <u>A077121</u> | <u>002</u> | Jul 29, 2005 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>10MG</u> | <u>A090334</u> | <u>001</u> | Feb 18, 2010 |
| <u>AB</u> | | | <u>20MG</u> | <u>A090334</u> | <u>002</u> | Feb 18, 2010 |
| <u>AB</u> | | NORTHSTAR HLTHCARE | <u>10MG</u> | <u>A078401</u> | <u>002</u> | Sep 18, 2009 |
| <u>AB</u> | | | <u>20MG</u> | <u>A078401</u> | <u>001</u> | Sep 18, 2009 |
| <u>AB</u> | | OXFORD PHARMS | <u>10MG</u> | <u>A077088</u> | <u>002</u> | Oct 31, 2007 |
| <u>AB</u> | | | <u>20MG</u> | <u>A077088</u> | <u>001</u> | Oct 31, 2007 |
| <u>AB</u> | | RUBICON RES PVT LTD | <u>10MG</u> | <u>A209102</u> | <u>002</u> | Nov 28, 2017 |
| <u>AB</u> | | | <u>20MG</u> | <u>A209102</u> | <u>003</u> | Nov 28, 2017 |
| <u>AB</u> | | SUN PHARM INDS INC | <u>10MG</u> | <u>A077862</u> | <u>001</u> | Aug 14, 2006 |
| <u>AB</u> | | | <u>20MG</u> | <u>A077862</u> | <u>002</u> | Aug 14, 2006 |
| <u>AB</u> | | USL PHARMA | <u>10MG</u> | <u>A074584</u> | <u>001</u> | Aug 19, 1996 |
| <u>AB</u> | | | <u>20MG</u> | <u>A074584</u> | <u>002</u> | Aug 19, 1996 |
| <u>AB</u> | | VINTAGE PHARMS | <u>10MG</u> | <u>A077068</u> | <u>002</u> | Aug 30, 2005 |
| <u>AB</u> | | | <u>20MG</u> | <u>A077068</u> | <u>001</u> | Aug 30, 2005 |
| <u>AB</u> | | ZYDUS WORLDWIDE | <u>10MG</u> | <u>A211659</u> | <u>001</u> | Nov 23, 2018 |
| <u>AB</u> | | | <u>20MG</u> | <u>A211659</u> | <u>002</u> | Nov 23, 2018 |
| | | RUBICON RES PVT LTD | 5MG | A209102 | 001 | Nov 28, 2017 |

BALOXAVIR MARBOXIL

TABLET; ORAL

XOFLUZA

| | | | | | |
|------------|---------------|------|---------|-----|--------------|
| <u>+</u> | GENENTECH INC | 20MG | N210854 | 001 | Oct 24, 2018 |
| <u>+</u> ! | | 40MG | N210854 | 002 | Oct 24, 2018 |

BALSALAZIDE DISODIUM

CAPSULE; ORAL

BALSALAZIDE DISODIUM

| | | | | | | |
|-----------|--|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | APOTEX INC | <u>750MG</u> | <u>A077883</u> | <u>001</u> | Dec 28, 2007 |
| <u>AB</u> | | MYLAN | <u>750MG</u> | <u>A077807</u> | <u>001</u> | Dec 28, 2007 |
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>750MG</u> | <u>A077806</u> | <u>001</u> | Dec 28, 2007 |

COLAZAL

| | | | | | | |
|-----------|------------|---------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | <u>+</u> ! | VALEANT PHARMS INTL | <u>750MG</u> | <u>N020610</u> | <u>001</u> | Jul 18, 2000 |
|-----------|------------|---------------------|--------------|----------------|------------|--------------|

TABLET; ORAL

BALSALAZIDE DISODIUM

| | | | | | | |
|-----------|--|---------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | PAR PHARM INC | <u>1.1GM</u> | <u>A206336</u> | <u>001</u> | Sep 08, 2015 |
|-----------|--|---------------|--------------|----------------|------------|--------------|

GIAZO

| | | | | | | |
|-----------|------------|---------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | <u>+</u> ! | VALEANT PHARMS INTL | <u>1.1GM</u> | <u>N022205</u> | <u>001</u> | Feb 03, 2012 |
|-----------|------------|---------------------|--------------|----------------|------------|--------------|

BARICITINIB

TABLET; ORAL

OLUMIANT

| | | | | | |
|------------|------------------|-----|---------|-----|--------------|
| <u>+</u> ! | ELI LILLY AND CO | 2MG | N207924 | 001 | May 31, 2018 |
|------------|------------------|-----|---------|-----|--------------|

BIARIUM SULFATE

FOR SUSPENSION; ORAL

E-Z-HD

| | | | | | |
|------------|--------|-----------------|---------|-----|--------------|
| <u>+</u> ! | BRACCO | 98% (334GM/BOT) | N208036 | 001 | Jan 11, 2016 |
|------------|--------|-----------------|---------|-----|--------------|

E-Z-PAQUE

| | | | | | |
|------------|--------|-----------------|---------|-----|--------------|
| <u>+</u> ! | BRACCO | 96% (169GM/BOT) | N208036 | 002 | Apr 07, 2017 |
|------------|--------|-----------------|---------|-----|--------------|

PASTE; ORAL

VARIBAR PUDDING

| | | | | | |
|--|--------|-----|---------|-----|--------------|
| | BRACCO | 40% | N208844 | 001 | Oct 14, 2016 |
|--|--------|-----|---------|-----|--------------|

SUSPENSION; ORAL

LIQUID E-Z-PAQUE

| | | | | | |
|------------|--------|-----------------|---------|-----|--------------|
| <u>+</u> ! | BRACCO | 60% (213GM/BOT) | N208143 | 003 | Mar 01, 2017 |
|------------|--------|-----------------|---------|-----|--------------|

READI-CAT 2

| | | | | | |
|------------|--------|--------------|---------|-----|--------------|
| <u>+</u> ! | BRACCO | 2% (9GM/BOT) | N208143 | 001 | Jan 15, 2016 |
|------------|--------|--------------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

BARIUM SULFATE

SUSPENSION; ORAL

READI-CAT 2 SMOOTHIES

+! BRACCO 2% (9GM/BOT) N208143 002 Jan 15, 2016

TAGITOL V

+! BRACCO 40% (8GM/BOT) N208143 005 Aug 04, 2017

VARIBAR HONEY

+! BRACCO 40% (100GM/250ML) N208143 007 Mar 26, 2018

VARIBAR NECTAR

+! BRACCO 40% (96GM/240ML) N208143 004 Jul 07, 2017

VARIBAR THIN HONEY

+! BRACCO 40% (100GM/250ML) N208143 006 Jan 23, 2018

BAZEDOXIFENE ACETATE; ESTROGENS, CONJUGATED

TABLET; ORAL

DUAVEE

+! WYETH PHARMS EQ 20MG BASE;0.45MG N022247 001 Oct 03, 2013

BECLOMETHASONE DIPROPIONATE

AEROSOL, METERED; INHALATION

QVAR REDIHALER

+ NORTON WATERFORD 0.04MG/INH N207921 001 Aug 03, 2017

+ 0.08MG/INH N207921 002 Aug 03, 2017

AEROSOL, METERED; NASAL

QNASL

+ TEVA BRANDED PHARM 0.04MG/ACTUATION N202813 002 Dec 17, 2014

+! 0.08MG/ACTUATION N202813 001 Mar 23, 2012

BECLOMETHASONE DIPROPIONATE MONOHYDRATE

SPRAY, METERED; NASAL

BECONASE AQ

+! GLAXOSMITHKLINE EQ 0.042MG DIPROP/SPRAY N019389 001 Jul 27, 1987

BEDAQUILINE FUMARATE

TABLET; ORAL

SIRTURO

+! JANSSEN THERAP EQ 100MG BASE N204384 001 Dec 28, 2012

BELINOSTAT

POWDER; INTRAVENOUS

BELEODAQ

+! SPECTRUM PHARMS 500MG/VIAL N206256 001 Jul 03, 2014

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE

| | | | | |
|-----------|----------------------|-------------|--------------------|--------------|
| <u>AB</u> | AMNEAL PHARMS | <u>5MG</u> | <u>A076820 001</u> | Feb 03, 2006 |
| <u>AB</u> | | <u>10MG</u> | <u>A076820 002</u> | Feb 03, 2006 |
| <u>AB</u> | | <u>20MG</u> | <u>A076820 003</u> | Feb 03, 2006 |
| <u>AB</u> | | <u>40MG</u> | <u>A076820 004</u> | Feb 03, 2006 |
| <u>AB</u> | APOTEX INC | <u>5MG</u> | <u>A077128 001</u> | Mar 08, 2006 |
| <u>AB</u> | | <u>10MG</u> | <u>A077128 002</u> | Mar 08, 2006 |
| <u>AB</u> | | <u>20MG</u> | <u>A077128 003</u> | Mar 08, 2006 |
| <u>AB</u> | | <u>40MG</u> | <u>A077128 004</u> | Mar 08, 2006 |
| <u>AB</u> | AUROBINDO PHARMA | <u>10MG</u> | <u>A078212 001</u> | May 22, 2008 |
| <u>AB</u> | | <u>20MG</u> | <u>A078212 002</u> | May 22, 2008 |
| <u>AB</u> | | <u>40MG</u> | <u>A078212 003</u> | May 22, 2008 |
| <u>AB</u> | CASI PHARMS INC | <u>5MG</u> | <u>A076402 001</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>10MG</u> | <u>A076402 002</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>20MG</u> | <u>A076402 003</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>40MG</u> | <u>A076402 004</u> | Feb 11, 2004 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>5MG</u> | <u>A076333 001</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>10MG</u> | <u>A076333 002</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>20MG</u> | <u>A076333 003</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>40MG</u> | <u>A076333 004</u> | Feb 11, 2004 |
| <u>AB</u> | MYLAN | <u>5MG</u> | <u>A076430 001</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>10MG</u> | <u>A076430 002</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>20MG</u> | <u>A076430 003</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>40MG</u> | <u>A076430 004</u> | Feb 11, 2004 |
| <u>AB</u> | PRINSTON INC | <u>5MG</u> | <u>A076118 001</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>10MG</u> | <u>A076118 002</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>20MG</u> | <u>A076118 003</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>40MG</u> | <u>A076118 004</u> | Feb 11, 2004 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>5MG</u> | <u>A076344 001</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>10MG</u> | <u>A076344 002</u> | Feb 11, 2004 |

PRESCRIPTION DRUG PRODUCT LIST

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE

| | | | | |
|-----------|------------------|-------------|--------------------|--------------|
| <u>AB</u> | | <u>20MG</u> | <u>A076344 003</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>40MG</u> | <u>A076344 004</u> | Feb 11, 2004 |
| <u>AB</u> | TEVA | <u>5MG</u> | <u>A076211 001</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>10MG</u> | <u>A076211 002</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>20MG</u> | <u>A076211 003</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>40MG</u> | <u>A076211 004</u> | Feb 11, 2004 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>5MG</u> | <u>A078848 001</u> | May 23, 2008 |
| <u>AB</u> | | <u>10MG</u> | <u>A078848 002</u> | May 23, 2008 |
| <u>AB</u> | | <u>20MG</u> | <u>A078848 003</u> | May 23, 2008 |
| <u>AB</u> | | <u>40MG</u> | <u>A078848 004</u> | May 23, 2008 |

LOTENSIN

| | | | | | |
|-----------|---|-------------------------|-------------|--------------------|--------------|
| <u>AB</u> | + | US PHARMS HOLDINGS I | <u>5MG</u> | <u>N019851 001</u> | Jun 25, 1991 |
| <u>AB</u> | + | | <u>10MG</u> | <u>N019851 002</u> | Jun 25, 1991 |
| <u>AB</u> | + | | <u>20MG</u> | <u>N019851 003</u> | Jun 25, 1991 |
| <u>AB</u> | + | | <u>40MG</u> | <u>N019851 004</u> | Jun 25, 1991 |

BENAZEPRIL HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

| | | | | |
|-----------|------------|---------------------|--------------------|--------------|
| <u>AB</u> | APOTEX INC | <u>5MG; 6.25MG</u> | <u>A078794 001</u> | Aug 21, 2014 |
| <u>AB</u> | | <u>10MG; 12.5MG</u> | <u>A078794 002</u> | Aug 21, 2014 |
| <u>AB</u> | | <u>20MG; 12.5MG</u> | <u>A078794 003</u> | Aug 21, 2014 |
| <u>AB</u> | | <u>20MG; 25MG</u> | <u>A078794 004</u> | Aug 21, 2014 |
| <u>AB</u> | MYLAN | <u>5MG; 6.25MG</u> | <u>A076688 001</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>10MG; 12.5MG</u> | <u>A076688 002</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>20MG; 12.5MG</u> | <u>A076688 003</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>20MG; 25MG</u> | <u>A076688 004</u> | Feb 11, 2004 |
| <u>AB</u> | SANDOZ | <u>5MG; 6.25MG</u> | <u>A076631 001</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>10MG; 12.5MG</u> | <u>A076631 002</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>20MG; 12.5MG</u> | <u>A076631 003</u> | Feb 11, 2004 |
| <u>AB</u> | | <u>20MG; 25MG</u> | <u>A076631 004</u> | Feb 11, 2004 |

LOTENSIN HCT

| | | | | | |
|-----------|---|-------------------------|---------------------|--------------------|--------------|
| <u>AB</u> | + | US PHARMS HOLDINGS I | <u>10MG; 12.5MG</u> | <u>N020033 002</u> | May 19, 1992 |
| <u>AB</u> | + | | <u>20MG; 12.5MG</u> | <u>N020033 004</u> | May 19, 1992 |
| <u>AB</u> | + | | <u>20MG; 25MG</u> | <u>N020033 003</u> | May 19, 1992 |

BENDAMUSTINE HYDROCHLORIDE

POWDER; IV (INFUSION)

BENDAMUSTINE HYDROCHLORIDE

| | | | | |
|-----------|---------------------|-------------------|--------------------|--------------|
| <u>AP</u> | ACCORD HLTHCARE | <u>25MG/VIAL</u> | <u>A205574 001</u> | May 19, 2016 |
| <u>AP</u> | | <u>100MG/VIAL</u> | <u>A205574 002</u> | May 19, 2016 |
| <u>AP</u> | BRECKENRIDGE PHARM | <u>25MG/VIAL</u> | <u>A205447 001</u> | Dec 29, 2016 |
| <u>AP</u> | | <u>100MG/VIAL</u> | <u>A205447 002</u> | Dec 29, 2016 |
| <u>AP</u> | GLENMARK PHARMS LTD | <u>25MG/VIAL</u> | <u>A204771 001</u> | Mar 24, 2016 |
| <u>AP</u> | | <u>100MG/VIAL</u> | <u>A204771 002</u> | Mar 24, 2016 |
| <u>AP</u> | MYLAN LABS LTD | <u>25MG/VIAL</u> | <u>A204104 001</u> | Apr 26, 2018 |
| <u>AP</u> | | <u>100MG/VIAL</u> | <u>A204104 002</u> | Apr 26, 2018 |

TREANDA

| | | | | | |
|-----------|---|----------|-------------------|--------------------|--------------|
| <u>AP</u> | + | CEPHALON | <u>25MG/VIAL</u> | <u>N022249 002</u> | May 01, 2009 |
| <u>AP</u> | + | | <u>100MG/VIAL</u> | <u>N022249 001</u> | Mar 20, 2008 |

SOLUTION; IV (INFUSION)

BELRAPZO

+! EAGLE PHARMS 100MG/4ML (25MG/ML) N205580 001 May 15, 2018

BENDEKA

+! EAGLE PHARMS 100MG/4ML (25MG/ML) N208194 001 Dec 07, 2015

BENDROFLUMETHIAZIDE; NADOLOL

TABLET; ORAL

CORZIDE

| | | | | | |
|-----------|---|-----------------|------------------|--------------------|--------------|
| <u>AB</u> | + | KING PHARMS LLC | <u>5MG; 40MG</u> | <u>N018647 001</u> | May 25, 1983 |
| <u>AB</u> | + | | <u>5MG; 80MG</u> | <u>N018647 002</u> | May 25, 1983 |

NADOLOL AND BENDROFLUMETHIAZIDE

| | | | | |
|-----------|------------|------------------|--------------------|--------------|
| <u>AB</u> | IMPAX LABS | <u>5MG; 40MG</u> | <u>A077833 001</u> | Mar 30, 2007 |
| <u>AB</u> | | <u>5MG; 80MG</u> | <u>A077833 002</u> | Mar 30, 2007 |

PRESCRIPTION DRUG PRODUCT LIST

BENOXINATE HYDROCHLORIDE; FLUORESCEIN SODIUM

SOLUTION/DROPS;OPHTHALMIC

ALTAFLUOR BENOX

+! ALTAIRE PHARMS INC 0.4%;0.25%

N208582 001 Dec 14, 2017

BENZNIDAZOLE

TABLET;ORAL

BENZNIDAZOLE

+ CHEMO RESEARCH SL 12.5MG

N209570 001 Aug 29, 2017

+! 100MG

N209570 002 Aug 29, 2017

BENZONATATE

CAPSULE;ORAL

BENZONATATEAA AIPING PHARM INC100MGA210562 001 Nov 09, 2018AA 150MGA210562 002 Nov 09, 2018AA 200MGA210562 003 Nov 09, 2018AA APOTEX INC 100MGA091310 001 Jan 16, 2015AA 200MGA091310 002 Jan 16, 2015AA BIONPHARMA INC 100MGA081297 001 Jan 29, 1993AA 200MGA081297 002 Oct 30, 2007AA CSPC NBP PHARM CO 100MGA202765 002 Aug 25, 2017AA 200MGA202765 001 Jul 31, 2015AA MIKART 100MGA040851 001 Nov 09, 2009AA 150MGA040851 002 Nov 09, 2009AA 200MGA040851 003 Nov 09, 2009AA ORIT LABS LLC 100MGA040682 001 Jul 30, 2007AA 200MGA040682 002 Jul 30, 2007AA PURACAP PHARM LLC 100MGA206948 001 Dec 19, 2018AA 200MGA206948 002 Dec 19, 2018AA STRIDES PHARMA 100MGA091133 001 Jul 30, 2015AA 200MGA091133 002 Jul 30, 2015AA SUN PHARM INDS INC 100MGA040587 001 Mar 19, 2008AA 200MGA040587 002 Mar 19, 2008AA ! THEPHARMANETWORK 100MGA040627 001 Mar 30, 2007

LLC

AA ! 150MGA201209 001 Sep 24, 2014AA ! 200MGA040749 001 Jul 25, 2007AA ZYDUS PHARMS USA 100MGA040597 001 Jun 08, 2007AA 200MGA040597 002 Jun 08, 2007TESSALONAA + PFIZER 100MGN011210 001BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL;TOPICAL

ACANYAAB +! DOW PHARM 2.5%;EQ 1.2% BASEN050819 001 Oct 23, 2008BENZACLINAB +! VALEANT BERMUDA 5%;EQ 1% BASEN050756 001 Dec 21, 2000CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDEAB ACTAVIS LABS UT INC 2.5%;EQ 1.2% BASEA205128 001 Jun 19, 2015AB MYLAN PHARMS INC 5%;EQ 1% BASEA065443 001 Aug 11, 2009AB PERRIGO ISRAEL 5%;EQ 1% BASEA202440 001 Sep 21, 2015AB 5%;1.2%A090979 001 Jun 26, 2012AB TARO 3.75%;EQ 1.2% BASEA208683 001 Jun 05, 2018AB 5%;EQ 1% BASEA208776 001 May 25, 2018AB TARO PHARMS 5%;1.2%A206218 001 Dec 15, 2017AB TOLMAR 5%;EQ 1% BASEA204087 001 Jun 27, 2017AB 5%;1.2%A203688 001 Aug 25, 2016AB ZYDUS PHARMS USA 5%;1.2%A210794 001 Dec 28, 2018DUACAB +! STIEFEL 5%;1.2%N050741 001 Aug 26, 2002ONEXTONAB +! DOW PHARM 3.75%;EQ 1.2% BASEN050819 002 Nov 24, 2014BENZOYL PEROXIDE; ERYTHROMYCIN

GEL;TOPICAL

BENZAMYCINAB +! VALEANT INTL 5%;3%N050557 001 Oct 26, 1984ERYTHROMYCIN AND BENZOYL PEROXIDEAB LYNE 5%;3%A065385 001 Sep 18, 2015AB TOLMAR 5%;3%A065112 001 Mar 29, 2004

PRESCRIPTION DRUG PRODUCT LIST

BENZOYL PEROXIDE; ERYTHROMYCIN

GEL; TOPICAL

AKTIPAK

+! CUTANEA

5%;3%

N050769 001 Nov 27, 2000

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL

BENZPHETAMINE HYDROCHLORIDE

| | | | | | |
|-----------|-------------------|-------------|----------------|------------|--------------|
| <u>AA</u> | EMCURE PHARMS LTD | <u>50MG</u> | <u>A202061</u> | <u>001</u> | Jan 27, 2012 |
| <u>AA</u> | EPIC PHARMA LLC | <u>50MG</u> | <u>A090346</u> | <u>001</u> | Dec 15, 2015 |
| <u>AA</u> | ! KVK TECH | <u>50MG</u> | <u>A090968</u> | <u>001</u> | Jul 20, 2010 |
| <u>AA</u> | MIKART | <u>50MG</u> | <u>A090473</u> | <u>002</u> | Sep 15, 2010 |
| <u>AA</u> | SPECGX LLC | <u>50MG</u> | <u>A040773</u> | <u>001</u> | Apr 25, 2007 |
| <u>AA</u> | TWI PHARMS | <u>50MG</u> | <u>A040578</u> | <u>001</u> | Apr 17, 2006 |
| | MIKART | 25MG | A090473 | 001 | Sep 15, 2010 |

BENZTROPINE MESYLATE

INJECTABLE; INJECTION

BENZTROPINE MESYLATE

| | | | | | |
|-----------|--------------------|---------------|----------------|------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>1MG/ML</u> | <u>A090233</u> | <u>001</u> | Jul 28, 2009 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>1MG/ML</u> | <u>A090287</u> | <u>001</u> | Aug 31, 2009 |
| <u>AP</u> | LUITPOLD | <u>1MG/ML</u> | <u>A091152</u> | <u>001</u> | Mar 29, 2010 |
| <u>AP</u> | NAVINTA LLC | <u>1MG/ML</u> | <u>A091525</u> | <u>001</u> | Feb 05, 2013 |

COGENTIN

| | | | | | |
|-----------|---------------------|---------------|----------------|------------|--|
| <u>AP</u> | +! OAK PHARMS AKORN | <u>1MG/ML</u> | <u>N012015</u> | <u>001</u> | |
|-----------|---------------------|---------------|----------------|------------|--|

TABLET; ORAL

BENZTROPINE MESYLATE

| | | | | | |
|-----------|--------------------|--------------|----------------|------------|--------------|
| <u>AA</u> | ASPEN GLOBAL INC | <u>0.5MG</u> | <u>A204713</u> | <u>001</u> | Apr 14, 2015 |
| <u>AA</u> | | <u>1MG</u> | <u>A204713</u> | <u>002</u> | Apr 14, 2015 |
| <u>AA</u> | | <u>2MG</u> | <u>A204713</u> | <u>003</u> | Apr 14, 2015 |
| <u>AA</u> | EPIC PHARMA LLC | <u>0.5MG</u> | <u>A072264</u> | <u>001</u> | Feb 27, 1989 |
| <u>AA</u> | | <u>1MG</u> | <u>A072265</u> | <u>001</u> | Feb 27, 1989 |
| <u>AA</u> | | <u>2MG</u> | <u>A072266</u> | <u>001</u> | Feb 27, 1989 |
| <u>AA</u> | INVAGEN PHARMS | <u>0.5MG</u> | <u>A090294</u> | <u>001</u> | Mar 29, 2010 |
| <u>AA</u> | | <u>1MG</u> | <u>A090294</u> | <u>002</u> | Mar 29, 2010 |
| <u>AA</u> | | <u>2MG</u> | <u>A090294</u> | <u>003</u> | Mar 29, 2010 |
| <u>AA</u> | LEADING PHARMA LLC | <u>0.5MG</u> | <u>A090168</u> | <u>001</u> | Nov 28, 2012 |
| <u>AA</u> | | <u>1MG</u> | <u>A090168</u> | <u>002</u> | Nov 28, 2012 |
| <u>AA</u> | | <u>2MG</u> | <u>A090168</u> | <u>003</u> | Nov 28, 2012 |
| <u>AA</u> | PLIVA | <u>0.5MG</u> | <u>A089058</u> | <u>001</u> | Aug 10, 1988 |
| <u>AA</u> | | <u>1MG</u> | <u>A089059</u> | <u>001</u> | Aug 10, 1988 |
| <u>AA</u> | | <u>2MG</u> | <u>A089060</u> | <u>001</u> | Aug 10, 1988 |
| <u>AA</u> | ! USL PHARMA | <u>0.5MG</u> | <u>A040103</u> | <u>001</u> | Dec 12, 1996 |
| <u>AA</u> | ! | <u>1MG</u> | <u>A040103</u> | <u>002</u> | Dec 12, 1996 |
| <u>AA</u> | ! | <u>2MG</u> | <u>A040103</u> | <u>003</u> | Dec 12, 1996 |
| <u>AA</u> | VINTAGE | <u>0.5MG</u> | <u>A040715</u> | <u>001</u> | Aug 27, 2007 |
| <u>AA</u> | | <u>1MG</u> | <u>A040715</u> | <u>002</u> | Aug 27, 2007 |
| <u>AA</u> | | <u>2MG</u> | <u>A040715</u> | <u>003</u> | Aug 27, 2007 |
| | OXFORD PHARMS | 0.5MG | A040706 | 002 | Feb 14, 2008 |
| | | 1MG | A040706 | 003 | Feb 14, 2008 |

BENZYL ALCOHOL

LOTION; TOPICAL

ULESFIA

+! SHIONOGI INC

5%

N022129 001 Apr 09, 2009

BENZYPENICILLOYL POLYLYSINE

INJECTABLE; INJECTION

PRE-PEN

+! ALLERQUEST

60UMOLAR

N050114 001

BEPOTASTINE BESILATE

SOLUTION/DROPS; OPHTHALMIC

BEPREVE

+! BAUSCH AND LOMB INC

1.5%

N022288 001 Sep 08, 2009

BERACTANT

SUSPENSION; INTRATRACHEAL

SURVANTA

+! ABBVIE

25MG/ML

N020032 001 Jul 01, 1991

PRESCRIPTION DRUG PRODUCT LIST

BESIFLOXACIN HYDROCHLORIDE

SUSPENSION/DROPS;OPHTHALMIC

BESIVANCE

+! BAUSCH AND LOMB EQ 0.6% BASE N022308 001 May 28, 2009

BETAINE

FOR SOLUTION;ORAL

CYSTADANE

+! ORPHAN EUROPE 1GM/SCOOPFUL N020576 001 Oct 25, 1996

BETAMETHASONE ACETATE; BETAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE**AB** LUITPOLD 3MG/ML;EQ 3MG BASE/ML **A090747 001** Jul 31, 2009CELESTONE SOLUSPAN**AB** +! MERCK SHARP DOHME 3MG/ML;EQ 3MG BASE/ML **N014602 001**BETAMETHASONE DIPROPIONATE

CREAM;TOPICAL

BETAMETHASONE DIPROPIONATE**AB** ACTAVIS MID EQ 0.05% BASE **A070885 001** Feb 03, 1987

ATLANTIC

AB +! FOUGERA PHARMS EQ 0.05% BASE **N019137 001** Jun 26, 1984**AB** G AND W LABS INC EQ 0.05% BASE **A210217 001** Oct 12, 2018**AB** TARO EQ 0.05% BASE **A073552 001** Apr 30, 1992**AB** ZYDUS PHARMS USA EQ 0.05% BASE **A208885 001** Jan 11, 2019

INC

CREAM, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATE**AB** ANDA REPOSITORY EQ 0.05% BASE **A076603 001** Jan 23, 2004**AB** FOUGERA PHARMS EQ 0.05% BASE **A076215 001** Dec 09, 2003**AB** GLENMARK GENERICS EQ 0.05% BASE **A078930 001** Sep 23, 2008**AB** PERRIGO NEW YORK EQ 0.05% BASE **A076592 001** Dec 09, 2003**AB** TARO EQ 0.05% BASE **A076543 001** Dec 09, 2003DIPROLENE AF**AB** +! MERCK SHARP DOHME EQ 0.05% BASE **N019555 001** Apr 27, 1987

GEL, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATE**AB** ! FOUGERA PHARMS EQ 0.05% BASE **A075276 001** May 13, 2003**AB** TARO EQ 0.05% BASE **A076508 001** Dec 02, 2003

LOTION;TOPICAL

BETAMETHASONE DIPROPIONATE**AB** ACTAVIS MID EQ 0.05% BASE **A070281 001** Jul 31, 1985

ATLANTIC

AB ! FOUGERA PHARMS INC EQ 0.05% BASE **A070275 001** Aug 12, 1985**AB** G AND W LABS INC EQ 0.05% BASE **A071467 001** Aug 10, 1987**AB** HI-TECH PHARMACAL EQ 0.05% BASE **A209896 001** Feb 06, 2018**AB** PERRIGO NEW YORK EQ 0.05% BASE **A072538 001** Jan 31, 1990

LOTION, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATE**AB** FOUGERA PHARMS EQ 0.05% BASE **A077111 001** May 21, 2007**AB** ! TARO EQ 0.05% BASE **A077477 001** May 21, 2007**AB** TELIGENT PHARMA INC EQ 0.05% BASE **A206389 001** Feb 13, 2018

OINTMENT;TOPICAL

BETAMETHASONE DIPROPIONATE**AB** ACTAVIS MID EQ 0.05% BASE **A071012 001** Feb 03, 1987

ATLANTIC

AB +! FOUGERA PHARMS INC EQ 0.05% BASE **N019141 001** Sep 04, 1984**AB** TARO EQ 0.05% BASE **A074271 001** Sep 15, 1994

OINTMENT, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATE**AB** ACTAVIS MID EQ 0.05% BASE **A074304 001** Aug 31, 1995

ATLANTIC

AB FOUGERA PHARMS EQ 0.05% BASE **A075373 001** Jun 22, 1999**AB** TARO EQ 0.05% BASE **A076753 001** Oct 12, 2004**AB** TELIGENT PHARMA INC EQ 0.05% BASE **A206118 001** Nov 09, 2017DIPROLENE**AB** +! MERCK SHARP DOHME EQ 0.05% BASE **N018741 001** Jul 27, 1983

SPRAY;TOPICAL

SERNIVO

+! PROMIUS PHARMA LLC EQ 0.05% BASE/SPRAY N208079 001 Feb 05, 2016

PRESCRIPTION DRUG PRODUCT LIST

BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE

AEROSOL, FOAM; TOPICAL

ENSTILAR

+! LEO PHARMA AS 0.064%;0.005% N207589 001 Oct 16, 2015

BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE

OINTMENT; TOPICAL

CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE**AB** PERRIGO ISRAEL 0.064%;0.005% **A200174 001** Dec 12, 2014**AB** TOLMAR 0.064%;0.005% **A201615 001** Jan 14, 2013TACLONEX**AB** +! LEO PHARMA AS 0.064%;0.005% **N021852 001** Jan 09, 2006

SUSPENSION; TOPICAL

TACLONEX

+! LEO PHARMA AS 0.064%;0.005% N022185 001 May 09, 2008

BETAMETHASONE DIPROPIONATE; CLOTRIMAZOLE

CREAM; TOPICAL

CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE**AB** ACTAVIS MID EQ 0.05% BASE;1% **A076002 001** Aug 02, 2002

ATLANTIC

AB FOUGERA PHARMS EQ 0.05% BASE;1% **A075502 001** Jun 05, 2001**AB** GLENMARK PHARMS EQ 0.05% BASE;1% **A202894 001** Oct 30, 2015**AB** TARO EQ 0.05% BASE;1% **A075673 001** May 29, 2001LOTRISONE**AB** +! MERCK SHARP DOHME EQ 0.05% BASE;1% **N018827 001** Jul 10, 1984

LOTION; TOPICAL

CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE**AB** FOUGERA PHARMS EQ 0.05% BASE;1% **A076516 001** Jun 16, 2005**AB** TARO EQ 0.05% BASE;1% **A076493 001** Jul 28, 2004LOTRISONE**AB** +! MERCK SHARP DOHME EQ 0.05% BASE;1% **N020010 001** Dec 08, 2000BETAMETHASONE VALERATE

AEROSOL, FOAM; TOPICAL

BETAMETHASONE VALERATE**AB** PERRIGO UK FINCO 0.12% **A078337 001** Nov 26, 2012**AB** RICONPHARMA LLC 0.12% **A207144 001** May 24, 2017**AB** TARO PHARM 0.12% **A208204 001** May 24, 2017LUXIQ**AB** +! MYLAN PHARMS INC 0.12% **N020934 001** Feb 28, 1999

CREAM; TOPICAL

BETA-VAL**AB** G AND W LABS INC EQ 0.1% BASE **N018642 001** Mar 24, 1983BETAMETHASONE VALERATE**AB** +! FOUGERA PHARMS INC EQ 0.1% BASE **N018861 001** Aug 31, 1983DERMABET**AB** TARO EQ 0.1% BASE **A072041 001** Jan 06, 1988VALNAC**AB** ACTAVIS MID EQ 0.1% BASE **A070050 001** Oct 10, 1984

ATLANTIC

LOTION; TOPICAL

BETAMETHASONE VALERATE**AB** +! FOUGERA PHARMS INC EQ 0.1% BASE **N018866 001** Aug 31, 1983**AB** STI PHARMA LLC EQ 0.1% BASE **A070052 001** Jul 31, 1985

OINTMENT; TOPICAL

BETA-VAL**AB** G AND W LABS INC EQ 0.1% BASE **A070069 001** Dec 19, 1985BETAMETHASONE VALERATE**AB** ACTAVIS MID EQ 0.1% BASE **A070051 001** Oct 10, 1984

ATLANTIC

AB +! FOUGERA PHARMS INC EQ 0.1% BASE **N018865 001** Aug 31, 1983BETAXOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

BETAXOLOL HYDROCHLORIDE**AT** AKORN EQ 0.5% BASE **A075386 001** Jun 30, 2000**AT** MEDIMETRIKS PHARMS EQ 0.5% BASE **A075630 001** Apr 12, 2001**AT** WOCKHARDT EQ 0.5% BASE **A078694 001** Nov 16, 2009BETOPTIC**AT** +! SANDOZ INC EQ 0.5% BASE **N019270 001** Aug 30, 1985

SUSPENSION/DROPS; OPHTHALMIC

BETOPTIC S

+! NOVARTIS PHARMS EQ 0.25% BASE N019845 001 Dec 29, 1989

CORP

PRESCRIPTION DRUG PRODUCT LIST

BETAXOLOL HYDROCHLORIDE

TABLET; ORAL

BETAXOLOL HYDROCHLORIDE

| | | | | |
|-----------|-------------|-------------|--------------------|--------------|
| <u>AB</u> | EPIC PHARMA | <u>10MG</u> | <u>A075541 001</u> | Oct 22, 1999 |
| <u>AB</u> | ! | <u>20MG</u> | <u>A075541 002</u> | Oct 22, 1999 |
| <u>AB</u> | KVK TECH | <u>10MG</u> | <u>A078962 001</u> | Jun 27, 2008 |
| <u>AB</u> | | <u>20MG</u> | <u>A078962 002</u> | Jun 27, 2008 |

BETHANECHOL CHLORIDE

TABLET; ORAL

BETHANECHOL CHLORIDE

| | | | | |
|-------------------|-------------------|-------------|--------------------|--------------|
| <u>AA</u> | AMNEAL PHARM | <u>5MG</u> | <u>A040855 001</u> | Nov 21, 2007 |
| <u>AA</u> | | <u>10MG</u> | <u>A040855 002</u> | Nov 21, 2007 |
| <u>AA</u> | | <u>25MG</u> | <u>A040855 003</u> | Nov 21, 2007 |
| <u>AA</u> | | <u>50MG</u> | <u>A040855 004</u> | Nov 21, 2007 |
| <u>AA</u> | ECI PHARMS LLC | <u>5MG</u> | <u>A040725 001</u> | Oct 26, 2007 |
| <u>AA</u> | | <u>10MG</u> | <u>A040726 001</u> | Oct 26, 2007 |
| <u>AA</u> | | <u>25MG</u> | <u>A040727 001</u> | Oct 26, 2007 |
| <u>AA</u> | | <u>50MG</u> | <u>A040728 001</u> | Oct 26, 2007 |
| <u>AA</u> | HERITAGE PHARMA | <u>5MG</u> | <u>A091256 001</u> | May 04, 2010 |
| <u>AA</u> | | <u>10MG</u> | <u>A091256 002</u> | May 04, 2010 |
| <u>AA</u> | | <u>25MG</u> | <u>A091256 003</u> | May 04, 2010 |
| <u>AA</u> | | <u>50MG</u> | <u>A091256 004</u> | May 04, 2010 |
| <u>AA</u> | LANNETT CO INC | <u>5MG</u> | <u>A040677 002</u> | Mar 27, 2008 |
| <u>AA</u> | | <u>10MG</u> | <u>A040677 003</u> | Mar 27, 2008 |
| <u>AA</u> | | <u>25MG</u> | <u>A040677 004</u> | Mar 27, 2008 |
| <u>AA</u> | | <u>50MG</u> | <u>A040677 001</u> | Mar 27, 2008 |
| <u>AA</u> | UPSHER SMITH LABS | <u>5MG</u> | <u>A040633 001</u> | Jun 01, 2005 |
| <u>AA</u> | | <u>10MG</u> | <u>A040634 001</u> | Jun 01, 2005 |
| <u>AA</u> | | <u>25MG</u> | <u>A040635 001</u> | Jun 01, 2005 |
| <u>AA</u> | | <u>50MG</u> | <u>A040636 001</u> | Jun 01, 2005 |
| <u>AA</u> | WOCKHARDT | <u>5MG</u> | <u>A040532 001</u> | Sep 29, 2003 |
| <u>AA</u> | | <u>10MG</u> | <u>A040533 001</u> | Sep 29, 2003 |
| <u>AA</u> | | <u>25MG</u> | <u>A040534 001</u> | Sep 29, 2003 |
| <u>AA</u> | | <u>50MG</u> | <u>A040518 001</u> | Sep 29, 2003 |
| <u>DUVOID</u> | | | | |
| <u>AA</u> | BI-COASTAL PHARMA | <u>10MG</u> | <u>A086262 001</u> | |
| <u>AA</u> | | <u>25MG</u> | <u>A086263 001</u> | |
| <u>AA</u> | | <u>50MG</u> | <u>A085882 003</u> | |
| <u>URECHOLINE</u> | | | | |
| <u>AA</u> | ! ODYSSEY PHARMS | <u>5MG</u> | <u>A089095 001</u> | Dec 19, 1985 |
| <u>AA</u> | ! | <u>10MG</u> | <u>A088440 001</u> | May 29, 1984 |
| <u>AA</u> | ! | <u>25MG</u> | <u>A088441 001</u> | May 29, 1984 |
| <u>AA</u> | ! | <u>50MG</u> | <u>A089096 001</u> | Dec 19, 1985 |

BETRIXABAN

CAPSULE; ORAL

BEVYXXA

| | | | | |
|---|--------------------|------|-------------|--------------|
| + | PORTOLA PHARMS INC | 40MG | N208383 001 | Jun 23, 2017 |
| + | ! | 80MG | N208383 002 | Jun 23, 2017 |

BEXAROTENE

CAPSULE; ORAL

BEXAROTENE

| | | | | |
|-----------------|-----------------------|-------------|--------------------|--------------|
| <u>AB</u> | AMERIGEN PHARMS LTD | <u>75MG</u> | <u>A209861 001</u> | May 08, 2018 |
| <u>AB</u> | AMNEAL PHARMS NY | <u>75MG</u> | <u>A210105 001</u> | Sep 04, 2018 |
| <u>AB</u> | BIONPHARMA INC | <u>75MG</u> | <u>A203174 001</u> | Aug 12, 2014 |
| <u>AB</u> | UPSHER SMITH LABS | <u>75MG</u> | <u>A209886 001</u> | Jul 25, 2018 |
| <u>TARGETIN</u> | | | | |
| <u>AB</u> | +! VALEANT LUXEMBOURG | <u>75MG</u> | <u>N021055 001</u> | Dec 29, 1999 |
| GEL; TOPICAL | | | | |
| TARGETIN | | | | |
| | +! VALEANT LUXEMBOURG | 1% | N021056 001 | Jun 28, 2000 |

BICALUTAMIDE

TABLET; ORAL

BICALUTAMIDE

| | | | | |
|-----------|-------------------------|-------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>50MG</u> | <u>A078917 001</u> | Jul 06, 2009 |
| <u>AB</u> | APOTEX INC | <u>50MG</u> | <u>A200274 001</u> | May 21, 2015 |
| <u>AB</u> | BOSCOGEN | <u>50MG</u> | <u>A091011 001</u> | Jun 10, 2015 |
| <u>AB</u> | FRESENIUS KABI ONCOL | <u>50MG</u> | <u>A079045 001</u> | May 13, 2010 |
| <u>AB</u> | MYLAN | <u>50MG</u> | <u>A079185 001</u> | Jul 06, 2009 |
| <u>AB</u> | SANDOZ | <u>50MG</u> | <u>A078575 001</u> | Jul 06, 2009 |
| <u>AB</u> | SUN PHARMA GLOBAL | <u>50MG</u> | <u>A079110 001</u> | Jul 06, 2009 |

PRESCRIPTION DRUG PRODUCT LIST

BICALUTAMIDE

TABLET;ORAL

BICALUTAMIDE

| | | | | |
|-----------|-------------------------|-------------|--------------------|--------------|
| AB | TEVA | 50MG | A076932 001 | Jul 06, 2009 |
| AB | WATSON LABS TEVA | 50MG | A078634 001 | Aug 28, 2009 |
| AB | ZYDUS PHARMS USA INC | 50MG | A079089 001 | Jul 06, 2009 |

CASODEX

| | | | | |
|-----------|-------------------|-------------|--------------------|--------------|
| AB | +! ANI PHARMS INC | 50MG | N020498 001 | Oct 04, 1995 |
|-----------|-------------------|-------------|--------------------|--------------|

BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

BIKTARVY

| | | | | |
|----|---------------------|---------------------------------|-------------|--------------|
| +! | GILEAD SCIENCES INC | EQ 50MG BASE;200MG;EQ 25MG BASE | N210251 001 | Feb 07, 2018 |
|----|---------------------|---------------------------------|-------------|--------------|

BIMATOPROST

SOLUTION/DROPS;OPHTHALMIC

BIMATOPROST

| | | | | |
|-----------|-------------------|--------------|--------------------|--------------|
| AT | APOTEX INC | 0.03% | A090449 001 | Jul 20, 2015 |
| AT | HI-TECH PHARMA CO | 0.03% | A203299 001 | Nov 08, 2018 |
| AT | ! LUPIN LTD | 0.03% | A203991 001 | Feb 20, 2015 |
| AT | SANDOZ INC | 0.03% | A202565 001 | May 05, 2015 |

LUMIGAN

| | | | | |
|----|----------|-------|-------------|--------------|
| +! | ALLERGAN | 0.01% | N022184 001 | Aug 31, 2010 |
|----|----------|-------|-------------|--------------|

SOLUTION/DROPS;TOPICAL

BIMATOPROST

| | | | | |
|-----------|-------------------|--------------|--------------------|--------------|
| AT | APOTEX INC | 0.03% | A201894 001 | Dec 01, 2014 |
| AT | HI-TECH PHARMACAL | 0.03% | A203051 001 | Oct 09, 2018 |
| AT | SANDOZ INC | 0.03% | A202719 001 | Apr 19, 2016 |

LATISSE

| | | | | |
|-----------|-------------|--------------|--------------------|--------------|
| AT | +! ALLERGAN | 0.03% | N022369 001 | Dec 24, 2008 |
|-----------|-------------|--------------|--------------------|--------------|

BINIMETINIB

TABLET;ORAL

MEKTOVI

| | | | | |
|----|---------------------|------|-------------|--------------|
| +! | ARRAY BIOPHARMA INC | 15MG | N210498 001 | Jun 27, 2018 |
|----|---------------------|------|-------------|--------------|

BISMUTH SUBCITRATE POTASSIUM; METRONIDAZOLE; TETRACYCLINE

CAPSULE;ORAL

PYLERA

| | | | | |
|----|--------------------|-------------------|-------------|--------------|
| +! | ALLERGAN SALES LLC | 140MG;125MG;125MG | N050786 001 | Sep 28, 2006 |
|----|--------------------|-------------------|-------------|--------------|

BISMUTH SUBSALICYLATE; METRONIDAZOLE; TETRACYCLINE HYDROCHLORIDE

TABLET, CHEWABLE, TABLET, CAPSULE;ORAL

BISMUTH SUBSALICYLATE, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE

| | | | | |
|---|------------------|------------------------------------------------------|-------------|--------------|
| ! | AILEX PHARMS LLC | 262.4MG, N/A, N/A;N/A, 250MG, N/A;N/A, N/A, 500MG | A202584 001 | Nov 30, 2018 |
|---|------------------|------------------------------------------------------|-------------|--------------|

BISOPROLOL FUMARATE

TABLET;ORAL

BISOPROLOL FUMARATE

| | | | | |
|-----------|-------------------------|-------------|--------------------|--------------|
| AB | AUROBINDO PHARMA | 5MG | A077910 001 | Dec 27, 2006 |
| AB | | 10MG | A077910 002 | Dec 27, 2006 |
| AB | CASI PHARMS INC | 5MG | A075643 001 | Nov 16, 2000 |
| AB | | 10MG | A075643 002 | Nov 16, 2000 |
| AB | MYLAN | 5MG | A075831 001 | Dec 14, 2005 |
| AB | ! | 10MG | A075831 002 | Dec 14, 2005 |
| AB | ORIT LABS LLC | 5MG | A204891 001 | Jan 11, 2017 |
| AB | | 10MG | A204891 002 | Jan 11, 2017 |
| AB | TEVA PHARMS | 5MG | A075644 001 | Jun 26, 2001 |
| AB | | 10MG | A075644 002 | Jun 26, 2001 |
| AB | UNICHEM PHARMS (USA) | 5MG | A078635 001 | Aug 18, 2009 |
| AB | | 10MG | A078635 002 | Aug 18, 2009 |

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

| | | | | |
|-----------|---------|---------------------|--------------------|--------------|
| AB | MYLAN | 2.5MG;6.25MG | A075768 001 | Sep 25, 2000 |
| AB | | 5MG;6.25MG | A075768 002 | Sep 25, 2000 |
| AB | | 10MG;6.25MG | A075768 003 | Sep 25, 2000 |
| AB | SANDOZ | 2.5MG;6.25MG | A075579 001 | Sep 25, 2000 |
| AB | | 5MG;6.25MG | A075579 002 | Sep 25, 2000 |
| AB | | 10MG;6.25MG | A075579 003 | Sep 25, 2000 |
| AB | UNICHEM | 2.5MG;6.25MG | A079106 001 | Jul 28, 2010 |
| AB | | 5MG;6.25MG | A079106 002 | Jul 28, 2010 |

PRESCRIPTION DRUG PRODUCT LIST

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

| | | | | | |
|-----------|-------------|--------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | | <u>10MG; 6.25MG</u> | <u>A079106 003</u> | Jul 28, 2010 |
| | <u>ZIAC</u> | | | | |
| <u>AB</u> | + | TEVA BRANDED PHARM | <u>2.5MG; 6.25MG</u> | <u>N020186 003</u> | Mar 26, 1993 |
| <u>AB</u> | + | | <u>5MG; 6.25MG</u> | <u>N020186 001</u> | Mar 26, 1993 |
| <u>AB</u> | + | ! | <u>10MG; 6.25MG</u> | <u>N020186 002</u> | Mar 26, 1993 |

BIVALIRUDIN

INJECTABLE; INTRAVENOUS

ANGIOMAX

| | | | | | |
|-----------|---|------------|-------------------|--------------------|--------------|
| <u>AP</u> | + | SANDOZ INC | <u>250MG/VIAL</u> | <u>N020873 001</u> | Dec 15, 2000 |
|-----------|---|------------|-------------------|--------------------|--------------|

BIVALIRUDIN

| | | | | | |
|-----------|--|----------------------|-------------------|--------------------|--------------|
| <u>AP</u> | | ACCORD HLTHCARE | <u>250MG/VIAL</u> | <u>A206551 001</u> | Nov 22, 2017 |
| <u>AP</u> | | APOTEX INC | <u>250MG/VIAL</u> | <u>A204876 001</u> | Jul 06, 2017 |
| <u>AP</u> | | AUROBINDO PHARMA LTD | <u>250MG/VIAL</u> | <u>A205962 001</u> | Jul 27, 2018 |
| <u>AP</u> | | CIPLA | <u>250MG/VIAL</u> | <u>A091602 001</u> | Jul 16, 2018 |
| <u>AP</u> | | DR REDDYS LABS LTD | <u>250MG/VIAL</u> | <u>A201577 001</u> | May 26, 2017 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>250MG/VIAL</u> | <u>A090189 001</u> | Oct 28, 2016 |
| <u>AP</u> | | HOSPIRA INC | <u>250MG/VIAL</u> | <u>A090811 001</u> | Jul 14, 2015 |
| <u>AP</u> | | | <u>250MG/VIAL</u> | <u>A090816 001</u> | Jul 14, 2015 |
| <u>AP</u> | | MYLAN INSTITUTIONAL | <u>250MG/VIAL</u> | <u>A202471 001</u> | Jun 01, 2018 |

SOLUTION; INTRAVENOUS

BIVALIRUDIN IN 0.9% SODIUM CHLORIDE

| | | | | | |
|--|---|----------------------|----------------------|-------------|--------------|
| | + | BAXTER HLTHCARE CORP | 250MG/50ML (5MG/ML) | N208374 001 | Dec 21, 2017 |
| | + | | 500MG/100ML (5MG/ML) | N208374 002 | Dec 21, 2017 |

BLEOMYCIN SULFATE

INJECTABLE; INJECTION

BLEOMYCIN SULFATE

| | | | | | |
|-----------|---|----------------------|------------------------------|--------------------|--------------|
| <u>AP</u> | ! | FRESENIUS KABI USA | <u>EQ 15 UNITS BASE/VIAL</u> | <u>A065185 001</u> | Jan 28, 2008 |
| <u>AP</u> | ! | | <u>EQ 30 UNITS BASE/VIAL</u> | <u>A065185 002</u> | Jan 28, 2008 |
| <u>AP</u> | | HONG KONG | <u>EQ 15 UNITS BASE/VIAL</u> | <u>A205030 001</u> | Apr 20, 2018 |
| <u>AP</u> | | | <u>EQ 30 UNITS BASE/VIAL</u> | <u>A205030 002</u> | Apr 20, 2018 |
| <u>AP</u> | | HOSPIRA | <u>EQ 15 UNITS BASE/VIAL</u> | <u>A065031 001</u> | Mar 10, 2000 |
| <u>AP</u> | | | <u>EQ 30 UNITS BASE/VIAL</u> | <u>A065031 002</u> | Mar 10, 2000 |
| <u>AP</u> | | TEVA PHARMS USA | <u>EQ 15 UNITS BASE/VIAL</u> | <u>A065033 001</u> | Jun 27, 2000 |
| <u>AP</u> | | | <u>EQ 30 UNITS BASE/VIAL</u> | <u>A065033 002</u> | Jun 27, 2000 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>EQ 15 UNITS BASE/VIAL</u> | <u>A065042 002</u> | Oct 17, 2001 |
| <u>AP</u> | | | <u>EQ 30 UNITS BASE/VIAL</u> | <u>A065042 001</u> | Oct 17, 2001 |

BORTEZOMIB

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

VELCADE

| | | | | | |
|--|---|-------------------|------------|-------------|--------------|
| | + | MILLENNIUM PHARMS | 3.5MG/VIAL | N021602 001 | May 13, 2003 |
|--|---|-------------------|------------|-------------|--------------|

POWDER; INTRAVENOUS

BORTEZOMIB

| | | | | | |
|--|--|--------------------|------------|-------------|--------------|
| | | FRESENIUS KABI USA | 3.5MG/VIAL | N205004 001 | Nov 06, 2017 |
|--|--|--------------------|------------|-------------|--------------|

POWDER; INTRAVENOUS, SUBCUTANEOUS

BORTEZOMIB

| | | | | | |
|--|---|-------------|----------|-------------|--------------|
| | + | HOSPIRA INC | 1MG/VIAL | N209191 002 | Dec 28, 2018 |
|--|---|-------------|----------|-------------|--------------|

BOSENTAN

TABLET; ORAL

TRACLEER

| | | | | | |
|--|---|---------------------|--------|-------------|--------------|
| | + | ACTELION PHARMS LTD | 62.5MG | N021290 001 | Nov 20, 2001 |
|--|---|---------------------|--------|-------------|--------------|

| | | | | | |
|--|---|---|-------|-------------|--------------|
| | + | ! | 125MG | N021290 002 | Nov 20, 2001 |
|--|---|---|-------|-------------|--------------|

TABLET, FOR SUSPENSION; ORAL

TRACLEER

| | | | | | |
|--|---|-----------------|------|-------------|--------------|
| | + | ACTELION PHARMS | 32MG | N209279 001 | Sep 05, 2017 |
|--|---|-----------------|------|-------------|--------------|

BOSUTINIB MONOHYDRATE

TABLET; ORAL

BOSULIF

| | | | | | |
|--|---|-------------|---------------|-------------|--------------|
| | + | PF PRISM CV | EQ 100MG BASE | N203341 001 | Sep 04, 2012 |
|--|---|-------------|---------------|-------------|--------------|

| | | | | | |
|--|---|--|---------------|-------------|--------------|
| | + | | EQ 400MG BASE | N203341 003 | Oct 27, 2017 |
|--|---|--|---------------|-------------|--------------|

| | | | | | |
|--|---|--|---------------|-------------|--------------|
| | + | | EQ 500MG BASE | N203341 002 | Sep 04, 2012 |
|--|---|--|---------------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

BRETYLIUM TOSYLATE

INJECTABLE; INJECTION

BRETYLIUM TOSYLATE

! ACADEMIC PHARMS INC 50MG/ML

A204386 001 Dec 21, 2018

BREXPIPIRAZOLE

TABLET; ORAL

REXULTI

+ OTSUKA PHARM CO LTD 0.25MG

N205422 001 Jul 10, 2015

+ 0.5MG

N205422 002 Jul 10, 2015

+ 1MG

N205422 003 Jul 10, 2015

+! 2MG

N205422 004 Jul 10, 2015

+ 3MG

N205422 005 Jul 10, 2015

+ 4MG

N205422 006 Jul 10, 2015

BRIGATINIB

TABLET; ORAL

ALUNBRIG

+ ARIAD 30MG

N208772 001 Apr 28, 2017

+ 90MG

N208772 002 Apr 28, 2017

+! 180MG

N208772 003 Oct 02, 2017

BRIMONIDINE TARTRATE

GEL; TOPICAL

MIRVASO

+! GALDERMA LABS LP EQ 0.33% BASE

N204708 001 Aug 23, 2013

SOLUTION/DROPS; OPHTHALMIC

ALPHAGAN P**AT** +! ALLERGAN **0.15%****N021262 001** Mar 16, 2001BRIMONIDINE TARTRATE**AT** AKORN **0.2%****A076439 001** Mar 14, 2006**AT** ! BAUSCH AND LOMB **0.2%****A076260 001** May 28, 2003**AT** INDOCO REMEDIES **0.2%****A091691 001** Nov 18, 2014**AT** SANDOZ INC **0.2%****A076254 001** Sep 16, 2003**AT** **0.2%****A078075 001** Jan 30, 2008QOLIANA**AT** SANDOZ INC **0.15%****N021764 001** May 22, 2006

ALPHAGAN P

+! ALLERGAN 0.1%

N021770 001 Aug 19, 2005

BRIMONIDINE TARTRATE; BRINZOLAMIDE

SUSPENSION/DROPS; OPHTHALMIC

SIMBRINZA

+! NOVARTIS PHARMS 0.2%; 1%

N204251 001 Apr 19, 2013

CORP

BRIMONIDINE TARTRATE; TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

COMBIGAN

+! ALLERGAN 0.2%; EQ 0.5% BASE

N021398 001 Oct 30, 2007

BRINZOLAMIDE

SUSPENSION/DROPS; OPHTHALMIC

AZOPT

+! NOVARTIS PHARMS 1%

N020816 001 Apr 01, 1998

CORP

BRIVARACETAM

SOLUTION; INTRAVENOUS

BRIVIACT

+! UCB INC 50MG/5ML (10MG/ML)

N205837 001 May 12, 2016

SOLUTION; ORAL

BRIVIACT

+! UCB INC 10MG/ML

N205838 001 May 12, 2016

TABLET; ORAL

BRIVIACT

+ UCB INC 10MG

N205836 001 May 12, 2016

+ 25MG

N205836 002 May 12, 2016

+ 50MG

N205836 003 May 12, 2016

+ 75MG

N205836 004 May 12, 2016

+! 100MG

N205836 005 May 12, 2016

PRESCRIPTION DRUG PRODUCT LIST

BROMFENAC SODIUM

SOLUTION/DROPS;OPHTHALMIC

BROMFENAC SODIUM

! HI-TECH PHARMACAL EQ 0.09% ACID

A203395 001 Jan 22, 2014

BROMSITE

+! SUN PHARMA GLOBAL EQ 0.075% ACID

N206911 001 Apr 08, 2016

PROLENSA

+! BAUSCH AND LOMB EQ 0.07% ACID

N203168 001 Apr 05, 2013

BROMOCRIPTINE MESYLATE

CAPSULE;ORAL

BROMOCRIPTINE MESYLATE**AB** ! MYLAN **EQ 5MG BASE****A077226 001** Apr 04, 2005**AB** ZYDUS PHARMS USA **EQ 5MG BASE****A078899 001** Jul 30, 2008

INC

PARLODEL**AB** + US PHARMS HOLDINGS **EQ 5MG BASE****N017962 002** Mar 01, 1982

I

TABLET;ORAL

BROMOCRIPTINE MESYLATE**AB** MYLAN **EQ 2.5MG BASE****A076962 001** Sep 24, 2004**AB** ! PADDOCK LLC **EQ 2.5MG BASE****A077646 001** Oct 01, 2008**AB** SANDOZ INC **EQ 2.5MG BASE****A074631 001** Jan 13, 1998PARLODEL**AB** + US PHARMS HOLDINGS **EQ 2.5MG BASE****N017962 001**

I

CYCLOSET

+! VEROSCIENCE EQ 0.8MG BASE

N020866 001 May 05, 2009

BROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SYRUP;ORAL

BROMFED-DM**AA** ! WOCKHARDT BIO AG **2MG/5ML;10MG/5ML;30MG/5ML****A088811 001** Jun 07, 1985BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE**AA** ACELLA PHARMS LLC **2MG/5ML;10MG/5ML;30MG/5ML****A203375 001** Sep 20, 2016**AA** MAYNE PHARMA INC **2MG/5ML;10MG/5ML;30MG/5ML****A207676 001** Dec 04, 2018**AA** PADDOCK LLC **2MG/5ML;10MG/5ML;30MG/5ML****A205292 001** Jul 15, 2014**AA** TARO PHARM **2MG/5ML;10MG/5ML;30MG/5ML****A205112 001** Feb 27, 2017**AA** VINTAGE PHARMS **2MG/5ML;10MG/5ML;30MG/5ML****A202940 001** Jul 21, 2014BUDESONIDE

AEROSOL, FOAM;RECTAL

UCERIS

+! VALEANT PHARMS INTL 2MG/ACTUATION

N205613 001 Oct 07, 2014

CAPSULE;ORAL

BUDESONIDE**AB** ALVOGEN MALTA **3MG****A206724 001** Nov 23, 2016**AB** AMNEAL PHARMS **3MG****A206200 001** Jul 31, 2017**AB** BARR LABS DIV TEVA **3MG****A090379 001** Apr 02, 2014**AB** MAYNE PHARMA **3MG****A206623 001** Apr 08, 2016**AB** MYLAN **3MG****A090410 001** May 16, 2011**AB** RISING PHARMS **3MG****A207367 001** Apr 07, 2017**AB** SCIECURE PHARMA INC **3MG****A209041 001** Sep 28, 2017**AB** ZYDUS PHARMS USA **3MG****A206134 001** May 04, 2017

INC

ENTOCORT EC**AB** +! PERRIGO PHARMA INTL **3MG****N021324 001** Oct 02, 2001

POWDER, METERED;INHALATION

PULMICORT FLEXHALER

+ ASTRAZENECA 0.08MG/INH

N021949 001 Jul 12, 2006

+! 0.16MG/INH

N021949 002 Jul 12, 2006

SUSPENSION; INHALATION

BUDESONIDE**AN** APOTEX INC **0.25MG/2ML****A078202 001** Mar 30, 2009**AN** **0.5MG/2ML****A078202 002** Mar 30, 2009**AN** CIPLA **0.25MG/2ML****A205710 001** Nov 16, 2017**AN** **0.5MG/2ML****A205710 002** Nov 16, 2017**AN** **1MG/2ML****A205710 003** Nov 16, 2017**AN** IMPAX LABS INC **0.25MG/2ML****A078404 001** Jul 31, 2012**AN** **0.5MG/2ML****A078404 002** Jul 31, 2012**AN** LUPIN ATLANTIS **0.5MG/2ML****A210897 001** Nov 09, 2018**AN** SANDOZ INC **0.25MG/2ML****A201966 003** Sep 27, 2013**AN** **0.5MG/2ML****A201966 002** Sep 27, 2013**AN** **1MG/2ML****A201966 001** Sep 27, 2013**AN** TEVA PHARMS **0.25MG/2ML****A077519 001** Nov 18, 2008

PRESCRIPTION DRUG PRODUCT LIST

BUDESONIDE

SUSPENSION; INHALATION

BUDESONIDE

| | | | | | |
|-----------|-----------------|------------------|----------------|------------|--------------|
| <u>AN</u> | | <u>0.5MG/2ML</u> | <u>A077519</u> | <u>002</u> | Nov 18, 2008 |
| <u>AN</u> | TEVA PHARMS USA | <u>1MG/2ML</u> | <u>A204548</u> | <u>001</u> | Mar 08, 2016 |

PULMICORT RESPULES

| | | | | | | |
|-----------|---|--------------------|-------------------|----------------|------------|--------------|
| <u>AN</u> | + | ASTRAZENECA PHARMS | <u>0.25MG/2ML</u> | <u>N020929</u> | <u>001</u> | Aug 08, 2000 |
| <u>AN</u> | + | | <u>0.5MG/2ML</u> | <u>N020929</u> | <u>002</u> | Aug 08, 2000 |
| <u>AN</u> | + | ! | <u>1MG/2ML</u> | <u>N020929</u> | <u>003</u> | Aug 08, 2000 |

TABLET, EXTENDED RELEASE; ORAL

BUDESONIDE

| | | | | | |
|-----------|---------------------|------------|----------------|------------|--------------|
| <u>AB</u> | ACTAVIS LABS FL INC | <u>9MG</u> | <u>A205457</u> | <u>001</u> | Jul 03, 2018 |
|-----------|---------------------|------------|----------------|------------|--------------|

UCERIS

| | | | | | | | |
|-----------|---|---|---------------------|------------|----------------|------------|--------------|
| <u>AB</u> | + | ! | VALEANT PHARMS INTL | <u>9MG</u> | <u>N203634</u> | <u>001</u> | Jan 14, 2013 |
|-----------|---|---|---------------------|------------|----------------|------------|--------------|

BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE

AEROSOL, METERED; INHALATION

SYMBICORT

| | | | | | | |
|---|---|-------------|-------------------------|---------|-----|--------------|
| + | ! | ASTRAZENECA | 0.08MG/INH;0.0045MG/INH | N021929 | 001 | Jul 21, 2006 |
| + | ! | | 0.16MG/INH;0.0045MG/INH | N021929 | 002 | Jul 21, 2006 |

BUMETANIDE

INJECTABLE; INJECTION

BUMETANIDE

| | | | | | | |
|-----------|---|-------------------------|------------------|----------------|------------|--------------|
| <u>AP</u> | | HOSPIRA | <u>0.25MG/ML</u> | <u>A074332</u> | <u>001</u> | Oct 31, 1994 |
| <u>AP</u> | ! | WEST-WARD PHARMS INT | <u>0.25MG/ML</u> | <u>A079196</u> | <u>001</u> | Apr 30, 2008 |

TABLET; ORAL

BUMETANIDE

| | | | | | | |
|-----------|---|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | AMNEAL PHARMS CO | <u>0.5MG</u> | <u>A209724</u> | <u>001</u> | Oct 18, 2017 |
| <u>AB</u> | | | <u>1MG</u> | <u>A209724</u> | <u>002</u> | Oct 18, 2017 |
| <u>AB</u> | | | <u>2MG</u> | <u>A209724</u> | <u>003</u> | Oct 18, 2017 |
| <u>AB</u> | | IVAX SUB TEVA PHARMS | <u>0.5MG</u> | <u>A074225</u> | <u>001</u> | Apr 24, 1995 |
| <u>AB</u> | | | <u>1MG</u> | <u>A074225</u> | <u>002</u> | Apr 24, 1995 |
| <u>AB</u> | | | <u>2MG</u> | <u>A074225</u> | <u>003</u> | Apr 24, 1995 |
| <u>AB</u> | | SANDOZ | <u>0.5MG</u> | <u>A074700</u> | <u>001</u> | Nov 21, 1996 |
| <u>AB</u> | | | <u>1MG</u> | <u>A074700</u> | <u>002</u> | Nov 21, 1996 |
| <u>AB</u> | ! | | <u>2MG</u> | <u>A074700</u> | <u>003</u> | Nov 21, 1996 |
| <u>AB</u> | | UPSHER SMITH LABS | <u>0.5MG</u> | <u>A209916</u> | <u>001</u> | Jan 23, 2018 |
| <u>AB</u> | | | <u>1MG</u> | <u>A209916</u> | <u>002</u> | Jan 23, 2018 |
| <u>AB</u> | | | <u>2MG</u> | <u>A209916</u> | <u>003</u> | Jan 23, 2018 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>0.5MG</u> | <u>A202900</u> | <u>001</u> | Apr 30, 2018 |
| <u>AB</u> | | | <u>1MG</u> | <u>A202900</u> | <u>002</u> | Apr 30, 2018 |
| <u>AB</u> | | | <u>2MG</u> | <u>A202900</u> | <u>003</u> | Apr 30, 2018 |

BUMEX

| | | | | | | |
|-----------|---|----------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | VALIDUS PHARMS | <u>0.5MG</u> | <u>N018225</u> | <u>002</u> | Feb 28, 1983 |
| <u>AB</u> | + | | <u>1MG</u> | <u>N018225</u> | <u>001</u> | Feb 28, 1983 |
| <u>AB</u> | + | | <u>2MG</u> | <u>N018225</u> | <u>003</u> | Jun 14, 1985 |

BUPIVACAINE

INJECTABLE, LIPOSOMAL; INJECTION

EXPAREL

| | | | | | | |
|---|---|-------------------|------------------------|---------|-----|--------------|
| + | ! | PACIRA PHARMS INC | 133MG/10ML (13.3MG/ML) | N022496 | 001 | Oct 28, 2011 |
| + | ! | | 266MG/20ML (13.3MG/ML) | N022496 | 002 | Oct 28, 2011 |

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|-------------------------|--------------|----------------|------------|--------------|
| <u>AP</u> | | AUROBINDO PHARMA LTD | <u>0.25%</u> | <u>A207183</u> | <u>001</u> | May 13, 2016 |
| <u>AP</u> | | | <u>0.5%</u> | <u>A207183</u> | <u>002</u> | May 13, 2016 |
| <u>AP</u> | | HOSPIRA | <u>0.25%</u> | <u>A070583</u> | <u>001</u> | Feb 17, 1987 |
| <u>AP</u> | | | <u>0.25%</u> | <u>A070586</u> | <u>001</u> | Mar 03, 1987 |
| <u>AP</u> | | | <u>0.25%</u> | <u>A070590</u> | <u>001</u> | Feb 17, 1987 |
| <u>AP</u> | | | <u>0.25%</u> | <u>N018053</u> | <u>002</u> | |
| <u>AP</u> | | | <u>0.5%</u> | <u>A070584</u> | <u>001</u> | Feb 17, 1986 |
| <u>AP</u> | | | <u>0.5%</u> | <u>A070597</u> | <u>001</u> | Mar 03, 1987 |
| <u>AP</u> | | | <u>0.5%</u> | <u>A070609</u> | <u>001</u> | Mar 03, 1987 |
| <u>AP</u> | | | <u>0.5%</u> | <u>N018053</u> | <u>001</u> | |
| <u>AP</u> | | | <u>0.75%</u> | <u>A070585</u> | <u>001</u> | Mar 03, 1987 |
| <u>AP</u> | | | <u>0.75%</u> | <u>N018053</u> | <u>003</u> | |
| <u>AP</u> | | MYLAN ASI | <u>0.25%</u> | <u>A091503</u> | <u>001</u> | Oct 18, 2011 |
| <u>AP</u> | | | <u>0.5%</u> | <u>A091503</u> | <u>002</u> | Oct 18, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>0.25%</u> | <u>A203895 001</u> | Nov 05, 2013 |
| <u>AP</u> | | <u>0.5%</u> | <u>A203895 002</u> | Nov 05, 2013 |
| <u>AP</u> | | <u>0.75%</u> | <u>A203895 003</u> | Nov 05, 2013 |
| <u>AP</u> | MYLAN ASI | <u>0.25%</u> | <u>A091487 002</u> | Oct 18, 2011 |
| <u>AP</u> | | <u>0.5%</u> | <u>A091487 001</u> | Oct 18, 2011 |
| <u>AP</u> | | <u>0.75%</u> | <u>A091487 003</u> | Oct 18, 2011 |

MARCAINE HYDROCHLORIDE

| | | | | |
|-----------|------------|--------------|--------------------|--|
| <u>AP</u> | +! HOSPIRA | <u>0.25%</u> | <u>N016964 001</u> | |
| <u>AP</u> | +! | <u>0.5%</u> | <u>N016964 006</u> | |

MARCAINE HYDROCHLORIDE PRESERVATIVE FREE

| | | | | |
|-----------|------------|--------------|--------------------|--|
| <u>AP</u> | +! HOSPIRA | <u>0.25%</u> | <u>N016964 012</u> | |
| <u>AP</u> | +! | <u>0.5%</u> | <u>N016964 005</u> | |
| <u>AP</u> | +! | <u>0.75%</u> | <u>N016964 009</u> | |

SENSORCAINE

| | | | | |
|-----------|--------------------|--------------|--------------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>0.25%</u> | <u>A070552 001</u> | May 21, 1986 |
| <u>AP</u> | | <u>0.25%</u> | <u>N018304 001</u> | |
| <u>AP</u> | | <u>0.5%</u> | <u>A070553 001</u> | May 21, 1986 |
| <u>AP</u> | | <u>0.5%</u> | <u>N018304 002</u> | |
| <u>AP</u> | | <u>0.75%</u> | <u>A070554 001</u> | May 21, 1986 |
| <u>AP</u> | | <u>0.75%</u> | <u>N018304 003</u> | |

INJECTABLE; SPINAL

BUPIVACAINE HYDROCHLORIDE

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| <u>AP</u> | BAXTER HLTHCARE CORP | <u>0.75%</u> | <u>A207266 001</u> | Jul 25, 2016 |
| <u>AP</u> | HOSPIRA | <u>0.75%</u> | <u>A071810 001</u> | Dec 11, 1987 |

MARCAINE

| | | | | |
|-----------|------------|--------------|--------------------|--------------|
| <u>AP</u> | +! HOSPIRA | <u>0.75%</u> | <u>N018692 001</u> | May 04, 1984 |
|-----------|------------|--------------|--------------------|--------------|

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

| | | | | |
|-----------|-----------|-------------------------|--------------------|--------------|
| <u>AP</u> | ! HOSPIRA | <u>0.5%;0.005MG/ML</u> | <u>A071168 001</u> | Jun 16, 1988 |
| <u>AP</u> | | <u>0.5%;0.005MG/ML</u> | <u>A071170 001</u> | Jun 16, 1988 |
| | ! | <u>0.25%;0.005MG/ML</u> | <u>A071165 001</u> | Jun 16, 1988 |
| | | <u>0.25%;0.005MG/ML</u> | <u>A071167 001</u> | Jun 16, 1988 |

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

| | | | | |
|-----------|-----------|-------------------------|--------------------|--------------|
| <u>AP</u> | SEPTODONT | <u>0.5%;0.0091MG/ML</u> | <u>A077250 001</u> | Sep 27, 2006 |
|-----------|-----------|-------------------------|--------------------|--------------|

BUPIVACAINE HYDROCHLORIDE W/EPINEPHRINE

| | | | | |
|-----------|------------|-------------------------|--------------------|--------------|
| <u>AP</u> | +! HOSPIRA | <u>0.5%;0.0091MG/ML</u> | <u>N022046 001</u> | Jul 13, 1983 |
|-----------|------------|-------------------------|--------------------|--------------|

MARCAINE HYDROCHLORIDE W/ EPINEPHRINE

| | | | | |
|-----------|------------|--------------------------|--------------------|--|
| <u>AP</u> | +! HOSPIRA | <u>0.25%;0.0091MG/ML</u> | <u>N016964 004</u> | |
| <u>AP</u> | +! | <u>0.5%;0.0091MG/ML</u> | <u>N016964 008</u> | |

MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE

| | | | | |
|-----------|------------|--------------------------|--------------------|--|
| <u>AP</u> | +! HOSPIRA | <u>0.25%;0.0091MG/ML</u> | <u>N016964 013</u> | |
| <u>AP</u> | +! | <u>0.5%;0.0091MG/ML</u> | <u>N016964 007</u> | |
| <u>AP</u> | +! | <u>0.75%;0.0091MG/ML</u> | <u>N016964 010</u> | |

SENSORCAINE

| | | | | |
|-----------|--------------------|--------------------------|--------------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>0.25%;0.0091MG/ML</u> | <u>A070966 001</u> | Oct 13, 1987 |
| <u>AP</u> | | <u>0.25%;0.0091MG/ML</u> | <u>A070967 001</u> | Oct 13, 1987 |
| <u>AP</u> | | <u>0.5%;0.0091MG/ML</u> | <u>A070968 001</u> | Oct 13, 1987 |
| <u>AP</u> | | <u>0.5%;0.0091MG/ML</u> | <u>N018304 004</u> | Sep 02, 1983 |
| <u>AP</u> | | <u>0.75%;0.0091MG/ML</u> | <u>N018304 005</u> | Sep 02, 1983 |

BUPRENORPHINE

FILM, EXTENDED RELEASE; TRANSDERMAL

BUPRENORPHINE

| | | | | |
|-----------|------------------|-----------------|--------------------|--------------|
| <u>AB</u> | WATSON LABS TEVA | <u>5MCG/HR</u> | <u>A204937 001</u> | Nov 20, 2018 |
| <u>AB</u> | | <u>10MCG/HR</u> | <u>A204937 002</u> | Nov 20, 2018 |
| <u>AB</u> | | <u>15MCG/HR</u> | <u>A204937 003</u> | Nov 20, 2018 |
| <u>AB</u> | | <u>20MCG/HR</u> | <u>A204937 004</u> | Nov 20, 2018 |

BUTRANS

| | | | | |
|-----------|--------------------|------------------|--------------------|--------------|
| <u>AB</u> | + PURDUE PHARMA LP | <u>5MCG/HR</u> | <u>N021306 001</u> | Jun 30, 2010 |
| <u>AB</u> | + | <u>10MCG/HR</u> | <u>N021306 002</u> | Jun 30, 2010 |
| <u>AB</u> | + | <u>15MCG/HR</u> | <u>N021306 004</u> | Jul 25, 2013 |
| <u>AB</u> | +! | <u>20MCG/HR</u> | <u>N021306 003</u> | Jun 30, 2010 |
| | + | <u>7.5MCG/HR</u> | <u>N021306 005</u> | Jun 30, 2014 |

PRESCRIPTION DRUG PRODUCT LIST

BUPRENORPHINE

SOLUTION, EXTENDED RELEASE; SUBCUTANEOUS

SUBLOCADE

| | | | | | |
|---|--------------|---------------------------|---------|-----|--------------|
| + | INDIVIOR INC | 100MG/0.5ML (100MG/0.5ML) | N209819 | 001 | Nov 30, 2017 |
| + | ! | 300MG/1.5ML (200MG/ML) | N209819 | 002 | Nov 30, 2017 |

BUPRENORPHINE HYDROCHLORIDE

FILM; BUCCAL

BELBUCA

| | | | | | |
|---|------|-----------------|---------|-----|--------------|
| + | BDSI | EQ 0.075MG BASE | N207932 | 001 | Oct 23, 2015 |
| + | | EQ 0.15MG BASE | N207932 | 002 | Oct 23, 2015 |
| + | | EQ 0.3MG BASE | N207932 | 003 | Oct 23, 2015 |
| + | | EQ 0.45MG BASE | N207932 | 004 | Oct 23, 2015 |
| + | | EQ 0.6MG BASE | N207932 | 005 | Oct 23, 2015 |
| + | | EQ 0.75MG BASE | N207932 | 006 | Oct 23, 2015 |
| + | ! | EQ 0.9MG BASE | N207932 | 007 | Oct 23, 2015 |

IMPLANT; IMPLANTATION

PROBUPHINE

| | | | | | |
|---|--------------|----------------------|---------|-----|--------------|
| + | TITAN PHARMS | EQ 80MG BASE/IMPLANT | N204442 | 001 | May 26, 2016 |
|---|--------------|----------------------|---------|-----|--------------|

INJECTABLE; INJECTION

BUPRENEX

| | | | | | |
|-----------|---|--------------|--------------------------------|-----------------------|-------------------|
| AP | + | INDIVIOR INC | <u>EQ 0.3MG BASE/ML</u> | <u>N018401</u> | <u>001</u> |
|-----------|---|--------------|--------------------------------|-----------------------|-------------------|

BUPRENORPHINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|----------------------|--------------------------------|-----------------------|-------------------|--------------|
| AP | | HOSPIRA | <u>EQ 0.3MG BASE/ML</u> | <u>A074137</u> | <u>001</u> | Jun 03, 1996 |
| AP | | LUITPOLD | <u>EQ 0.3MG BASE/ML</u> | <u>A078331</u> | <u>001</u> | Mar 27, 2007 |
| AP | | PAR STERILE PRODUCTS | <u>EQ 0.3MG BASE/ML</u> | <u>A206586</u> | <u>001</u> | Jul 28, 2015 |
| AP | | WEST-WARD PHARMS INT | <u>EQ 0.3MG BASE/ML</u> | <u>A076931</u> | <u>001</u> | Mar 02, 2005 |

TABLET; SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE

| | | | | | | |
|-----------|---|----------------------|---------------------------|-----------------------|-------------------|--------------|
| AB | | ACTAVIS ELIZABETH | <u>EQ 2MG BASE</u> | <u>A090819</u> | <u>001</u> | Feb 19, 2015 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A090819</u> | <u>002</u> | Feb 19, 2015 |
| AB | | BARR | <u>EQ 2MG BASE</u> | <u>A090360</u> | <u>001</u> | May 07, 2010 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A090360</u> | <u>002</u> | May 07, 2010 |
| AB | | CASI PHARMS INC | <u>EQ 2MG BASE</u> | <u>A090279</u> | <u>001</u> | Jun 10, 2015 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A090279</u> | <u>002</u> | Jun 10, 2015 |
| AB | | ETHYPHARM | <u>EQ 2MG BASE</u> | <u>A090622</u> | <u>001</u> | Sep 24, 2010 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A090622</u> | <u>002</u> | Sep 24, 2010 |
| AB | | MYLAN PHARMS INC | <u>EQ 2MG BASE</u> | <u>A201066</u> | <u>001</u> | Mar 06, 2015 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A201066</u> | <u>002</u> | Mar 06, 2015 |
| AB | | RHODES PHARMS | <u>EQ 2MG BASE</u> | <u>A207276</u> | <u>001</u> | Mar 27, 2017 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A207276</u> | <u>002</u> | Mar 27, 2017 |
| AB | | SUN PHARM INDS LTD | <u>EQ 2MG BASE</u> | <u>A201760</u> | <u>001</u> | Jan 29, 2016 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A201760</u> | <u>002</u> | Jan 29, 2016 |
| AB | | WEST-WARD PHARMS INT | <u>EQ 2MG BASE</u> | <u>A078633</u> | <u>001</u> | Oct 08, 2009 |
| AB | ! | | <u>EQ 8MG BASE</u> | <u>A078633</u> | <u>002</u> | Oct 08, 2009 |

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE

FILM; BUCCAL

BUNAVAIL

| | | | | | |
|---|------|-----------------------------|---------|-----|--------------|
| + | BDSI | EQ 2.1MG BASE;EQ 0.3MG BASE | N205637 | 001 | Jun 06, 2014 |
| + | | EQ 4.2MG BASE;EQ 0.7MG BASE | N205637 | 002 | Jun 06, 2014 |
| + | ! | EQ 6.3MG BASE;EQ 1MG BASE | N205637 | 003 | Jun 06, 2014 |

FILM; BUCCAL, SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

| | | | | | | |
|-----------|--|--------------------|-----------------------------------------|-----------------------|-------------------|--------------|
| AB | | DR REDDYS LABS SA | <u>EQ 2MG BASE;EQ 0.5MG BASE</u> | <u>A205299</u> | <u>001</u> | Jun 14, 2018 |
| AB | | | <u>EQ 4MG BASE;EQ 1MG BASE</u> | <u>A205806</u> | <u>001</u> | Jun 14, 2018 |
| AB | | | <u>EQ 8MG BASE;EQ 2MG BASE</u> | <u>A205299</u> | <u>002</u> | Jun 14, 2018 |
| AB | | | <u>EQ 12MG BASE;EQ 3MG BASE</u> | <u>A205806</u> | <u>002</u> | Jun 14, 2018 |
| AB | | MYLAN TECHNOLOGIES | <u>EQ 8MG BASE;EQ 2MG BASE</u> | <u>A207607</u> | <u>001</u> | Jun 14, 2018 |
| AB | | | <u>EQ 12MG BASE;EQ 3MG BASE</u> | <u>A207607</u> | <u>002</u> | Jun 14, 2018 |

SUBOXONE

| | | | | | | |
|-----------|---|--------------|-----------------------------------------|-----------------------|-------------------|--------------|
| AB | + | INDIVIOR INC | <u>EQ 2MG BASE;EQ 0.5MG BASE</u> | <u>N022410</u> | <u>001</u> | Aug 30, 2010 |
| AB | + | | <u>EQ 4MG BASE;EQ 1MG BASE</u> | <u>N022410</u> | <u>003</u> | Aug 10, 2012 |
| AB | + | | <u>EQ 8MG BASE;EQ 2MG BASE</u> | <u>N022410</u> | <u>002</u> | Aug 30, 2010 |
| AB | + | ! | <u>EQ 12MG BASE;EQ 3MG BASE</u> | <u>N022410</u> | <u>004</u> | Aug 10, 2012 |

FILM; SUBLINGUAL

CASSIPA

| | | | | | |
|---|-----------------|--------------------------|---------|-----|--------------|
| + | TEVA PHARMS USA | EQ 16MG BASE;EQ 4MG BASE | N208042 | 001 | Sep 07, 2018 |
|---|-----------------|--------------------------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE

TABLET;SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|----------------------------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS ELIZABETH | <u>EQ 2MG BASE;EQ 0.5MG BASE</u> | <u>A091422 001</u> | Feb 22, 2013 |
| <u>AB</u> | ! | <u>EQ 8MG BASE;EQ 2MG BASE</u> | <u>A091422 002</u> | Feb 22, 2013 |
| <u>AB</u> | AMNEAL PHARMS | <u>EQ 2MG BASE;EQ 0.5MG BASE</u> | <u>A203136 001</u> | Feb 22, 2013 |
| <u>AB</u> | | <u>EQ 8MG BASE;EQ 2MG BASE</u> | <u>A203136 002</u> | Feb 22, 2013 |
| <u>AB</u> | ETHYPHARM USA CORP | <u>EQ 2MG BASE;EQ 0.5MG BASE</u> | <u>A204431 001</u> | Oct 16, 2015 |
| <u>AB</u> | | <u>EQ 8MG BASE;EQ 2MG BASE</u> | <u>A204431 002</u> | Oct 16, 2015 |
| <u>AB</u> | LANNETT CO INC | <u>EQ 2MG BASE;EQ 0.5MG BASE</u> | <u>A205022 001</u> | Sep 19, 2016 |
| <u>AB</u> | | <u>EQ 8MG BASE;EQ 2MG BASE</u> | <u>A205022 002</u> | Sep 19, 2016 |
| <u>AB</u> | SPECGX LLC | <u>EQ 2MG BASE;EQ 0.5MG BASE</u> | <u>A207000 001</u> | Dec 13, 2017 |
| <u>AB</u> | | <u>EQ 8MG BASE;EQ 2MG BASE</u> | <u>A207000 002</u> | Dec 13, 2017 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>EQ 2MG BASE;EQ 0.5MG BASE</u> | <u>A201633 001</u> | Aug 05, 2016 |
| <u>AB</u> | | <u>EQ 8MG BASE;EQ 2MG BASE</u> | <u>A201633 002</u> | Aug 05, 2016 |
| <u>AB</u> | TEVA PHARMS USA | <u>EQ 2MG BASE;EQ 0.5MG BASE</u> | <u>A091149 001</u> | Sep 08, 2014 |
| <u>AB</u> | | <u>EQ 8MG BASE;EQ 2MG BASE</u> | <u>A091149 002</u> | Sep 08, 2014 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>EQ 2MG BASE;EQ 0.5MG BASE</u> | <u>A203326 001</u> | Jun 27, 2014 |
| <u>AB</u> | | <u>EQ 8MG BASE;EQ 2MG BASE</u> | <u>A203326 002</u> | Jun 27, 2014 |
| | ZUBSOLV | | | |
| + | OREXO US INC | EQ 0.7MG BASE;EQ 0.18MG BASE | N204242 006 | Oct 04, 2016 |
| + | | EQ 1.4MG BASE;EQ 0.36MG BASE | N204242 001 | Jul 03, 2013 |
| + | | EQ 2.9MG BASE;EQ 0.71MG BASE | N204242 005 | Jun 04, 2015 |
| + | | EQ 5.7MG BASE;EQ 1.4MG BASE | N204242 002 | Jul 03, 2013 |
| + | | EQ 8.6MG BASE;EQ 2.1MG BASE | N204242 003 | Dec 11, 2014 |
| + | ! | EQ 11.4MG BASE;EQ 2.9MG BASE | N204242 004 | Dec 11, 2014 |

BUPROPION HYDROBROMIDE

TABLET, EXTENDED RELEASE;ORAL

APLENZIN

| | | | | |
|---|-------------------------|-------|-------------|--------------|
| + | VALEANT PHARMS NORTH | 174MG | N022108 001 | Apr 23, 2008 |
| + | | 348MG | N022108 002 | Apr 23, 2008 |
| + | ! | 522MG | N022108 003 | Apr 23, 2008 |

BUPROPION HYDROCHLORIDE

TABLET;ORAL

BUPROPION HYDROCHLORIDE

| | | | | |
|-----------|--------------------|--------------|--------------------|--------------|
| <u>AB</u> | ALEMBIC PHARMS LTD | <u>75MG</u> | <u>A203013 001</u> | Jun 08, 2018 |
| <u>AB</u> | | <u>100MG</u> | <u>A203013 002</u> | Jun 08, 2018 |
| <u>AB</u> | APOTEX INC | <u>75MG</u> | <u>A076143 001</u> | Jan 17, 2006 |
| <u>AB</u> | ! | <u>100MG</u> | <u>A076143 002</u> | Jan 17, 2006 |
| <u>AB</u> | HERITAGE PHARMA | <u>75MG</u> | <u>A206975 001</u> | Aug 19, 2016 |
| <u>AB</u> | | <u>100MG</u> | <u>A206975 002</u> | Aug 19, 2016 |
| <u>AB</u> | INVAGEN PHARMS | <u>75MG</u> | <u>A207389 001</u> | Sep 18, 2017 |
| <u>AB</u> | | <u>100MG</u> | <u>A207389 002</u> | Sep 18, 2017 |
| <u>AB</u> | MYLAN | <u>75MG</u> | <u>A075491 001</u> | Apr 17, 2000 |
| <u>AB</u> | | <u>100MG</u> | <u>A075491 002</u> | Apr 17, 2000 |
| <u>AB</u> | SANDOZ | <u>75MG</u> | <u>A075584 001</u> | Feb 07, 2000 |
| <u>AB</u> | | <u>100MG</u> | <u>A075584 002</u> | Feb 07, 2000 |

TABLET, EXTENDED RELEASE;ORAL

BUPROPION HYDROCHLORIDE

| | | | | |
|------------|---------------------|--------------|--------------------|--------------|
| <u>AB1</u> | ACTAVIS LABS FL INC | <u>100MG</u> | <u>A079095 001</u> | Mar 24, 2009 |
| <u>AB1</u> | | <u>150MG</u> | <u>A079095 002</u> | Mar 24, 2009 |
| <u>AB1</u> | | <u>200MG</u> | <u>A079095 003</u> | Mar 24, 2009 |
| <u>AB1</u> | ANCHEN PHARMS | <u>100MG</u> | <u>A091459 001</u> | Jun 09, 2011 |
| <u>AB1</u> | | <u>150MG</u> | <u>A091459 002</u> | Jun 09, 2011 |
| <u>AB1</u> | | <u>200MG</u> | <u>A091459 003</u> | Jun 09, 2011 |
| <u>AB1</u> | IMPAX LABS | <u>100MG</u> | <u>A075913 001</u> | Jan 28, 2004 |
| <u>AB1</u> | | <u>150MG</u> | <u>A075913 002</u> | Mar 22, 2004 |
| <u>AB1</u> | | <u>200MG</u> | <u>A076711 001</u> | Dec 03, 2004 |
| <u>AB1</u> | INVAGEN PHARMS | <u>100MG</u> | <u>A206674 001</u> | Feb 09, 2016 |
| <u>AB1</u> | | <u>150MG</u> | <u>A206674 002</u> | Feb 09, 2016 |
| <u>AB1</u> | | <u>200MG</u> | <u>A206674 003</u> | Feb 09, 2016 |
| <u>AB1</u> | JUBILANT GENERICS | <u>100MG</u> | <u>A202774 001</u> | Oct 11, 2013 |
| <u>AB1</u> | | <u>150MG</u> | <u>A202774 002</u> | Oct 11, 2013 |
| <u>AB1</u> | | <u>200MG</u> | <u>A202774 003</u> | Oct 11, 2013 |
| <u>AB1</u> | MYLAN | <u>100MG</u> | <u>A090325 001</u> | Apr 08, 2010 |
| <u>AB1</u> | | <u>150MG</u> | <u>A090325 002</u> | Apr 08, 2010 |
| <u>AB1</u> | | <u>200MG</u> | <u>A090325 003</u> | Apr 08, 2010 |
| <u>AB1</u> | PRINSTON INC | <u>100MG</u> | <u>A202304 001</u> | May 26, 2015 |
| <u>AB1</u> | | <u>150MG</u> | <u>A202304 002</u> | May 26, 2015 |
| <u>AB1</u> | | <u>200MG</u> | <u>A202304 003</u> | May 26, 2015 |

PRESCRIPTION DRUG PRODUCT LIST

BUPROPION HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

BUPROPION HYDROCHLORIDE

| | | | | |
|------------|--------------------|--------------|--------------------|--------------|
| <u>AB1</u> | SANDOZ | <u>100MG</u> | <u>A075932 001</u> | Nov 25, 2003 |
| <u>AB1</u> | | <u>150MG</u> | <u>A075932 002</u> | Mar 22, 2004 |
| <u>AB1</u> | | <u>200MG</u> | <u>A075932 003</u> | Jun 22, 2005 |
| <u>AB1</u> | SCIEGEN PHARMS INC | <u>100MG</u> | <u>A205794 001</u> | Mar 01, 2016 |
| <u>AB1</u> | | <u>150MG</u> | <u>A205794 002</u> | Mar 01, 2016 |
| <u>AB1</u> | | <u>200MG</u> | <u>A205794 003</u> | Mar 01, 2016 |
| <u>AB1</u> | SUN PHARMA GLOBAL | <u>100MG</u> | <u>A078866 001</u> | Apr 06, 2010 |
| <u>AB1</u> | | <u>150MG</u> | <u>A078866 002</u> | Apr 06, 2010 |
| <u>AB1</u> | | <u>200MG</u> | <u>A078866 003</u> | Apr 06, 2010 |
| <u>AB1</u> | TORRENT PHARMS LTD | <u>100MG</u> | <u>A203969 001</u> | Oct 31, 2014 |
| <u>AB1</u> | | <u>150MG</u> | <u>A203969 002</u> | Oct 31, 2014 |
| <u>AB1</u> | | <u>200MG</u> | <u>A203969 003</u> | Oct 31, 2014 |
| <u>AB1</u> | WATSON LABS INC | <u>100MG</u> | <u>A077455 001</u> | Jul 19, 2010 |
| <u>AB1</u> | | <u>150MG</u> | <u>A077455 002</u> | Mar 12, 2008 |
| <u>AB1</u> | | <u>200MG</u> | <u>A077455 003</u> | Jul 19, 2010 |
| <u>AB1</u> | YICHANG HUMANWELL | <u>100MG</u> | <u>A211347 001</u> | Oct 16, 2018 |
| <u>AB1</u> | | <u>150MG</u> | <u>A211347 002</u> | Oct 16, 2018 |
| <u>AB1</u> | | <u>200MG</u> | <u>A211347 003</u> | Oct 16, 2018 |

WELLBUTRIN SR

| | | | | |
|------------|-------------------|--------------|--------------------|--------------|
| <u>AB1</u> | + GLAXOSMITHKLINE | <u>100MG</u> | <u>N020358 002</u> | Oct 04, 1996 |
| <u>AB1</u> | + | <u>150MG</u> | <u>N020358 003</u> | Oct 04, 1996 |
| <u>AB1</u> | + | <u>200MG</u> | <u>N020358 004</u> | Jun 14, 2002 |

BUPROPION HYDROCHLORIDE

| | | | | |
|------------|---------------------|--------------|--------------------|--------------|
| <u>AB2</u> | ACTAVIS LABS FL INC | <u>150MG</u> | <u>A079094 001</u> | Mar 24, 2009 |
| <u>AB2</u> | ANCHEN PHARMS | <u>150MG</u> | <u>A091520 001</u> | Jun 09, 2011 |
| <u>AB2</u> | IMPAX LABS | <u>150MG</u> | <u>A075914 001</u> | May 27, 2004 |
| <u>AB2</u> | JUBILANT GENERICS | <u>150MG</u> | <u>A202775 001</u> | Oct 11, 2013 |
| <u>AB2</u> | MYLAN | <u>150MG</u> | <u>A090941 001</u> | May 03, 2010 |
| <u>AB2</u> | SANDOZ INC | <u>150MG</u> | <u>A077475 001</u> | Mar 12, 2008 |
| <u>AB2</u> | SCIEGEN PHARMS INC | <u>150MG</u> | <u>A206122 001</u> | Aug 17, 2016 |

ZYBAN

| | | | | |
|------------|--------------------|--------------|--------------------|--------------|
| <u>AB2</u> | +! GLAXOSMITHKLINE | <u>150MG</u> | <u>N020711 003</u> | May 14, 1997 |
|------------|--------------------|--------------|--------------------|--------------|

BUPROPION HYDROCHLORIDE

| | | | | |
|------------|----------------------|--------------|--------------------|--------------|
| <u>AB3</u> | ACCORD HLTHCARE | <u>150MG</u> | <u>A210497 001</u> | Oct 31, 2018 |
| <u>AB3</u> | | <u>300MG</u> | <u>A210497 002</u> | Oct 31, 2018 |
| <u>AB3</u> | ACTAVIS LABS FL INC | <u>150MG</u> | <u>A077715 001</u> | Nov 26, 2008 |
| <u>AB3</u> | ANBISON LAB | <u>150MG</u> | <u>A207224 001</u> | Jun 30, 2017 |
| <u>AB3</u> | | <u>300MG</u> | <u>A207224 002</u> | Jun 30, 2017 |
| <u>AB3</u> | ANCHEN PHARMS | <u>150MG</u> | <u>A077284 001</u> | Dec 14, 2006 |
| <u>AB3</u> | | <u>300MG</u> | <u>A077284 002</u> | Dec 14, 2006 |
| <u>AB3</u> | IMPAX LABS | <u>150MG</u> | <u>A077415 001</u> | Nov 26, 2008 |
| <u>AB3</u> | INVAGEN PHARMS | <u>150MG</u> | <u>A206556 001</u> | Aug 26, 2016 |
| <u>AB3</u> | | <u>300MG</u> | <u>A206556 002</u> | Aug 26, 2016 |
| <u>AB3</u> | JUBILANT GENERICS | <u>150MG</u> | <u>A207459 001</u> | Jun 30, 2017 |
| <u>AB3</u> | | <u>300MG</u> | <u>A207459 002</u> | Jun 30, 2017 |
| <u>AB3</u> | LUPIN LTD | <u>150MG</u> | <u>A090693 001</u> | Apr 06, 2017 |
| <u>AB3</u> | | <u>300MG</u> | <u>A090693 002</u> | Apr 06, 2017 |
| <u>AB3</u> | MYLAN | <u>150MG</u> | <u>A090942 001</u> | Jul 14, 2010 |
| <u>AB3</u> | | <u>300MG</u> | <u>A090942 002</u> | Jul 14, 2010 |
| <u>AB3</u> | SCIEGEN PHARMS INC | <u>150MG</u> | <u>A207479 001</u> | Apr 12, 2017 |
| <u>AB3</u> | | <u>300MG</u> | <u>A207479 002</u> | Apr 12, 2017 |
| <u>AB3</u> | SINOTHERAPEUTICS INC | <u>150MG</u> | <u>A208652 001</u> | Aug 21, 2017 |
| <u>AB3</u> | | <u>300MG</u> | <u>A208652 002</u> | Aug 21, 2017 |
| <u>AB3</u> | SUN PHARMA GLOBAL | <u>150MG</u> | <u>A200695 001</u> | Dec 18, 2014 |
| <u>AB3</u> | TWI PHARMS | <u>150MG</u> | <u>A210081 001</u> | Nov 03, 2017 |
| <u>AB3</u> | | <u>300MG</u> | <u>A210081 002</u> | Nov 03, 2017 |
| <u>AB3</u> | WATSON LABS INC | <u>150MG</u> | <u>A077285 001</u> | Nov 26, 2008 |
| <u>AB3</u> | | <u>300MG</u> | <u>A077285 002</u> | Aug 15, 2008 |
| <u>AB3</u> | WOCKHARDT LTD | <u>150MG</u> | <u>A202189 001</u> | Nov 21, 2012 |
| <u>AB3</u> | YICHANG HUMANWELL | <u>150MG</u> | <u>A210015 001</u> | Jun 14, 2018 |
| <u>AB3</u> | | <u>300MG</u> | <u>A210015 002</u> | Jun 14, 2018 |
| <u>AB3</u> | ZYDUS PHARMS USA INC | <u>150MG</u> | <u>A201567 002</u> | Jul 23, 2018 |
| <u>AB3</u> | | <u>300MG</u> | <u>A201567 001</u> | Jan 17, 2014 |

WELLBUTRIN XL

| | | | | |
|------------|----------------|--------------|--------------------|--------------|
| <u>AB3</u> | + VALEANT INTL | <u>150MG</u> | <u>N021515 001</u> | Aug 28, 2003 |
| <u>AB3</u> | + | <u>300MG</u> | <u>N021515 002</u> | Aug 28, 2003 |

PRESCRIPTION DRUG PRODUCT LIST

BUPROPION HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

FORFIVO XL

+! ALVOGEN

450MG

N022497 001 Nov 10, 2011

BUPROPION HYDROCHLORIDE; NALTREXONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CONTRAVE

+! NALPROPION

90MG;8MG

N200063 001 Sep 10, 2014

BUSPIRONE HYDROCHLORIDE

TABLET;ORAL

BUSPIRONE HYDROCHLORIDE

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>5MG</u> | <u>A202557 001</u> | Dec 30, 2014 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A202557 002</u> | Dec 30, 2014 |
| <u>AB</u> | | <u>10MG</u> | <u>A202557 003</u> | Dec 30, 2014 |
| <u>AB</u> | | <u>15MG</u> | <u>A202557 004</u> | Dec 30, 2014 |
| <u>AB</u> | | <u>30MG</u> | <u>A202557 005</u> | Dec 30, 2014 |
| <u>AB</u> | AMNEAL PHARMS CO | <u>5MG</u> | <u>A208829 001</u> | May 24, 2017 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A208829 002</u> | May 24, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A208829 003</u> | May 24, 2017 |
| <u>AB</u> | | <u>15MG</u> | <u>A208829 004</u> | May 24, 2017 |
| <u>AB</u> | | <u>30MG</u> | <u>A208829 005</u> | May 24, 2017 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>5MG</u> | <u>A078246 001</u> | Feb 27, 2009 |
| <u>AB</u> | | <u>10MG</u> | <u>A078246 002</u> | Feb 27, 2009 |
| <u>AB</u> | | <u>15MG</u> | <u>A078246 003</u> | Feb 27, 2009 |
| <u>AB</u> | | <u>30MG</u> | <u>A078246 004</u> | Feb 27, 2009 |
| <u>AB</u> | HERITAGE PHARMA | <u>5MG</u> | <u>A204582 001</u> | Sep 18, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A204582 002</u> | Sep 18, 2015 |
| <u>AB</u> | | <u>15MG</u> | <u>A204582 003</u> | Sep 18, 2015 |
| <u>AB</u> | | <u>30MG</u> | <u>A204582 004</u> | Sep 18, 2015 |
| <u>AB</u> | IMPAX LABS INC | <u>5MG</u> | <u>A074253 001</u> | Mar 28, 2001 |
| <u>AB</u> | | <u>10MG</u> | <u>A074253 002</u> | Mar 28, 2001 |
| <u>AB</u> | | <u>15MG</u> | <u>A074253 003</u> | Mar 13, 2002 |
| <u>AB</u> | INVENTIA HLTHCARE | <u>5MG</u> | <u>A209696 001</u> | May 03, 2018 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A209696 002</u> | May 03, 2018 |
| <u>AB</u> | | <u>10MG</u> | <u>A209696 003</u> | May 03, 2018 |
| <u>AB</u> | | <u>15MG</u> | <u>A209696 004</u> | May 03, 2018 |
| <u>AB</u> | | <u>30MG</u> | <u>A209696 005</u> | May 03, 2018 |
| <u>AB</u> | MYLAN | <u>5MG</u> | <u>A076008 003</u> | Mar 01, 2002 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A075467 002</u> | Mar 28, 2001 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A076008 002</u> | Jul 08, 2013 |
| <u>AB</u> | | <u>10MG</u> | <u>A076008 004</u> | Mar 01, 2002 |
| <u>AB</u> | | <u>15MG</u> | <u>A076008 005</u> | Mar 28, 2001 |
| <u>AB</u> | | <u>30MG</u> | <u>A076008 001</u> | Jun 28, 2001 |
| <u>AB</u> | OXFORD PHARMS | <u>30MG</u> | <u>A078302 001</u> | Dec 17, 2007 |
| <u>AB</u> | STRIDES PHARMA | <u>5MG</u> | <u>A202330 001</u> | Aug 25, 2014 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A202330 005</u> | Feb 17, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A202330 002</u> | Aug 25, 2014 |
| <u>AB</u> | | <u>15MG</u> | <u>A202330 003</u> | Aug 25, 2014 |
| <u>AB</u> | | <u>30MG</u> | <u>A202330 004</u> | Aug 25, 2014 |
| <u>AB</u> | TEVA | <u>5MG</u> | <u>A075022 001</u> | Feb 28, 2002 |
| <u>AB</u> | | <u>10MG</u> | <u>A075022 002</u> | Feb 28, 2002 |
| <u>AB</u> | ! | <u>15MG</u> | <u>A075022 003</u> | Feb 28, 2002 |
| <u>AB</u> | | <u>30MG</u> | <u>A075022 004</u> | Mar 25, 2004 |
| <u>AB</u> | YILING PHARM LTD | <u>5MG</u> | <u>A202087 001</u> | Dec 16, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A202087 002</u> | Dec 16, 2015 |
| <u>AB</u> | | <u>15MG</u> | <u>A202087 003</u> | Dec 16, 2015 |
| <u>AB</u> | | <u>30MG</u> | <u>A202087 004</u> | Dec 16, 2015 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>5MG</u> | <u>A078888 001</u> | Feb 07, 2014 |
| <u>AB</u> | | <u>10MG</u> | <u>A078888 002</u> | Feb 07, 2014 |
| <u>AB</u> | | <u>15MG</u> | <u>A078888 003</u> | Feb 07, 2014 |
| <u>AB</u> | | <u>30MG</u> | <u>A078888 004</u> | Feb 07, 2014 |

BUSULFAN

INJECTABLE; INJECTION

BUSULFAN

| | | | | |
|-----------|------------------|---------------|--------------------|--------------|
| <u>AP</u> | ACTAVIS LLC | <u>6MG/ML</u> | <u>A205139 001</u> | Dec 08, 2017 |
| <u>AP</u> | AMNEAL PHARMS CO | <u>6MG/ML</u> | <u>A209580 001</u> | Dec 18, 2017 |
| <u>AP</u> | HOSPIRA INC | <u>6MG/ML</u> | <u>A205672 001</u> | Jul 31, 2018 |
| <u>AP</u> | LUITPOLD | <u>6MG/ML</u> | <u>A202259 001</u> | Dec 22, 2015 |
| <u>AP</u> | MYLAN LABS LTD | <u>6MG/ML</u> | <u>A205184 001</u> | Jul 13, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

BUSULFAN

INJECTABLE; INJECTION

BUSULFAN

| | | | | | |
|-----------|-------------------|---------------|----------------|------------|--------------|
| <u>AP</u> | NEXUS PHARMS | <u>6MG/ML</u> | <u>A207794</u> | <u>001</u> | Jan 14, 2019 |
| <u>AP</u> | PHARMASCIENCE INC | <u>6MG/ML</u> | <u>A207050</u> | <u>001</u> | Mar 24, 2017 |
| <u>AP</u> | SANDOZ INC | <u>6MG/ML</u> | <u>A205106</u> | <u>001</u> | Sep 21, 2018 |

BUSULFEX

| | | | | | |
|-----------|-----------------|---------------|----------------|------------|--------------|
| <u>AP</u> | +! OTSUKA PHARM | <u>6MG/ML</u> | <u>N020954</u> | <u>001</u> | Feb 04, 1999 |
|-----------|-----------------|---------------|----------------|------------|--------------|

MYLERAN

| | | | | | |
|-----------|------------------|---------------|----------------|------------|--------------|
| <u>AP</u> | ASPEN GLOBAL INC | <u>6MG/ML</u> | <u>A208536</u> | <u>001</u> | Nov 20, 2017 |
|-----------|------------------|---------------|----------------|------------|--------------|

TABLET; ORAL

MYLERAN

| | | | | | |
|----|--------------|-----|---------|-----|--|
| +! | ASPEN GLOBAL | 2MG | N009386 | 001 | |
|----|--------------|-----|---------|-----|--|

BUTABARBITAL SODIUM

TABLET; ORAL

BUTISOL SODIUM

| | | | | | |
|----|---------------------|------|---------|-----|--|
| +! | MYLAN SPECIALITY LP | 30MG | N000793 | 004 | |
|----|---------------------|------|---------|-----|--|

BUTENAFINE HYDROCHLORIDE

CREAM; TOPICAL

MENTAX

| | | | | | |
|----|-------|----|---------|-----|--------------|
| +! | MYLAN | 1% | N020524 | 001 | Oct 18, 1996 |
|----|-------|----|---------|-----|--------------|

BUTOCONAZOLE NITRATE

CREAM; VAGINAL

GYNAZOLE-1

| | | | | | |
|---|----------------|----|---------|-----|--------------|
| ! | PERRIGO ISRAEL | 2% | A200923 | 001 | May 18, 2012 |
|---|----------------|----|---------|-----|--------------|

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

BUTORPHANOL TARTRATE

| | | | | | |
|-----------|-------------------------|---------------|----------------|------------|--------------|
| <u>AP</u> | HIKMA FARMACEUTICA | <u>1MG/ML</u> | <u>A078400</u> | <u>001</u> | May 01, 2009 |
| <u>AP</u> | | <u>2MG/ML</u> | <u>A078400</u> | <u>002</u> | May 01, 2009 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>2MG/ML</u> | <u>A075046</u> | <u>001</u> | Aug 12, 1998 |

BUTORPHANOL TARTRATE PRESERVATIVE FREE

| | | | | | |
|-----------|-------------------------|---------------|----------------|------------|--------------|
| <u>AP</u> | ! HOSPIRA | <u>1MG/ML</u> | <u>A074626</u> | <u>001</u> | Jan 23, 1997 |
| <u>AP</u> | ! | <u>2MG/ML</u> | <u>A074626</u> | <u>002</u> | Jan 23, 1997 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>1MG/ML</u> | <u>A075045</u> | <u>001</u> | Aug 12, 1998 |
| <u>AP</u> | | <u>2MG/ML</u> | <u>A075045</u> | <u>002</u> | Aug 12, 1998 |

SPRAY, METERED; NASAL

BUTORPHANOL TARTRATE

| | | | | | |
|-----------|-------------------------|------------------|----------------|------------|--------------|
| <u>AB</u> | APOTEX INC | <u>1MG/SPRAY</u> | <u>A075499</u> | <u>001</u> | Dec 04, 2002 |
| <u>AB</u> | ! MYLAN | <u>1MG/SPRAY</u> | <u>A075759</u> | <u>001</u> | Aug 08, 2001 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>1MG/SPRAY</u> | <u>A075824</u> | <u>001</u> | Mar 12, 2002 |

CABAZITAXEL

SOLUTION; INTRAVENOUS

JEVTANA KIT

| | | | | | |
|----|-------------------|----------------------|---------|-----|--------------|
| +! | SANOFI AVENTIS US | 60MG/1.5ML (40MG/ML) | N201023 | 001 | Jun 17, 2010 |
|----|-------------------|----------------------|---------|-----|--------------|

CABERGOLINE

TABLET; ORAL

CABERGOLINE

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ACTAVIS LABS FL INC | <u>0.5MG</u> | <u>A078035</u> | <u>001</u> | Apr 21, 2008 |
| <u>AB</u> | INGENUS PHARMS LLC | <u>0.5MG</u> | <u>A204735</u> | <u>001</u> | Aug 01, 2018 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>0.5MG</u> | <u>A077750</u> | <u>001</u> | Mar 07, 2007 |
| <u>AB</u> | MYLAN PHARMS INC | <u>0.5MG</u> | <u>A202947</u> | <u>001</u> | Dec 02, 2013 |
| <u>AB</u> | ! PAR PHARM | <u>0.5MG</u> | <u>A076310</u> | <u>001</u> | Dec 29, 2005 |

CABOZANTINIB S-MALATE

CAPSULE; ORAL

COMETRIQ

| | | | | | |
|----|----------|--------------|---------|-----|--------------|
| +! | EXELIXIS | EQ 20MG BASE | N203756 | 001 | Nov 29, 2012 |
| + | | EQ 80MG BASE | N203756 | 002 | Nov 29, 2012 |

TABLET; ORAL

CABOMETYX

| | | | | | |
|----|--------------|--------------|---------|-----|--------------|
| + | EXELIXIS INC | EQ 20MG BASE | N208692 | 001 | Apr 25, 2016 |
| + | | EQ 40MG BASE | N208692 | 002 | Apr 25, 2016 |
| +! | | EQ 60MG BASE | N208692 | 003 | Apr 25, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

CAFFEINE CITRATE

SOLUTION; INTRAVENOUS

CAFCTT

| | | | | | |
|-----------|------------|-------------------------|-------------------------------------------|--------------------|--------------|
| AP | + ! | WEST-WARD PHARMS INT | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>N020793 001</u> | Sep 21, 1999 |
|-----------|------------|-------------------------|-------------------------------------------|--------------------|--------------|

CAFFEINE CITRATE

| | | | | | |
|-----------|--|-------------------------|-------------------------------------------|--------------------|--------------|
| AP | | AUROBINDO PHARMA LTD | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>A205013 001</u> | Sep 22, 2015 |
| AP | | EXELA PHARMA SCIENCE | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>A077233 001</u> | Sep 21, 2006 |
| AP | | FRESENIUS KABI USA | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>A077997 001</u> | Jul 20, 2007 |
| AP | | LUITPOLD | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>A077906 001</u> | May 15, 2007 |
| AP | | MICRO LABS | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>A207400 001</u> | Dec 14, 2017 |
| AP | | SAGENT PHARMS | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>A090827 001</u> | Aug 29, 2012 |
| AP | | SUN PHARMA GLOBAL | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>A090077 001</u> | Sep 30, 2009 |

SOLUTION; ORAL

CAFCTT

| | | | | | |
|-----------|------------|-------------------------|-------------------------------------------|--------------------|--------------|
| AA | + ! | WEST-WARD PHARMS INT | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>N020793 002</u> | Apr 12, 2000 |
|-----------|------------|-------------------------|-------------------------------------------|--------------------|--------------|

CAFFEINE CITRATE

| | | | | | |
|-----------|--|-------------------------|-------------------------------------------|--------------------|--------------|
| AA | | EXELA PHARMA SCS LLC | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>A077304 001</u> | Sep 21, 2006 |
| AA | | FRESENIUS KABI USA | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>A078002 001</u> | Jan 31, 2008 |
| AA | | LUITPOLD | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>A090064 001</u> | Nov 20, 2009 |
| AA | | SAGENT PHARMS | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>A091102 001</u> | Aug 29, 2012 |
| AA | | SUN PHARMA GLOBAL | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>A090357 001</u> | Sep 30, 2009 |

CAFFEINE; ERGOTAMINE TARTRATE

SUPPOSITORY; RECTAL

MIGERGOT

! HZNP

100MG; 2MG

A086557 001 Oct 04, 1983

TABLET; ORAL

CAFERGOT

| | | | | | |
|-----------|----------|--------|-------------------|--------------------|--|
| AA | ! | SANDOZ | <u>100MG; 1MG</u> | <u>A084294 001</u> | |
|-----------|----------|--------|-------------------|--------------------|--|

ERGOTAMINE TARTRATE AND CAFFEINE

| | | | | | |
|-----------|--|-------------------|-------------------|--------------------|--------------|
| AA | | HIKMA INTL PHARMS | <u>100MG; 1MG</u> | <u>A040510 001</u> | Sep 17, 2004 |
| AA | | MIKART | <u>100MG; 1MG</u> | <u>A040590 001</u> | Sep 16, 2005 |

CALCIFEDIOL

CAPSULE, EXTENDED RELEASE; ORAL

RAYALDEE

+! OPKO IRELAND GLOBAL 0.03MG

N208010 001 Jun 17, 2016

CALCIPOTRIENE

AEROSOL, FOAM; TOPICAL

SORILUX

+! MAYNE PHARMA 0.005%

N022563 001 Oct 06, 2010

CREAM; TOPICAL

CALCIPOTRIENE

| | | | | | |
|-----------|--|-----------------|---------------|--------------------|--------------|
| AB | | GLENMARK PHARMS | <u>0.005%</u> | <u>A205772 001</u> | Jun 09, 2015 |
| AB | | TOLMAR | <u>0.005%</u> | <u>A200935 001</u> | May 30, 2012 |

DOVONEX

| | | | | | |
|-----------|------------|---------------|---------------|--------------------|--------------|
| AB | + ! | LEO PHARMA AS | <u>0.005%</u> | <u>N020554 001</u> | Jul 22, 1996 |
|-----------|------------|---------------|---------------|--------------------|--------------|

OINTMENT; TOPICAL

CALCIPOTRIENE

! GLENMARK PHARMS INC 0.005%

A090633 001 Mar 24, 2010

SOLUTION; TOPICAL

CALCIPOTRIENE

| | | | | | |
|-----------|----------|------------------|---------------|--------------------|--------------|
| AT | | FOUGERA PHARMS | <u>0.005%</u> | <u>A078305 001</u> | May 06, 2008 |
| AT | | G AND W LABS INC | <u>0.005%</u> | <u>A078468 001</u> | Mar 24, 2011 |
| AT | | HI TECH PHARMA | <u>0.005%</u> | <u>A077579 001</u> | Nov 19, 2009 |
| AT | | NOVEL LABS INC | <u>0.005%</u> | <u>A207163 001</u> | Dec 26, 2017 |
| AT | ! | TOLMAR | <u>0.005%</u> | <u>A077029 001</u> | Nov 20, 2009 |

CALCITONIN SALMON

INJECTABLE; INJECTION

MIACALCIN

+! MYLAN IRELAND LTD 200 IU/ML

N017808 002 Mar 29, 1991

SPRAY, METERED; NASAL

CALCITONIN-SALMON

| | | | | | |
|-----------|----------|------------|---------------------|--------------------|--------------|
| AB | ! | APOTEX INC | <u>200 IU/SPRAY</u> | <u>A076396 001</u> | Nov 17, 2008 |
| AB | | PAR PHARM | <u>200 IU/SPRAY</u> | <u>A076979 001</u> | Jun 08, 2009 |

PRESCRIPTION DRUG PRODUCT LIST

CALCITRIOL

CAPSULE; ORAL

CALCITRIOL

| | | | | |
|-----------|-------------------------|----------------|--------------------|--------------|
| <u>AB</u> | AMNEAL PHARMS | <u>0.25MCG</u> | <u>A203289 002</u> | Jun 14, 2017 |
| <u>AB</u> | | <u>0.5MCG</u> | <u>A203289 001</u> | Jun 14, 2017 |
| <u>AB</u> | BIONPHARMA INC | <u>0.25MCG</u> | <u>A091174 001</u> | May 24, 2013 |
| <u>AB</u> | | <u>0.5MCG</u> | <u>A091174 002</u> | May 24, 2013 |
| <u>AB</u> | STRIDES PHARMA | <u>0.25MCG</u> | <u>A091356 001</u> | Dec 12, 2014 |
| <u>AB</u> | | <u>0.5MCG</u> | <u>A091356 002</u> | Dec 12, 2014 |
| <u>AB</u> | TEVA | <u>0.25MCG</u> | <u>A075765 001</u> | Oct 12, 2001 |
| <u>AB</u> | | <u>0.5MCG</u> | <u>A075765 002</u> | Oct 12, 2001 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>0.25MCG</u> | <u>A076917 001</u> | Mar 27, 2006 |

ROCALTROL

| | | | | |
|-----------|------------------|----------------|--------------------|--|
| <u>AB</u> | + VALIDUS PHARMS | <u>0.25MCG</u> | <u>N018044 001</u> | |
| <u>AB</u> | +! | <u>0.5MCG</u> | <u>N018044 002</u> | |

INJECTABLE; INJECTION

CALCITRIOL

! AKORN

0.001MG/ML

A078066 001 Jan 29, 2008

OINTMENT; TOPICAL

VECTICAL

+! GALDERMA LABS LP

3MCG/GM

N022087 001 Jan 23, 2009

SOLUTION; ORAL

CALCITRIOL

| | | | | |
|-----------|-------------------------|----------------|--------------------|--------------|
| <u>AA</u> | INVATECH PHARMA | <u>1MCG/ML</u> | <u>A209798 001</u> | Nov 21, 2018 |
| <u>AA</u> | WEST-WARD PHARMS INT | <u>1MCG/ML</u> | <u>A076242 001</u> | Jul 18, 2003 |

ROCALTROL

| | | | | |
|-----------|-------------------|----------------|--------------------|--------------|
| <u>AA</u> | +! VALIDUS PHARMS | <u>1MCG/ML</u> | <u>N021068 001</u> | Nov 20, 1998 |
|-----------|-------------------|----------------|--------------------|--------------|

CALCIUM ACETATE

CAPSULE; ORAL

CALCIUM ACETATE

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | AMNEAL PHARMS | <u>667MG</u> | <u>A201658 001</u> | Oct 06, 2014 |
| <u>AB</u> | CHARTWELL RX | <u>667MG</u> | <u>A091312 001</u> | Jun 01, 2012 |
| <u>AB</u> | HERITAGE PHARMS INC | <u>667MG</u> | <u>A202315 001</u> | Jun 29, 2015 |
| <u>AB</u> | INVAGEN PHARMS | <u>667MG</u> | <u>A203135 001</u> | Feb 07, 2013 |
| <u>AB</u> | LOTUS PHARM CO LTD | <u>667MG</u> | <u>A203298 001</u> | Jul 26, 2016 |
| <u>AB</u> | LUPIN LTD | <u>667MG</u> | <u>A202127 001</u> | Jul 09, 2015 |
| <u>AB</u> | NOSTRUM LABS INC | <u>667MG</u> | <u>A203179 001</u> | Oct 26, 2015 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>667MG</u> | <u>A077728 001</u> | Feb 26, 2008 |

PHOSLO GELCAPS

| | | | | |
|-----------|--------------------|--------------|--------------------|--------------|
| <u>AB</u> | +! FRESENIUS MEDCL | <u>667MG</u> | <u>N021160 003</u> | Apr 02, 2001 |
|-----------|--------------------|--------------|--------------------|--------------|

SOLUTION; ORAL

PHOSLYRA

+! FRESENIUS MEDCL

667MG/5ML

N022581 001 Apr 18, 2011

TABLET; ORAL

CALCIUM ACETATE

| | | | | |
|-----------|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | HERITAGE PHARMS INC | <u>667MG</u> | <u>A202885 001</u> | Jan 22, 2015 |
| <u>AB</u> | INVAGEN PHARMS | <u>667MG</u> | <u>A202420 001</u> | Feb 05, 2013 |
| <u>AB</u> | ! PADDOCK LLC | <u>667MG</u> | <u>A091561 001</u> | Apr 13, 2011 |

CALCIUM CHLORIDE

INJECTABLE; INJECTION

CALCIUM CHLORIDE 10%

| | | | | |
|-----------|---------------------|-----------------|--------------------|--------------|
| <u>AP</u> | INTL MEDICATION SYS | <u>100MG/ML</u> | <u>A203477 001</u> | May 09, 2018 |
| <u>AP</u> | LUITPOLD | <u>100MG/ML</u> | <u>A209088 001</u> | Jul 27, 2017 |

CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER

| | | | | |
|-----------|------------|-----------------|--------------------|--------------|
| <u>AP</u> | +! HOSPIRA | <u>100MG/ML</u> | <u>N021117 001</u> | Jan 28, 2000 |
|-----------|------------|-----------------|--------------------|--------------|

CALCIUM CHLORIDE; DEXTROSE; GLUTATHIONE DISULFIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

BSS PLUS

+! ALCON

0.154MG/ML; 0.92MG/ML; 0.184MG/ML; 0.2MG/ML; 0.38MG/ML; 2.1MG/ML; 7.14MG/ML; 0.42MG/ML

N018469 001

PRESCRIPTION DRUG PRODUCT LIST

CALCIUM CHLORIDE; DEXTROSE; LACTIC ACID; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PRISMASOL B22GK 4/0 IN PLASTIC CONTAINER

| | | | | |
|----|----------------------|-------------------------------------------------------------------------------------------------------|-------------|--------------|
| +! | BAXTER HLTHCARE CORP | N/A/1000ML;20GM/1000ML;5.4GM/1000ML;3.05GM/1000ML;0.314GM/1000ML;2.21GM/1000ML;7.07GM/1000ML (5000ML) | N021703 011 | Oct 10, 2008 |
|----|----------------------|-------------------------------------------------------------------------------------------------------|-------------|--------------|

PRISMASOL BGK 0/2.5 IN PLASTIC CONTAINER

| | | | | |
|----|----------------------|------------------------------------------------------------------------------------------------------|-------------|--------------|
| +! | BAXTER HLTHCARE CORP | 3.68GM/1000ML;20GM/1000ML;5.4GM/1000ML;3.05GM/1000ML;N/A/1000ML;3.09GM/1000ML;6.46GM/1000ML (5000ML) | N021703 006 | Oct 25, 2006 |
|----|----------------------|------------------------------------------------------------------------------------------------------|-------------|--------------|

PRISMASOL BGK 2/0 IN PLASTIC CONTAINER

| | | | | |
|----|----------------------|-------------------------------------------------------------------------------------------------------|-------------|--------------|
| +! | BAXTER HLTHCARE CORP | N/A/1000ML;20GM/1000ML;5.4GM/1000ML;2.03GM/1000ML;0.157GM/1000ML;3.09GM/1000ML;6.46GM/1000ML (5000ML) | N021703 002 | Oct 25, 2006 |
|----|----------------------|-------------------------------------------------------------------------------------------------------|-------------|--------------|

PRISMASOL BGK 2/3.5 IN PLASTIC CONTAINER

| | | | | |
|----|----------------------|----------------------------------------------------------------------------------------------------------|-------------|--------------|
| +! | BAXTER HLTHCARE CORP | 5.15GM/1000ML;20GM/1000ML;5.4GM/1000ML;2.03GM/1000ML;0.157GM/1000ML;3.09GM/1000ML;6.46GM/1000ML (5000ML) | N021703 003 | Oct 25, 2006 |
|----|----------------------|----------------------------------------------------------------------------------------------------------|-------------|--------------|

PRISMASOL BGK 4/0/1.2 IN PLASTIC CONTAINER

| | | | | |
|----|----------------------|-------------------------------------------------------------------------------------------------------|-------------|--------------|
| +! | BAXTER HLTHCARE CORP | N/A/1000ML;20GM/1000ML;5.4GM/1000ML;2.44GM/1000ML;0.314GM/1000ML;3.09GM/1000ML;6.46GM/1000ML (5000ML) | N021703 015 | Oct 10, 2008 |
|----|----------------------|-------------------------------------------------------------------------------------------------------|-------------|--------------|

PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER

| | | | | |
|----|----------------------|----------------------------------------------------------------------------------------------------------|-------------|--------------|
| +! | BAXTER HLTHCARE CORP | 3.68GM/1000ML;20GM/1000ML;5.4GM/1000ML;3.05GM/1000ML;0.314GM/1000ML;3.09GM/1000ML;6.46GM/1000ML (5000ML) | N021703 004 | Oct 25, 2006 |
|----|----------------------|----------------------------------------------------------------------------------------------------------|-------------|--------------|

PRISMASOL BK 0/0/1.2 IN PLASTIC CONTAINER

| | | | | |
|----|----------------------|--------------------------------------------------------------------------------------------------|-------------|--------------|
| +! | BAXTER HLTHCARE CORP | N/A/1000ML;N/A/1000ML;5.4GM/1000ML;2.44GM/1000ML;N/A/1000ML;3.09GM/1000ML;6.46GM/1000ML (5000ML) | N021703 014 | Oct 10, 2008 |
|----|----------------------|--------------------------------------------------------------------------------------------------|-------------|--------------|

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DELFLX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--------------|
| AT | FRESENIUS MEDCL | <u>25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;5.67MG/100ML;392MG/100ML</u> | N018883 001 | Nov 30, 1984 |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--------------|

DELFLX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--------------|
| AT | FRESENIUS MEDCL | <u>25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;5.38MG/100ML;448MG/100ML</u> | N018883 004 | Nov 30, 1984 |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--------------|

DELFLX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--------------|
| AT | FRESENIUS MEDCL | <u>18.4MG/100ML;1.5GM/100ML;5.08MG/100ML;5.38MG/100ML;448MG/100ML</u> | N020171 001 | Aug 19, 1992 |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--------------|

DELFLX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--------------|
| AT | FRESENIUS MEDCL | <u>25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5.67MG/100ML;392MG/100ML</u> | N018883 002 | Nov 30, 1984 |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--------------|

DELFLX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--------------|
| AT | FRESENIUS MEDCL | <u>25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;5.38MG/100ML;448MG/100ML</u> | N018883 005 | Nov 30, 1984 |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--------------|

DELFLX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--------------|
| AT | FRESENIUS MEDCL | <u>18.4MG/100ML;2.5GM/100ML;5.08MG/100ML;5.38MG/100ML;448MG/100ML</u> | N020171 002 | Aug 19, 1992 |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--------------|

DELFLX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|------------------------------------------------------------------------|--------------------|--------------|
| AT | FRESENIUS MEDCL | <u>25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;5.67MG/100ML;392MG/100ML</u> | N018883 003 | Nov 30, 1984 |
|-----------|-----------------|------------------------------------------------------------------------|--------------------|--------------|

DELFLX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|------------------------------------------------------------------------|--------------------|--------------|
| AT | FRESENIUS MEDCL | <u>25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;5.38MG/100ML;448MG/100ML</u> | N018883 006 | Nov 30, 1984 |
|-----------|-----------------|------------------------------------------------------------------------|--------------------|--------------|

DELFLX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|------------------------------------------------------------------------|--------------------|--------------|
| AT | FRESENIUS MEDCL | <u>18.4MG/100ML;4.25GM/100ML;5.08MG/100ML;5.38MG/100ML;448MG/100ML</u> | N020171 003 | Aug 19, 1992 |
|-----------|-----------------|------------------------------------------------------------------------|--------------------|--------------|

DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--------------|
| AT | BAXTER HLTHCARE | <u>18.3MG/100ML;1.5GM/100ML;5.08MG/100ML;5.38MG/100ML;448MG/100ML</u> | N020183 001 | Dec 04, 1992 |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--------------|

DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--|
| AT | BAXTER HLTHCARE | <u>18.3MG/100ML;1.5GM/100ML;5.08MG/100ML;5.38MG/100ML;448MG/100ML</u> | N017512 004 | |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--|

| | | | | |
|-----------|--|-----------------------------------------------------------------------|--------------------|--------------|
| AT | | <u>25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;5.38MG/100ML;448MG/100ML</u> | N020163 001 | Dec 04, 1992 |
|-----------|--|-----------------------------------------------------------------------|--------------------|--------------|

DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--|
| AT | BAXTER HLTHCARE | <u>25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;5.38MG/100ML;448MG/100ML</u> | N017512 005 | |
|-----------|-----------------|-----------------------------------------------------------------------|--------------------|--|

| | | | | |
|-----------|--|-----------------------------------------------------------------------|--------------------|--------------|
| AT | | <u>25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;5.38MG/100ML;448MG/100ML</u> | N020163 002 | Dec 04, 1992 |
|-----------|--|-----------------------------------------------------------------------|--------------------|--------------|

DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|------------------------------------------------------------------------|--------------------|--|
| AT | BAXTER HLTHCARE | <u>25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;5.38MG/100ML;448MG/100ML</u> | N017512 006 | |
|-----------|-----------------|------------------------------------------------------------------------|--------------------|--|

| | | | | |
|-----------|--|------------------------------------------------------------------------|--------------------|--------------|
| AT | | <u>25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;5.38MG/100ML;448MG/100ML</u> | N020163 003 | Dec 04, 1992 |
|-----------|--|------------------------------------------------------------------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER538MG/100ML; 448MG/100ML

DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 18.3MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5.38MG/100ML; 448MG/100ML N020183 002 Dec 04, 1992

DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 18.3MG/100ML; 3.5GM/100ML; 5.08MG/100ML; 5.38MG/100ML; 448MG/100ML N020183 003 Dec 04, 1992

DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

BAXTER HLTHCARE 18.3MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5.38MG/100ML; 448MG/100ML N020183 004 Dec 04, 1992

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INTRATHECAL

ELLIOTTS B SOLUTION

+! LUKARE MEDICAL LLC 0.2MG/ML; 0.8MG/ML; 0.3MG/ML; 0.3MG/ML; 1.9MG/ML; 7.3MG/ML; 0.2MG/ML N020577 001 Sep 27, 1996

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINERAP + ICU MEDICAL INC 20MG/100ML; 5GM/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML N017608 001DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINERAP B BRAUN 20MG/100ML; 5GM/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML N019634 003 Feb 24, 1988LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINERAP BAXTER HLTHCARE 20MG/100ML; 5GM/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML N016679 001POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINERAP BAXTER HLTHCARE 20MG/100ML; 5GM/100ML; 254MG/100ML; 600MG/100ML; 310MG/100ML N019367 006 Apr 05, 1985POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINERAP BAXTER HLTHCARE 20MG/100ML; 5GM/100ML; 179MG/100ML; 600MG/100ML; 310MG/100ML N019367 004 Apr 05, 1985AP 20MG/100ML; 5GM/100ML; 328MG/100ML; 600MG/100ML; 310MG/100ML N019367 005 Apr 05, 1985AP + ICU MEDICAL INC 20MG/100ML; 5GM/100ML; 179MG/100ML; 600MG/100ML; 310MG/100ML N019685 002 Oct 17, 1988POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINERAP BAXTER HLTHCARE 20MG/100ML; 5GM/100ML; 254MG/100ML; 600MG/100ML; 310MG/100ML N019367 007 Apr 05, 1985POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINERAP BAXTER HLTHCARE 20MG/100ML; 5GM/100ML; 328MG/100ML; 600MG/100ML; 310MG/100ML N019367 008 Apr 05, 1985

DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER

B BRAUN 10MG/100ML; 2.5GM/100ML; 15MG/100ML; 300MG/100ML; 160MG/100ML N019634 001 Feb 24, 1988

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

BAXTER HLTHCARE 20MG/100ML; 5GM/100ML; 105MG/100ML; 600MG/100ML; 310MG/100ML N019367 002 Apr 05, 1985

20MG/100ML; 5GM/100ML; 179MG/100ML; 600MG/100ML; 310MG/100ML N019367 003 Apr 05, 1985

POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

BAXTER HLTHCARE 20MG/100ML; 5GM/100ML; 105MG/100ML; 600MG/100ML; 310MG/100ML N019367 001 Apr 05, 1985

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TPN ELECTROLYTES IN PLASTIC CONTAINER

+! HOSPIRA 16.5MG/ML; 25.4MG/ML; 74.6MG/ML; 121MG/ML; 16.1MG/ML N018895 001 Jul 20, 1984

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

SOLUTION; IRRIGATION

BALANCED SALTAT AKORN 0.48MG/ML; 0.3MG/ML; 0.75MG/ML; 3.9MG/ML; 6.4MG/ML; 1.7MG/ML A075503 001 Sep 27, 2006AT B BRAUN 0.48MG/ML; 0.3MG/ML; 0.75MG/ML; 3.9MG/ML; 6.4MG/ML; 1.7MG/ML A091387 001 Feb 03, 2010BSSAT +! ALCON 0.48MG/ML; 0.3MG/ML; 0.75MG/ML; 3.9MG/ML; 6.4MG/ML; 1.7MG/ML N020742 001 Dec 10, 1997

PRESCRIPTION DRUG PRODUCT LIST

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

INJECTABLE; INJECTION

PHOXILLUM B2K 4/0 IN PLASTIC CONTAINER

| | | | | |
|----|----------------------|--------------------------------------------------------------------------------------------------|-------------|--------------|
| +! | BAXTER HLTHCARE CORP | N/A/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 2.21GM/1000ML; 6.95GM/1000ML; 0.187GM/1000ML (5000ML) | N207026 002 | Jan 13, 2015 |
|----|----------------------|--------------------------------------------------------------------------------------------------|-------------|--------------|

PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER

| | | | | |
|----|----------------------|--------------------------------------------------------------------------------------------|-------------|--------------|
| +! | BAXTER HLTHCARE CORP | 3.68GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML; 6.34GM/1000ML; 0.187GM/1000ML | N207026 001 | Jan 13, 2015 |
|----|----------------------|--------------------------------------------------------------------------------------------|-------------|--------------|

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

SOLUTION; PERFUSION, CARDIAC

CARDIOPLEGIC IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|-----------------------------------------------------------------------|---------------------------|--------------|
| AT | BAXTER HLTHCARE | <u>17.6MG/100ML; 325.3MG/100ML; 119.3MG/100ML; 643MG/100ML</u> | <u>A075323 001</u> | Apr 21, 2000 |
|-----------|-----------------|-----------------------------------------------------------------------|---------------------------|--------------|

PLEGISOL IN PLASTIC CONTAINER

| | | | | |
|-----------|------------|-----------------------------------------------------------------------|---------------------------|--------------|
| AT | +! HOSPIRA | <u>17.6MG/100ML; 325.3MG/100ML; 119.3MG/100ML; 643MG/100ML</u> | <u>N018608 001</u> | Feb 26, 1982 |
|-----------|------------|-----------------------------------------------------------------------|---------------------------|--------------|

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

RINGER'S IN PLASTIC CONTAINER

| | | | | |
|-----------|---------|---------------------------------------------------|---------------------------|--------------|
| AP | B BRAUN | <u>33MG/100ML; 30MG/100ML; 860MG/100ML</u> | <u>N020002 001</u> | Apr 17, 1992 |
|-----------|---------|---------------------------------------------------|---------------------------|--------------|

| | | | | |
|-----------|-----------------|---------------------------------------------------|---------------------------|--|
| AP | BAXTER HLTHCARE | <u>33MG/100ML; 30MG/100ML; 860MG/100ML</u> | <u>N016693 001</u> | |
|-----------|-----------------|---------------------------------------------------|---------------------------|--|

| | | | | |
|-----------|-----------------|---------------------------------------------------|---------------------------|--|
| AP | ICU MEDICAL INC | <u>33MG/100ML; 30MG/100ML; 860MG/100ML</u> | <u>N018251 001</u> | |
|-----------|-----------------|---------------------------------------------------|---------------------------|--|

SOLUTION; IRRIGATION

RINGER'S IN PLASTIC CONTAINER

| | | | | |
|-----------|---------|---------------------------------------------------|---------------------------|--|
| AT | B BRAUN | <u>33MG/100ML; 30MG/100ML; 860MG/100ML</u> | <u>N018156 001</u> | |
|-----------|---------|---------------------------------------------------|---------------------------|--|

| | | | | |
|-----------|-----------------|---------------------------------------------------|---------------------------|--------------|
| AT | BAXTER HLTHCARE | <u>33MG/100ML; 30MG/100ML; 860MG/100ML</u> | <u>N018495 001</u> | Feb 19, 1982 |
|-----------|-----------------|---------------------------------------------------|---------------------------|--------------|

| | | | | |
|-----------|-----------------|---------------------------------------------------|---------------------------|--|
| AT | ICU MEDICAL INC | <u>33MG/100ML; 30MG/100ML; 860MG/100ML</u> | <u>N017635 001</u> | |
|-----------|-----------------|---------------------------------------------------|---------------------------|--|

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

LACTATED RINGER'S IN PLASTIC CONTAINER

| | | | | |
|-----------|---------|----------------------------------------------------------------|---------------------------|--------------|
| AP | B BRAUN | <u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u> | <u>N019632 001</u> | Feb 29, 1988 |
|-----------|---------|----------------------------------------------------------------|---------------------------|--------------|

| | | | | |
|-----------|--------------------|----------------------------------------------------------------|---------------------------|--|
| AP | +! BAXTER HLTHCARE | <u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u> | <u>N016682 001</u> | |
|-----------|--------------------|----------------------------------------------------------------|---------------------------|--|

| | | | | |
|-----------|-----------------|----------------------------------------------------------------|---------------------------|--|
| AP | ICU MEDICAL INC | <u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u> | <u>N017641 001</u> | |
|-----------|-----------------|----------------------------------------------------------------|---------------------------|--|

SOLUTION; IRRIGATION

LACTATED RINGER'S IN PLASTIC CONTAINER

| | | | | |
|-----------|------------|----------------------------------------------------------------|---------------------------|--------------|
| AT | +! B BRAUN | <u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u> | <u>N018681 001</u> | Dec 27, 1982 |
|-----------|------------|----------------------------------------------------------------|---------------------------|--------------|

| | | | | |
|-----------|-----------------|----------------------------------------------------------------|---------------------------|--------------|
| AT | BAXTER HLTHCARE | <u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u> | <u>N018494 001</u> | Feb 19, 1982 |
|-----------|-----------------|----------------------------------------------------------------|---------------------------|--------------|

| | | | | |
|-----------|----|----------------------------------------------------------------|---------------------------|--------------|
| AT | +! | <u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u> | <u>N018921 001</u> | Apr 03, 1984 |
|-----------|----|----------------------------------------------------------------|---------------------------|--------------|

| | | | | |
|-----------|--------------------|----------------------------------------------------------------|---------------------------|--------------|
| AT | +! ICU MEDICAL INC | <u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u> | <u>N019416 001</u> | Jan 17, 1986 |
|-----------|--------------------|----------------------------------------------------------------|---------------------------|--------------|

CALCIUM GLUCONATE

SOLUTION; INTRAVENOUS

CALCIUM GLUCONATE

| | | | | |
|----|--------------------|---------------------|-------------|--------------|
| +! | FRESENIUS KABI USA | 1GM/10ML (100MG/ML) | N208418 001 | Jun 15, 2017 |
|----|--------------------|---------------------|-------------|--------------|

| | | | | |
|----|--|---------------------|-------------|--------------|
| +! | | 5GM/50ML (100MG/ML) | N208418 002 | Jun 15, 2017 |
|----|--|---------------------|-------------|--------------|

| | | | | |
|----|--|-----------------------|-------------|--------------|
| +! | | 10GM/100ML (100MG/ML) | N208418 003 | Jun 15, 2017 |
|----|--|-----------------------|-------------|--------------|

CALCIUM GLUCONATE IN SODIUM CHLORIDE

| | | | | |
|----|-----------------|--------------------|-------------|--------------|
| +! | HQ SPCLT PHARMA | 1GM/50ML (20MG/ML) | N210906 001 | Oct 29, 2018 |
|----|-----------------|--------------------|-------------|--------------|

| | | | | |
|----|--|---------------------|-------------|--------------|
| +! | | 2GM/100ML (20MG/ML) | N210906 002 | Oct 29, 2018 |
|----|--|---------------------|-------------|--------------|

CALFACTANT

SUSPENSION; INTRATRACHEAL

INFASURF PRESERVATIVE FREE

| | | | | |
|----|-----|---------|-------------|--------------|
| +! | ONY | 35MG/ML | N020521 001 | Jul 01, 1998 |
|----|-----|---------|-------------|--------------|

CANAGLIFLOZIN

TABLET; ORAL

INVOKANA

| | | | | |
|---|----------------|-------|-------------|--------------|
| + | JANSSEN PHARMS | 100MG | N204042 001 | Mar 29, 2013 |
|---|----------------|-------|-------------|--------------|

| | | | | |
|----|--|-------|-------------|--------------|
| +! | | 300MG | N204042 002 | Mar 29, 2013 |
|----|--|-------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

INVOKAMET

| | | | | | |
|---|----------------|--------------|---------|-----|--------------|
| + | JANSSEN PHARMS | 50MG; 500MG | N204353 | 001 | Aug 08, 2014 |
| + | | 50MG; 1GM | N204353 | 002 | Aug 08, 2014 |
| + | | 150MG; 500MG | N204353 | 003 | Aug 08, 2014 |
| + | ! | 150MG; 1GM | N204353 | 004 | Aug 08, 2014 |

TABLET, EXTENDED RELEASE; ORAL

INVOKAMET XR

| | | | | | |
|---|----------------|--------------|---------|-----|--------------|
| + | JANSSEN PHARMS | 50MG; 500MG | N205879 | 001 | Sep 20, 2016 |
| + | | 50MG; 1GM | N205879 | 002 | Sep 20, 2016 |
| + | | 150MG; 500MG | N205879 | 003 | Sep 20, 2016 |
| + | ! | 150MG; 1GM | N205879 | 004 | Sep 20, 2016 |

CANDESARTAN CILEXETIL

TABLET; ORAL

ATACAND

| | | | | | | |
|-----------|---|----------------|-------------|----------------|------------|--------------|
| <u>AB</u> | + | ANI PHARMS INC | <u>4MG</u> | <u>N020838</u> | <u>001</u> | Jun 04, 1998 |
| <u>AB</u> | + | | <u>8MG</u> | <u>N020838</u> | <u>002</u> | Jun 04, 1998 |
| <u>AB</u> | + | | <u>16MG</u> | <u>N020838</u> | <u>003</u> | Jun 04, 1998 |
| <u>AB</u> | + | ! | <u>32MG</u> | <u>N020838</u> | <u>004</u> | Jun 04, 1998 |

CANDESARTAN CILEXETIL

| | | | | | | |
|-----------|--|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | ALEMBIC PHARMS LTD | <u>4MG</u> | <u>A210302</u> | <u>001</u> | Dec 04, 2018 |
| <u>AB</u> | | | <u>8MG</u> | <u>A210302</u> | <u>002</u> | Dec 04, 2018 |
| <u>AB</u> | | | <u>16MG</u> | <u>A210302</u> | <u>003</u> | Dec 04, 2018 |
| <u>AB</u> | | | <u>32MG</u> | <u>A209119</u> | <u>001</u> | Jun 20, 2017 |
| <u>AB</u> | | MACLEODS PHARMS LTD | <u>4MG</u> | <u>A203813</u> | <u>001</u> | Dec 05, 2016 |
| <u>AB</u> | | | <u>8MG</u> | <u>A203813</u> | <u>002</u> | Dec 05, 2016 |
| <u>AB</u> | | | <u>16MG</u> | <u>A203813</u> | <u>003</u> | Dec 05, 2016 |
| <u>AB</u> | | | <u>32MG</u> | <u>A203813</u> | <u>004</u> | Dec 05, 2016 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>4MG</u> | <u>A078702</u> | <u>001</u> | May 03, 2013 |
| <u>AB</u> | | | <u>8MG</u> | <u>A078702</u> | <u>002</u> | May 03, 2013 |
| <u>AB</u> | | | <u>16MG</u> | <u>A078702</u> | <u>003</u> | May 03, 2013 |
| <u>AB</u> | | | <u>32MG</u> | <u>A078702</u> | <u>004</u> | May 03, 2013 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>4MG</u> | <u>A091390</u> | <u>001</u> | Aug 23, 2017 |
| <u>AB</u> | | | <u>8MG</u> | <u>A091390</u> | <u>002</u> | Aug 23, 2017 |
| <u>AB</u> | | | <u>16MG</u> | <u>A091390</u> | <u>003</u> | Aug 23, 2017 |
| <u>AB</u> | | | <u>32MG</u> | <u>A091390</u> | <u>004</u> | Aug 23, 2017 |

CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ATACAND HCT

| | | | | | | |
|-----------|---|----------------|---------------------|----------------|------------|--------------|
| <u>AB</u> | + | ANI PHARMS INC | <u>16MG; 12.5MG</u> | <u>N021093</u> | <u>001</u> | Sep 05, 2000 |
| <u>AB</u> | + | | <u>32MG; 12.5MG</u> | <u>N021093</u> | <u>002</u> | Sep 05, 2000 |
| <u>AB</u> | + | ! | <u>32MG; 25MG</u> | <u>N021093</u> | <u>003</u> | May 16, 2008 |

CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE

| | | | | | | |
|-----------|--|-------------------------|---------------------|----------------|------------|--------------|
| <u>AB</u> | | DR REDDYS LABS LTD | <u>16MG; 12.5MG</u> | <u>A202965</u> | <u>001</u> | Jun 03, 2013 |
| <u>AB</u> | | | <u>32MG; 12.5MG</u> | <u>A202965</u> | <u>002</u> | Jun 03, 2013 |
| <u>AB</u> | | | <u>32MG; 25MG</u> | <u>A202965</u> | <u>003</u> | Jun 03, 2013 |
| <u>AB</u> | | MACLEODS PHARMS LTD | <u>16MG; 12.5MG</u> | <u>A204100</u> | <u>001</u> | Feb 27, 2015 |
| <u>AB</u> | | | <u>32MG; 12.5MG</u> | <u>A204100</u> | <u>002</u> | Feb 27, 2015 |
| <u>AB</u> | | | <u>32MG; 25MG</u> | <u>A204100</u> | <u>003</u> | Feb 27, 2015 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>16MG; 12.5MG</u> | <u>A090704</u> | <u>001</u> | Dec 04, 2012 |
| <u>AB</u> | | | <u>32MG; 12.5MG</u> | <u>A090704</u> | <u>002</u> | Dec 04, 2012 |
| <u>AB</u> | | | <u>32MG; 25MG</u> | <u>A090704</u> | <u>003</u> | Dec 04, 2012 |
| <u>AB</u> | | PRINSTON INC | <u>16MG; 12.5MG</u> | <u>A207455</u> | <u>001</u> | Apr 11, 2018 |
| <u>AB</u> | | | <u>32MG; 12.5MG</u> | <u>A207455</u> | <u>002</u> | Apr 11, 2018 |
| <u>AB</u> | | | <u>32MG; 25MG</u> | <u>A207455</u> | <u>003</u> | Apr 11, 2018 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>16MG; 12.5MG</u> | <u>A203466</u> | <u>001</u> | Nov 27, 2017 |
| <u>AB</u> | | | <u>32MG; 12.5MG</u> | <u>A203466</u> | <u>002</u> | Nov 27, 2017 |
| <u>AB</u> | | | <u>32MG; 25MG</u> | <u>A203466</u> | <u>003</u> | Nov 27, 2017 |

CANGRELOR

POWDER; INTRAVENOUS

KENGREAL

| | | | | | |
|---|----------------|-----------|---------|-----|--------------|
| + | CHIESI USA INC | 50MG/VIAL | N204958 | 001 | Jun 22, 2015 |
|---|----------------|-----------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

CANNABIDIOL

SOLUTION; ORAL

EPIDIOLEX

+! GW RES LTD

100MG/ML

N210365 001 Sep 28, 2018

CAPECITABINE

TABLET; ORAL

CAPECITABINE

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>150MG</u> | <u>A202593 001</u> | Apr 23, 2015 |
| <u>AB</u> | | <u>500MG</u> | <u>A202593 002</u> | Apr 23, 2015 |
| <u>AB</u> | ALKEM LABS LTD | <u>150MG</u> | <u>A207652 001</u> | Nov 24, 2017 |
| <u>AB</u> | | <u>500MG</u> | <u>A207652 002</u> | Nov 24, 2017 |
| <u>AB</u> | AMNEAL PHARMS | <u>150MG</u> | <u>A204741 001</u> | Feb 28, 2017 |
| <u>AB</u> | | <u>500MG</u> | <u>A204741 002</u> | Feb 28, 2017 |
| <u>AB</u> | EUGIA PHARMA | <u>150MG</u> | <u>A210604 001</u> | Apr 17, 2018 |
| <u>AB</u> | | <u>500MG</u> | <u>A210604 002</u> | Apr 17, 2018 |
| <u>AB</u> | MSN LABS PVT LTD | <u>150MG</u> | <u>A209365 001</u> | Jul 02, 2018 |
| <u>AB</u> | | <u>500MG</u> | <u>A209365 002</u> | Jul 02, 2018 |
| <u>AB</u> | MYLAN PHARMS INC | <u>150MG</u> | <u>A090943 001</u> | Aug 08, 2014 |
| <u>AB</u> | | <u>500MG</u> | <u>A090943 002</u> | Aug 08, 2014 |
| <u>AB</u> | SHILPA MEDICARE LTD | <u>150MG</u> | <u>A207456 001</u> | Dec 12, 2016 |
| <u>AB</u> | | <u>500MG</u> | <u>A207456 002</u> | Dec 12, 2016 |
| <u>AB</u> | TEVA PHARMS USA | <u>150MG</u> | <u>A091649 001</u> | Sep 16, 2013 |
| <u>AB</u> | | <u>500MG</u> | <u>A091649 002</u> | Sep 16, 2013 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>150MG</u> | <u>A200483 001</u> | Jul 14, 2016 |
| <u>AB</u> | | <u>500MG</u> | <u>A200483 002</u> | Jul 14, 2016 |
| | <u>XELODA</u> | | | |
| <u>AB</u> | + HOFFMANN LA ROCHE | <u>150MG</u> | <u>N020896 001</u> | Apr 30, 1998 |
| <u>AB</u> | +! | <u>500MG</u> | <u>N020896 002</u> | Apr 30, 1998 |

CAPREOMYCIN SULFATE

INJECTABLE; INJECTION

CAPASTAT SULFATEAP +! AKORNEQ 1GM BASE/VIALN050095 001CAPREOMYCIN SULFATEAP HISUN PHARMEQ 1GM BASE/VIALA204796 001 Oct 18, 2018

HANGZHOU

AP MYLAN LABS LTDEQ 1GM BASE/VIALA202634 001 Nov 27, 2017CAPSAICIN

PATCH; TOPICAL

QUTENZA

+! AVERITAS

8%

N022395 001 Nov 16, 2009

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

| | | | | |
|-----------|-------------------|---------------|--------------------|--------------|
| <u>AB</u> | HIKMA INTL PHARMS | <u>12.5MG</u> | <u>A074505 001</u> | Feb 13, 1996 |
| <u>AB</u> | | <u>25MG</u> | <u>A074505 002</u> | Feb 13, 1996 |
| <u>AB</u> | | <u>50MG</u> | <u>A074505 003</u> | Feb 13, 1996 |
| <u>AB</u> | | <u>100MG</u> | <u>A074505 004</u> | Feb 13, 1996 |
| <u>AB</u> | MYLAN PHARMS INC | <u>12.5MG</u> | <u>A074434 001</u> | Feb 13, 1996 |
| <u>AB</u> | | <u>25MG</u> | <u>A074434 002</u> | Feb 13, 1996 |
| <u>AB</u> | | <u>50MG</u> | <u>A074434 003</u> | Feb 13, 1996 |
| <u>AB</u> | ! | <u>100MG</u> | <u>A074434 004</u> | Feb 13, 1996 |
| <u>AB</u> | PRINSTON INC | <u>12.5MG</u> | <u>A074477 001</u> | Feb 13, 1996 |
| <u>AB</u> | | <u>25MG</u> | <u>A074477 002</u> | Feb 13, 1996 |
| <u>AB</u> | | <u>50MG</u> | <u>A074477 003</u> | Feb 13, 1996 |
| <u>AB</u> | | <u>100MG</u> | <u>A074477 004</u> | Feb 13, 1996 |
| <u>AB</u> | TEVA | <u>12.5MG</u> | <u>A074322 001</u> | Feb 13, 1996 |
| <u>AB</u> | | <u>25MG</u> | <u>A074322 002</u> | Feb 13, 1996 |
| <u>AB</u> | | <u>50MG</u> | <u>A074322 003</u> | Feb 13, 1996 |
| <u>AB</u> | | <u>100MG</u> | <u>A074322 004</u> | Feb 13, 1996 |
| <u>AB</u> | WATSON LABS | <u>12.5MG</u> | <u>A074386 001</u> | May 23, 1996 |
| <u>AB</u> | | <u>25MG</u> | <u>A074386 002</u> | May 23, 1996 |
| <u>AB</u> | | <u>50MG</u> | <u>A074386 003</u> | May 23, 1996 |
| <u>AB</u> | | <u>100MG</u> | <u>A074386 004</u> | May 23, 1996 |
| <u>AB</u> | WOCKHARDT LTD | <u>12.5MG</u> | <u>A074532 001</u> | Mar 28, 1997 |
| <u>AB</u> | | <u>25MG</u> | <u>A074532 002</u> | Mar 28, 1997 |
| <u>AB</u> | | <u>50MG</u> | <u>A074532 003</u> | Mar 28, 1997 |
| <u>AB</u> | | <u>100MG</u> | <u>A074532 004</u> | Mar 28, 1997 |

PRESCRIPTION DRUG PRODUCT LIST

CAPTOPRIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CAPTOPRIL AND HYDROCHLOROTHIAZIDE

| | | | | | |
|---|-------|------------|---------|-----|--------------|
| | MYLAN | 25MG; 15MG | A074896 | 001 | Dec 29, 1997 |
| ! | | 25MG; 25MG | A074896 | 002 | Dec 29, 1997 |
| ! | | 50MG; 15MG | A074896 | 004 | Dec 29, 1997 |
| | | 50MG; 25MG | A074896 | 003 | Dec 29, 1997 |

CARBACHOL

SOLUTION; INTRAOCULAR

MIOSTAT

| | | | | | |
|---|-------|-------|---------|-----|--|
| + | ALCON | 0.01% | N016968 | 001 | |
|---|-------|-------|---------|-----|--|

CARBAMAZEPINE

CAPSULE, EXTENDED RELEASE; ORAL

CARBAMAZEPINE

| | | | | | |
|-----------|-------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | APOTEX INC | <u>100MG</u> | <u>A078986</u> | <u>001</u> | Nov 25, 2011 |
| <u>AB</u> | | <u>200MG</u> | <u>A078986</u> | <u>002</u> | Nov 25, 2011 |
| <u>AB</u> | | <u>300MG</u> | <u>A078986</u> | <u>003</u> | Nov 25, 2011 |
| <u>AB</u> | MYLAN IRELAND LTD | <u>100MG</u> | <u>A076697</u> | <u>001</u> | May 20, 2011 |
| <u>AB</u> | | <u>200MG</u> | <u>A076697</u> | <u>002</u> | May 20, 2011 |
| <u>AB</u> | | <u>300MG</u> | <u>A076697</u> | <u>003</u> | May 20, 2011 |
| <u>AB</u> | TARO | <u>100MG</u> | <u>A201106</u> | <u>001</u> | Jun 21, 2013 |
| <u>AB</u> | | <u>200MG</u> | <u>A201106</u> | <u>002</u> | Jun 21, 2013 |
| <u>AB</u> | | <u>300MG</u> | <u>A201106</u> | <u>003</u> | Jun 21, 2013 |
| <u>AB</u> | TEVA PHARMS | <u>100MG</u> | <u>A078592</u> | <u>001</u> | Sep 20, 2012 |
| <u>AB</u> | | <u>200MG</u> | <u>A078592</u> | <u>002</u> | Sep 20, 2012 |
| <u>AB</u> | | <u>300MG</u> | <u>A078592</u> | <u>003</u> | Sep 20, 2012 |

CARBATROL

| | | | | | | |
|-----------|---|-------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | SHIRE | <u>100MG</u> | <u>N020712</u> | <u>003</u> | Sep 30, 1997 |
| <u>AB</u> | + | | <u>200MG</u> | <u>N020712</u> | <u>001</u> | Sep 30, 1997 |
| <u>AB</u> | + | ! | <u>300MG</u> | <u>N020712</u> | <u>002</u> | Sep 30, 1997 |

EQUETRO

| | | | | | |
|---|----------------|-------|---------|-----|--------------|
| + | VALIDUS PHARMS | 100MG | N021710 | 001 | Dec 10, 2004 |
| + | | 200MG | N021710 | 002 | Dec 10, 2004 |
| + | ! | 300MG | N021710 | 003 | Dec 10, 2004 |

SUSPENSION; ORAL

CARBAMAZEPINE

| | | | | | | |
|-----------|------------------|------------------|------------------|----------------|--------------|--------------|
| <u>AB</u> | WOCKHARDT BIO AG | <u>100MG/5ML</u> | <u>A075714</u> | <u>001</u> | Jun 05, 2002 | |
| | <u>TEGRETOL</u> | | | | | |
| <u>AB</u> | + | NOVARTIS | <u>100MG/5ML</u> | <u>N018927</u> | <u>001</u> | Dec 18, 1987 |
| | <u>TERIL</u> | | | | | |
| <u>AB</u> | TARO PHARM | <u>100MG/5ML</u> | <u>A076729</u> | <u>001</u> | Sep 20, 2004 | |

TABLET; ORAL

CARBAMAZEPINE

| | | | | | |
|-----------|----------------|--------------|----------------|------------|--------------|
| <u>AB</u> | APOTEX INC | <u>200MG</u> | <u>A075948</u> | <u>001</u> | Feb 27, 2002 |
| <u>AB</u> | TARO | <u>200MG</u> | <u>A074649</u> | <u>001</u> | Oct 03, 1996 |
| <u>AB</u> | TORRENT PHARMS | <u>200MG</u> | <u>A077272</u> | <u>002</u> | Dec 07, 2005 |

EPITOL

| | | | | | |
|-----------|------|--------------|----------------|------------|--------------|
| <u>AB</u> | TEVA | <u>200MG</u> | <u>A070541</u> | <u>001</u> | Sep 17, 1986 |
|-----------|------|--------------|----------------|------------|--------------|

TEGRETOL

| | | | | | |
|-----------|---|----------|--------------|----------------|------------|
| <u>AB</u> | + | NOVARTIS | <u>200MG</u> | <u>N016608</u> | <u>001</u> |
|-----------|---|----------|--------------|----------------|------------|

CARBAMAZEPINE

| | | | | | |
|--|----------------|-------|---------|-----|--------------|
| | TORRENT PHARMS | 100MG | A077272 | 001 | Dec 07, 2005 |
| | | 300MG | A077272 | 003 | Dec 07, 2005 |
| | | 400MG | A077272 | 004 | Dec 07, 2005 |

TABLET, CHEWABLE; ORAL

CARBAMAZEPINE

| | | | | | |
|-----------|-----------------|--------------|----------------|------------|--------------|
| <u>AB</u> | TARO PHARM INDS | <u>100MG</u> | <u>A075687</u> | <u>001</u> | Oct 24, 2000 |
| <u>AB</u> | TORRENT PHARMS | <u>100MG</u> | <u>A075712</u> | <u>001</u> | Jul 05, 2001 |

EPITOL

| | | | | | |
|-----------|------|--------------|----------------|------------|--------------|
| <u>AB</u> | TEVA | <u>100MG</u> | <u>A073524</u> | <u>001</u> | Jul 29, 1992 |
|-----------|------|--------------|----------------|------------|--------------|

TEGRETOL

| | | | | | |
|-----------|---|----------|--------------|----------------|------------|
| <u>AB</u> | + | NOVARTIS | <u>100MG</u> | <u>N018281</u> | <u>001</u> |
|-----------|---|----------|--------------|----------------|------------|

CARBAMAZEPINE

| | | | | | |
|---|-----------------|-------|---------|-----|--------------|
| ! | TARO PHARM INDS | 200MG | A075687 | 002 | Jul 29, 2002 |
|---|-----------------|-------|---------|-----|--------------|

TABLET, EXTENDED RELEASE; ORAL

CARBAMAZEPINE

| | | | | | |
|-----------|------|--------------|----------------|------------|--------------|
| <u>AB</u> | TARO | <u>100MG</u> | <u>A078115</u> | <u>001</u> | Mar 31, 2009 |
| <u>AB</u> | | <u>200MG</u> | <u>A078115</u> | <u>002</u> | Mar 31, 2009 |
| <u>AB</u> | | <u>400MG</u> | <u>A078115</u> | <u>003</u> | Mar 31, 2009 |

TEGRETOL-XR

| | | | | | | |
|-----------|---|----------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | NOVARTIS | <u>100MG</u> | <u>N020234</u> | <u>001</u> | Mar 25, 1996 |
|-----------|---|----------|--------------|----------------|------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

CARBAMAZEPINE

TABLET, EXTENDED RELEASE;ORAL

TEGRETOL-XR

| | | | | |
|-----------|---|--------------|--------------------|--------------|
| AB | + | 200MG | N020234 002 | Mar 25, 1996 |
| AB | + | 400MG | N020234 003 | Mar 25, 1996 |

CARBIDOPA

TABLET;ORAL

CARBIDOPA

| | | | | |
|-----------|-------------------------|-------------|--------------------|--------------|
| AB | ALVOGEN MALTA | 25MG | A204291 001 | Jan 08, 2016 |
| AB | AMERIGEN PHARMS LTD | 25MG | A203261 001 | Mar 10, 2014 |
| AB | EDENBRIDGE PHARMS | 25MG | A205304 001 | Feb 17, 2016 |
| AB | NOVEL LABS INC | 25MG | A204763 001 | Oct 20, 2017 |
| AB | ZYDUS PHARMS USA INC | 25MG | A209910 001 | May 07, 2018 |

LODOSYN

| | | | | |
|-----------|---|------|-------------|--------------------|
| AB | + | ATON | 25MG | N017830 001 |
|-----------|---|------|-------------|--------------------|

CARBIDOPA; ENTACAPONE; LEVODOPA

TABLET;ORAL

CARBIDOPA, LEVODOPA AND ENTACAPONE

| | | | | |
|-----------|-------------------|----------------------------|--------------------|--------------|
| AB | SUN PHARMA GLOBAL | 25MG;200MG;100MG | A079085 001 | May 10, 2012 |
| AB | | 37.5MG;200MG;150MG | A079085 002 | May 10, 2012 |
| AB | WOCKHARDT LTD | 12.5MG;200MG;50MG | A090786 001 | Nov 20, 2012 |
| AB | | 18.75MG;200MG;75MG | A090833 001 | Nov 20, 2012 |
| AB | | 25MG;200MG;100MG | A090833 002 | Nov 20, 2012 |
| AB | | 31.25MG;200MG;125MG | A090833 003 | Nov 20, 2012 |
| AB | | 37.5MG;200MG;150MG | A090833 004 | Nov 20, 2012 |
| AB | | 50MG;200MG;200MG | A090833 005 | Nov 20, 2012 |

STALEVO 100

| | | | | | |
|-----------|---|--------------|-------------------------|--------------------|--------------|
| AB | + | ORION PHARMA | 25MG;200MG;100MG | N021485 002 | Jun 11, 2003 |
|-----------|---|--------------|-------------------------|--------------------|--------------|

STALEVO 125

| | | | | | |
|-----------|---|--------------|----------------------------|--------------------|--------------|
| AB | + | ORION PHARMA | 31.25MG;200MG;125MG | N021485 006 | Aug 29, 2008 |
|-----------|---|--------------|----------------------------|--------------------|--------------|

STALEVO 150

| | | | | | |
|-----------|---|--------------|---------------------------|--------------------|--------------|
| AB | + | ORION PHARMA | 37.5MG;200MG;150MG | N021485 003 | Jun 11, 2003 |
|-----------|---|--------------|---------------------------|--------------------|--------------|

STALEVO 200

| | | | | | |
|-----------|---|--------------|-------------------------|--------------------|--------------|
| AB | + | ORION PHARMA | 50MG;200MG;200MG | N021485 004 | Aug 02, 2007 |
|-----------|---|--------------|-------------------------|--------------------|--------------|

STALEVO 50

| | | | | | |
|-----------|---|--------------|--------------------------|--------------------|--------------|
| AB | + | ORION PHARMA | 12.5MG;200MG;50MG | N021485 001 | Jun 11, 2003 |
|-----------|---|--------------|--------------------------|--------------------|--------------|

STALEVO 75

| | | | | | |
|-----------|---|--------------|---------------------------|--------------------|--------------|
| AB | + | ORION PHARMA | 18.75MG;200MG;75MG | N021485 005 | Aug 29, 2008 |
|-----------|---|--------------|---------------------------|--------------------|--------------|

CARBIDOPA; LEVODOPA

CAPSULE, EXTENDED RELEASE;ORAL

RYTARY

| | | | | | |
|--|---|----------------|---------------|-------------|--------------|
| | + | IMPAX LABS INC | 23.75MG;95MG | N203312 001 | Jan 07, 2015 |
| | + | | 36.25MG;145MG | N203312 002 | Jan 07, 2015 |
| | + | | 48.75MG;195MG | N203312 003 | Jan 07, 2015 |
| | + | | 61.25MG;245MG | N203312 004 | Jan 07, 2015 |

SUSPENSION;ENTERAL

DUOPA

| | | | | | |
|--|---|------------|-------------------|-------------|--------------|
| | + | ABBVIE INC | 4.63MG/ML;20MG/ML | N203952 001 | Jan 09, 2015 |
|--|---|------------|-------------------|-------------|--------------|

TABLET;ORAL

CARBIDOPA AND LEVODOPA

| | | | | |
|-----------|-------------------|-------------------|--------------------|--------------|
| AB | ACTAVIS ELIZABETH | 10MG;100MG | A074260 001 | Sep 03, 1993 |
| AB | | 25MG;100MG | A074260 002 | Sep 03, 1993 |
| AB | | 25MG;250MG | A074260 003 | Sep 03, 1993 |
| AB | APOTEX INC | 10MG;100MG | A077120 001 | Jun 02, 2008 |
| AB | | 25MG;100MG | A077120 002 | Jun 02, 2008 |
| AB | | 25MG;250MG | A077120 003 | Jun 02, 2008 |
| AB | MAYNE PHARMA | 10MG;100MG | A073618 001 | Aug 28, 1992 |
| AB | | 25MG;100MG | A073589 001 | Aug 28, 1992 |
| AB | | 25MG;250MG | A073607 001 | Aug 28, 1992 |
| AB | MYLAN | 10MG;100MG | A090324 001 | Sep 28, 2009 |
| AB | | 25MG;100MG | A090324 002 | Sep 28, 2009 |
| AB | | 25MG;250MG | A090324 003 | Sep 28, 2009 |
| AB | SUN PHARM INDS | 10MG;100MG | A078536 001 | Oct 28, 2008 |
| AB | | 25MG;100MG | A078536 002 | Oct 28, 2008 |
| AB | | 25MG;250MG | A078536 003 | Oct 28, 2008 |

SINEMET

| | | | | |
|-----------|---|-------------------|-------------------|--------------------|
| AB | + | MERCK SHARP DOHME | 10MG;100MG | N017555 001 |
| AB | + | | 25MG;100MG | N017555 003 |
| AB | + | | 25MG;250MG | N017555 002 |

PRESCRIPTION DRUG PRODUCT LIST

CARBIDOPA; LEVODOPA

TABLET, EXTENDED RELEASE;ORAL

CARBIDOPA AND LEVODOPA

| | | | | |
|-----------|-----------------|-------------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>25MG;100MG</u> | <u>A202323 001</u> | Feb 08, 2013 |
| <u>AB</u> | | <u>50MG;200MG</u> | <u>A202323 002</u> | Feb 08, 2013 |
| <u>AB</u> | APOTEX | <u>25MG;100MG</u> | <u>A076212 001</u> | Jun 16, 2004 |
| <u>AB</u> | | <u>50MG;200MG</u> | <u>A076212 002</u> | Jun 16, 2004 |
| <u>AB</u> | IMPAX LABS | <u>25MG;100MG</u> | <u>A076521 001</u> | May 14, 2004 |
| <u>AB</u> | | <u>50MG;200MG</u> | <u>A076521 002</u> | May 14, 2004 |
| <u>AB</u> | MYLAN | <u>25MG;100MG</u> | <u>A075091 002</u> | Apr 21, 2000 |
| <u>AB</u> | | <u>50MG;200MG</u> | <u>A075091 001</u> | Sep 30, 1999 |
| <u>AB</u> | SUN PHARM INDS | <u>25MG;100MG</u> | <u>A077828 001</u> | Aug 23, 2007 |
| <u>AB</u> | | <u>50MG;200MG</u> | <u>A077828 002</u> | Aug 23, 2007 |

SINEMET CR

| | | | | |
|-----------|---------------------|-------------------|--------------------|--------------|
| <u>AB</u> | + MERCK SHARP DOHME | <u>25MG;100MG</u> | <u>N019856 002</u> | Dec 24, 1992 |
| <u>AB</u> | +! | <u>50MG;200MG</u> | <u>N019856 001</u> | May 30, 1991 |

TABLET, ORALLY DISINTEGRATING;ORAL

CARBIDOPA AND LEVODOPA

| | | | | |
|-----------|-------------------|-------------------|--------------------|--------------|
| <u>AB</u> | MYLAN | <u>10MG;100MG</u> | <u>A078893 001</u> | Sep 18, 2008 |
| <u>AB</u> | | <u>25MG;100MG</u> | <u>A078893 002</u> | Sep 18, 2008 |
| <u>AB</u> | ! | <u>25MG;250MG</u> | <u>A078893 003</u> | Sep 18, 2008 |
| <u>AB</u> | SUN PHARMA GLOBAL | <u>10MG;100MG</u> | <u>A078690 001</u> | Jul 31, 2009 |
| <u>AB</u> | | <u>25MG;100MG</u> | <u>A078690 002</u> | Jul 31, 2009 |
| <u>AB</u> | | <u>25MG;250MG</u> | <u>A078690 003</u> | Jul 31, 2009 |

CARBINOXAMINE MALEATE

SOLUTION;ORAL

CARBINOXAMINE MALEATE

| | | | | |
|-----------|----------------|----------------|--------------------|--------------|
| <u>AA</u> | ! MIKART | <u>4MG/5ML</u> | <u>A040458 001</u> | Apr 25, 2003 |
| <u>AA</u> | VINTAGE PHARMS | <u>4MG/5ML</u> | <u>A040814 001</u> | Feb 26, 2008 |

SUSPENSION, EXTENDED RELEASE;ORAL

KARBINAL ER

| | | | | |
|----|-----------------|---------|-------------|--------------|
| +! | TRIS PHARMA INC | 4MG/5ML | N022556 001 | Mar 28, 2013 |
|----|-----------------|---------|-------------|--------------|

TABLET;ORAL

CARBINOXAMINE MALEATE

| | | | | |
|-----------|----------------------|------------|--------------------|--------------|
| <u>AA</u> | INVAGEN PHARMS | <u>4MG</u> | <u>A090435 001</u> | Apr 15, 2010 |
| <u>AA</u> | ! MIKART | <u>4MG</u> | <u>A040442 001</u> | Mar 19, 2003 |
| <u>AA</u> | MISSION PHARMACAL CO | <u>4MG</u> | <u>A090756 001</u> | May 27, 2011 |
| <u>AA</u> | VINTAGE PHARMS | <u>4MG</u> | <u>A040639 002</u> | May 30, 2008 |
| | MIKART | 6MG | A207484 001 | May 31, 2016 |

CARBOPLATIN

INJECTABLE;IV (INFUSION)

CARBOPLATIN

| | | | | |
|-----------|--------------------|-----------------------------|--------------------|--------------|
| <u>AP</u> | ACCORD HLTHCARE | <u>50MG/5ML (10MG/ML)</u> | <u>A206775 001</u> | Feb 09, 2017 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A206775 002</u> | Feb 09, 2017 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A206775 003</u> | Feb 09, 2017 |
| <u>AP</u> | | <u>600MG/60ML (10MG/ML)</u> | <u>A206775 004</u> | Feb 09, 2017 |
| <u>AP</u> | AKORN | <u>50MG/5ML (10MG/ML)</u> | <u>A090475 001</u> | Jul 29, 2009 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A090475 002</u> | Jul 29, 2009 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A090475 003</u> | Jul 29, 2009 |
| <u>AP</u> | | <u>600MG/60ML (10MG/ML)</u> | <u>A091268 002</u> | Jul 28, 2010 |
| <u>AP</u> | CIPLA LTD | <u>50MG/5ML (10MG/ML)</u> | <u>A077861 001</u> | Jan 18, 2007 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A077861 002</u> | Jan 18, 2007 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A077861 003</u> | Jan 18, 2007 |
| <u>AP</u> | | <u>600MG/60ML (10MG/ML)</u> | <u>A077861 004</u> | Jan 18, 2007 |
| <u>AP</u> | EUGIA PHARMA | <u>50MG/5ML (10MG/ML)</u> | <u>A205487 001</u> | Mar 28, 2016 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A205487 002</u> | Mar 28, 2016 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A205487 003</u> | Mar 28, 2016 |
| <u>AP</u> | FRESENIUS KABI USA | <u>50MG/5ML (10MG/ML)</u> | <u>A077432 001</u> | Sep 29, 2006 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A077432 002</u> | Sep 29, 2006 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A077432 003</u> | Sep 29, 2006 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A077247 003</u> | Oct 21, 2004 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A077266 003</u> | Feb 15, 2006 |
| <u>AP</u> | | <u>600MG/60ML (10MG/ML)</u> | <u>A077266 004</u> | Feb 15, 2006 |
| <u>AP</u> | GLAND PHARMA LTD | <u>50MG/5ML (10MG/ML)</u> | <u>A207324 001</u> | Feb 15, 2017 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A207324 002</u> | Feb 15, 2017 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A207324 003</u> | Feb 15, 2017 |
| <u>AP</u> | | <u>600MG/60ML (10MG/ML)</u> | <u>A207324 004</u> | Feb 15, 2017 |
| <u>AP</u> | HOSPIRA | <u>50MG/5ML (10MG/ML)</u> | <u>A076517 001</u> | Oct 14, 2004 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A076517 002</u> | Oct 14, 2004 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A076517 003</u> | Oct 14, 2004 |
| <u>AP</u> | | <u>600MG/60ML (10MG/ML)</u> | <u>A077059 001</u> | Nov 23, 2004 |

PRESCRIPTION DRUG PRODUCT LIST

CARBOPLATIN

INJECTABLE; IV (INFUSION)

CARBOPLATIN

| | | | | |
|-----------|-------------------------|-----------------------------|--------------------|--------------|
| <u>AP</u> | INGENUS PHARMS LLC | <u>50MG/5ML (10MG/ML)</u> | <u>A208487 001</u> | Apr 26, 2017 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A208487 002</u> | Apr 26, 2017 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A208487 003</u> | Apr 26, 2017 |
| <u>AP</u> | | <u>600MG/60ML (10MG/ML)</u> | <u>A208487 004</u> | Apr 26, 2017 |
| <u>AP</u> | MYLAN INSTITUTIONAL | <u>50MG/5ML (10MG/ML)</u> | <u>A077998 001</u> | Apr 24, 2007 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A077998 002</u> | Apr 24, 2007 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A077998 003</u> | Apr 24, 2007 |
| <u>AP</u> | MYLAN LABS LTD | <u>50MG/5ML (10MG/ML)</u> | <u>A091063 001</u> | Nov 09, 2011 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A091063 002</u> | Nov 09, 2011 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A091063 003</u> | Nov 09, 2011 |
| <u>AP</u> | | <u>600MG/60ML (10MG/ML)</u> | <u>A091063 004</u> | Nov 09, 2011 |
| <u>AP</u> | NANJING KING-FRIEND | <u>50MG/5ML (10MG/ML)</u> | <u>A077096 001</u> | Jun 14, 2005 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A077096 002</u> | Jun 14, 2005 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A077096 003</u> | Jun 14, 2005 |
| <u>AP</u> | | <u>600MG/60ML (10MG/ML)</u> | <u>A077096 004</u> | Jun 03, 2013 |
| <u>AP</u> | ! PHARMACHEMIE BV | <u>50MG/5ML (10MG/ML)</u> | <u>A077269 001</u> | Oct 14, 2004 |
| <u>AP</u> | ! | <u>150MG/15ML (10MG/ML)</u> | <u>A077269 002</u> | Oct 14, 2004 |
| <u>AP</u> | ! | <u>450MG/45ML (10MG/ML)</u> | <u>A077269 003</u> | Oct 14, 2004 |
| <u>AP</u> | ! | <u>600MG/60ML (10MG/ML)</u> | <u>A077269 004</u> | Dec 28, 2007 |
| <u>AP</u> | PLIVA LACHEMA | <u>50MG/5ML (10MG/ML)</u> | <u>A078631 001</u> | Dec 02, 2008 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A078631 002</u> | Dec 02, 2008 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A078631 003</u> | Dec 02, 2008 |
| <u>AP</u> | | <u>600MG/60ML (10MG/ML)</u> | <u>A078631 004</u> | Dec 02, 2008 |
| <u>AP</u> | SANDOZ INC | <u>50MG/5ML (10MG/ML)</u> | <u>A078280 001</u> | May 08, 2008 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A078280 002</u> | May 08, 2008 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A078280 003</u> | May 08, 2008 |
| <u>AP</u> | SUN PHARMA GLOBAL | <u>50MG/5ML (10MG/ML)</u> | <u>A077926 001</u> | Sep 19, 2008 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A077926 002</u> | Sep 19, 2008 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A077926 003</u> | Sep 19, 2008 |
| <u>AP</u> | ! TEVA PHARMS USA | <u>50MG/5ML (10MG/ML)</u> | <u>A077139 001</u> | Sep 21, 2005 |
| <u>AP</u> | ! | <u>150MG/15ML (10MG/ML)</u> | <u>A077139 002</u> | Sep 21, 2005 |
| <u>AP</u> | ! | <u>450MG/45ML (10MG/ML)</u> | <u>A077139 003</u> | Sep 21, 2005 |
| <u>AP</u> | ! | <u>600MG/60ML (10MG/ML)</u> | <u>A077139 004</u> | Sep 21, 2005 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>50MG/5ML (10MG/ML)</u> | <u>A077244 001</u> | Oct 15, 2004 |
| <u>AP</u> | | <u>150MG/15ML (10MG/ML)</u> | <u>A077244 002</u> | Oct 15, 2004 |
| <u>AP</u> | | <u>450MG/45ML (10MG/ML)</u> | <u>A077244 003</u> | Oct 15, 2004 |
| <u>AP</u> | | <u>600MG/60ML (10MG/ML)</u> | <u>A077244 004</u> | Jan 20, 2006 |

CARBOPROST TROMETHAMINE

INJECTABLE; INJECTION

HEMABATE

+! PHARMACIA AND
UPJOHN

EQ 0.25MG BASE/ML

N017989 001

CARFILZOMIB

POWDER; INTRAVENOUS

KYPROLIS

+ ONYX THERAP

10MG/VIAL

N202714 003 Jun 07, 2018

+

30MG/VIAL

N202714 002 Jun 03, 2016

+!

60MG/VIAL

N202714 001 Jul 20, 2012

CARGLUMIC ACID

TABLET; ORAL

CARBAGLU

+! ORPHAN EUROPE

200MG

N022562 001 Mar 18, 2010

CARIPRAZINE HYDROCHLORIDE

CAPSULE; ORAL

VRAYLAR

+ ALLERGAN SALES LLC

EQ 1.5MG BASE

N204370 001 Sep 17, 2015

+

EQ 3MG BASE

N204370 002 Sep 17, 2015

+

EQ 4.5MG BASE

N204370 003 Sep 17, 2015

+!

EQ 6MG BASE

N204370 004 Sep 17, 2015

CARISOPRODOL

TABLET; ORAL

CARISOPRODOL

| | | | | |
|-----------|-------------------|--------------|--------------------|--------------|
| <u>AA</u> | ACCELRX LABS | <u>350MG</u> | <u>A040576 001</u> | Jun 07, 2005 |
| <u>AA</u> | AUROBINDO PHARMA | <u>350MG</u> | <u>A040792 001</u> | Aug 06, 2009 |
| <u>AA</u> | HIKMA INTL PHARMS | <u>350MG</u> | <u>A040124 001</u> | Jan 24, 1996 |
| <u>AA</u> | NATCO PHARMA LTD | <u>350MG</u> | <u>A090988 001</u> | Oct 28, 2014 |

PRESCRIPTION DRUG PRODUCT LIST

CARISOPRODOL

TABLET; ORAL

CARISOPRODOL

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AA</u> | NOVAST LABS | <u>350MG</u> | <u>A040823</u> | <u>001</u> | Oct 22, 2008 |
| <u>AA</u> | ORIENT PHARMA CO LTD | <u>350MG</u> | <u>A205085</u> | <u>001</u> | Oct 28, 2014 |
| <u>AA</u> | SCIEGEN PHARMS INC | <u>350MG</u> | <u>A203374</u> | <u>001</u> | Jan 27, 2014 |
| <u>AA</u> | STRIDES PHARMA | <u>350MG</u> | <u>A205513</u> | <u>002</u> | Nov 12, 2015 |
| <u>AA</u> | SUN PHARM INDUSTRIES | <u>350MG</u> | <u>A089346</u> | <u>001</u> | Oct 17, 1991 |
| <u>AA</u> | VINTAGE PHARMS | <u>350MG</u> | <u>A040245</u> | <u>001</u> | Sep 08, 1997 |
| <u>AA</u> | WATSON LABS | <u>350MG</u> | <u>A087499</u> | <u>001</u> | Apr 20, 1982 |
| <u>AA</u> | WILSHIRE PHARMS INC | <u>350MG</u> | <u>A205126</u> | <u>002</u> | Jul 08, 2015 |

SOMA

| | | | | | |
|-----------|----------|---------------------|--------------|----------------|------------|
| <u>AA</u> | <u>+</u> | MYLAN SPECIALITY LP | <u>350MG</u> | <u>N011792</u> | <u>001</u> |
|-----------|----------|---------------------|--------------|----------------|------------|

CARISOPRODOL

| | | | | | |
|-----------|---------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA | <u>250MG</u> | <u>A040792</u> | <u>002</u> | Nov 08, 2016 |
| <u>AB</u> | NOSTRUM LABS INC | <u>250MG</u> | <u>A207237</u> | <u>001</u> | May 11, 2017 |
| <u>AB</u> | STRIDES PHARMA | <u>250MG</u> | <u>A205513</u> | <u>001</u> | Nov 12, 2015 |
| <u>AB</u> | WILSHIRE PHARMS INC | <u>250MG</u> | <u>A205126</u> | <u>001</u> | Jul 08, 2015 |

SOMA

| | | | | | | |
|-----------|----------|---------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | <u>+</u> | MYLAN SPECIALITY LP | <u>250MG</u> | <u>N011792</u> | <u>004</u> | Sep 13, 2007 |
|-----------|----------|---------------------|--------------|----------------|------------|--------------|

CARMUSTINE

IMPLANT; INTRACRANIAL

GLIADEL

| | | | | | |
|----------|------------------|-------|---------|-----|--------------|
| <u>+</u> | ARBOR PHARMS LLC | 7.7MG | N020637 | 001 | Sep 23, 1996 |
|----------|------------------|-------|---------|-----|--------------|

INJECTABLE; INJECTION

BICNU

| | | | | | |
|-----------|----------|-------------------|-------------------|----------------|------------|
| <u>AP</u> | <u>+</u> | EMCURE PHARMS LTD | <u>100MG/VIAL</u> | <u>N017422</u> | <u>001</u> |
|-----------|----------|-------------------|-------------------|----------------|------------|

CARMUSTINE

| | | | | | |
|-----------|------------------|-------------------|----------------|------------|--------------|
| <u>AP</u> | AMNEAL PHARMS CO | <u>100MG/VIAL</u> | <u>A211229</u> | <u>001</u> | Oct 16, 2018 |
| <u>AP</u> | NAVINTA LLC | <u>100MG/VIAL</u> | <u>A210179</u> | <u>001</u> | Sep 11, 2018 |

CARTEOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CARTEOLOL HYDROCHLORIDE

| | | | | | | |
|-----------|-----------------|------------|----------------|----------------|--------------|--------------|
| <u>AT</u> | BAUSCH AND LOMB | <u>1%</u> | <u>A075546</u> | <u>001</u> | Jan 20, 2000 | |
| <u>AT</u> | <u>!</u> | SANDOZ INC | <u>1%</u> | <u>A075476</u> | <u>001</u> | Jan 03, 2000 |

CARVEDILOL

TABLET; ORAL

CARVEDILOL

| | | | | | |
|-----------|---------------------|----------------|----------------|------------|--------------|
| <u>AB</u> | APOTEX INC | <u>3.125MG</u> | <u>A078165</u> | <u>001</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>6.25MG</u> | <u>A078165</u> | <u>002</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A078165</u> | <u>003</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A078165</u> | <u>004</u> | Sep 05, 2007 |
| <u>AB</u> | AUROBINDO PHARMA | <u>3.125MG</u> | <u>A078332</u> | <u>001</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>6.25MG</u> | <u>A078332</u> | <u>002</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A078332</u> | <u>003</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A078332</u> | <u>004</u> | Sep 05, 2007 |
| <u>AB</u> | BEXIMCO USA | <u>3.125MG</u> | <u>A078384</u> | <u>001</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>6.25MG</u> | <u>A078384</u> | <u>002</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A078384</u> | <u>003</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A078384</u> | <u>004</u> | Sep 05, 2007 |
| <u>AB</u> | CHARTWELL MOLECULAR | <u>3.125MG</u> | <u>A077474</u> | <u>001</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>6.25MG</u> | <u>A077474</u> | <u>002</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A077474</u> | <u>003</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A077474</u> | <u>004</u> | Sep 05, 2007 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>3.125MG</u> | <u>A076649</u> | <u>001</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>6.25MG</u> | <u>A076649</u> | <u>002</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A076649</u> | <u>003</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A076649</u> | <u>004</u> | Sep 05, 2007 |
| <u>AB</u> | GLENMARK GENERICS | <u>3.125MG</u> | <u>A078251</u> | <u>001</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>6.25MG</u> | <u>A078251</u> | <u>002</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A078251</u> | <u>003</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A078251</u> | <u>004</u> | Sep 05, 2007 |
| <u>AB</u> | LUPIN | <u>3.125MG</u> | <u>A078217</u> | <u>001</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>6.25MG</u> | <u>A078217</u> | <u>002</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A078217</u> | <u>003</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A078217</u> | <u>004</u> | Sep 05, 2007 |
| <u>AB</u> | MYLAN | <u>3.125MG</u> | <u>A077316</u> | <u>001</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>6.25MG</u> | <u>A077316</u> | <u>002</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A077316</u> | <u>003</u> | Sep 05, 2007 |

PRESCRIPTION DRUG PRODUCT LIST

CARVEDILOL

TABLET; ORAL

CARVEDILOL

| | | | | | |
|-----------|-------------------------|----------------|----------------|------------|--------------|
| <u>AB</u> | | <u>25MG</u> | <u>A077316</u> | <u>004</u> | Sep 05, 2007 |
| <u>AB</u> | SANDOZ | <u>3.125MG</u> | <u>A078227</u> | <u>001</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>6.25MG</u> | <u>A078227</u> | <u>002</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A078227</u> | <u>003</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A078227</u> | <u>004</u> | Sep 05, 2007 |
| <u>AB</u> | SUN PHARM INDS INC | <u>3.125MG</u> | <u>A077346</u> | <u>004</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>6.25MG</u> | <u>A077346</u> | <u>001</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A077346</u> | <u>002</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A077346</u> | <u>003</u> | Sep 05, 2007 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>3.125MG</u> | <u>A076989</u> | <u>001</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>6.25MG</u> | <u>A076989</u> | <u>002</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A076989</u> | <u>003</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A076989</u> | <u>004</u> | Sep 05, 2007 |
| <u>AB</u> | TARO | <u>3.125MG</u> | <u>A077780</u> | <u>001</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>6.25MG</u> | <u>A077780</u> | <u>002</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A077780</u> | <u>003</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A077780</u> | <u>004</u> | Sep 05, 2007 |
| <u>AB</u> | TEVA | <u>3.125MG</u> | <u>A076373</u> | <u>001</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>6.25MG</u> | <u>A076373</u> | <u>002</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A076373</u> | <u>003</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A076373</u> | <u>004</u> | Sep 05, 2007 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>3.125MG</u> | <u>A077614</u> | <u>004</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>6.25MG</u> | <u>A077614</u> | <u>001</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A077614</u> | <u>002</u> | Sep 05, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A077614</u> | <u>003</u> | Sep 05, 2007 |
| | <u>COREG</u> | | | | |
| <u>AB</u> | + SMITHKLINE BEECHAM | <u>3.125MG</u> | <u>N020297</u> | <u>004</u> | May 29, 1997 |
| <u>AB</u> | + | <u>6.25MG</u> | <u>N020297</u> | <u>003</u> | Sep 14, 1995 |
| <u>AB</u> | +! | <u>12.5MG</u> | <u>N020297</u> | <u>002</u> | Sep 14, 1995 |
| <u>AB</u> | + | <u>25MG</u> | <u>N020297</u> | <u>001</u> | Sep 14, 1995 |

CARVEDILOL PHOSPHATE

CAPSULE, EXTENDED RELEASE; ORAL

CARVEDILOL PHOSPHATE

| | | | | | |
|-----------|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | IMPAX LABS INC | <u>10MG</u> | <u>A204717</u> | <u>001</u> | May 07, 2018 |
| <u>AB</u> | | <u>20MG</u> | <u>A204717</u> | <u>002</u> | May 07, 2018 |
| <u>AB</u> | | <u>40MG</u> | <u>A204717</u> | <u>003</u> | May 07, 2018 |
| <u>AB</u> | | <u>80MG</u> | <u>A204717</u> | <u>004</u> | May 07, 2018 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>10MG</u> | <u>A090132</u> | <u>001</u> | Oct 25, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A090132</u> | <u>002</u> | Oct 25, 2017 |
| <u>AB</u> | | <u>40MG</u> | <u>A090132</u> | <u>003</u> | Oct 25, 2017 |
| <u>AB</u> | | <u>80MG</u> | <u>A090132</u> | <u>004</u> | Oct 25, 2017 |
| | <u>COREG CR</u> | | | | |
| <u>AB</u> | + SMITHKLINE BEECHAM | <u>10MG</u> | <u>N022012</u> | <u>001</u> | Oct 20, 2006 |
| <u>AB</u> | + | <u>20MG</u> | <u>N022012</u> | <u>002</u> | Oct 20, 2006 |
| <u>AB</u> | +! | <u>40MG</u> | <u>N022012</u> | <u>003</u> | Oct 20, 2006 |
| <u>AB</u> | + | <u>80MG</u> | <u>N022012</u> | <u>004</u> | Oct 20, 2006 |

CASPOFUNGIN ACETATE

POWDER; INTRAVENOUS

CANCIDAS

| | | | | | |
|-----------|----------|------------------|----------------|------------|--------------|
| <u>AP</u> | +! MERCK | <u>50MG/VIAL</u> | <u>N021227</u> | <u>001</u> | Jan 26, 2001 |
| <u>AP</u> | +! | <u>70MG/VIAL</u> | <u>N021227</u> | <u>002</u> | Jan 26, 2001 |

CASPOFUNGIN ACETATE

| | | | | | |
|-----------|--------------------|------------------|----------------|------------|--------------|
| <u>AP</u> | CIPLA | <u>50MG/VIAL</u> | <u>A209489</u> | <u>001</u> | Jul 12, 2018 |
| <u>AP</u> | | <u>70MG/VIAL</u> | <u>A209489</u> | <u>002</u> | Jul 12, 2018 |
| <u>AP</u> | FRESENIUS KABI USA | <u>50MG/VIAL</u> | <u>N206110</u> | <u>001</u> | Dec 30, 2016 |
| <u>AP</u> | | <u>70MG/VIAL</u> | <u>N206110</u> | <u>002</u> | Dec 30, 2016 |
| <u>AP</u> | GLAND PHARMA LTD | <u>50MG/VIAL</u> | <u>A207092</u> | <u>001</u> | Sep 29, 2017 |
| <u>AP</u> | | <u>70MG/VIAL</u> | <u>A207092</u> | <u>002</u> | Sep 29, 2017 |
| <u>AP</u> | MYLAN LABS LTD | <u>50MG/VIAL</u> | <u>A207650</u> | <u>001</u> | Sep 29, 2017 |
| <u>AP</u> | | <u>70MG/VIAL</u> | <u>A207650</u> | <u>002</u> | Sep 29, 2017 |
| <u>AP</u> | SANDOZ INC | <u>50MG/VIAL</u> | <u>A200833</u> | <u>001</u> | Jun 28, 2018 |
| <u>AP</u> | | <u>70MG/VIAL</u> | <u>A200833</u> | <u>002</u> | Jun 28, 2018 |
| <u>AP</u> | XELLIA PHARMS APS | <u>50MG/VIAL</u> | <u>A205923</u> | <u>001</u> | Jul 02, 2018 |
| <u>AP</u> | | <u>70MG/VIAL</u> | <u>A205923</u> | <u>002</u> | Jul 02, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

CEFACTOR

CAPSULE; ORAL

CEFACTOR

| | | | | |
|-----------|-----------------|----------------------|--------------------|--------------|
| <u>AB</u> | HIKMA | <u>EQ 250MG BASE</u> | <u>A065350 001</u> | Apr 03, 2007 |
| <u>AB</u> | ! | <u>EQ 500MG BASE</u> | <u>A065350 002</u> | Apr 03, 2007 |
| <u>AB</u> | YUNG SHIN PHARM | <u>EQ 250MG BASE</u> | <u>A065146 001</u> | Jan 22, 2004 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065146 002</u> | Jan 22, 2004 |

FOR SUSPENSION; ORAL

CEFACTOR

| | | | | |
|--|-----------------|-------------------|-------------|--------------|
| | YUNG SHIN PHARM | EQ 125MG BASE/5ML | A065412 001 | Feb 17, 2012 |
| | | EQ 187MG BASE/5ML | A065412 002 | Feb 17, 2012 |
| | | EQ 250MG BASE/5ML | A065412 003 | Feb 17, 2012 |
| | ! | EQ 375MG BASE/5ML | A065412 004 | Feb 17, 2012 |

TABLET, EXTENDED RELEASE; ORAL

CEFACTOR

| | | | | |
|--|------|---------------|-------------|--------------|
| | TEVA | EQ 375MG BASE | A065058 001 | Sep 04, 2002 |
| | ! | EQ 500MG BASE | A065058 002 | Sep 04, 2002 |

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE; ORAL

CEFADROXIL

| | | | | |
|-----------|------------------|----------------------|--------------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA | <u>EQ 500MG BASE</u> | <u>A065352 001</u> | Jan 25, 2007 |
| <u>AB</u> | HIKMA | <u>EQ 500MG BASE</u> | <u>A065311 001</u> | Feb 07, 2006 |
| <u>AB</u> | LUPIN | <u>EQ 500MG BASE</u> | <u>A065392 001</u> | May 29, 2007 |
| <u>AB</u> | ORCHID HLTHCARE | <u>EQ 500MG BASE</u> | <u>A065309 001</u> | Sep 18, 2006 |
| <u>AB</u> | ! | <u>EQ 500MG BASE</u> | <u>A065282 001</u> | Jan 20, 2006 |

FOR SUSPENSION; ORAL

CEFADROXIL

| | | | | |
|-----------|-----------------|--------------------------|--------------------|--------------|
| <u>AB</u> | AUROBINDO | <u>EQ 250MG BASE/5ML</u> | <u>A065349 001</u> | Apr 25, 2013 |
| <u>AB</u> | | <u>EQ 500MG BASE/5ML</u> | <u>A065349 002</u> | Apr 25, 2013 |
| <u>AB</u> | HIKMA PHARMS | <u>EQ 250MG BASE/5ML</u> | <u>A091036 001</u> | Nov 28, 2012 |
| <u>AB</u> | | <u>EQ 500MG BASE/5ML</u> | <u>A091036 002</u> | Nov 28, 2012 |
| <u>AB</u> | LUPIN | <u>EQ 250MG BASE/5ML</u> | <u>A065396 001</u> | Feb 21, 2008 |
| <u>AB</u> | ! | <u>EQ 500MG BASE/5ML</u> | <u>A065396 002</u> | Feb 21, 2008 |
| <u>AB</u> | ORCHID HLTHCARE | <u>EQ 250MG BASE/5ML</u> | <u>A065307 002</u> | Oct 16, 2006 |
| <u>AB</u> | | <u>EQ 500MG BASE/5ML</u> | <u>A065307 003</u> | Oct 16, 2006 |

TABLET; ORAL

CEFADROXIL

| | | | | |
|-----------|-----------------|--------------------|--------------------|--------------|
| <u>AB</u> | HIKMA | <u>EQ 1GM BASE</u> | <u>A065260 001</u> | Mar 30, 2006 |
| <u>AB</u> | ORCHID HLTHCARE | <u>EQ 1GM BASE</u> | <u>A065301 001</u> | Sep 18, 2006 |
| | ! | EQ 1GM BASE | A062774 001 | Apr 08, 1987 |

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

CEFAZOLIN SODIUM

| | | | | | |
|-----------|----------------------------|---------------------------|--------------------|--------------|--------------|
| <u>AP</u> | ACS DOBFAR | <u>EQ 500MG BASE/VIAL</u> | <u>A065303 001</u> | Oct 22, 2008 | |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A065303 002</u> | Oct 22, 2008 | |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A065306 001</u> | Oct 22, 2008 | |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>EQ 500MG BASE/VIAL</u> | <u>A065047 001</u> | Sep 18, 2001 | |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A065047 002</u> | Sep 18, 2001 | |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A065143 001</u> | Oct 18, 2004 | |
| <u>AP</u> | ! | <u>EQ 500MG BASE/VIAL</u> | <u>A065226 001</u> | Apr 21, 2005 | |
| <u>AP</u> | ! | <u>EQ 1GM BASE/VIAL</u> | <u>A065226 002</u> | Apr 21, 2005 | |
| <u>AP</u> | ! | <u>EQ 1GM BASE/VIAL</u> | <u>A065244 001</u> | Aug 12, 2005 | |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A201654 001</u> | Feb 03, 2016 | |
| <u>AP</u> | ! | <u>EQ 10GM BASE/VIAL</u> | <u>A065247 001</u> | Aug 12, 2005 | |
| <u>AP</u> | QILU PHARM CO LTD | <u>EQ 1GM BASE/VIAL</u> | <u>A203661 001</u> | Dec 28, 2015 | |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A209217 001</u> | Oct 17, 2018 | |
| <u>AP</u> | SANDOZ | <u>EQ 500MG BASE/VIAL</u> | <u>A062831 001</u> | Dec 09, 1988 | |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A062831 002</u> | Dec 09, 1988 | |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A065345 001</u> | May 09, 2007 | |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A062831 003</u> | Sep 25, 1992 | |
| | ANCEF IN PLASTIC CONTAINER | | | | |
| | ! | BAXTER HLTHCARE | EQ 20MG BASE/ML | A063002 002 | Mar 28, 1991 |
| | | CEFAZOLIN AND DEXTROSE | | | |
| | +! | B BRAUN | EQ 1GM BASE/VIAL | N050779 002 | Jul 27, 2000 |
| | + | | EQ 2GM BASE/VIAL | N050779 003 | Jan 13, 2012 |
| | | CEFAZOLIN SODIUM | | | |
| | ! | ACS DOBFAR | EQ 20GM BASE/VIAL | A065306 002 | Aug 18, 2014 |
| | ! | SAMSON MEDCL | EQ 100GM BASE/VIAL | A065141 001 | Nov 29, 2006 |
| | ! | | EQ 300GM BASE/VIAL | A065141 002 | Nov 29, 2006 |

PRESCRIPTION DRUG PRODUCT LIST

CEFAZOLIN SODIUM

SOLUTION;INTRAVENOUS

CEFAZOLIN IN PLASTIC CONTAINER

| | | | | |
|-----------------|-------------------------------------|---------|-----|--------------|
| BAXTER HLTHCARE | EQ 2GM BASE/100ML (EQ 20MG BASE/ML) | N207131 | 001 | Aug 07, 2015 |
| CORP | | | | |

CEFDINIR

CAPSULE;ORAL

CEFDINIR

| | | | | | |
|-----------|------------------|--------------|----------------|------------|--------------|
| AB | AUROBINDO PHARMA | 300MG | A065434 | 001 | Jan 07, 2008 |
| AB | LUPIN | 300MG | A065264 | 001 | May 19, 2006 |
| AB | ORCHID HLTHCARE | 300MG | A065418 | 001 | Jul 18, 2007 |
| AB | ! SANDOZ | 300MG | A065330 | 001 | Apr 06, 2007 |
| AB | TEVA PHARMS | 300MG | A065368 | 001 | May 09, 2007 |

FOR SUSPENSION;ORAL

CEFDINIR

| | | | | | |
|-----------|------------------|------------------|----------------|------------|--------------|
| AB | AUROBINDO PHARMA | 125MG/5ML | A065473 | 001 | Dec 14, 2007 |
| AB | | 250MG/5ML | A065473 | 002 | Dec 14, 2007 |
| AB | LUPIN | 125MG/5ML | A065259 | 001 | May 31, 2006 |
| AB | | 250MG/5ML | A065259 | 002 | May 07, 2007 |
| AB | ORCHID HLTHCARE | 125MG/5ML | A065429 | 001 | Jul 18, 2007 |
| AB | | 250MG/5ML | A065429 | 002 | Jul 18, 2007 |
| AB | SANDOZ | 125MG/5ML | A065337 | 001 | Apr 06, 2007 |
| AB | ! | 250MG/5ML | A065337 | 002 | Apr 06, 2007 |
| AB | TEVA PHARMS | 125MG/5ML | A065332 | 001 | May 04, 2007 |
| AB | | 250MG/5ML | A065332 | 002 | May 04, 2007 |

CEFEPIME HYDROCHLORIDE

INJECTABLE;INJECTION

CEFEPIME HYDROCHLORIDE

| | | | | | |
|-----------|-------------------|---------------------------|----------------|------------|--------------|
| AP | ACS DOBFAR | EQ 1GM BASE/VIAL | A065441 | 001 | Mar 20, 2008 |
| AP | | EQ 2GM BASE/VIAL | A065441 | 002 | Mar 20, 2008 |
| AP | HOSPIRA INC | EQ 500MG BASE/VIAL | A065369 | 001 | Jun 18, 2007 |
| AP | | EQ 1GM BASE/VIAL | A065369 | 002 | Jun 18, 2007 |
| AP | | EQ 1GM BASE/VIAL | A202268 | 001 | Jul 30, 2012 |
| AP | | EQ 2GM BASE/VIAL | A065369 | 003 | Jun 18, 2007 |
| AP | | EQ 2GM BASE/VIAL | A202268 | 002 | Jul 30, 2012 |
| AP | QILU PHARM CO LTD | EQ 500MG BASE/VIAL | A203704 | 001 | Feb 01, 2016 |
| AP | | EQ 1GM BASE/VIAL | A203704 | 002 | Feb 01, 2016 |
| AP | | EQ 2GM BASE/VIAL | A203704 | 003 | Feb 01, 2016 |
| AP | SAGENT PHARMS | EQ 1GM BASE/VIAL | A091048 | 001 | Jan 04, 2017 |
| AP | | EQ 2GM BASE/VIAL | A091048 | 002 | Jan 04, 2017 |

MAXIPIME

| | | | | | |
|-----------|----------------|---------------------------|----------------|------------|--------------|
| AP | +! HOSPIRA INC | EQ 500MG BASE/VIAL | N050679 | 001 | Jan 18, 1996 |
| AP | +! | EQ 1GM BASE/VIAL | N050679 | 002 | Jan 18, 1996 |
| AP | +! | EQ 2GM BASE/VIAL | N050679 | 003 | Jan 18, 1996 |

CEFEPIME AND DEXTROSE IN DUPLEX CONTAINER

| | | | | |
|---------|------------------|---------|-----|--------------|
| B BRAUN | EQ 1GM BASE/VIAL | N050821 | 001 | May 06, 2010 |
| | EQ 2GM BASE/VIAL | N050821 | 002 | May 06, 2010 |

CEFEPIME IN PLASTIC CONTAINER

| | | | | | |
|----|-----------------|-------------------------------------|---------|-----|--------------|
| +! | BAXTER HLTHCARE | EQ 1GM BASE/50ML (EQ 20MG BASE/ML) | N050817 | 001 | Aug 05, 2008 |
| +! | | EQ 2GM BASE/100ML (EQ 20MG BASE/ML) | N050817 | 002 | Aug 05, 2008 |

POWDER;INTRAVENOUS

CEFEPIME HYDROCHLORIDE IN PLASTIC CONTAINER

| | | | | |
|--------------|---------------|---------|-----|--------------|
| SAMSON MEDCL | EQ 100GM BASE | A209408 | 001 | Aug 21, 2018 |
|--------------|---------------|---------|-----|--------------|

CEFIXIME

CAPSULE;ORAL

CEFIXIME

| | | | | | |
|-----------|----------------|--------------|----------------|------------|--------------|
| AB | ALKEM LABS LTD | 400MG | A210574 | 001 | Oct 09, 2018 |
| AB | +! LUPIN LTD | 400MG | N203195 | 001 | Jun 01, 2012 |

FOR SUSPENSION;ORAL

CEFIXIME

| | | | | | |
|-----------|----------------------|------------------|----------------|------------|--------------|
| AB | AUROBINDO PHARMA LTD | 100MG/5ML | A204835 | 001 | Apr 14, 2015 |
| AB | | 200MG/5ML | A204835 | 002 | Apr 14, 2015 |
| AB | BELCHER PHARMS LLC | 100MG/5ML | A206938 | 001 | Feb 06, 2017 |
| AB | | 200MG/5ML | A206938 | 002 | Feb 06, 2017 |
| AB | | 500MG/5ML | A206939 | 001 | Feb 06, 2017 |
| AB | SANDOZ INC | 100MG/5ML | A206144 | 001 | Nov 17, 2017 |
| AB | | 200MG/5ML | A206144 | 002 | Nov 17, 2017 |

SUPRAX

| | | | | | |
|-----------|--------------|------------------|----------------|------------|--------------|
| AB | +! LUPIN LTD | 500MG/5ML | N202091 | 001 | Feb 20, 2013 |
|-----------|--------------|------------------|----------------|------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

CEFIXIME

FOR SUSPENSION; ORAL

SUPRAX

| | | | | |
|------------------------|--------------|------------------|--------------------|--------------|
| <u>AB</u> | LUPIN PHARMS | <u>100MG/5ML</u> | <u>A065129 001</u> | Feb 23, 2004 |
| <u>AB</u> | | <u>200MG/5ML</u> | <u>A065355 001</u> | Apr 10, 2007 |
| TABLET; ORAL | | | | |
| SUPRAX | | | | |
| ! | LUPIN PHARMS | 400MG | A065130 001 | Feb 12, 2004 |
| TABLET, CHEWABLE; ORAL | | | | |
| SUPRAX | | | | |
| | LUPIN LTD | 100MG | A065380 001 | Oct 25, 2010 |
| | | 150MG | A065380 002 | Oct 25, 2010 |
| ! | | 200MG | A065380 003 | Oct 25, 2010 |

CEFOTAXIME SODIUM

INJECTABLE; INJECTION

CEFOTAXIME

| | | | | | |
|-----------|---|-----------|---------------------------|--------------------|--------------|
| <u>AP</u> | ! | HIKMA | <u>EQ 500MG BASE/VIAL</u> | <u>A065072 001</u> | Nov 20, 2002 |
| <u>AP</u> | ! | | <u>EQ 1GM BASE/VIAL</u> | <u>A065072 002</u> | Nov 20, 2002 |
| <u>AP</u> | ! | | <u>EQ 2GM BASE/VIAL</u> | <u>A065072 003</u> | Nov 20, 2002 |
| <u>AP</u> | ! | | <u>EQ 10GM BASE/VIAL</u> | <u>A065071 001</u> | Nov 20, 2002 |
| <u>AP</u> | | WOCKHARDT | <u>EQ 1GM BASE/VIAL</u> | <u>A065197 001</u> | Aug 29, 2006 |

CEFOTAXIME SODIUM

| | | | | | |
|-----------|--|-------------|---------------------------|--------------------|--------------|
| <u>AP</u> | | HOSPIRA INC | <u>EQ 500MG BASE/VIAL</u> | <u>A065290 001</u> | Aug 11, 2006 |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A065290 002</u> | Aug 11, 2006 |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A065293 001</u> | Aug 10, 2006 |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A203132 001</u> | Feb 19, 2016 |
| <u>AP</u> | | | <u>EQ 2GM BASE/VIAL</u> | <u>A065290 003</u> | Aug 11, 2006 |
| <u>AP</u> | | | <u>EQ 2GM BASE/VIAL</u> | <u>A065293 002</u> | Aug 10, 2006 |
| <u>AP</u> | | | <u>EQ 2GM BASE/VIAL</u> | <u>A203132 002</u> | Feb 19, 2016 |
| <u>AP</u> | | | <u>EQ 10GM BASE/VIAL</u> | <u>A065292 001</u> | Aug 10, 2006 |
| <u>AP</u> | | WOCKHARDT | <u>EQ 500MG BASE/VIAL</u> | <u>A065197 002</u> | Jun 20, 2008 |
| <u>AP</u> | | | <u>EQ 2GM BASE/VIAL</u> | <u>A065197 003</u> | Jun 20, 2008 |

CEFOTETAN DISODIUM

INJECTABLE; INJECTION

CEFOTAN

| | | | | | |
|-----------|---|----------|-------------------------|--------------------|--------------|
| <u>AP</u> | + | TELIGENT | <u>EQ 1GM BASE/VIAL</u> | <u>N050588 001</u> | Dec 27, 1985 |
| <u>AP</u> | + | | <u>EQ 2GM BASE/VIAL</u> | <u>N050588 002</u> | Dec 27, 1985 |

CEFOTETAN

| | | | | | |
|-----------|---|-------------------------|--------------------------|--------------------|--------------|
| <u>AP</u> | ! | FRESENIUS KABI USA | <u>EQ 1GM BASE/VIAL</u> | <u>A065374 001</u> | Aug 09, 2007 |
| <u>AP</u> | ! | | <u>EQ 2GM BASE/VIAL</u> | <u>A065374 002</u> | Aug 09, 2007 |
| <u>AP</u> | ! | | <u>EQ 10GM BASE/VIAL</u> | <u>A065375 001</u> | Aug 09, 2007 |
| <u>AP</u> | | HIKMA FARMACEUTICA | <u>EQ 1GM BASE/VIAL</u> | <u>A091031 001</u> | Oct 26, 2011 |
| <u>AP</u> | | | <u>EQ 2GM BASE/VIAL</u> | <u>A091031 002</u> | Oct 26, 2011 |
| <u>AP</u> | | WEST-WARD PHARM CORP | <u>EQ 10GM BASE/VIAL</u> | <u>A091030 001</u> | Oct 26, 2011 |

CEFOTETAN AND DEXTROSE IN DUPLEX CONTAINER

| | | | | | |
|---|---|---------|------------------|-------------|--------------|
| + | ! | B BRAUN | EQ 1GM BASE/VIAL | N065430 001 | Aug 09, 2007 |
| + | ! | | EQ 2GM BASE/VIAL | N065430 002 | Aug 09, 2007 |

CEFOXITIN SODIUM

INJECTABLE; INJECTION

CEFOXITIN

| | | | | | | |
|---------------------------------------------------|---|-------------------------|--------------------------|-------------------------|--------------------|--------------|
| <u>AP</u> | ! | ACS DOBFAR | <u>EQ 1GM BASE/VIAL</u> | <u>A065414 001</u> | Jun 12, 2009 | |
| <u>AP</u> | ! | | <u>EQ 2GM BASE/VIAL</u> | <u>A065414 002</u> | Jun 12, 2009 | |
| <u>AP</u> | ! | | <u>EQ 10GM BASE/VIAL</u> | <u>A065415 001</u> | May 19, 2010 | |
| <u>AP</u> | | HIKMA FARMACEUTICA | <u>EQ 1GM BASE/VIAL</u> | <u>A065238 001</u> | Mar 12, 2010 | |
| <u>AP</u> | | | <u>EQ 2GM BASE/VIAL</u> | <u>A065238 002</u> | Mar 12, 2010 | |
| <u>AP</u> | | | <u>EQ 10GM BASE/VIAL</u> | <u>A065239 001</u> | Mar 02, 2010 | |
| <u>AP</u> | | HOSPIRA INC | <u>EQ 1GM BASE/VIAL</u> | <u>A065313 001</u> | Jan 23, 2006 | |
| <u>AP</u> | | | <u>EQ 2GM BASE/VIAL</u> | <u>A065313 002</u> | Jan 23, 2006 | |
| <u>AP</u> | | | <u>EQ 10GM BASE/VIAL</u> | <u>A065312 001</u> | Feb 13, 2006 | |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>EQ 1GM BASE/VIAL</u> | <u>A065051 001</u> | Sep 11, 2000 | |
| <u>AP</u> | | | <u>EQ 2GM BASE/VIAL</u> | <u>A065051 002</u> | Sep 11, 2000 | |
| <u>AP</u> | | | <u>EQ 10GM BASE/VIAL</u> | <u>A065050 001</u> | Sep 11, 2000 | |
| <u>CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER</u> | | | | | | |
| <u>AP</u> | + | ! | B BRAUN | <u>EQ 1GM BASE/VIAL</u> | <u>N065214 001</u> | Mar 10, 2006 |
| <u>AP</u> | + | ! | | <u>EQ 2GM BASE/VIAL</u> | <u>N065214 002</u> | Mar 10, 2006 |

MEFOXIN IN PLASTIC CONTAINER

| | | | | | |
|---|--|---------------------|-----------------|-------------|--------------|
| ! | | MYLAN INSTITUTIONAL | EQ 20MG BASE/ML | A063182 001 | Jan 25, 1993 |
| ! | | | EQ 40MG BASE/ML | A063182 002 | Jan 25, 1993 |

PRESCRIPTION DRUG PRODUCT LIST

CEFOXITIN SODIUM

POWDER; INTRAVENOUS

CEFOXITIN IN PLASTIC CONTAINER

SAMSON MEDCL

EQ 100GM BASE

A200938 001 Nov 16, 2015

CEFPODOXIME PROXETIL

FOR SUSPENSION; ORAL

CEFPODOXIME PROXETILAB AUROBINDO PHARMA
LTDEQ 50MG BASE/5MLA065409 001 Jun 08, 2007AB EQ 100MG BASE/5ML A065409 002 Jun 08, 2007AB SANDOZ EQ 50MG BASE/5ML A090031 001 Jan 14, 2009AB ! EQ 100MG BASE/5ML A090031 002 Jan 14, 2009

TABLET; ORAL

CEFPODOXIME PROXETILAB AUROBINDO PHARMAEQ 100MG BASEA065370 001 Jun 11, 2007AB EQ 200MG BASE A065370 002 Jun 11, 2007AB ORCHID HLTHCARE EQ 100MG BASE A065388 001 Nov 14, 2007AB EQ 200MG BASE A065388 002 Nov 14, 2007AB SANDOZ EQ 100MG BASE A065462 001 May 28, 2008AB ! EQ 200MG BASE A065462 002 May 28, 2008CEFPROZIL

FOR SUSPENSION; ORAL

CEFPROZILAB APOTEX INC125MG/5MLA065351 001 Feb 29, 2012AB 250MG/5ML A065351 002 Feb 29, 2012AB AUROBINDO PHARMA 125MG/5ML A065381 001 Jan 30, 2007AB 250MG/5ML A065381 002 Jan 30, 2007AB LUPIN 125MG/5ML A065261 001 Dec 19, 2005AB ! 250MG/5ML A065261 002 Dec 19, 2005AB ORCHID HLTHCARE 125MG/5ML A065284 002 Dec 30, 2005AB 250MG/5ML A065284 001 Dec 30, 2005AB SANDOZ 125MG/5ML A065257 001 Dec 08, 2005AB 250MG/5ML A065257 002 Dec 08, 2005AB TEVA PHARMS 125MG/5ML A065236 001 Dec 08, 2005AB 250MG/5ML A065236 002 Dec 08, 2005

TABLET; ORAL

CEFPROZILAB APOTEX INC250MGA065327 001 Mar 26, 2008AB 500MG A065327 002 Mar 26, 2008AB AUROBINDO PHARMA 250MG A065340 001 May 24, 2007AB LTD 500MG A065340 002 May 24, 2007AB CASI PHARMS INC 250MG A065235 001 Nov 14, 2005AB 500MG A065235 002 Nov 14, 2005AB LUPIN 250MG A065276 001 Dec 08, 2005AB ! 500MG A065276 002 Dec 08, 2005AB ORCHID HLTHCARE 250MG A065267 001 Dec 19, 2005AB 500MG A065267 002 Dec 19, 2005AB TEVA 250MG A065208 001 Dec 06, 2005AB 500MG A065208 002 Dec 06, 2005AB WOCKHARDT 250MG A065428 001 Jun 14, 2007AB 500MG A065428 002 Jun 14, 2007CEFTAROLINE FOSAMIL

POWDER; INTRAVENOUS

TEFLARO

+ ALLERGAN SALES LLC

400MG/VIAL

N200327 001 Oct 29, 2010

+!

600MG/VIAL

N200327 002 Oct 29, 2010

CEFTAZIDIME

INJECTABLE; INJECTION

CEFTAZIDIMEAP ACS DOBFAR1GM/VIALA062640 002 Nov 20, 1985AP 2GM/VIAL A062640 003 Nov 20, 1985AP 6GM/VIAL A062640 004 Feb 03, 1992AP WOCKHARDT 1GM/VIAL A065196 001 Oct 15, 2008FORTAZAP +! TELIGENT500MG/VIALN050578 001 Jul 19, 1985AP +! 1GM/VIAL N050578 002 Jul 19, 1985AP +! 2GM/VIAL N050578 003 Jul 19, 1985AP +! 6GM/VIAL N050578 004 Jul 19, 1985

PRESCRIPTION DRUG PRODUCT LIST

CEFTAZIDIME

INJECTABLE; INJECTION

TAZICEF

| | | | | |
|-----------|-----------------------------------|-------------------|--------------------|--------------|
| <u>AP</u> | HOSPIRA | <u>500MG/VIAL</u> | <u>A062662 001</u> | Mar 06, 1986 |
| <u>AP</u> | | <u>1GM/VIAL</u> | <u>A062662 002</u> | Mar 06, 1986 |
| <u>AP</u> | | <u>1GM/VIAL</u> | <u>A064032 001</u> | Oct 31, 1993 |
| <u>AP</u> | | <u>2GM/VIAL</u> | <u>A062662 003</u> | Mar 06, 1986 |
| <u>AP</u> | | <u>2GM/VIAL</u> | <u>A064032 002</u> | Oct 31, 1993 |
| <u>AP</u> | | <u>6GM/VIAL</u> | <u>A062662 004</u> | Mar 06, 1986 |
| | CEFTAZIDIME IN DEXTROSE CONTAINER | | | |
| | + B BRAUN | EQ 1GM BASE | N050823 001 | Jun 13, 2011 |
| | +! | EQ 2GM BASE | N050823 002 | Jun 13, 2011 |

CEFTOLOZANE SULFATE; TAZOBACTAM SODIUM

POWDER; INTRAVENOUS

ZERBAXA

| | | | | |
|--|----------------------|-------------------------------------|-------------|--------------|
| | +! CUBIST PHARMS LLC | EQ 1GM BASE/VIAL;EQ 0.5GM BASE/VIAL | N206829 001 | Dec 19, 2014 |
|--|----------------------|-------------------------------------|-------------|--------------|

CEFTRIAZONE SODIUM

INJECTABLE; INJECTION

CEFTRIAZONE

| | | | | |
|-----------|-----------------------------------------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | ACS DOBFAR | <u>EQ 500MG BASE/VIAL</u> | <u>A065329 001</u> | Jul 24, 2008 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A065329 002</u> | Jul 24, 2008 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A065329 003</u> | Jul 24, 2008 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A065328 001</u> | Jul 24, 2008 |
| <u>AP</u> | HOSPIRA INC | <u>EQ 10GM BASE/VIAL</u> | <u>A065232 001</u> | Aug 02, 2005 |
| <u>AP</u> | QILU PHARM CO LTD | <u>EQ 10GM BASE/VIAL</u> | <u>A209218 001</u> | Oct 17, 2018 |
| <u>AP</u> | ! SANDOZ | <u>EQ 10GM BASE/VIAL</u> | <u>A065168 001</u> | May 17, 2005 |
| <u>AP</u> | ! SANDOZ INC | <u>EQ 1GM BASE/VIAL</u> | <u>A065204 001</u> | May 03, 2005 |
| <u>AP</u> | ! SANDOZ INC | <u>EQ 2GM BASE/VIAL</u> | <u>A065204 002</u> | May 03, 2005 |
| <u>AP</u> | WOCKHARDT | <u>EQ 1GM BASE/VIAL</u> | <u>A065180 001</u> | May 12, 2006 |
| | <u>CEFTRIAZONE AND DEXTROSE IN DUPLIX CONTAINER</u> | | | |
| <u>AP</u> | +! B BRAUN | <u>EQ 1GM BASE/VIAL</u> | <u>N050796 001</u> | Apr 20, 2005 |
| <u>AP</u> | +! | <u>EQ 2GM BASE/VIAL</u> | <u>N050796 002</u> | Apr 20, 2005 |
| | <u>CEFTRIAZONE SODIUM</u> | | | |
| <u>AP</u> | ASTRAL | <u>EQ 10GM BASE/VIAL</u> | <u>A091117 001</u> | Jan 20, 2017 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>EQ 10GM BASE/VIAL</u> | <u>A090701 001</u> | Oct 04, 2017 |
| | CEFTRIAZONE | | | |
| | SAMSON MEDCL | EQ 100GM BASE/VIAL | A090057 001 | Apr 25, 2014 |
| | CEFTRIAZONE IN PLASTIC CONTAINER | | | |
| | ! BAXTER HLTHCARE | EQ 20MG BASE/ML | A065224 001 | Aug 23, 2005 |
| | ! | EQ 40MG BASE/ML | A065224 002 | Aug 23, 2005 |

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFTRIAZONE

| | | | | |
|-----------|--------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | AKORN INC | <u>EQ 250MG BASE/VIAL</u> | <u>A065305 001</u> | Jan 11, 2008 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A065305 002</u> | Jan 11, 2008 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A065305 003</u> | Jan 11, 2008 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A065305 004</u> | Jan 11, 2008 |
| <u>AP</u> | ASTRAL | <u>EQ 250MG BASE/VIAL</u> | <u>A091049 001</u> | Jun 11, 2018 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A091049 002</u> | Jun 11, 2018 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A091049 003</u> | Jun 11, 2018 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A091049 004</u> | Jun 11, 2018 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>EQ 250MG BASE/VIAL</u> | <u>A065342 001</u> | Jan 10, 2008 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A065342 002</u> | Jan 10, 2008 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A065342 003</u> | Jan 10, 2008 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A065342 004</u> | Jan 10, 2008 |
| <u>AP</u> | HOSPIRA INC | <u>EQ 250MG BASE/VIAL</u> | <u>A065230 001</u> | Aug 02, 2005 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A065230 002</u> | Aug 02, 2005 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A065230 003</u> | Aug 02, 2005 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A065230 004</u> | Aug 02, 2005 |
| <u>AP</u> | LUPIN | <u>EQ 250MG BASE/VIAL</u> | <u>A065125 001</u> | Sep 30, 2003 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A065125 002</u> | Sep 30, 2003 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A065125 003</u> | Sep 30, 2003 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A065125 004</u> | Sep 30, 2003 |
| <u>AP</u> | QILU PHARM CO LTD | <u>EQ 250MG BASE/VIAL</u> | <u>A203702 001</u> | Jun 29, 2016 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A203702 002</u> | Jun 29, 2016 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A203702 003</u> | Jun 29, 2016 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A203702 004</u> | Jun 29, 2016 |
| <u>AP</u> | ! SANDOZ | <u>EQ 250MG BASE/VIAL</u> | <u>A065169 001</u> | May 09, 2005 |
| <u>AP</u> | ! SANDOZ | <u>EQ 500MG BASE/VIAL</u> | <u>A065169 002</u> | May 09, 2005 |
| <u>AP</u> | ! SANDOZ | <u>EQ 1GM BASE/VIAL</u> | <u>A065169 003</u> | May 09, 2005 |
| <u>AP</u> | ! SANDOZ | <u>EQ 2GM BASE/VIAL</u> | <u>A065169 004</u> | May 09, 2005 |
| <u>AP</u> | WOCKHARDT | <u>EQ 250MG BASE/VIAL</u> | <u>A065391 001</u> | Apr 12, 2007 |

PRESCRIPTION DRUG PRODUCT LIST

CEFTRIAZONE SODIUM

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFTRIAZONE

| | | | | |
|-----------|--|---------------------------|--------------------|--------------|
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A065391 002</u> | Apr 12, 2007 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A065391 003</u> | Apr 12, 2007 |

CEFUROXIME AXETIL

TABLET; ORAL

CEFUROXIME AXETIL

| | | | | |
|-----------|----------------------|----------------------|--------------------|--------------|
| <u>AB</u> | ALKEM LABS LTD | <u>EQ 250MG BASE</u> | <u>A065496 001</u> | Jun 07, 2010 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065496 002</u> | Jun 07, 2010 |
| <u>AB</u> | ANI PHARMS INC | <u>EQ 250MG BASE</u> | <u>A065190 001</u> | Oct 18, 2004 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065190 002</u> | Oct 18, 2004 |
| <u>AB</u> | APOTEX | <u>EQ 250MG BASE</u> | <u>A065069 001</u> | Oct 02, 2002 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065069 002</u> | Oct 02, 2002 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 125MG BASE</u> | <u>A065308 001</u> | Mar 29, 2006 |
| <u>AB</u> | | <u>EQ 250MG BASE</u> | <u>A065308 002</u> | Mar 29, 2006 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065308 003</u> | Mar 29, 2006 |
| <u>AB</u> | LUPIN | <u>EQ 250MG BASE</u> | <u>A065135 001</u> | Jul 25, 2003 |
| <u>AB</u> | ! | <u>EQ 500MG BASE</u> | <u>A065135 002</u> | Jul 25, 2003 |
| <u>AB</u> | ORCHID HLTHCARE | <u>EQ 125MG BASE</u> | <u>A065359 001</u> | Feb 15, 2008 |
| <u>AB</u> | | <u>EQ 250MG BASE</u> | <u>A065359 002</u> | Feb 15, 2008 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065359 003</u> | Feb 15, 2008 |
| <u>AB</u> | WOCKHARDT | <u>EQ 125MG BASE</u> | <u>A065166 001</u> | Jul 29, 2005 |
| <u>AB</u> | | <u>EQ 250MG BASE</u> | <u>A065166 002</u> | Jul 29, 2005 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065166 003</u> | Jul 29, 2005 |

CEFUROXIME SODIUM

INJECTABLE; INJECTION

CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER

| | | | | |
|-----------|------------|---------------------------|--------------------|--------------|
| <u>AP</u> | +! B BRAUN | <u>EQ 750MG BASE/VIAL</u> | <u>N050780 001</u> | Feb 21, 2001 |
| <u>AP</u> | +! | <u>EQ 1.5GM BASE/VIAL</u> | <u>N050780 002</u> | Feb 21, 2001 |

CEFUROXIME SODIUM

| | | | | |
|-----------|--------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | ACS DOBFAR SPA | <u>EQ 1.5GM BASE/VIAL</u> | <u>A064125 002</u> | May 30, 1997 |
| <u>AP</u> | | <u>EQ 7.5GM BASE/VIAL</u> | <u>A064124 001</u> | May 30, 1997 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>EQ 1.5GM BASE/VIAL</u> | <u>A065048 002</u> | Jan 09, 2004 |
| <u>AP</u> | | <u>EQ 7.5GM BASE/VIAL</u> | <u>A065046 001</u> | Jan 09, 2004 |
| <u>AP</u> | HOSPIRA INC | <u>EQ 1.5GM BASE/VIAL</u> | <u>A065483 002</u> | Oct 15, 2008 |
| <u>AP</u> | | <u>EQ 1.5GM BASE/VIAL</u> | <u>A065503 001</u> | Oct 15, 2008 |
| <u>AP</u> | | <u>EQ 7.5GM BASE/VIAL</u> | <u>A065484 001</u> | Oct 15, 2008 |

ZINACEF

| | | | | |
|-----------|-------------|---------------------------|--------------------|--------------|
| <u>AP</u> | +! TELIGENT | <u>EQ 1.5GM BASE/VIAL</u> | <u>N050558 003</u> | Oct 19, 1983 |
| <u>AP</u> | +! | <u>EQ 7.5GM BASE/VIAL</u> | <u>N050558 004</u> | Oct 23, 1986 |

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFUROXIME SODIUM

| | | | | |
|-----------|--------------------|---------------------------|--------------------|--------------|
| <u>AB</u> | ACS DOBFAR SPA | <u>EQ 750MG BASE/VIAL</u> | <u>A064125 001</u> | May 30, 1997 |
| <u>AB</u> | HIKMA FARMACEUTICA | <u>EQ 750MG BASE/VIAL</u> | <u>A065048 001</u> | Jan 09, 2004 |

ZINACEF

| | | | | |
|-----------|-------------|---------------------------|--------------------|--------------|
| <u>AB</u> | +! TELIGENT | <u>EQ 750MG BASE/VIAL</u> | <u>N050558 002</u> | Oct 19, 1983 |
|-----------|-------------|---------------------------|--------------------|--------------|

CEFUROXIME SODIUM

| | | | | |
|-----------|-------------|---------------------------|--------------------|--------------|
| <u>AP</u> | HOSPIRA INC | <u>EQ 750MG BASE/VIAL</u> | <u>A065483 001</u> | Oct 15, 2008 |
|-----------|-------------|---------------------------|--------------------|--------------|

CELECOXIB

CAPSULE; ORAL

CELEBREX

| | | | | |
|-----------|-------------|--------------|--------------------|--------------|
| <u>AB</u> | + GD SEARLE | <u>50MG</u> | <u>N020998 004</u> | Dec 15, 2006 |
| <u>AB</u> | + | <u>100MG</u> | <u>N020998 001</u> | Dec 31, 1998 |
| <u>AB</u> | + | <u>200MG</u> | <u>N020998 002</u> | Dec 31, 1998 |
| <u>AB</u> | +! | <u>400MG</u> | <u>N020998 003</u> | Aug 29, 2002 |

CELECOXIB

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| <u>AB</u> | ALEMBIC PHARMS LTD | <u>50MG</u> | <u>A204519 001</u> | Aug 21, 2015 |
| <u>AB</u> | | <u>100MG</u> | <u>A204519 002</u> | Aug 21, 2015 |
| <u>AB</u> | | <u>200MG</u> | <u>A204519 003</u> | Aug 21, 2015 |
| <u>AB</u> | | <u>400MG</u> | <u>A204519 004</u> | Aug 21, 2015 |
| <u>AB</u> | AMNEAL PHARMS | <u>50MG</u> | <u>A208833 001</u> | May 31, 2018 |
| <u>AB</u> | | <u>100MG</u> | <u>A208833 002</u> | May 31, 2018 |
| <u>AB</u> | | <u>200MG</u> | <u>A208833 003</u> | May 31, 2018 |
| <u>AB</u> | | <u>400MG</u> | <u>A208833 004</u> | May 31, 2018 |
| <u>AB</u> | APOTEX INC | <u>50MG</u> | <u>A204197 001</u> | Jun 02, 2015 |
| <u>AB</u> | | <u>100MG</u> | <u>A204197 002</u> | Jun 02, 2015 |
| <u>AB</u> | | <u>200MG</u> | <u>A204197 003</u> | Jun 02, 2015 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>50MG</u> | <u>A206827 001</u> | Feb 01, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

CELECOXIB

CAPSULE; ORAL

CELECOXIB

| | | | | |
|-----------|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | | <u>100MG</u> | <u>A206827 002</u> | Feb 01, 2016 |
| <u>AB</u> | | <u>200MG</u> | <u>A206827 003</u> | Feb 01, 2016 |
| <u>AB</u> | | <u>400MG</u> | <u>A206827 004</u> | Feb 01, 2016 |
| <u>AB</u> | CIPLA | <u>50MG</u> | <u>A207446 001</u> | Sep 23, 2015 |
| <u>AB</u> | | <u>100MG</u> | <u>A207446 002</u> | Sep 23, 2015 |
| <u>AB</u> | | <u>200MG</u> | <u>A207446 003</u> | Sep 23, 2015 |
| <u>AB</u> | | <u>400MG</u> | <u>A207446 004</u> | Sep 23, 2015 |
| <u>AB</u> | CSPC OUYI PHARM CO | <u>50MG</u> | <u>A210071 001</u> | Jan 23, 2018 |
| <u>AB</u> | | <u>100MG</u> | <u>A210071 002</u> | Jan 23, 2018 |
| <u>AB</u> | | <u>200MG</u> | <u>A210071 003</u> | Jan 23, 2018 |
| <u>AB</u> | JUBILANT GENERICS | <u>50MG</u> | <u>A207061 001</u> | Apr 04, 2017 |
| <u>AB</u> | | <u>100MG</u> | <u>A207061 002</u> | Apr 04, 2017 |
| <u>AB</u> | | <u>200MG</u> | <u>A207061 003</u> | Apr 04, 2017 |
| <u>AB</u> | | <u>400MG</u> | <u>A207061 004</u> | Apr 04, 2017 |
| <u>AB</u> | LUPIN LTD | <u>50MG</u> | <u>A202240 001</u> | Oct 29, 2014 |
| <u>AB</u> | | <u>100MG</u> | <u>A202240 002</u> | Jun 09, 2015 |
| <u>AB</u> | | <u>200MG</u> | <u>A202240 003</u> | Jun 09, 2015 |
| <u>AB</u> | | <u>400MG</u> | <u>A202240 004</u> | Jun 09, 2015 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>50MG</u> | <u>A204590 001</u> | Mar 16, 2016 |
| <u>AB</u> | | <u>100MG</u> | <u>A204590 002</u> | Mar 16, 2016 |
| <u>AB</u> | | <u>200MG</u> | <u>A204590 003</u> | Mar 16, 2016 |
| <u>AB</u> | | <u>400MG</u> | <u>A204590 004</u> | Mar 16, 2016 |
| <u>AB</u> | MICRO LABS | <u>50MG</u> | <u>A204776 001</u> | Apr 30, 2018 |
| <u>AB</u> | | <u>100MG</u> | <u>A204776 002</u> | Apr 30, 2018 |
| <u>AB</u> | | <u>200MG</u> | <u>A204776 003</u> | Apr 30, 2018 |
| <u>AB</u> | | <u>400MG</u> | <u>A204776 004</u> | Apr 30, 2018 |
| <u>AB</u> | MYLAN PHARMS INC | <u>50MG</u> | <u>A078857 001</u> | May 30, 2014 |
| <u>AB</u> | | <u>100MG</u> | <u>A078857 002</u> | Feb 11, 2015 |
| <u>AB</u> | | <u>200MG</u> | <u>A078857 003</u> | Feb 11, 2015 |
| <u>AB</u> | | <u>400MG</u> | <u>A078857 004</u> | Feb 11, 2015 |
| <u>AB</u> | TEVA | <u>50MG</u> | <u>A076898 001</u> | May 30, 2014 |
| <u>AB</u> | | <u>100MG</u> | <u>A076898 002</u> | May 30, 2014 |
| <u>AB</u> | | <u>200MG</u> | <u>A076898 003</u> | May 30, 2014 |
| <u>AB</u> | | <u>400MG</u> | <u>A076898 004</u> | May 30, 2014 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>50MG</u> | <u>A207677 001</u> | Dec 23, 2015 |
| <u>AB</u> | | <u>100MG</u> | <u>A207677 002</u> | Dec 23, 2015 |
| <u>AB</u> | | <u>200MG</u> | <u>A207677 003</u> | Dec 23, 2015 |
| <u>AB</u> | | <u>400MG</u> | <u>A207677 004</u> | Dec 23, 2015 |
| <u>AB</u> | WATSON LABS INC | <u>50MG</u> | <u>A200562 001</u> | Feb 11, 2015 |
| <u>AB</u> | | <u>100MG</u> | <u>A200562 002</u> | Feb 11, 2015 |
| <u>AB</u> | | <u>200MG</u> | <u>A200562 003</u> | Feb 11, 2015 |
| <u>AB</u> | | <u>400MG</u> | <u>A200562 004</u> | Feb 11, 2015 |

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

| | | | | |
|---------------|----------------------|----------------------|----------------------|---------------------------------|
| <u>AB</u> | ALKEM LABS LTD | <u>EQ 250MG BASE</u> | <u>A090836 001</u> | Dec 20, 2010 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A090836 002</u> | Dec 20, 2010 |
| <u>AB</u> | | <u>EQ 750MG BASE</u> | <u>A090836 004</u> | Mar 29, 2013 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 250MG BASE</u> | <u>A065253 001</u> | Nov 16, 2005 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065253 002</u> | Nov 16, 2005 |
| <u>AB</u> | BELCHER PHARMS | <u>EQ 250MG BASE</u> | <u>A062713 001</u> | Jul 15, 1988 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A062713 002</u> | Jul 15, 1988 |
| <u>AB</u> | HIKMA | <u>EQ 250MG BASE</u> | <u>A065215 001</u> | Jan 24, 2006 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065215 002</u> | Jan 24, 2006 |
| <u>AB</u> | LUPIN | <u>EQ 250MG BASE</u> | <u>A065229 001</u> | Nov 25, 2005 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065229 002</u> | Nov 25, 2005 |
| <u>AB</u> | ORCHID HLTHCARE | <u>EQ 250MG BASE</u> | <u>A065248 001</u> | Jun 28, 2005 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065248 002</u> | Jun 28, 2005 |
| <u>AB</u> | SUN PHARM INDS (IN) | <u>EQ 250MG BASE</u> | <u>A062791 001</u> | Jun 11, 1987 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A062791 002</u> | Jun 11, 1987 |
| <u>AB</u> | TEVA | <u>EQ 250MG BASE</u> | <u>A062702 001</u> | Feb 13, 1987 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A062702 002</u> | Feb 13, 1987 |
| <u>AB</u> | YUNG SHIN PHARM | <u>EQ 250MG BASE</u> | <u>A065152 001</u> | Feb 24, 2005 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A065152 002</u> | Feb 24, 2005 |
| <u>KEFLEX</u> | | | | |
| <u>AB</u> | + | <u>PRAGMA</u> | <u>EQ 250MG BASE</u> | <u>N050405 002</u> |
| <u>AB</u> | + | | <u>EQ 500MG BASE</u> | <u>N050405 003</u> |
| <u>AB</u> | + | | <u>EQ 750MG BASE</u> | <u>N050405 005</u> May 12, 2006 |

PRESCRIPTION DRUG PRODUCT LIST

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

| | | | |
|----------------|---------------|-------------|--------------|
| ALKEM LABS LTD | EQ 333MG BASE | A090836 003 | Mar 29, 2013 |
|----------------|---------------|-------------|--------------|

FOR SUSPENSION; ORAL

CEPHALEXIN

| | | | | |
|-----------|-----------------|--------------------------|--------------------|--------------|
| AB | LUPIN | EQ 125MG BASE/5ML | A065234 001 | Aug 17, 2005 |
| AB | | EQ 250MG BASE/5ML | A065234 002 | Aug 17, 2005 |
| AB | ORCHID HLTHCARE | EQ 125MG BASE/5ML | A065326 001 | Jul 10, 2006 |
| AB | | EQ 250MG BASE/5ML | A065326 002 | Jul 10, 2006 |
| AB | TEVA | EQ 125MG BASE/5ML | A062703 001 | Feb 13, 1987 |
| AB | ! | EQ 250MG BASE/5ML | A062703 002 | Feb 13, 1987 |
| AB | YUNG SHIN PHARM | EQ 125MG BASE/5ML | A065336 001 | Jul 25, 2007 |
| AB | | EQ 250MG BASE/5ML | A065336 002 | Jul 25, 2007 |

TABLET; ORAL

CEPHALEXIN

| | | | |
|------|---------------|-------------|--------------|
| TEVA | EQ 250MG BASE | A063023 001 | Jan 12, 1989 |
|------|---------------|-------------|--------------|

| | | | |
|---|---------------|-------------|--------------|
| ! | EQ 500MG BASE | A063024 001 | Jan 12, 1989 |
|---|---------------|-------------|--------------|

CERITINIB

CAPSULE; ORAL

ZYKADIA

| | | | |
|--------------------|-------|-------------|--------------|
| +! NOVARTIS PHARMS | 150MG | N205755 001 | Apr 29, 2014 |
|--------------------|-------|-------------|--------------|

CORP

CETIRIZINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

ZERVIAE

| | | | |
|--------------------|---------------|-------------|--------------|
| +! EYEVANCE PHARMS | EQ 0.24% BASE | N208694 001 | May 30, 2017 |
|--------------------|---------------|-------------|--------------|

SYRUP; ORAL

CETIRIZINE HYDROCHLORIDE

| | | | | |
|-----------|--------------------|----------------|--------------------|--------------|
| AA | AMNEAL PHARMS | 5MG/5ML | A090766 001 | Oct 07, 2009 |
| AA | ANDA REPOSITORY | 5MG/5ML | A090191 001 | Nov 12, 2009 |
| AA | BIO PHARM INC | 5MG/5ML | A078870 001 | Apr 27, 2009 |
| AA | BRECKENRIDGE PHARM | 5MG/5ML | A078488 001 | Oct 06, 2008 |
| AA | LANNETT CO INC | 5MG/5ML | A078496 001 | Sep 25, 2009 |
| AA | | 5MG/5ML | A078876 001 | May 11, 2012 |
| AA | ! PERRIGO R AND D | 5MG/5ML | A078398 001 | Jun 17, 2008 |
| AA | TARO | 5MG/5ML | A076601 001 | Jun 20, 2008 |
| AA | TEVA PHARMS | 5MG/5ML | A077279 001 | May 27, 2008 |

CETRORELIX

INJECTABLE; INJECTION

CETROTIDE

| | | | |
|-------------------|-------------------|-------------|--------------|
| +! EMD SERONO INC | EQ 0.25MG BASE/ML | N021197 001 | Aug 11, 2000 |
|-------------------|-------------------|-------------|--------------|

CEVIMELINE HYDROCHLORIDE

CAPSULE; ORAL

CEVIMELINE HYDROCHLORIDE

| | | | | |
|-----------|------------------|-------------|--------------------|--------------|
| AB | NOVEL LABS INC | 30MG | A204746 001 | Dec 30, 2016 |
| AB | RISING PHARMS | 30MG | A203775 001 | Jun 04, 2014 |
| AB | WEST-WARD PHARMS | 30MG | A091591 001 | Jul 08, 2013 |

EVOXAC

| | | | | |
|-----------|-----------------------|-------------|--------------------|--------------|
| AB | +! DAIICHI SANKYO INC | 30MG | N020989 002 | Jan 11, 2000 |
|-----------|-----------------------|-------------|--------------------|--------------|

CHENODIOL

TABLET; ORAL

CHENODIOL

| | | | |
|-----------------|-------|-------------|--------------|
| ! NEXGEN PHARMA | 250MG | A091019 001 | Oct 22, 2009 |
|-----------------|-------|-------------|--------------|

CHLORAMBUCIL

TABLET; ORAL

LEUKERAN

| | | | |
|---------------------|-----|-------------|--|
| +! ASPEN GLOBAL INC | 2MG | N010669 002 | |
|---------------------|-----|-------------|--|

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE; INJECTION

CHLORAMPHENICOL SODIUM SUCCINATE

| | | | |
|----------------------|------------------|-------------|--------------|
| ! FRESENIUS KABI USA | EQ 1GM BASE/VIAL | A062365 001 | Aug 25, 1982 |
|----------------------|------------------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE

| | | | | | |
|----------------|--------------------|-------------|----------------|------------|--|
| <u>AB</u> | BARR | <u>5MG</u> | <u>A084768</u> | <u>001</u> | |
| <u>AB</u> | | <u>10MG</u> | <u>A083116</u> | <u>001</u> | |
| <u>AB</u> | | <u>25MG</u> | <u>A084769</u> | <u>001</u> | |
| <u>LIBRIUM</u> | | | | | |
| <u>AB</u> | VALEANT PHARM INTL | <u>5MG</u> | <u>A085461</u> | <u>001</u> | |
| <u>AB</u> | | <u>10MG</u> | <u>A085472</u> | <u>001</u> | |
| <u>AB</u> | ! | <u>25MG</u> | <u>A085475</u> | <u>001</u> | |

CHLORDIAZEPOXIDE HYDROCHLORIDE; CLIDINIUM BROMIDE

CAPSULE; ORAL

LIBRAX

+! VALEANT PHARMS 5MG; 2.5MG N012750 001

CHLORHEXIDINE GLUCONATE

SOLUTION; DENTAL

CHLORHEXIDINE GLUCONATE

| | | | | | |
|------------------|-------------------------|--------------|----------------|------------|--------------|
| <u>AT</u> | HI TECH PHARMA | <u>0.12%</u> | <u>A074356</u> | <u>001</u> | May 07, 1996 |
| <u>AT</u> | LYNE | <u>0.12%</u> | <u>A074291</u> | <u>001</u> | Dec 28, 1995 |
| <u>AT</u> | TEVA | <u>0.12%</u> | <u>A074522</u> | <u>001</u> | Dec 15, 1995 |
| <u>AT</u> | WOCKHARDT BIO AG | <u>0.12%</u> | <u>A075006</u> | <u>001</u> | Mar 03, 2004 |
| <u>AT</u> | XTRIMUM | <u>0.12%</u> | <u>A077789</u> | <u>001</u> | Jun 18, 2009 |
| <u>PAROEX</u> | | | | | |
| <u>AT</u> | SUNSTAR AMERICAS | <u>0.12%</u> | <u>A076434</u> | <u>001</u> | Nov 29, 2005 |
| <u>PERIDEX</u> | | | | | |
| <u>AT</u> | +! 3M | <u>0.12%</u> | <u>N019028</u> | <u>001</u> | Aug 13, 1986 |
| <u>PERIOGARD</u> | | | | | |
| <u>AT</u> | COLGATE PALMOLIVE CO | <u>0.12%</u> | <u>A073695</u> | <u>001</u> | Jan 14, 1994 |
| <u>AT</u> | COLGATE-PALMOLIVE CO | <u>0.12%</u> | <u>A203212</u> | <u>001</u> | Jan 28, 2016 |
| TABLET; DENTAL | | | | | |
| PERIOCHIP | | | | | |
| | +! DEXCEL PHARMA | 2.5MG | N020774 | 001 | May 15, 1998 |

CHLOROPROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHLOROPROCAINE HYDROCHLORIDE

| | | | | | |
|-----------------------|-------------------------|--------------------|----------------|------------|--------------|
| <u>AP</u> | HOSPIRA | <u>2%</u> | <u>A087447</u> | <u>001</u> | Apr 16, 1982 |
| <u>AP</u> | | <u>3%</u> | <u>A087446</u> | <u>001</u> | Apr 16, 1982 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>2%</u> | <u>A040273</u> | <u>001</u> | Sep 09, 1998 |
| <u>AP</u> | | <u>3%</u> | <u>A040273</u> | <u>002</u> | Sep 09, 1998 |
| <u>NESACAINE</u> | | | | | |
| <u>AP</u> | + FRESENIUS KABI USA | <u>2%</u> | <u>N009435</u> | <u>002</u> | |
| <u>NESACAINE-MPF</u> | | | | | |
| <u>AP</u> | +! FRESENIUS KABI USA | <u>2%</u> | <u>N009435</u> | <u>006</u> | May 02, 1996 |
| <u>AP</u> | +! | <u>3%</u> | <u>N009435</u> | <u>007</u> | May 02, 1996 |
| NESACAINE | | | | | |
| | +! FRESENIUS KABI USA | 1% | N009435 | 001 | |
| SOLUTION; INTRATHECAL | | | | | |
| CLOROTEKAL | | | | | |
| | + B BRAUN MEDICAL INC | 50MG/5ML (10MG/ML) | N208791 | 001 | Sep 26, 2017 |

CHLOROQUINE PHOSPHATE

TABLET; ORAL

CHLOROQUINE PHOSPHATE

| | | | | | |
|-----------|---|------------------|----------------------|----------------|------------|
| <u>AA</u> | ! | HIKMA PHARMS | <u>EQ 150MG BASE</u> | <u>A083082</u> | <u>001</u> |
| <u>AA</u> | | | <u>EQ 300MG BASE</u> | <u>A083082</u> | <u>002</u> |
| <u>AA</u> | | IPCA LABS LTD | <u>EQ 150MG BASE</u> | <u>A090610</u> | <u>001</u> |
| <u>AA</u> | | | <u>EQ 300MG BASE</u> | <u>A090249</u> | <u>001</u> |
| <u>AA</u> | | NATCO PHARMA LTD | <u>EQ 150MG BASE</u> | <u>A091621</u> | <u>001</u> |
| <u>AA</u> | ! | | <u>EQ 300MG BASE</u> | <u>A090612</u> | <u>001</u> |

CHLOROTHIAZIDE

SUSPENSION; ORAL

DIURIL

+! SALIX PHARMS 250MG/5ML N011870 001

TABLET; ORAL

CHLOROTHIAZIDE

! MYLAN 250MG A084217 002
500MG A084217 001

PRESCRIPTION DRUG PRODUCT LIST

CHLOROTHIAZIDE SODIUM

INJECTABLE; INJECTION

CHLOROTHIAZIDE SODIUM

| | | | | |
|-----------|---------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 500MG BASE/VIAL</u> | <u>A090896 001</u> | Oct 16, 2009 |
| <u>AP</u> | LUITPOLD | <u>EQ 500MG BASE/VIAL</u> | <u>A202561 001</u> | Apr 22, 2013 |
| <u>AP</u> | MYLAN INSTITUTIONAL | <u>EQ 500MG BASE/VIAL</u> | <u>A202493 001</u> | Jun 18, 2014 |
| <u>AP</u> | SAGENT PHARMS | <u>EQ 500MG BASE/VIAL</u> | <u>A202462 001</u> | May 29, 2015 |
| <u>AP</u> | SUN PHARMA GLOBAL | <u>EQ 500MG BASE/VIAL</u> | <u>A091546 001</u> | Jul 26, 2011 |

DIURIL

| | | | | |
|-----------|----|------------------|---------------------------|--------------------|
| <u>AP</u> | +! | OAK PHARMS AKORN | <u>EQ 500MG BASE/VIAL</u> | <u>N011145 005</u> |
|-----------|----|------------------|---------------------------|--------------------|

CHLORPHENIRAMINE MALEATE; CODEINE PHOSPHATE

TABLET, EXTENDED RELEASE; ORAL

TUXARIN ER

MAINPOINTE

8MG; 54.3MG

N206323 001 Jun 22, 2015

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND CHLORPHENIRAMINE MALEATE

| | | | | |
|-----------|-------------------|-------------------------|--------------------|--------------|
| <u>AA</u> | ACELLA PHARMS LLC | <u>4MG/5ML; 5MG/5ML</u> | <u>A206891 001</u> | Jun 09, 2017 |
|-----------|-------------------|-------------------------|--------------------|--------------|

VITUZ

| | | | | | |
|-----------|----|---------------|-------------------------|--------------------|--------------|
| <u>AA</u> | +! | CYPRESS PHARM | <u>4MG/5ML; 5MG/5ML</u> | <u>N204307 001</u> | Feb 20, 2013 |
|-----------|----|---------------|-------------------------|--------------------|--------------|

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

| | | | | |
|-----------|------------------|-----------------------------------|--------------------|--------------|
| <u>AA</u> | MAYNE PHARMA INC | <u>4MG/5ML; 5MG/5ML; 60MG/5ML</u> | <u>A205657 001</u> | Aug 03, 2015 |
|-----------|------------------|-----------------------------------|--------------------|--------------|

| | | | | |
|-----------|-------------|-----------------------------------|--------------------|--------------|
| <u>AA</u> | PADDOCK LLC | <u>4MG/5ML; 5MG/5ML; 60MG/5ML</u> | <u>A204627 001</u> | Apr 29, 2014 |
|-----------|-------------|-----------------------------------|--------------------|--------------|

ZUTRIPRO

| | | | | | |
|-----------|----|---------------|-----------------------------------|--------------------|--------------|
| <u>AA</u> | +! | CYPRESS PHARM | <u>4MG/5ML; 5MG/5ML; 60MG/5ML</u> | <u>N022439 001</u> | Jun 08, 2011 |
|-----------|----|---------------|-----------------------------------|--------------------|--------------|

CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL

TUZISTRA XR

+!

EQ 2.8MG BASE/5ML; EQ 14.7MG BASE/5ML

N207768 001 Apr 30, 2015

CHLORPHENIRAMINE POLISTIREX; HYDROCODONE POLISTIREX

CAPSULE, EXTENDED RELEASE; ORAL

TUSSICAPS

ECR PHARMA

EQ 4MG MALEATE; EQ 5MG BITARTRATE

A077273 002 Sep 24, 2007

!

EQ 8MG MALEATE; EQ 10MG BITARTRATE

A077273 001 Sep 24, 2007

SUSPENSION, EXTENDED RELEASE; ORAL

HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX

| | | | | |
|-----------|-----------------|---------------------------------------------------|--------------------|--------------|
| <u>AB</u> | TRIS PHARMA INC | <u>EQ 8MG MALEATE/5ML; EQ 10MG BITARTRATE/5ML</u> | <u>A091632 001</u> | Oct 01, 2010 |
|-----------|-----------------|---------------------------------------------------|--------------------|--------------|

HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX

| | | | | | |
|-----------|---|-----------------|---------------------------------------------------|--------------------|--------------|
| <u>AB</u> | ! | NEOS THERAP INC | <u>EQ 8MG MALEATE/5ML; EQ 10MG BITARTRATE/5ML</u> | <u>A091671 001</u> | Jun 29, 2012 |
|-----------|---|-----------------|---------------------------------------------------|--------------------|--------------|

CHLORPROMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHLORPROMAZINE HYDROCHLORIDE

!

WEST-WARD PHARMS 25MG/ML

A083329 001

INT

TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

| | | | | |
|-----------|------------------|--------------|--------------------|--------------|
| <u>AB</u> | AMNEAL PHARMS CO | <u>10MG</u> | <u>A209755 001</u> | Sep 10, 2018 |
| <u>AB</u> | | <u>25MG</u> | <u>A209755 002</u> | Sep 10, 2018 |
| <u>AB</u> | | <u>50MG</u> | <u>A209755 003</u> | Sep 10, 2018 |
| <u>AB</u> | | <u>100MG</u> | <u>A209755 004</u> | Sep 10, 2018 |
| <u>AB</u> | | <u>200MG</u> | <u>A209755 005</u> | Sep 10, 2018 |
| <u>AB</u> | USL PHARMA | <u>10MG</u> | <u>A083386 001</u> | |
| <u>AB</u> | ! | <u>25MG</u> | <u>A084112 001</u> | |
| <u>AB</u> | | <u>50MG</u> | <u>A084113 001</u> | |
| <u>AB</u> | ! | <u>100MG</u> | <u>A084114 001</u> | |
| <u>AB</u> | | <u>200MG</u> | <u>A084115 001</u> | |

CHLORPROPAMIDE

TABLET; ORAL

CHLORPROPAMIDE

MYLAN

100MG

A088549 002 Jun 01, 1984

!

250MG

A088549 001 Jun 01, 1984

PRESCRIPTION DRUG PRODUCT LIST

CHLORTHALIDONE

TABLET; ORAL

CHLORTHALIDONE

| | | | | | |
|-----------|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | APPCO PHARMA LLC | <u>25MG</u> | <u>A210742</u> | <u>001</u> | Oct 12, 2018 |
| <u>AB</u> | | <u>50MG</u> | <u>A210742</u> | <u>002</u> | Oct 12, 2018 |
| <u>AB</u> | MYLAN | <u>25MG</u> | <u>A086831</u> | <u>002</u> | |
| <u>AB</u> | ! | <u>50MG</u> | <u>A086831</u> | <u>001</u> | |
| <u>AB</u> | RICONPHARMA LLC | <u>25MG</u> | <u>A206904</u> | <u>001</u> | Mar 30, 2017 |
| <u>AB</u> | | <u>50MG</u> | <u>A206904</u> | <u>002</u> | Mar 30, 2017 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>25MG</u> | <u>A089286</u> | <u>002</u> | Jul 21, 1986 |
| <u>AB</u> | | <u>50MG</u> | <u>A089286</u> | <u>001</u> | Jul 21, 1986 |
| <u>AB</u> | UMEDICA LABS PVT LTD | <u>25MG</u> | <u>A207222</u> | <u>001</u> | May 24, 2018 |
| <u>AB</u> | | <u>50MG</u> | <u>A207222</u> | <u>002</u> | May 24, 2018 |

CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE

| | | | | | |
|---|-------------|-------|---------|-----|--------------|
| ! | MIKART | 250MG | A207483 | 001 | Jun 24, 2016 |
| | | 375MG | A040861 | 001 | Jun 01, 2010 |
| ! | | 750MG | A040861 | 002 | Jun 01, 2010 |
| ! | WATSON LABS | 500MG | A089859 | 001 | May 04, 1988 |

CHOLESTYRAMINE

POWDER; ORAL

CHOLESTYRAMINE

| | | | | | |
|-----------|-------------------------|------------------------------|----------------|------------|--------------|
| <u>AB</u> | ANI PHARMS INC | <u>EQ 4GM RESIN/PACKET</u> | <u>A074554</u> | <u>001</u> | Oct 02, 1996 |
| <u>AB</u> | | <u>EQ 4GM RESIN/SCOOPFUL</u> | <u>A074554</u> | <u>002</u> | Oct 02, 1996 |
| <u>AB</u> | PAR PHARM | <u>EQ 4GM RESIN/PACKET</u> | <u>A077204</u> | <u>001</u> | Aug 26, 2005 |
| <u>AB</u> | | <u>EQ 4GM RESIN/SCOOPFUL</u> | <u>A077204</u> | <u>002</u> | Aug 26, 2005 |
| <u>AB</u> | ! | <u>EQ 4GM RESIN/PACKET</u> | <u>A074557</u> | <u>001</u> | Aug 15, 1996 |
| <u>AB</u> | | <u>EQ 4GM RESIN/SCOOPFUL</u> | <u>A074557</u> | <u>002</u> | Aug 15, 1996 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 4GM RESIN/SCOOPFUL</u> | <u>A202901</u> | <u>001</u> | Jul 02, 2018 |

CHOLESTYRAMINE LIGHT

| | | | | | |
|-----------|-------------------------|------------------------------|----------------|------------|--------------|
| <u>AB</u> | PAR PHARM | <u>EQ 4GM RESIN/PACKET</u> | <u>A077203</u> | <u>001</u> | Aug 26, 2005 |
| <u>AB</u> | | <u>EQ 4GM RESIN/SCOOPFUL</u> | <u>A077203</u> | <u>002</u> | Aug 26, 2005 |
| <u>AB</u> | ! | <u>EQ 4GM RESIN/PACKET</u> | <u>A074558</u> | <u>001</u> | Aug 15, 1996 |
| <u>AB</u> | | <u>EQ 4GM RESIN/SCOOPFUL</u> | <u>A074558</u> | <u>002</u> | Aug 15, 1996 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 4GM RESIN/SCOOPFUL</u> | <u>A202902</u> | <u>001</u> | Apr 25, 2017 |

PREVALITE

| | | | | | |
|-----------|-------------------|------------------------------|----------------|------------|--------------|
| <u>AB</u> | UPSHER SMITH LABS | <u>EQ 4GM RESIN/PACKET</u> | <u>A073263</u> | <u>001</u> | Feb 22, 1996 |
| <u>AB</u> | | <u>EQ 4GM RESIN/SCOOPFUL</u> | <u>A073263</u> | <u>002</u> | Oct 30, 1997 |

CHOLIC ACID

CAPSULE; ORAL

CHOLBAM

| | | | | | |
|---|------|-------|---------|-----|--------------|
| + | RTRX | 50MG | N205750 | 001 | Mar 17, 2015 |
| + | ! | 250MG | N205750 | 002 | Mar 17, 2015 |

CHOLINE C-11

INJECTABLE; INTRAVENOUS

CHOLINE C-11

| | | | | | |
|-----------|----------------------------------------|------------------------------------|----------------|------------|--------------|
| <u>AP</u> | GLOBAL ISOTOPES LLC | <u>4-33.1mCi/ML</u> | <u>A206319</u> | <u>001</u> | Nov 13, 2015 |
| <u>AP</u> | + | <u>4-33.1mCi/ML</u> | <u>N203155</u> | <u>001</u> | Sep 12, 2012 |
| <u>AP</u> | UCSF RODIOPHARM | <u>4-33.1mCi/ML</u> | <u>A208444</u> | <u>001</u> | Nov 20, 2017 |
| <u>AP</u> | WA UNIV SCH MED UNIV TX MD ANDERSON | <u>4-33.1mCi/ML</u> 4-100mCi/ML | <u>A208413</u> | <u>001</u> | Jan 10, 2017 |
| | | | A205690 | 001 | Oct 29, 2015 |

CHOLINE FENOFIBRATE

CAPSULE, DELAYED RELEASE; ORAL

FENOFIBRIC ACID

| | | | | | |
|-----------|--------------------|---------------------------------|----------------|------------|--------------|
| <u>AB</u> | ACTAVIS ELIZABETH | <u>EQ 45MG FENOFIBRIC ACID</u> | <u>A200920</u> | <u>001</u> | Oct 07, 2015 |
| <u>AB</u> | | <u>EQ 135MG FENOFIBRIC ACID</u> | <u>A200920</u> | <u>002</u> | Oct 07, 2015 |
| <u>AB</u> | ALEMbic PHARMS LTD | <u>EQ 45MG FENOFIBRIC ACID</u> | <u>A208705</u> | <u>001</u> | May 12, 2017 |
| <u>AB</u> | | <u>EQ 135MG FENOFIBRIC ACID</u> | <u>A208705</u> | <u>002</u> | May 12, 2017 |
| <u>AB</u> | ANCHEN PHARMS | <u>EQ 45MG FENOFIBRIC ACID</u> | <u>A201573</u> | <u>002</u> | Jul 18, 2013 |
| <u>AB</u> | | <u>EQ 135MG FENOFIBRIC ACID</u> | <u>A201573</u> | <u>001</u> | Jul 18, 2013 |
| <u>AB</u> | IMPAX LABS INC | <u>EQ 45MG FENOFIBRIC ACID</u> | <u>A200264</u> | <u>001</u> | Sep 07, 2016 |
| <u>AB</u> | | <u>EQ 135MG FENOFIBRIC ACID</u> | <u>A200264</u> | <u>002</u> | Sep 07, 2016 |
| <u>AB</u> | LUPIN LTD | <u>EQ 45MG FENOFIBRIC ACID</u> | <u>A200750</u> | <u>001</u> | Dec 04, 2013 |
| <u>AB</u> | | <u>EQ 135MG FENOFIBRIC ACID</u> | <u>A200750</u> | <u>002</u> | Dec 04, 2013 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 45MG FENOFIBRIC ACID</u> | <u>A200913</u> | <u>001</u> | Mar 25, 2013 |
| <u>AB</u> | | <u>EQ 135MG FENOFIBRIC ACID</u> | <u>A200913</u> | <u>002</u> | Mar 25, 2013 |

PRESCRIPTION DRUG PRODUCT LIST

CHOLINE FENOFIBRATE

CAPSULE, DELAYED RELEASE;ORAL

TRILIPIX

| | | | | | |
|-----------|---|--------|---------------------------------|--------------------|--------------|
| AB | + | ABBVIE | EQ 45MG FENOFIBRIC ACID | N022224 001 | Dec 15, 2008 |
| AB | + | ! | EQ 135MG FENOFIBRIC ACID | N022224 002 | Dec 15, 2008 |

CHORIOGONADOTROPIN ALFA

INJECTABLE;SUBCUTANEOUS

OVIDREL

| | | | | | |
|---|---|------------|------------------|-------------|--------------|
| + | ! | EMD SERONO | EQ 0.25MG /0.5ML | N021149 002 | Oct 06, 2003 |
|---|---|------------|------------------|-------------|--------------|

CHROMIC CHLORIDE

INJECTABLE;INJECTION

CHROMIC CHLORIDE IN PLASTIC CONTAINER

| | | | | | |
|---|---|---------|------------------------|-------------|--------------|
| + | ! | HOSPIRA | EQ 0.004MG CHROMIUM/ML | N018961 001 | Jun 26, 1986 |
|---|---|---------|------------------------|-------------|--------------|

CICLESONIDE

AEROSOL, METERED;INHALATION

ALVESCO

| | | | | | |
|---|---|--------------------|------------|-------------|--------------|
| + | ! | ASTRAZENECA PHARMS | 0.08MG/INH | N021658 002 | Jan 10, 2008 |
|---|---|--------------------|------------|-------------|--------------|

| | | | | | |
|---|---|--|------------|-------------|--------------|
| + | ! | | 0.16MG/INH | N021658 003 | Jan 10, 2008 |
|---|---|--|------------|-------------|--------------|

AEROSOL, METERED;NASAL

ZETONNA

| | | | | | |
|---|---|--------------------|-------------|-------------|--------------|
| + | ! | ASTRAZENECA PHARMS | 0.037MG/INH | N202129 001 | Jan 20, 2012 |
|---|---|--------------------|-------------|-------------|--------------|

SPRAY, METERED;NASAL

OMNARIS

| | | | | | |
|---|---|--------------------|------------|-------------|--------------|
| + | ! | ASTRAZENECA PHARMS | 0.05MG/INH | N022004 001 | Oct 20, 2006 |
|---|---|--------------------|------------|-------------|--------------|

CICLOPIROX

CREAM;TOPICAL

CICLOPIROX

| | | | | | |
|-----------|--|------------------|--------------|--------------------|--------------|
| AB | | FOUGERA PHARMS | 0.77% | A076435 001 | Dec 29, 2004 |
| AB | | G AND W LABS INC | 0.77% | A078463 001 | Dec 20, 2010 |
| AB | | GLENMARK PHARMS | 0.77% | A090273 001 | Nov 10, 2009 |
| AB | | PERRIGO NEW YORK | 0.77% | A077364 001 | Mar 03, 2006 |
| AB | | TARO | 0.77% | A076790 001 | Apr 12, 2005 |

LOPROX

| | | | | | | |
|-----------|---|---|--------------------|--------------|--------------------|--------------|
| AB | + | ! | MEDIMETRIKS PHARMS | 0.77% | N018748 001 | Dec 30, 1982 |
|-----------|---|---|--------------------|--------------|--------------------|--------------|

GEL;TOPICAL

CICLOPIROX

| | | | | | | |
|-----------|---|---|-------------------|--------------|--------------------|--------------|
| AB | + | ! | CNTY LINE PHARMS | 0.77% | N020519 001 | Jul 21, 1997 |
| AB | | | FOUGERA PHARMS | 0.77% | A077896 001 | Jun 10, 2008 |
| AB | | | GLENMARK GENERICS | 0.77% | A091595 001 | Feb 29, 2012 |
| AB | | | PADDOCK LLC | 0.77% | A078266 001 | Jan 07, 2009 |

SHAMPOO;TOPICAL

CICLOPIROX

| | | | | | | |
|-----------|--|--|----------------------|-----------|--------------------|--------------|
| AT | | | ACTAVIS MID ATLANTIC | 1% | A090490 001 | Nov 24, 2009 |
| AT | | | FOUGERA PHARMS | 1% | A090146 001 | May 25, 2010 |
| AT | | | PERRIGO CO | 1% | A078594 001 | Feb 16, 2010 |
| AT | | | TARO | 1% | A090269 001 | Feb 23, 2011 |
| AT | | | TELIGENT PHARMA INC | 1% | A209975 001 | Apr 05, 2018 |

LOPROX

| | | | | | | |
|-----------|---|---|---------|-----------|--------------------|--------------|
| AT | + | ! | MEDICIS | 1% | N021159 001 | Feb 28, 2003 |
|-----------|---|---|---------|-----------|--------------------|--------------|

SOLUTION;TOPICAL

CICLOPIROX

| | | | | | | |
|-----------|--|--|----------------------|-----------|--------------------|--------------|
| AT | | | ACTAVIS MID ATLANTIC | 8% | A078046 001 | Sep 18, 2007 |
| AT | | | AKORN | 8% | A078975 001 | Feb 17, 2010 |
| AT | | | APOTEX INC | 8% | A078172 001 | Sep 18, 2007 |
| AT | | | G AND W LABS | 8% | A078233 001 | Sep 18, 2007 |
| AT | | | HI TECH PHARMA | 8% | A078270 001 | Sep 18, 2007 |
| AT | | | INGENUS PHARMS LLC | 8% | A078124 001 | Sep 18, 2007 |
| AT | | | PERRIGO NEW YORK | 8% | A077623 001 | Sep 18, 2007 |
| AT | | | TARO PHARM INDS | 8% | A078144 001 | Sep 18, 2007 |
| AT | | | TOLMAR | 8% | A077687 001 | Sep 18, 2007 |

PENLAC

| | | | | | | |
|-----------|---|---|-----------------|-----------|--------------------|--------------|
| AT | + | ! | VALEANT BERMUDA | 8% | N021022 001 | Dec 17, 1999 |
|-----------|---|---|-----------------|-----------|--------------------|--------------|

SUSPENSION;TOPICAL

CICLOPIROX

| | | | | | | |
|-----------|--|--|------------------|--------------|--------------------|--------------|
| AB | | | FOUGERA PHARMS | 0.77% | A076422 001 | Aug 06, 2004 |
| AB | | | PERRIGO NEW YORK | 0.77% | A077676 001 | Dec 15, 2006 |
| AB | | | TARO | 0.77% | A077092 001 | Aug 10, 2005 |

PRESCRIPTION DRUG PRODUCT LIST

CICLOPIROX

SUSPENSION; TOPICAL

LOPROX

| | | | | | |
|-----------|------------|--------------------|--------------|--------------------|--------------|
| AB | + ! | MEDIMETRIKS PHARMS | 0.77% | N019824 001 | Dec 30, 1988 |
|-----------|------------|--------------------|--------------|--------------------|--------------|

CIDOFOVIR

INJECTABLE; INJECTION

CIDOFOVIR

| | | | | | |
|-----------|--|-------------------|------------------------|--------------------|--------------|
| AP | | EMCURE PHARMS LTD | EQ 75MG BASE/ML | A202501 001 | Jul 26, 2012 |
|-----------|--|-------------------|------------------------|--------------------|--------------|

| | | | | | |
|-----------|----------|---------------------|------------------------|--------------------|--------------|
| AP | ! | MYLAN INSTITUTIONAL | EQ 75MG BASE/ML | A201276 001 | Jun 27, 2012 |
|-----------|----------|---------------------|------------------------|--------------------|--------------|

CILASTATIN SODIUM; IMPENEM

POWDER; INTRAVENOUS

IMPENEM AND CILASTATIN

| | | | | | |
|-----------|--|------------|---------------------------------------|--------------------|--------------|
| AP | | ACS DOBFAR | EQ 500MG BASE/VIAL; 500MG/VIAL | A090577 002 | Dec 21, 2011 |
|-----------|--|------------|---------------------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------------|---------------------------------------|--------------------|--------------|
| AP | | HOSPIRA INC | EQ 500MG BASE/VIAL; 500MG/VIAL | A090825 002 | Nov 16, 2011 |
|-----------|--|-------------|---------------------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|---------------------------------------|--------------------|--------------|
| AP | | | EQ 500MG BASE/VIAL; 500MG/VIAL | A091007 001 | Nov 16, 2011 |
|-----------|--|--|---------------------------------------|--------------------|--------------|

PRIMAXIN

| | | | | | |
|-----------|------------|-------|---------------------------------------|--------------------|--------------|
| AP | + ! | MERCK | EQ 500MG BASE/VIAL; 500MG/VIAL | N050587 002 | Nov 26, 1985 |
|-----------|------------|-------|---------------------------------------|--------------------|--------------|

IMPENEM AND CILASTATIN

| | | | | | |
|----------|--|------------|---------------------------------------|--------------------|--------------|
| ! | | ACS DOBFAR | EQ 250MG BASE/VIAL; 250MG/VIAL | A090577 001 | Dec 21, 2011 |
|----------|--|------------|---------------------------------------|--------------------|--------------|

CILOSTAZOL

TABLET; ORAL

CILOSTAZOL

| | | | | | |
|-----------|--|------------|-------------|--------------------|--------------|
| AB | | APOTEX INC | 50MG | A077030 001 | Dec 10, 2004 |
|-----------|--|------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 100MG | A077030 002 | Dec 10, 2004 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------------|-------------|--------------------|--------------|
| AB | | BRECKENRIDGE PHARM | 50MG | A077708 001 | Sep 28, 2009 |
|-----------|--|--------------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 100MG | A077708 002 | Sep 28, 2009 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|-----------------|-------------|--------------------|--------------|
| AB | | CASI PHARMS INC | 50MG | A077310 001 | Nov 08, 2005 |
|-----------|--|-----------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 100MG | A077021 001 | Nov 23, 2004 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------|-------------|--------------------|--------------|
| AB | | CHARTWELL RX | 50MG | A077722 001 | Sep 24, 2012 |
|-----------|--|--------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 100MG | A077831 001 | Sep 24, 2012 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------------------|-------------|--------------------|--------------|
| AB | | FRONTIDA BIOPHARM | 50MG | A077208 002 | Mar 29, 2006 |
|-----------|--|-------------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 100MG | A077208 001 | Mar 29, 2006 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|----------|------|-------------|--------------------|--------------|
| AB | ! | TEVA | 50MG | A077027 001 | Nov 24, 2004 |
|-----------|----------|------|-------------|--------------------|--------------|

| | | | | | |
|-----------|----------|--|--------------|--------------------|--------------|
| AB | ! | | 100MG | A077027 002 | Nov 24, 2004 |
|-----------|----------|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------------|-------------|--------------------|--------------|
| AB | | WEST-WARD PHARMS | 50MG | A077024 001 | May 17, 2005 |
|-----------|--|------------------|-------------|--------------------|--------------|

INT

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 100MG | A077024 002 | May 17, 2005 |
|-----------|--|--|--------------|--------------------|--------------|

CIMETIDINE

TABLET; ORAL

CIMETIDINE

| | | | | | |
|-----------|--|--------|--------------|--------------------|--------------|
| AB | | APOTEX | 200MG | A074890 001 | Dec 18, 1998 |
|-----------|--|--------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 300MG | A074890 002 | Dec 18, 1998 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 400MG | A074890 003 | Dec 18, 1998 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 800MG | A074890 004 | Dec 18, 1998 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------|--------------|--------------------|--------------|
| AB | | MYLAN | 200MG | A074246 001 | May 17, 1994 |
|-----------|--|-------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 300MG | A074246 002 | May 17, 1994 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 400MG | A074246 003 | May 17, 1994 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|----------|--|--------------|--------------------|--------------|
| AB | ! | | 800MG | A074246 004 | May 17, 1994 |
|-----------|----------|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------|--------------|--------------------|--------------|
| AB | | PLIVA | 800MG | A074566 001 | Feb 27, 1997 |
|-----------|--|-------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|------|--------------|--------------------|--------------|
| AB | | TEVA | 200MG | A074151 001 | May 17, 1994 |
|-----------|--|------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 300MG | A074151 002 | May 17, 1994 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 400MG | A074151 003 | May 17, 1994 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 800MG | A074463 001 | May 17, 1994 |
|-----------|--|--|--------------|--------------------|--------------|

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

| | | | | | |
|--|--|-----------------|--------------------------|--------------------|--------------|
| | | DAVA PHARMS INC | EQ 300MG BASE/2ML | A074428 001 | Apr 25, 1996 |
|--|--|-----------------|--------------------------|--------------------|--------------|

SOLUTION; ORAL

CIMETIDINE HYDROCHLORIDE

| | | | | | |
|-----------|--|----------------|--------------------------|--------------------|--------------|
| AA | | ANI PHARMS INC | EQ 300MG BASE/5ML | A074610 001 | Sep 26, 1996 |
|-----------|--|----------------|--------------------------|--------------------|--------------|

| | | | | | |
|-----------|----------|----------------|--------------------------|--------------------|--------------|
| AA | ! | HI TECH PHARMA | EQ 300MG BASE/5ML | A074664 001 | Oct 28, 1997 |
|-----------|----------|----------------|--------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------------|--------------------------|--------------------|--------------|
| AA | | PHARM ASSOC | EQ 300MG BASE/5ML | A074553 001 | Jan 27, 1997 |
|-----------|--|-------------|--------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------------|--------------------------|--------------------|--------------|
| AA | | WOCKHARDT BIO AG | EQ 300MG BASE/5ML | A074757 001 | Oct 17, 1997 |
|-----------|--|------------------|--------------------------|--------------------|--------------|

CINACALCET HYDROCHLORIDE

TABLET; ORAL

CINACALCET HYDROCHLORIDE

| | | | | | |
|-----------|--|----------------------|---------------------|--------------------|--------------|
| AB | | AUROBINDO PHARMA LTD | EQ 30MG BASE | A206125 001 | Mar 08, 2018 |
|-----------|--|----------------------|---------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|---------------------|--------------------|--------------|
| AB | | | EQ 60MG BASE | A206125 002 | Mar 08, 2018 |
|-----------|--|--|---------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|---------------------|--------------------|--------------|
| AB | | | EQ 90MG BASE | A206125 003 | Mar 08, 2018 |
|-----------|--|--|---------------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

CINACALCET HYDROCHLORIDE

TABLET; ORAL

CINACALCET HYDROCHLORIDE

| | | | | |
|-----------------|---------------------|---------------------|--------------------|--------------|
| <u>AB</u> | CIPLA | <u>EQ 30MG BASE</u> | <u>A208915 001</u> | Mar 08, 2018 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A208915 002</u> | Mar 08, 2018 |
| <u>AB</u> | | <u>EQ 90MG BASE</u> | <u>A208915 003</u> | Mar 08, 2018 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 30MG BASE</u> | <u>A203422 001</u> | Oct 16, 2018 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A203422 002</u> | Oct 16, 2018 |
| <u>AB</u> | | <u>EQ 90MG BASE</u> | <u>A203422 003</u> | Oct 16, 2018 |
| <u>AB</u> | PIRAMAL HLTHCARE UK | <u>EQ 30MG BASE</u> | <u>A210207 001</u> | Aug 01, 2018 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A210207 002</u> | Aug 01, 2018 |
| <u>AB</u> | | <u>EQ 90MG BASE</u> | <u>A210207 003</u> | Aug 01, 2018 |
| <u>AB</u> | STRIDES PHARMA | <u>EQ 30MG BASE</u> | <u>A209226 001</u> | Apr 30, 2018 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A209226 002</u> | Apr 30, 2018 |
| <u>AB</u> | | <u>EQ 90MG BASE</u> | <u>A209226 003</u> | Apr 30, 2018 |
| <u>AB</u> | SUN PHARMA GLOBAL | <u>EQ 30MG BASE</u> | <u>A207008 001</u> | Oct 11, 2018 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A207008 002</u> | Oct 11, 2018 |
| <u>AB</u> | | <u>EQ 90MG BASE</u> | <u>A207008 003</u> | Oct 11, 2018 |
| <u>AB</u> | WATSON LABS TEVA | <u>EQ 30MG BASE</u> | <u>A204377 001</u> | Dec 27, 2018 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A204377 002</u> | Dec 27, 2018 |
| <u>AB</u> | | <u>EQ 90MG BASE</u> | <u>A204377 003</u> | Dec 27, 2018 |
| <u>SENSIPAR</u> | | | | |
| <u>AB</u> | + AMGEN | <u>EQ 30MG BASE</u> | <u>N021688 001</u> | Mar 08, 2004 |
| <u>AB</u> | + | <u>EQ 60MG BASE</u> | <u>N021688 002</u> | Mar 08, 2004 |
| <u>AB</u> | +! | <u>EQ 90MG BASE</u> | <u>N021688 003</u> | Mar 08, 2004 |

CIPROFLOXACIN

FOR SUSPENSION; ORAL

CIPRO

| | | | | |
|-----------|------------------|------------------|--------------------|--------------|
| <u>AB</u> | + BAYER HLTHCARE | <u>250MG/5ML</u> | <u>N020780 001</u> | Sep 26, 1997 |
| <u>AB</u> | +! | <u>500MG/5ML</u> | <u>N020780 002</u> | Sep 26, 1997 |

CIPROFLOXACIN

| | | | | |
|-----------|-----------|------------------|--------------------|--------------|
| <u>AB</u> | LUPIN LTD | <u>250MG/5ML</u> | <u>A200563 001</u> | Mar 05, 2014 |
| <u>AB</u> | | <u>500MG/5ML</u> | <u>A200563 002</u> | Mar 05, 2014 |

INJECTABLE; INJECTION

CIPROFLOXACIN

| | | | | |
|-----------|------------------------|-----------------------------|--------------------|--------------|
| <u>AP</u> | ! BAXTER HLTHCARE CORP | <u>200MG/20ML (10MG/ML)</u> | <u>A078062 001</u> | Apr 29, 2008 |
| <u>AP</u> | ! | <u>400MG/40ML (10MG/ML)</u> | <u>A078062 002</u> | Apr 29, 2008 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>200MG/20ML (10MG/ML)</u> | <u>A076717 001</u> | Dec 22, 2009 |
| <u>AP</u> | | <u>400MG/40ML (10MG/ML)</u> | <u>A076717 002</u> | Dec 22, 2009 |
| <u>AP</u> | HOSPIRA | <u>200MG/20ML (10MG/ML)</u> | <u>A077245 001</u> | Aug 28, 2006 |
| <u>AP</u> | | <u>400MG/40ML (10MG/ML)</u> | <u>A077245 002</u> | Aug 28, 2006 |

CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|-----------|----------------------|--------------------|--------------------|--------------|
| <u>AP</u> | BAXTER HLTHCARE CORP | <u>200MG/100ML</u> | <u>A078024 001</u> | Mar 18, 2008 |
| <u>AP</u> | | <u>400MG/200ML</u> | <u>A078024 002</u> | Mar 18, 2008 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>400MG/200ML</u> | <u>A078431 001</u> | Nov 18, 2009 |
| <u>AP</u> | ! HOSPIRA | <u>200MG/100ML</u> | <u>A077753 001</u> | Mar 18, 2008 |
| <u>AP</u> | ! | <u>400MG/200ML</u> | <u>A077753 002</u> | Mar 18, 2008 |
| <u>AP</u> | INFORLIFE | <u>200MG/100ML</u> | <u>A078252 001</u> | Mar 18, 2008 |
| <u>AP</u> | | <u>400MG/200ML</u> | <u>A078252 002</u> | Mar 18, 2008 |

INJECTABLE, SUSPENSION; OTIC

OTIPRIO

| | | | | |
|----|-------------|--------------|-------------|--------------|
| +! | OTONOMY INC | 6% (60MG/ML) | N207986 001 | Dec 10, 2015 |
|----|-------------|--------------|-------------|--------------|

CIPROFLOXACIN HYDROCHLORIDE

OINTMENT; OPHTHALMIC

CILOXAN

| | | | | |
|----|----------------------|--------------|-------------|--------------|
| +! | NOVARTIS PHARMS CORP | EQ 0.3% BASE | N020369 001 | Mar 30, 1998 |
|----|----------------------|--------------|-------------|--------------|

SOLUTION/DROPS; OPHTHALMIC

CILOXAN

| | | | | | |
|-----------|----|----------------------|---------------------|--------------------|--------------|
| <u>AT</u> | +! | NOVARTIS PHARMS CORP | <u>EQ 0.3% BASE</u> | <u>N019992 001</u> | Dec 31, 1990 |
|-----------|----|----------------------|---------------------|--------------------|--------------|

CIPROFLOXACIN HYDROCHLORIDE

| | | | | |
|-----------|--------------------|---------------------|--------------------|--------------|
| <u>AT</u> | AKORN INC | <u>EQ 0.3% BASE</u> | <u>A076555 001</u> | Dec 11, 2008 |
| <u>AT</u> | ALTAIRE PHARMS INC | <u>EQ 0.3% BASE</u> | <u>A204613 001</u> | May 03, 2018 |
| <u>AT</u> | FDC LTD | <u>EQ 0.3% BASE</u> | <u>A077568 001</u> | Jun 30, 2008 |
| <u>AT</u> | RISING PHARMS | <u>EQ 0.3% BASE</u> | <u>A077689 001</u> | Dec 13, 2006 |
| <u>AT</u> | TELGENT | <u>EQ 0.3% BASE</u> | <u>A076754 001</u> | Jun 09, 2004 |
| <u>AT</u> | WATSON LABS INC | <u>EQ 0.3% BASE</u> | <u>A076673 001</u> | Jan 21, 2005 |

PRESCRIPTION DRUG PRODUCT LISTCIPROFLOXACIN HYDROCHLORIDE

SOLUTION/DROPS;OTIC

CETRAXAL

+! WRASER PHARMS

EQ 0.2% BASE

N021918 001 May 01, 2009

TABLET;ORAL

CIPRO**AB** + BAYER HLTHCAREEQ 250MG BASEN019537 002 Oct 22, 1987**AB** +!EQ 500MG BASEN019537 003 Oct 22, 1987CIPROFLOXACIN HYDROCHLORIDE**AB** APOTEXEQ 250MG BASEA076896 001 Nov 04, 2004**AB**EQ 500MG BASEA076896 002 Nov 04, 2004**AB**EQ 750MG BASEA076896 003 Nov 04, 2004**AB** AUROBINDO PHARMAEQ 250MG BASEA077859 001 Apr 26, 2007**AB**EQ 500MG BASEA077859 002 Apr 26, 2007**AB**EQ 750MG BASEA077859 003 Apr 26, 2007**AB** CARLSBADEQ 250MG BASEA076126 002 Jun 09, 2004**AB**EQ 500MG BASEA076126 003 Jun 09, 2004**AB**EQ 750MG BASEA076126 004 Jun 09, 2004**AB** DR REDDYS LABS LTDEQ 100MG BASEA075593 002 Jun 09, 2004**AB**EQ 250MG BASEA075593 003 Jun 09, 2004**AB**EQ 500MG BASEA075593 004 Jun 09, 2004**AB**EQ 750MG BASEA075593 001 Jun 09, 2004**AB** HIKMAEQ 250MG BASEA076558 002 Jun 09, 2004**AB**EQ 500MG BASEA076558 003 Jun 09, 2004**AB**EQ 750MG BASEA076558 004 Jun 09, 2004**AB** IVAX SUB TEVAEQ 250MG BASEA076089 002 Jun 09, 2004

PHARMS

ABEQ 500MG BASEA076089 003 Jun 09, 2004**AB**EQ 750MG BASEA076089 004 Jun 09, 2004**AB** MYLANEQ 500MG BASEA075817 003 Jun 09, 2004**AB** TARO PHARMEQ 100MG BASEA076912 001 Feb 18, 2005**AB**EQ 250MG BASEA076912 002 Oct 06, 2004**AB**EQ 500MG BASEA076912 003 Oct 06, 2004**AB**EQ 750MG BASEA076912 004 Oct 06, 2004**AB** UNIQUE PHARM LABSEQ 250MG BASEA076639 001 Sep 10, 2004**AB**EQ 500MG BASEA076639 002 Sep 10, 2004**AB**EQ 750MG BASEA076639 003 Sep 10, 2004**AB** WATSON LABSEQ 100MG BASEA076794 001 Feb 10, 2005**AB**EQ 250MG BASEA076794 002 Jun 09, 2004**AB**EQ 500MG BASEA076794 003 Jun 09, 2004**AB**EQ 750MG BASEA076794 004 Jun 09, 2004**AB** YILING PHARM LTDEQ 250MG BASEA208921 001 Jun 22, 2018**AB**EQ 500MG BASEA208921 002 Jun 22, 2018CIPROFLOXACIN HYDROCHLORIDE; FLUOCINOLONE ACETONIDE

SOLUTION/DROPS;OTIC

OTOVEL

+! LABORATORIOS SALVAT EQ 0.3% BASE;0.025%

N208251 001 Apr 29, 2016

CIPROFLOXACIN HYDROCHLORIDE; HYDROCORTISONE

SUSPENSION/DROPS;OTIC

CIPRO HC

+! NOVARTIS PHARMS

EQ 0.2% BASE;1%

N020805 001 Feb 10, 1998

CORP

CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CIPROFLOXACIN EXTENDED RELEASE**AB** ANCHEN PHARMS212.6MG;EQ 287.5MG BASEA078166 002 Nov 27, 2007**AB**425.2MG;EQ 574.9MG BASEA078166 001 Nov 27, 2007**AB** DR REDDYS LABS LTD425.2MG;EQ 574.9MG BASEA077701 001 Mar 26, 2007**AB** ! MYLAN PHARMS INC212.6MG;EQ 287.5MG BASEA078183 001 Mar 22, 2007**AB** !425.2MG;EQ 574.9MG BASEA078183 002 Mar 22, 2007CIPROFLOXACIN; DEXAMETHASONE

SUSPENSION/DROPS;OTIC

CIPRODEX

+! NOVARTIS PHARMS

0.3%;0.1%

N021537 001 Jul 18, 2003

CORP

PRESCRIPTION DRUG PRODUCT LIST

CISATRACURIUM BESYLATE

INJECTABLE; INJECTION

CISATRACURIUM BESYLATE

| | | | | |
|-----------|---------------------|-----------------------|--------------------|--------------|
| <u>AP</u> | ACCORD HLTHCARE | <u>EQ 2MG BASE/ML</u> | <u>A205873 001</u> | Jun 16, 2017 |
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 2MG BASE/ML</u> | <u>A203183 001</u> | Feb 26, 2015 |
| <u>AP</u> | JIANGSU HENGRUI MED | <u>EQ 2MG BASE/ML</u> | <u>A209334 001</u> | Aug 30, 2017 |
| <u>AP</u> | SANDOZ INC | <u>EQ 2MG BASE/ML</u> | <u>A200159 001</u> | Feb 03, 2012 |

CISATRACURIUM BESYLATE PRESERVATIVE FREE

| | | | | |
|-----------|---------------------|------------------------|--------------------|--------------|
| <u>AP</u> | ACCORD HLTHCARE | <u>EQ 2MG BASE/ML</u> | <u>A205872 001</u> | Jun 16, 2017 |
| <u>AP</u> | | <u>EQ 10MG BASE/ML</u> | <u>A205872 002</u> | Jun 16, 2017 |
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 2MG BASE/ML</u> | <u>A203182 001</u> | Feb 26, 2015 |
| <u>AP</u> | | <u>EQ 10MG BASE/ML</u> | <u>A203182 002</u> | Feb 26, 2015 |
| <u>AP</u> | JIANGSU HENGRUI MED | <u>EQ 2MG BASE/ML</u> | <u>A204960 001</u> | Jan 27, 2017 |
| <u>AP</u> | | <u>EQ 10MG BASE/ML</u> | <u>A204960 002</u> | Sep 19, 2017 |
| <u>AP</u> | SANDOZ INC | <u>EQ 2MG BASE/ML</u> | <u>A200154 001</u> | Feb 03, 2012 |
| <u>AP</u> | | <u>EQ 10MG BASE/ML</u> | <u>A200154 002</u> | Feb 03, 2012 |

NIMBEX

| | | | | |
|-----------|-----------|-----------------------|--------------------|--------------|
| <u>AP</u> | +! ABBVIE | <u>EQ 2MG BASE/ML</u> | <u>N020551 001</u> | Dec 15, 1995 |
|-----------|-----------|-----------------------|--------------------|--------------|

NIMBEX PRESERVATIVE FREE

| | | | | |
|-----------|-----------|------------------------|--------------------|--------------|
| <u>AP</u> | +! ABBVIE | <u>EQ 2MG BASE/ML</u> | <u>N020551 003</u> | Dec 15, 1995 |
| | +! | <u>EQ 10MG BASE/ML</u> | <u>N020551 002</u> | Dec 15, 1995 |

CISPLATIN

INJECTABLE; INJECTION

CISPLATIN

| | | | | |
|-----------|-------------------------|---------------|--------------------|--------------|
| <u>AP</u> | ACCORD HLTHCARE | <u>1MG/ML</u> | <u>A206774 001</u> | Aug 18, 2015 |
| <u>AP</u> | ! FRESENIUS KABI USA | <u>1MG/ML</u> | <u>A074735 001</u> | Jul 16, 1999 |
| <u>AP</u> | GLAND PHARMA LTD | <u>1MG/ML</u> | <u>A207323 001</u> | Mar 17, 2017 |
| <u>AP</u> | + HQ SPCLT PHARMA | <u>1MG/ML</u> | <u>N018057 004</u> | Nov 08, 1988 |
| <u>AP</u> | MYLAN LABS LTD | <u>1MG/ML</u> | <u>A091062 001</u> | Apr 18, 2012 |
| <u>AP</u> | PHARMACHEMIE BV | <u>1MG/ML</u> | <u>A074656 001</u> | May 16, 2000 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>1MG/ML</u> | <u>A075036 001</u> | Nov 07, 2000 |

CITALOPRAM HYDROBROMIDE

SOLUTION; ORAL

CITALOPRAM HYDROBROMIDE

| | | | | |
|-----------|---------------------------|-------------------------|--------------------|--------------|
| <u>AA</u> | AUROBINDO PHARMA LTD | <u>EQ 10MG BASE/5ML</u> | <u>A077812 001</u> | Aug 28, 2006 |
| <u>AA</u> | HETERO LABS LTD III | <u>EQ 10MG BASE/5ML</u> | <u>A201450 001</u> | Dec 15, 2015 |
| <u>AA</u> | LANNETT CO INC | <u>EQ 10MG BASE/5ML</u> | <u>A077629 001</u> | Jun 15, 2006 |
| <u>AA</u> | ! WEST-WARD PHARMS INT | <u>EQ 10MG BASE/5ML</u> | <u>A077043 001</u> | Dec 13, 2004 |

TABLET; ORAL

CELEXA

| | | | | |
|-----------|----------------------|---------------------|--------------------|--------------|
| <u>AB</u> | + ALLERGAN SALES LLC | <u>EQ 10MG BASE</u> | <u>N020822 001</u> | Apr 27, 2000 |
| <u>AB</u> | + | <u>EQ 20MG BASE</u> | <u>N020822 002</u> | Jul 17, 1998 |
| <u>AB</u> | +! | <u>EQ 40MG BASE</u> | <u>N020822 003</u> | Jul 17, 1998 |

CITALOPRAM HYDROBROMIDE

| | | | | |
|-----------|---------------------|---------------------|--------------------|--------------|
| <u>AB</u> | AMNEAL PHARMS NY | <u>EQ 10MG BASE</u> | <u>A077289 001</u> | Nov 30, 2006 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077289 002</u> | Nov 30, 2006 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077289 003</u> | Nov 30, 2006 |
| <u>AB</u> | APOTEX INC | <u>EQ 10MG BASE</u> | <u>A077046 001</u> | Nov 24, 2004 |
| <u>AB</u> | AUROBINDO | <u>EQ 10MG BASE</u> | <u>A077031 001</u> | Oct 28, 2004 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077031 002</u> | Oct 28, 2004 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077031 003</u> | Oct 28, 2004 |
| <u>AB</u> | CHARTWELL MOLECULAR | <u>EQ 10MG BASE</u> | <u>A077044 001</u> | Nov 05, 2004 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077044 002</u> | Nov 05, 2004 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077044 003</u> | Nov 05, 2004 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 10MG BASE</u> | <u>A077038 001</u> | Oct 28, 2004 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077038 002</u> | Oct 28, 2004 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077038 003</u> | Oct 28, 2004 |
| <u>AB</u> | EPIC PHARMA | <u>EQ 10MG BASE</u> | <u>A077045 003</u> | Apr 29, 2005 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077045 002</u> | Apr 29, 2005 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077045 001</u> | Apr 29, 2005 |
| <u>AB</u> | G AND W LABS INC | <u>EQ 10MG BASE</u> | <u>A077048 001</u> | Nov 16, 2004 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077048 002</u> | Nov 16, 2004 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077048 003</u> | Nov 16, 2004 |
| <u>AB</u> | GLENMARK GENERICS | <u>EQ 10MG BASE</u> | <u>A077654 001</u> | Feb 27, 2009 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077654 002</u> | Feb 27, 2009 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077654 003</u> | Feb 27, 2009 |
| <u>AB</u> | INVAGEN PHARMS | <u>EQ 10MG BASE</u> | <u>A077534 001</u> | Oct 03, 2006 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077534 002</u> | Oct 03, 2006 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077534 003</u> | Oct 03, 2006 |

PRESCRIPTION DRUG PRODUCT LIST

CITALOPRAM HYDROBROMIDE

TABLET; ORAL

CITALOPRAM HYDROBROMIDE

| | | | | |
|-----------|--------------------|---------------------|--------------------|--------------|
| <u>AB</u> | JUBILANT GENERICS | <u>EQ 10MG BASE</u> | <u>A205407 001</u> | Dec 23, 2015 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A205407 002</u> | Dec 23, 2015 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A205407 003</u> | Dec 23, 2015 |
| <u>AB</u> | MYLAN | <u>EQ 10MG BASE</u> | <u>A077042 001</u> | Nov 05, 2004 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077042 002</u> | Nov 05, 2004 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077042 003</u> | Nov 05, 2004 |
| <u>AB</u> | PLIVA | <u>EQ 10MG BASE</u> | <u>A077232 001</u> | Oct 31, 2005 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077232 002</u> | Oct 31, 2005 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077232 003</u> | Oct 31, 2005 |
| <u>AB</u> | SUN PHARM INDS INC | <u>EQ 10MG BASE</u> | <u>A077032 001</u> | Nov 12, 2004 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077032 002</u> | Nov 12, 2004 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077032 003</u> | Nov 12, 2004 |
| <u>AB</u> | TORPHARM | <u>EQ 20MG BASE</u> | <u>A077046 002</u> | Nov 24, 2004 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077046 003</u> | Nov 24, 2004 |
| <u>AB</u> | TORRENT PHARMS | <u>EQ 10MG BASE</u> | <u>A078216 001</u> | Mar 27, 2007 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A078216 002</u> | Mar 27, 2007 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A078216 003</u> | Mar 27, 2007 |

CITRIC ACID; GLUCONOLACTONE; MAGNESIUM CARBONATE

SOLUTION; IRRIGATION

RENACIDIN

+! UNITED GUARDIAN 6.602GM/100ML; 198MG/100ML; 3.177GM/100ML N019481 001 Oct 02, 1990

CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE

FOR SOLUTION; ORAL

PREPOPIK

+! FERRING PHARMS INC 12GM/PACKET; 3.5GM/PACKET; 10MG/PACKET N202535 001 Jul 16, 2012

SOLUTION; ORAL

CLENPIQ

+! FERRING PHARMS INC 12GM/BOT; 3.5GM/BOT; 10MG/BOT N209589 001 Nov 28, 2017

CITRIC ACID; UREA C-13

FOR SOLUTION, TABLET, FOR SOLUTION; ORAL

IDKIT:HP

+! EXALENZ BIOSCIENCE N/A, 4GM; 75MG, N/A N021314 001 Dec 17, 2002

CLADRIBINE

INJECTABLE; INJECTION

CLADRIBINE

| | | | | | |
|-----------|---|-------------------------|---------------|--------------------|--------------|
| <u>AP</u> | ! | FRESENIUS KABI USA | <u>1MG/ML</u> | <u>A076571 001</u> | Apr 22, 2004 |
| <u>AP</u> | | MYLAN LABS LTD | <u>1MG/ML</u> | <u>A200510 001</u> | Oct 06, 2011 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>1MG/ML</u> | <u>A075405 001</u> | Feb 28, 2000 |

CLARITHROMYCIN

FOR SUSPENSION; ORAL

CLARITHROMYCIN

| | | | | | |
|-----------|---|--------|------------------|--------------------|--------------|
| <u>AB</u> | | SANDOZ | <u>125MG/5ML</u> | <u>A065283 002</u> | Sep 04, 2007 |
| <u>AB</u> | ! | | <u>250MG/5ML</u> | <u>A065283 003</u> | Sep 04, 2007 |

TABLET; ORAL

CLARITHROMYCIN

| | | | | | |
|-----------|---|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | | ALLIED PHARMA INC | <u>250MG</u> | <u>A202710 001</u> | Jun 10, 2013 |
| <u>AB</u> | | | <u>500MG</u> | <u>A202710 002</u> | Jun 10, 2013 |
| <u>AB</u> | | APOTEX CORP | <u>250MG</u> | <u>A065384 001</u> | Aug 20, 2007 |
| <u>AB</u> | | | <u>500MG</u> | <u>A065384 002</u> | Aug 20, 2007 |
| <u>AB</u> | ! | AUROBINDO | <u>250MG</u> | <u>A065489 001</u> | Jul 25, 2012 |
| <u>AB</u> | ! | | <u>500MG</u> | <u>A065489 002</u> | Jul 25, 2012 |
| <u>AB</u> | | HEC PHARM | <u>250MG</u> | <u>A203584 001</u> | Sep 28, 2015 |
| <u>AB</u> | | | <u>500MG</u> | <u>A203584 002</u> | Sep 28, 2015 |
| <u>AB</u> | | SANDOZ | <u>250MG</u> | <u>A065144 001</u> | Oct 18, 2005 |
| <u>AB</u> | | | <u>500MG</u> | <u>A065136 001</u> | Aug 25, 2005 |
| <u>AB</u> | | TEVA | <u>250MG</u> | <u>A065155 001</u> | May 31, 2005 |
| <u>AB</u> | | | <u>500MG</u> | <u>A065155 002</u> | May 31, 2005 |
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>250MG</u> | <u>A065178 002</u> | May 25, 2004 |
| <u>AB</u> | | | <u>500MG</u> | <u>A065178 001</u> | May 25, 2004 |
| <u>AB</u> | | WOCKHARDT | <u>250MG</u> | <u>A065266 001</u> | May 31, 2006 |
| <u>AB</u> | | | <u>500MG</u> | <u>A065266 002</u> | May 31, 2006 |

TABLET, EXTENDED RELEASE; ORAL

CLARITHROMYCIN

| | | | | | |
|-----------|--|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS LABS FL INC | <u>500MG</u> | <u>A065145 001</u> | Jun 24, 2004 |
| <u>AB</u> | | ALLIED | <u>500MG</u> | <u>A203243 001</u> | Feb 29, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

CLARITHROMYCIN

TABLET, EXTENDED RELEASE;ORAL

CLARITHROMYCIN

| | | | | |
|-----------|----------------|--------------|--------------------|--------------|
| AB | LUPIN LTD | 500MG | A202532 001 | Sep 15, 2015 |
| AB | ! MAYNE PHARMA | 500MG | A065154 001 | May 18, 2005 |
| AB | SUNSHINE LAKE | 500MG | A208987 001 | Jul 09, 2018 |

CLEMASTINE FUMARATE

SYRUP;ORAL

CLEMASTINE FUMARATE

! TEVA

EQ 0.5MG BASE/5ML

A073399 001 Jun 30, 1994

TABLET;ORAL

CLEMASTINE FUMARATE

! TEVA

2.68MG

A073283 001 Jan 31, 1992

CLEVIDIPINE

EMULSION;INTRAVENOUS

CLEVIPREX

+! CHIESI USA INC

25MG/50ML (0.5MG/ML)

N022156 001 Aug 01, 2008

+!

50MG/100ML (0.5MG/ML)

N022156 002 Aug 01, 2008

CLINDAMYCIN HYDROCHLORIDE

CAPSULE;ORAL

CLEOCIN HYDROCHLORIDE

| | | | | |
|-----------|---------------------------|----------------------|--------------------|--------------|
| AB | + PHARMACIA AND UPJOHN | EQ 75MG BASE | N050162 001 | |
| AB | + | EQ 150MG BASE | N050162 002 | |
| AB | +! | EQ 300MG BASE | N050162 003 | Apr 14, 1988 |

CLINDAMYCIN HYDROCHLORIDE

| | | | | |
|-----------|--------------------|----------------------|--------------------|--------------|
| AB | AUROBINDO PHARMA | EQ 150MG BASE | A065442 001 | Aug 26, 2009 |
| AB | | EQ 300MG BASE | A065442 002 | Aug 26, 2009 |
| AB | EPIC PHARMA LLC | EQ 150MG BASE | A065194 001 | Mar 22, 2004 |
| AB | | EQ 300MG BASE | A065194 002 | Mar 22, 2004 |
| AB | G AND W LABS INC | EQ 150MG BASE | A063029 001 | Sep 20, 1989 |
| AB | | EQ 300MG BASE | A063029 002 | Aug 05, 2005 |
| AB | LANNETT CO INC | EQ 75MG BASE | A065243 002 | Aug 12, 2005 |
| AB | | EQ 150MG BASE | A065243 003 | Aug 12, 2005 |
| AB | | EQ 300MG BASE | A065243 001 | Aug 12, 2005 |
| AB | MICRO LABS | EQ 75MG BASE | A207402 001 | Nov 05, 2018 |
| AB | | EQ 150MG BASE | A207402 002 | Nov 05, 2018 |
| AB | | EQ 300MG BASE | A207402 003 | Nov 05, 2018 |
| AB | SUN PHARM INDS LTD | EQ 150MG BASE | A065061 001 | Feb 02, 2001 |
| AB | | EQ 300MG BASE | A065061 002 | Feb 02, 2001 |
| AB | WATSON LABS | EQ 150MG BASE | A063083 001 | Jul 31, 1991 |
| AB | | EQ 300MG BASE | A063083 002 | Mar 18, 2003 |
| AB | ZYDUS PHARMS USA | EQ 75MG BASE | A065217 001 | Jan 31, 2005 |
| AB | | EQ 150MG BASE | A065217 002 | Jan 31, 2005 |
| AB | | EQ 300MG BASE | A065217 003 | Jan 31, 2005 |

CLINDAMYCIN PALMITATE HYDROCHLORIDE

FOR SOLUTION;ORAL

CLEOCIN

| | | | | |
|-----------|---------------------------|-------------------------|--------------------|--------------|
| AA | ! PHARMACIA AND UPJOHN | EQ 75MG BASE/5ML | A062644 001 | Apr 07, 1986 |
|-----------|---------------------------|-------------------------|--------------------|--------------|

CLINDAMYCIN PALMITATE HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|-------------------------|--------------------|--------------|
| AA | AMNEAL PHARMS | EQ 75MG BASE/5ML | A203513 001 | Mar 13, 2014 |
| AA | AUROBINDO PHARMA LTD | EQ 75MG BASE/5ML | A202409 001 | Apr 30, 2013 |
| AA | HERITAGE PHARMS INC | EQ 75MG BASE/5ML | A207047 001 | May 11, 2018 |
| AA | LYNE | EQ 75MG BASE/5ML | A201821 001 | Aug 28, 2012 |
| AA | MYLAN PHARMS INC | EQ 75MG BASE/5ML | A203063 001 | May 25, 2016 |
| AA | ORIT LABS LLC | EQ 75MG BASE/5ML | A206958 001 | May 05, 2017 |
| AA | PADDOCK LLC | EQ 75MG BASE/5ML | A090902 001 | Jul 07, 2010 |

CLINDAMYCIN PHOSPHATE

AEROSOL, FOAM;TOPICAL

CLINDAMYCIN PHOSPHATE

| | | | | |
|-----------|------------------|-----------|--------------------|--------------|
| AT | PERRIGO UK FINCO | 1% | A090785 001 | Mar 31, 2010 |
|-----------|------------------|-----------|--------------------|--------------|

EVOCLIN

| | | | | |
|-----------|---------------------|-----------|--------------------|--------------|
| AT | +! MYLAN PHARMS INC | 1% | N050801 001 | Oct 22, 2004 |
|-----------|---------------------|-----------|--------------------|--------------|

CREAM;VAGINAL

CLEOCIN

| | | | | |
|-----------|----------------------------|-------------------|--------------------|--------------|
| AB | +! PHARMACIA AND UPJOHN | EQ 2% BASE | N050680 002 | Mar 02, 1998 |
|-----------|----------------------------|-------------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

CLINDAMYCIN PHOSPHATE

CREAM; VAGINAL

CLINDAMYCIN PHOSPHATE

AB FOUGERA PHARMS **EQ 2% BASE** **A065139 001** Dec 27, 2004
 CLINDESSE
 +! PERRIGO PHARMA INTL **EQ 2% BASE** **N050793 001** Nov 30, 2004

GEL; TOPICAL

CLEOCIN T

AB +! PHARMACIA AND **EQ 1% BASE** **N050615 001** Jan 07, 1987
 UPJOHN

CLINDAMYCIN PHOSPHATE

AB FOUGERA PHARMS **EQ 1% BASE** **A064160 001** Jan 28, 2000
 CLINDAGEL
 BT +! PRECISION DERMAT **EQ 1% BASE** **N050782 001** Nov 27, 2000

INJECTABLE; INJECTION

CLEOCIN PHOSPHATE

AP PHARMACIA AND **EQ 150MG BASE/ML** **A062803 001** Oct 16, 1987
 UPJOHN

AP +! **EQ 150MG BASE/ML** **N050441 001**

CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER

AP +! PHARMACIA AND **EQ 6MG BASE/ML** **N050639 001** Aug 30, 1989
 UPJOHN

AP +! **EQ 12MG BASE/ML** **N050639 002** Aug 30, 1989

AP +! **EQ 18MG BASE/ML** **N050639 003** Apr 10, 1991

CLINDAMYCIN PHOSPHATE

AP ALVOGEN INC **EQ 150MG BASE/ML** **A062800 001** Jul 24, 1987

AP **EQ 150MG BASE/ML** **A062801 001** Jul 24, 1987

AP **EQ 150MG BASE/ML** **A062943 001** Sep 29, 1988

AP FRESENIUS KABI USA **EQ 150MG BASE/ML** **A065346 001** Mar 29, 2007

AP **EQ 150MG BASE/ML** **A065347 001** May 09, 2007

AP MYLAN LABS LTD **EQ 150MG BASE/ML** **A204748 001** Oct 10, 2017

AP **EQ 150MG BASE/ML** **A204749 001** Oct 10, 2017

AP SAGENT PHARMS **EQ 150MG BASE/ML** **A090108 001** Sep 30, 2011

AP **EQ 150MG BASE/ML** **A090109 001** Sep 30, 2011

AP WEST-WARD PHARMS **EQ 150MG BASE/ML** **A062889 001** Apr 25, 1988

INT

AP **EQ 150MG BASE/ML** **A065206 001** Sep 24, 2004

CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER

AP AKORN INC **EQ 6MG BASE/ML** **A203048 001** Apr 04, 2013

AP **EQ 12MG BASE/ML** **A203048 002** Apr 04, 2013

AP **EQ 18MG BASE/ML** **A203048 003** Apr 04, 2013

AP BAXTER HLTHCARE **EQ 6MG BASE/ML** **A208084 001** Jun 28, 2017

CORP

AP **EQ 12MG BASE/ML** **A208084 002** Jun 28, 2017

AP **EQ 18MG BASE/ML** **A208084 003** Jun 28, 2017

AP SANDOZ INC **EQ 6MG BASE/ML** **A201692 001** May 31, 2012

AP **EQ 12MG BASE/ML** **A201692 002** May 31, 2012

AP **EQ 18MG BASE/ML** **A201692 003** May 31, 2012

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%

+! ABRAXIS PHARM **EQ 900MG BASE/100ML** **N050635 001** Dec 22, 1989

LOTION; TOPICAL

CLEOCIN T

AB +! PHARMACIA AND **EQ 1% BASE** **N050600 001** May 31, 1989
 UPJOHN

CLINDAMYCIN PHOSPHATE

AB FOUGERA PHARMS **EQ 1% BASE** **A065067 001** Jan 31, 2002

SOLUTION; INTRAVENOUS

CLINDAMYCIN PHOSPHATE IN 0.9% SODIUM CHLORIDE

+! BAXTER HLTHCARE **EQ 300MG BASE/50ML (EQ 6MG BASE/ML)** **N208083 001** Apr 20, 2017

CORP

+! **EQ 600MG BASE/50ML (EQ 12MG BASE/ML)** **N208083 002** Apr 20, 2017

+! **EQ 900MG BASE/50ML (EQ 18MG BASE/ML)** **N208083 003** Apr 20, 2017

SOLUTION; TOPICAL

CLEOCIN T

AT +! PHARMACIA AND **EQ 1% BASE** **N050537 001**
 UPJOHN

CLINDA-DERM

AT PADDOCK LLC **EQ 1% BASE** **A063329 001** Sep 30, 1992

CLINDAMYCIN PHOSPHATE

AT FOUGERA PHARMS **EQ 1% BASE** **A065254 001** Feb 14, 2006

AT FOUGERA PHARMS INC **EQ 1% BASE** **A064159 001** Jun 05, 1997

AT G AND W LABS INC **EQ 1% BASE** **A062811 001** Sep 01, 1988

AT GLASSHOUSE PHARMS **EQ 1% BASE** **A209846 001** Feb 08, 2018

AT PERRIGO NEW YORK **EQ 1% BASE** **A064050 001** Nov 30, 1995

PRESCRIPTION DRUG PRODUCT LIST

CLINDAMYCIN PHOSPHATE

SOLUTION; TOPICAL

CLINDAMYCIN PHOSPHATE

| | | | | |
|-----------|-------------------------|-------------------|--------------------|--------------|
| AT | TARO PHARM INDS | EQ 1% BASE | A065184 001 | Mar 31, 2004 |
| AT | TELLIGENT PHARMA INC | EQ 1% BASE | A206945 001 | Dec 30, 2016 |
| AT | VINTAGE PHARMS | EQ 1% BASE | A203343 001 | May 29, 2015 |
| AT | ZYDUS PHARMS USA INC | EQ 1% BASE | A208767 001 | Jul 16, 2018 |

SUPPOSITORY; VAGINAL

CLEOCIN

| | | | | |
|---|-------------------------|-------|-------------|--------------|
| + | PHARMACIA AND UPJOHN | 100MG | N050767 001 | Aug 13, 1999 |
|---|-------------------------|-------|-------------|--------------|

SWAB; TOPICAL

CLEOCIN

| | | | | | |
|-----------|---|-------------------------|-------------------|--------------------|--------------|
| AT | + | PHARMACIA AND UPJOHN | EQ 1% BASE | N050537 002 | Feb 22, 1994 |
|-----------|---|-------------------------|-------------------|--------------------|--------------|

CLINDAMYCIN PHOSPHATE

| | | | | |
|-----------|------------------|-------------------|--------------------|--------------|
| AT | AKORN | EQ 1% BASE | A065513 001 | Jun 17, 2010 |
| AT | PERRIGO NEW YORK | EQ 1% BASE | A065049 001 | May 25, 2000 |

CLINDETS

| | | | | |
|-----------|------------|-------------------|--------------------|--------------|
| AT | PERRIGO CO | EQ 1% BASE | A064136 001 | Sep 30, 1996 |
|-----------|------------|-------------------|--------------------|--------------|

CLINDAMYCIN PHOSPHATE; TRETINOIN

GEL; TOPICAL

CLINDAMYCIN PHOSPHATE AND TRETINOIN

| | | | | |
|-----------|-------------------------|---------------------|--------------------|--------------|
| AB | ACTAVIS MID ATLANTIC | 1.2%; 0.025% | A202564 001 | Jun 12, 2015 |
|-----------|-------------------------|---------------------|--------------------|--------------|

ZIANA

| | | | | | |
|-----------|---|---------|---------------------|--------------------|--------------|
| AB | + | MEDICIS | 1.2%; 0.025% | N050802 001 | Nov 07, 2006 |
|-----------|---|---------|---------------------|--------------------|--------------|

VELTIN

| | | | | | |
|----|---|-----------------|--------------|-------------|--------------|
| BX | + | AQUA PHARMS LLC | 1.2%; 0.025% | N050803 001 | Jul 16, 2010 |
|----|---|-----------------|--------------|-------------|--------------|

CLOBAZAM

FILM; ORAL

SYMPAZAN

| | | | | |
|---|------------------|------|-------------|--------------|
| + | AQUESTIVE THERAP | 5MG | N210833 001 | Nov 01, 2018 |
| + | | 10MG | N210833 002 | Nov 01, 2018 |
| + | | 20MG | N210833 003 | Nov 01, 2018 |

SUSPENSION; ORAL

CLOBAZAM

| | | | | |
|-----------|-------------------------|-----------------|--------------------|--------------|
| AB | AMNEAL PHARMS LLC | 2.5MG/ML | A210039 001 | Oct 22, 2018 |
| AB | BIONPHARMA INC | 2.5MG/ML | A208819 001 | Oct 22, 2018 |
| AB | LUPIN LTD | 2.5MG/ML | A210546 001 | Dec 28, 2018 |
| AB | MYLAN PHARMS INC | 2.5MG/ML | A211259 001 | Oct 22, 2018 |
| AB | UPSHER SMITH LABS | 2.5MG/ML | A210569 001 | Oct 22, 2018 |
| AB | WEST-WARD PHARMS INT | 2.5MG/ML | A209715 001 | Oct 22, 2018 |

ONFI

| | | | | | |
|-----------|---|---------------------|-----------------|--------------------|--------------|
| AB | + | LUNDBECK PHARMS LLC | 2.5MG/ML | N203993 001 | Dec 14, 2012 |
|-----------|---|---------------------|-----------------|--------------------|--------------|

TABLET; ORAL

CLOBAZAM

| | | | | |
|-----------|-------------------------|-------------|--------------------|--------------|
| AB | AMNEAL PHARMS CO | 10MG | A209718 001 | Oct 22, 2018 |
| AB | | 20MG | A209718 002 | Oct 22, 2018 |
| AB | BIONPHARMA INC | 10MG | A208825 001 | Oct 22, 2018 |
| AB | | 20MG | A208825 002 | Oct 22, 2018 |
| AB | BRECKENRIDGE PHARM | 10MG | A209308 001 | Oct 22, 2018 |
| AB | | 20MG | A209308 002 | Oct 22, 2018 |
| AB | HETERO LABS LTD III | 10MG | A209795 001 | Oct 22, 2018 |
| AB | | 20MG | A209795 002 | Oct 22, 2018 |
| AB | LUPIN LTD | 10MG | A210545 001 | Dec 14, 2018 |
| AB | | 20MG | A210545 002 | Dec 14, 2018 |
| AB | PIRAMAL HLTHCARE UK | 10MG | A209808 001 | Oct 22, 2018 |
| AB | | 20MG | A209808 002 | Oct 22, 2018 |
| AB | TARO PHARM | 10MG | A209440 001 | Oct 22, 2018 |
| AB | | 20MG | A209440 002 | Oct 22, 2018 |
| AB | UPSHER SMITH LABS | 10MG | A209687 001 | Oct 22, 2018 |
| AB | | 20MG | A209687 002 | Oct 22, 2018 |
| AB | WEST-WARD PHARMS INT | 10MG | A208785 001 | Oct 22, 2018 |
| AB | | 20MG | A208785 002 | Oct 22, 2018 |
| AB | ZYDUS PHARMS USA INC | 10MG | A211449 001 | Oct 22, 2018 |
| AB | | 20MG | A211449 002 | Oct 22, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

CLOBAZAM

TABLET; ORAL

ONFI

| | | | | | | |
|-----------|---|---------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | + | LUNDBECK PHARMS LLC | <u>10MG</u> | <u>N202067</u> | <u>002</u> | Oct 21, 2011 |
| <u>AB</u> | + | ! | <u>20MG</u> | <u>N202067</u> | <u>003</u> | Oct 21, 2011 |

CLOBETASOL PROPIONATE

AEROSOL, FOAM; TOPICAL

CLOBETASOL PROPIONATE

| | | | | | | |
|------------|--|--------------------|--------------|----------------|------------|--------------|
| <u>AB1</u> | | INGENUS PHARMS LLC | <u>0.05%</u> | <u>A206805</u> | <u>001</u> | Jul 31, 2017 |
| <u>AB1</u> | | PERRIGO ISRAEL | <u>0.05%</u> | <u>A077763</u> | <u>001</u> | Mar 10, 2008 |
| <u>AB1</u> | | TARO PHARM | <u>0.05%</u> | <u>A208779</u> | <u>001</u> | Oct 04, 2018 |

OLUX

| | | | | | | | |
|------------|---|---|------------------|--------------|----------------|------------|--------------|
| <u>AB1</u> | + | ! | MYLAN PHARMS INC | <u>0.05%</u> | <u>N021142</u> | <u>001</u> | May 26, 2000 |
|------------|---|---|------------------|--------------|----------------|------------|--------------|

CLOBETASOL PROPIONATE

| | | | | | | |
|------------|--|----------------|--------------|----------------|------------|--------------|
| <u>AB2</u> | | PERRIGO ISRAEL | <u>0.05%</u> | <u>A201402</u> | <u>001</u> | Aug 14, 2012 |
|------------|--|----------------|--------------|----------------|------------|--------------|

OLUX E

| | | | | | | | |
|------------|---|---|------------------|--------------|----------------|------------|--------------|
| <u>AB2</u> | + | ! | MYLAN PHARMS INC | <u>0.05%</u> | <u>N022013</u> | <u>001</u> | Jan 12, 2007 |
|------------|---|---|------------------|--------------|----------------|------------|--------------|

CREAM; TOPICAL

CLOBETASOL PROPIONATE

| | | | | | | |
|------------|--|-------------------------|--------------|----------------|------------|--------------|
| <u>AB1</u> | | AMNEAL PHARMS LLC | <u>0.05%</u> | <u>A211256</u> | <u>001</u> | Dec 26, 2018 |
| <u>AB1</u> | | CHEMO RESEARCH SL | <u>0.05%</u> | <u>A210034</u> | <u>001</u> | Jun 15, 2018 |
| <u>AB1</u> | | FOUGERA PHARMS INC | <u>0.05%</u> | <u>A074392</u> | <u>001</u> | Sep 30, 1996 |
| <u>AB1</u> | | G AND W LABS INC | <u>0.05%</u> | <u>A074139</u> | <u>001</u> | Aug 03, 1994 |
| <u>AB1</u> | | GLENMARK PHARMS | <u>0.05%</u> | <u>A209095</u> | <u>001</u> | May 10, 2018 |
| <u>AB1</u> | | LUPIN LTD | <u>0.05%</u> | <u>A210208</u> | <u>001</u> | Jan 30, 2018 |
| <u>AB1</u> | | MYLAN PHARMS INC | <u>0.05%</u> | <u>A075338</u> | <u>001</u> | Feb 09, 2001 |
| <u>AB1</u> | | SOLARIS PHARMA CORP | <u>0.05%</u> | <u>A211401</u> | <u>001</u> | Jan 11, 2019 |
| <u>AB1</u> | | TARO | <u>0.05%</u> | <u>A074249</u> | <u>001</u> | Jul 08, 1996 |
| <u>AB1</u> | | TELIGENT PHARMA INC | <u>0.05%</u> | <u>A209974</u> | <u>001</u> | Apr 17, 2018 |
| <u>AB1</u> | | ZYDUS PHARMS USA INC | <u>0.05%</u> | <u>A211074</u> | <u>001</u> | Oct 15, 2018 |

CORMAX

| | | | | | | |
|------------|---|----------------|--------------|----------------|------------|--------------|
| <u>AB1</u> | ! | HI TECH PHARMA | <u>0.05%</u> | <u>A074220</u> | <u>001</u> | May 16, 1997 |
|------------|---|----------------|--------------|----------------|------------|--------------|

CLOBETASOL PROPIONATE (EMOLLIENT)

| | | | | | | |
|------------|---|---------------------|--------------|----------------|------------|--------------|
| <u>AB2</u> | ! | FOUGERA PHARMS | <u>0.05%</u> | <u>A075430</u> | <u>001</u> | May 26, 1999 |
| <u>AB2</u> | | TARO | <u>0.05%</u> | <u>A075633</u> | <u>001</u> | May 17, 2000 |
| <u>AB2</u> | | TELIGENT PHARMA INC | <u>0.05%</u> | <u>A209411</u> | <u>001</u> | Aug 21, 2017 |

EMBELINE E

| | | | | | | |
|------------|--|----------------|--------------|----------------|------------|--------------|
| <u>AB2</u> | | HI TECH PHARMA | <u>0.05%</u> | <u>A075325</u> | <u>001</u> | Dec 24, 1998 |
|------------|--|----------------|--------------|----------------|------------|--------------|

IMPOYZ

| | | | | | | | |
|--|---|---|---------------|---------------|----------------|------------|--------------|
| | + | ! | ENCORE DERMAT | <u>0.025%</u> | <u>N209483</u> | <u>001</u> | Nov 28, 2017 |
|--|---|---|---------------|---------------|----------------|------------|--------------|

GEL; TOPICAL

CLOBETASOL PROPIONATE

| | | | | | | |
|-----------|---|---------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ! | FOUGERA PHARMS | <u>0.05%</u> | <u>A075368</u> | <u>001</u> | Feb 15, 2000 |
| <u>AB</u> | | PERRIGO CO | <u>0.05%</u> | <u>A075027</u> | <u>001</u> | Oct 31, 1997 |
| <u>AB</u> | | TARO | <u>0.05%</u> | <u>A075279</u> | <u>001</u> | May 28, 1999 |
| <u>AB</u> | | TELIGENT PHARMA INC | <u>0.05%</u> | <u>A208881</u> | <u>001</u> | Mar 06, 2017 |

EMBELINE

| | | | | | | |
|-----------|--|----------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | HI TECH PHARMA | <u>0.05%</u> | <u>A076141</u> | <u>001</u> | Apr 12, 2002 |
|-----------|--|----------------|--------------|----------------|------------|--------------|

LOTION; TOPICAL

CLOBETASOL PROPIONATE

| | | | | | | |
|-----------|--|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | ACTAVIS MID ATLANTIC | <u>0.05%</u> | <u>A078223</u> | <u>001</u> | Dec 04, 2008 |
| <u>AB</u> | | HI-TECH PHARMACAL | <u>0.05%</u> | <u>A211348</u> | <u>001</u> | Oct 26, 2018 |
| <u>AB</u> | | LUPIN LTD | <u>0.05%</u> | <u>A209147</u> | <u>001</u> | Sep 22, 2017 |
| <u>AB</u> | | TARO | <u>0.05%</u> | <u>A200302</u> | <u>001</u> | Jul 02, 2012 |
| <u>AB</u> | | TELIGENT PHARMA INC | <u>0.05%</u> | <u>A208667</u> | <u>001</u> | Nov 29, 2016 |

CLOBEX

| | | | | | | | |
|-----------|---|---|------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | ! | GALDERMA LABS LP | <u>0.05%</u> | <u>N021535</u> | <u>001</u> | Jul 24, 2003 |
|-----------|---|---|------------------|--------------|----------------|------------|--------------|

OINTMENT; TOPICAL

CLOBETASOL PROPIONATE

| | | | | | | |
|-----------|---|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | AMNEAL PHARMS LLC | <u>0.05%</u> | <u>A210551</u> | <u>001</u> | Aug 21, 2018 |
| <u>AB</u> | | CHEMO RESEARCH SL | <u>0.05%</u> | <u>A209701</u> | <u>001</u> | Apr 17, 2018 |
| <u>AB</u> | ! | FOUGERA PHARMS | <u>0.05%</u> | <u>A074407</u> | <u>001</u> | Feb 23, 1996 |
| <u>AB</u> | | G AND W LABS INC | <u>0.05%</u> | <u>A074089</u> | <u>001</u> | Feb 16, 1994 |
| <u>AB</u> | | GLENMARK PHARMS | <u>0.05%</u> | <u>A208933</u> | <u>001</u> | Mar 20, 2017 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>0.05%</u> | <u>A075057</u> | <u>001</u> | Aug 12, 1998 |
| <u>AB</u> | | NOVEL LABS INC | <u>0.05%</u> | <u>A208841</u> | <u>001</u> | May 04, 2018 |
| <u>AB</u> | | TARO | <u>0.05%</u> | <u>A074248</u> | <u>001</u> | Jul 12, 1996 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>0.05%</u> | <u>A210199</u> | <u>001</u> | Oct 27, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

CLOBETASOL PROPIONATE

OINTMENT; TOPICAL

EMBELINE

AB HI TECH PHARMA 0.05% **A074221 001** Mar 31, 1995
SHAMPOO; TOPICAL

CLOBETASOL PROPIONATE

AB ACTAVIS MID 0.05% **A078854 001** Jun 07, 2011
ATLANTIC

AB HI-TECH PHARMACAL 0.05% **A209871 001** Oct 27, 2017

AB PERRIGO ISRAEL 0.05% **A090974 001** Aug 09, 2012

CLOBEX

AB +! GALDERMA LABS 0.05% **N021644 001** Feb 05, 2004
SOLUTION; TOPICAL

CLOBETASOL PROPIONATE

AT FOUGERA PHARMS 0.05% **A075391 001** Feb 08, 1999

AT G AND W LABS INC 0.05% **A074331 001** Dec 15, 1995

AT GLENMARK PHARMS LTD 0.05% **A210190 001** Apr 18, 2018

AT MACLEODS PHARMS LTD 0.05% **A209361 001** Oct 25, 2017

AT NOVEL LABS INC 0.05% **A206075 001** Nov 23, 2015

AT TARO 0.05% **A075224 001** Nov 16, 1998

AT 0.05% **A075363 001** Dec 29, 2000

AT TOLMAR 0.05% **A076977 001** Aug 05, 2005

AT WOCHKHARDT BIO AG 0.05% **A075205 001** Nov 13, 1998

EMBELINE

AT ! HI TECH PHARMA 0.05% **A074222 001** Dec 06, 1995
SPRAY; TOPICAL

CLOBETASOL PROPIONATE

AT AKORN 0.05% **A207218 001** Apr 28, 2017

AT GLENMARK PHARMS 0.05% **A209004 001** Mar 26, 2018

AT LUPIN LTD 0.05% **A208125 001** Mar 26, 2018

AT PADDOCK LLC 0.05% **A090898 001** Jun 16, 2011

AT TARO 0.05% **A208842 001** Mar 26, 2018

AT ZYDUS PHARMS USA 0.05% **A206378 001** Feb 16, 2017
INC

CLOBEX

AT +! GALDERMA LABS LP 0.05% **N021835 001** Oct 27, 2005

CLOCORTOLONE PIVALATE

CREAM; TOPICAL

CLODERM

+! EPI HLTH 0.1% N017765 001

CLOFARABINE

SOLUTION; INTRAVENOUS

CLOFARABINE

AP ABON PHARMS LLC 20MG/20ML (1MG/ML) **A204029 001** May 09, 2017

AP AMNEAL PHARMS CO 20MG/20ML (1MG/ML) **A208857 001** Nov 06, 2017

AP DR REDDYS LABS LTD 20MG/20ML (1MG/ML) **A205375 001** Nov 06, 2017

AP GLAND PHARMA LTD 20MG/20ML (1MG/ML) **A207831 001** Oct 31, 2018

AP HOSPIRA INC 20MG/20ML (1MG/ML) **A210283 001** Dec 27, 2018

AP INGENUS PHARMS LLC 20MG/20ML (1MG/ML) **A210270 001** Sep 14, 2018

AP MSN LABS PVT LTD 20MG/20ML (1MG/ML) **A209775 001** Dec 06, 2017

AP MYLAN LABS LTD 20MG/20ML (1MG/ML) **A208860 001** Nov 06, 2017

CLOLAR

AP +! GENZYME 20MG/20ML (1MG/ML) **N021673 001** Dec 28, 2004

CLOMIPHENE CITRATE

TABLET; ORAL

CLOMIPHENE CITRATE

! PAR PHARM 50MG A075528 001 Aug 30, 1999

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

ANAFRANIL

AB +! SPECGX LLC 25MG **N019906 001** Dec 29, 1989

AB + 50MG **N019906 002** Dec 29, 1989

AB + 75MG **N019906 003** Dec 29, 1989

CLOMIPRAMINE HYDROCHLORIDE

AB AMNEAL PHARMS CO 25MG **A208632 001** Oct 31, 2018

AB 50MG **A208632 002** Oct 31, 2018

AB 75MG **A208632 003** Oct 31, 2018

AB LUPIN LTD 25MG **A209294 001** Nov 21, 2018

AB 50MG **A209294 002** Nov 21, 2018

AB 75MG **A209294 003** Nov 21, 2018

AB MYLAN 25MG **A074947 001** Apr 30, 1998

PRESCRIPTION DRUG PRODUCT LIST

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

CLOMIPRAMINE HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|-------------|--------------------|--------------|
| <u>AB</u> | | <u>50MG</u> | <u>A074947 002</u> | Apr 30, 1998 |
| <u>AB</u> | | <u>75MG</u> | <u>A074947 003</u> | Apr 30, 1998 |
| <u>AB</u> | SANDOZ | <u>25MG</u> | <u>A074364 001</u> | Mar 29, 1996 |
| <u>AB</u> | | <u>25MG</u> | <u>A074953 001</u> | Jun 25, 1997 |
| <u>AB</u> | | <u>50MG</u> | <u>A074364 002</u> | Mar 29, 1996 |
| <u>AB</u> | | <u>50MG</u> | <u>A074953 002</u> | Jun 25, 1997 |
| <u>AB</u> | | <u>75MG</u> | <u>A074364 003</u> | Mar 29, 1996 |
| <u>AB</u> | | <u>75MG</u> | <u>A074953 003</u> | Jun 25, 1997 |
| <u>AB</u> | TARO | <u>25MG</u> | <u>A074694 001</u> | Dec 31, 1996 |
| <u>AB</u> | | <u>50MG</u> | <u>A074694 002</u> | Dec 31, 1996 |
| <u>AB</u> | | <u>75MG</u> | <u>A074694 003</u> | Dec 31, 1996 |
| <u>AB</u> | TEVA | <u>25MG</u> | <u>A074958 001</u> | Aug 26, 1997 |
| <u>AB</u> | | <u>50MG</u> | <u>A074958 002</u> | Aug 26, 1997 |
| <u>AB</u> | | <u>75MG</u> | <u>A074958 003</u> | Aug 26, 1997 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>25MG</u> | <u>A208961 001</u> | Dec 27, 2017 |
| <u>AB</u> | | <u>50MG</u> | <u>A208961 002</u> | Dec 27, 2017 |
| <u>AB</u> | | <u>75MG</u> | <u>A208961 003</u> | Dec 27, 2017 |

CLONAZEPAM

TABLET; ORAL

CLONAZEPAM

| | | | | |
|-----------|--------------------|--------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>0.5MG</u> | <u>A077147 001</u> | May 02, 2005 |
| <u>AB</u> | | <u>1MG</u> | <u>A077147 002</u> | May 02, 2005 |
| <u>AB</u> | | <u>2MG</u> | <u>A077147 003</u> | May 02, 2005 |
| <u>AB</u> | ACTAVIS ELIZABETH | <u>0.5MG</u> | <u>A074869 001</u> | Oct 31, 1996 |
| <u>AB</u> | | <u>1MG</u> | <u>A074869 002</u> | Oct 31, 1996 |
| <u>AB</u> | | <u>2MG</u> | <u>A074869 003</u> | Oct 31, 1996 |
| <u>AB</u> | MYLAN | <u>0.5MG</u> | <u>A075150 001</u> | Oct 05, 1998 |
| <u>AB</u> | | <u>1MG</u> | <u>A075150 002</u> | Oct 05, 1998 |
| <u>AB</u> | | <u>2MG</u> | <u>A075150 003</u> | Oct 05, 1998 |
| <u>AB</u> | PRINSTON INC | <u>0.5MG</u> | <u>A077856 001</u> | Jun 28, 2006 |
| <u>AB</u> | | <u>1MG</u> | <u>A077856 002</u> | Jun 28, 2006 |
| <u>AB</u> | | <u>2MG</u> | <u>A077856 003</u> | Jun 28, 2006 |
| <u>AB</u> | SANDOZ | <u>0.5MG</u> | <u>A074979 001</u> | Aug 29, 1997 |
| <u>AB</u> | | <u>1MG</u> | <u>A074979 002</u> | Aug 29, 1997 |
| <u>AB</u> | | <u>2MG</u> | <u>A074979 003</u> | Aug 29, 1997 |
| <u>AB</u> | SUN PHARM INDS INC | <u>0.5MG</u> | <u>A075423 001</u> | Apr 27, 2001 |
| <u>AB</u> | | <u>1MG</u> | <u>A075423 002</u> | Apr 27, 2001 |
| <u>AB</u> | | <u>2MG</u> | <u>A075423 003</u> | Apr 27, 2001 |
| <u>AB</u> | TEVA | <u>0.5MG</u> | <u>A074569 001</u> | Sep 10, 1996 |
| <u>AB</u> | | <u>1MG</u> | <u>A074569 002</u> | Sep 10, 1996 |
| <u>AB</u> | | <u>2MG</u> | <u>A074569 003</u> | Sep 10, 1996 |
| <u>AB</u> | WATSON LABS | <u>0.5MG</u> | <u>A074964 001</u> | Dec 30, 1997 |
| <u>AB</u> | | <u>1MG</u> | <u>A074964 002</u> | Dec 30, 1997 |
| <u>AB</u> | | <u>2MG</u> | <u>A074964 003</u> | Dec 30, 1997 |

KLONOPIN

| | | | | |
|-----------|---------|--------------|--------------------|--|
| <u>AB</u> | + ROCHE | <u>0.5MG</u> | <u>N017533 001</u> | |
| <u>AB</u> | +! | <u>1MG</u> | <u>N017533 002</u> | |
| <u>AB</u> | + | <u>2MG</u> | <u>N017533 003</u> | |

TABLET, ORALLY DISINTEGRATING; ORAL

CLONAZEPAM

| | | | | |
|-----------|--------------------|----------------|--------------------|--------------|
| <u>AB</u> | BARR | <u>0.125MG</u> | <u>A077194 001</u> | Aug 10, 2005 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A077194 002</u> | Aug 10, 2005 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A077194 003</u> | Aug 10, 2005 |
| <u>AB</u> | | <u>1MG</u> | <u>A077194 004</u> | Aug 10, 2005 |
| <u>AB</u> | | <u>2MG</u> | <u>A077194 005</u> | Aug 10, 2005 |
| <u>AB</u> | PAR PHARM | <u>0.125MG</u> | <u>A077171 001</u> | Aug 03, 2005 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A077171 002</u> | Aug 03, 2005 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A077171 003</u> | Aug 03, 2005 |
| <u>AB</u> | ! | <u>1MG</u> | <u>A077171 004</u> | Aug 03, 2005 |
| <u>AB</u> | | <u>2MG</u> | <u>A077171 005</u> | Aug 03, 2005 |
| <u>AB</u> | SUN PHARM INDS INC | <u>0.125MG</u> | <u>A078654 001</u> | Aug 27, 2014 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A078654 002</u> | Aug 27, 2014 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A078654 003</u> | Aug 27, 2014 |
| <u>AB</u> | | <u>1MG</u> | <u>A078654 004</u> | Aug 27, 2014 |
| <u>AB</u> | | <u>2MG</u> | <u>A078654 005</u> | Aug 27, 2014 |

PRESCRIPTION DRUG PRODUCT LIST

CLONIDINE

FILM, EXTENDED RELEASE; TRANSDERMAL

CATAPRES-TTS-1

| | | | | | |
|-----------|---|-------------------------|-------------------|--------------------|--------------|
| <u>AB</u> | + | BOEHRINGER INGELHEIM | <u>0.1MG/24HR</u> | <u>N018891 001</u> | Oct 10, 1984 |
|-----------|---|-------------------------|-------------------|--------------------|--------------|

CATAPRES-TTS-2

| | | | | | |
|-----------|---|-------------------------|-------------------|--------------------|--------------|
| <u>AB</u> | + | BOEHRINGER INGELHEIM | <u>0.2MG/24HR</u> | <u>N018891 002</u> | Oct 10, 1984 |
|-----------|---|-------------------------|-------------------|--------------------|--------------|

CATAPRES-TTS-3

| | | | | | |
|-----------|---|-------------------------|-------------------|--------------------|--------------|
| <u>AB</u> | + | BOEHRINGER INGELHEIM | <u>0.3MG/24HR</u> | <u>N018891 003</u> | Oct 10, 1984 |
|-----------|---|-------------------------|-------------------|--------------------|--------------|

CLONIDINE

| | | | | | |
|-----------|--|---------------------|-------------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS LABS UT INC | <u>0.1MG/24HR</u> | <u>A090873 001</u> | May 06, 2014 |
| <u>AB</u> | | | <u>0.2MG/24HR</u> | <u>A090873 002</u> | May 06, 2014 |
| <u>AB</u> | | | <u>0.3MG/24HR</u> | <u>A090873 003</u> | May 06, 2014 |
| <u>AB</u> | | AVEVA | <u>0.1MG/24HR</u> | <u>A076157 001</u> | Aug 18, 2009 |
| <u>AB</u> | | | <u>0.2MG/24HR</u> | <u>A076157 002</u> | Aug 18, 2009 |
| <u>AB</u> | | | <u>0.3MG/24HR</u> | <u>A076157 003</u> | Aug 18, 2009 |
| <u>AB</u> | | MAYNE PHARMA | <u>0.1MG/24HR</u> | <u>A079090 001</u> | Aug 20, 2010 |
| <u>AB</u> | | | <u>0.2MG/24HR</u> | <u>A079090 002</u> | Aug 20, 2010 |
| <u>AB</u> | | | <u>0.3MG/24HR</u> | <u>A079090 003</u> | Aug 20, 2010 |
| <u>AB</u> | | MYLAN TECHNOLOGIES | <u>0.1MG/24HR</u> | <u>A076166 001</u> | Jul 16, 2010 |
| <u>AB</u> | | | <u>0.2MG/24HR</u> | <u>A076166 002</u> | Jul 16, 2010 |
| <u>AB</u> | | | <u>0.3MG/24HR</u> | <u>A076166 003</u> | Jul 16, 2010 |

CLONIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CLONIDINE HYDROCHLORIDE

| | | | | | |
|-----------|--|-------------------------|----------------------------|--------------------|--------------|
| <u>AP</u> | | FRESENIUS KABI USA | <u>1MG/10ML (0.1MG/ML)</u> | <u>A200673 001</u> | Jul 08, 2011 |
| <u>AP</u> | | | <u>5MG/10ML (0.5MG/ML)</u> | <u>A200673 002</u> | Jul 08, 2011 |
| <u>AP</u> | | HIKMA FARMACEUTICA | <u>1MG/10ML (0.1MG/ML)</u> | <u>A200300 001</u> | Jan 26, 2011 |
| <u>AP</u> | | | <u>5MG/10ML (0.5MG/ML)</u> | <u>A200300 002</u> | Jan 26, 2011 |
| <u>AP</u> | | LUITPOLD | <u>1MG/10ML (0.1MG/ML)</u> | <u>A091104 001</u> | Oct 08, 2009 |
| <u>AP</u> | | | <u>5MG/10ML (0.5MG/ML)</u> | <u>A091104 002</u> | Oct 08, 2009 |
| <u>AP</u> | | X-GEN PHARMS INC | <u>1MG/10ML (0.1MG/ML)</u> | <u>A203167 001</u> | Oct 29, 2013 |
| <u>AP</u> | | | <u>5MG/10ML (0.5MG/ML)</u> | <u>A203167 002</u> | Oct 29, 2013 |
| <u>AP</u> | | ZYDUS PHARMS USA INC | <u>1MG/10ML (0.1MG/ML)</u> | <u>A202601 001</u> | Feb 20, 2014 |
| <u>AP</u> | | | <u>5MG/10ML (0.5MG/ML)</u> | <u>A202601 002</u> | Feb 20, 2014 |

DURACLON

| | | | | | |
|-----------|---|---------------------|----------------------------|--------------------|--------------|
| <u>AP</u> | + | MYLAN INSTITUTIONAL | <u>1MG/10ML (0.1MG/ML)</u> | <u>N020615 001</u> | Oct 02, 1996 |
| <u>AP</u> | + | | <u>5MG/10ML (0.5MG/ML)</u> | <u>N020615 002</u> | Apr 27, 1999 |

TABLET; ORAL

CATAPRES

| | | | | | |
|-----------|---|-------------------------|--------------|--------------------|--|
| <u>AB</u> | + | BOEHRINGER INGELHEIM | <u>0.1MG</u> | <u>N017407 001</u> | |
| <u>AB</u> | + | | <u>0.2MG</u> | <u>N017407 002</u> | |
| <u>AB</u> | + | | <u>0.3MG</u> | <u>N017407 003</u> | |

CLONIDINE HYDROCHLORIDE

| | | | | | |
|-----------|--|--------------------|--------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS ELIZABETH | <u>0.1MG</u> | <u>A070974 001</u> | Dec 16, 1986 |
| <u>AB</u> | | | <u>0.2MG</u> | <u>A070975 001</u> | Dec 16, 1986 |
| <u>AB</u> | | | <u>0.3MG</u> | <u>A070976 001</u> | Dec 16, 1986 |
| <u>AB</u> | | ALEMBIC PHARMS LTD | <u>0.1MG</u> | <u>A091368 001</u> | Dec 06, 2011 |
| <u>AB</u> | | | <u>0.2MG</u> | <u>A091368 002</u> | Dec 06, 2011 |
| <u>AB</u> | | | <u>0.3MG</u> | <u>A091368 003</u> | Dec 06, 2011 |
| <u>AB</u> | | FRONTIDA BIOPHARM | <u>0.1MG</u> | <u>A070925 001</u> | Sep 04, 1987 |
| <u>AB</u> | | | <u>0.2MG</u> | <u>A070924 001</u> | Sep 04, 1987 |
| <u>AB</u> | | | <u>0.3MG</u> | <u>A070923 001</u> | Sep 04, 1987 |
| <u>AB</u> | | IMPAX LABS | <u>0.1MG</u> | <u>A078099 001</u> | Aug 27, 2009 |
| <u>AB</u> | | | <u>0.2MG</u> | <u>A078099 002</u> | Aug 27, 2009 |
| <u>AB</u> | | | <u>0.3MG</u> | <u>A078099 003</u> | Aug 27, 2009 |
| <u>AB</u> | | MYLAN | <u>0.1MG</u> | <u>A070317 002</u> | Jul 09, 1987 |
| <u>AB</u> | | | <u>0.2MG</u> | <u>A070317 003</u> | Jun 09, 1987 |
| <u>AB</u> | | | <u>0.3MG</u> | <u>A070317 001</u> | Jun 09, 1987 |
| <u>AB</u> | | PRINSTON INC | <u>0.1MG</u> | <u>A077901 001</u> | Mar 09, 2007 |
| <u>AB</u> | | | <u>0.2MG</u> | <u>A077901 002</u> | Mar 09, 2007 |
| <u>AB</u> | | | <u>0.3MG</u> | <u>A077901 003</u> | Mar 09, 2007 |
| <u>AB</u> | | SUN PHARM INDS INC | <u>0.1MG</u> | <u>A090329 001</u> | Jul 03, 2014 |
| <u>AB</u> | | | <u>0.2MG</u> | <u>A090329 002</u> | Jul 03, 2014 |
| <u>AB</u> | | | <u>0.3MG</u> | <u>A090329 003</u> | Jul 03, 2014 |
| <u>AB</u> | | UNICHEM | <u>0.1MG</u> | <u>A078895 001</u> | Aug 26, 2009 |
| <u>AB</u> | | | <u>0.2MG</u> | <u>A078895 002</u> | Aug 26, 2009 |
| <u>AB</u> | | | <u>0.3MG</u> | <u>A078895 003</u> | Aug 26, 2009 |
| <u>AB</u> | | YUNG SHIN PHARM | <u>0.1MG</u> | <u>A202297 001</u> | Jun 13, 2013 |

PRESCRIPTION DRUG PRODUCT LIST

CLONIDINE HYDROCHLORIDE

TABLET;ORAL

CLONIDINE HYDROCHLORIDE

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| <u>AB</u> | | <u>0.2MG</u> | <u>A202297 002</u> | Jun 13, 2013 |
| <u>AB</u> | | <u>0.3MG</u> | <u>A202297 003</u> | Jun 13, 2013 |

TABLET, EXTENDED RELEASE;ORAL

CLONIDINE HYDROCHLORIDE

| | | | | |
|------------|--------------------|--------------|--------------------|--------------|
| <u>AB1</u> | ACTAVIS ELIZABETH | <u>0.1MG</u> | <u>A203320 001</u> | May 15, 2015 |
| <u>AB1</u> | AJANTA PHARMA LTD | <u>0.1MG</u> | <u>A209686 001</u> | Nov 20, 2017 |
| <u>AB1</u> | AMNEAL PHARMS NY | <u>0.1MG</u> | <u>A210052 001</u> | Nov 20, 2017 |
| <u>AB1</u> | ANCHEN PHARMS | <u>0.1MG</u> | <u>A202984 001</u> | Sep 30, 2013 |
| <u>AB1</u> | JUBILANT GENERICS | <u>0.1MG</u> | <u>A210338 001</u> | Jan 29, 2018 |
| <u>AB1</u> | LUPIN LTD | <u>0.1MG</u> | <u>A209285 001</u> | Oct 23, 2017 |
| <u>AB1</u> | MAYNE PHARMA INC | <u>0.1MG</u> | <u>A210680 001</u> | Apr 30, 2018 |
| <u>AB1</u> | UPSHER SMITH LABS | <u>0.1MG</u> | <u>A211433 001</u> | Oct 12, 2018 |
| <u>AB1</u> | XIAMEN LP PHARM CO | <u>0.1MG</u> | <u>A209757 001</u> | Nov 20, 2017 |

KAPVAY

| | | | | | |
|------------|------------|-------------------------|--------------|--------------------|--------------|
| <u>AB1</u> | <u>+</u> ! | CONCORDIA PHARMS INC | <u>0.1MG</u> | <u>N022331 003</u> | Sep 28, 2010 |
|------------|------------|-------------------------|--------------|--------------------|--------------|

CLONIDINE HYDROCHLORIDE

| | | | | | |
|------------|--|-------------------|--------------|--------------------|--------------|
| <u>AB2</u> | | ACTAVIS ELIZABETH | <u>0.1MG</u> | <u>A202792 001</u> | May 15, 2015 |
|------------|--|-------------------|--------------|--------------------|--------------|

CLOPIDOGREL BISULFATE

TABLET;ORAL

CLOPIDOGREL BISULFATE

| | | | | | |
|-----------|--|-------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | ACCORD HLTHCARE | <u>EQ 75MG BASE</u> | <u>A202925 001</u> | Mar 27, 2013 |
| <u>AB</u> | | | <u>EQ 300MG BASE</u> | <u>A202925 002</u> | Mar 27, 2013 |
| <u>AB</u> | | ACME LABS | <u>EQ 75MG BASE</u> | <u>A078004 001</u> | May 17, 2012 |
| <u>AB</u> | | AMNEAL PHARMS | <u>EQ 75MG BASE</u> | <u>A203751 001</u> | Apr 11, 2014 |
| <u>AB</u> | | APOTEX INC | <u>EQ 75MG BASE</u> | <u>A076274 001</u> | May 17, 2012 |
| <u>AB</u> | | | <u>EQ 300MG BASE</u> | <u>A076274 002</u> | Mar 04, 2014 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>EQ 75MG BASE</u> | <u>A090540 001</u> | May 17, 2012 |
| <u>AB</u> | | CSPC OUYI PHARM CO | <u>EQ 75MG BASE</u> | <u>A204359 001</u> | Feb 02, 2017 |
| <u>AB</u> | | DR REDDYS LABS INC | <u>EQ 75MG BASE</u> | <u>A076273 001</u> | Jan 14, 2008 |
| <u>AB</u> | | DR REDDYS LABS LTD | <u>EQ 300MG BASE</u> | <u>A091023 001</u> | May 17, 2012 |
| <u>AB</u> | | GATE PHARMS | <u>EQ 300MG BASE</u> | <u>A091216 001</u> | May 17, 2012 |
| <u>AB</u> | | MACLEODS PHARMS LTD | <u>EQ 75MG BASE</u> | <u>A202928 001</u> | Feb 10, 2014 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>EQ 75MG BASE</u> | <u>A077665 001</u> | May 17, 2012 |
| <u>AB</u> | | | <u>EQ 300MG BASE</u> | <u>A077665 002</u> | May 17, 2012 |
| <u>AB</u> | | PRINSTON INC | <u>EQ 75MG BASE</u> | <u>A206376 001</u> | May 07, 2018 |
| <u>AB</u> | | | <u>EQ 300MG BASE</u> | <u>A206376 002</u> | May 07, 2018 |
| <u>AB</u> | | SCIEGEN PHARMS INC | <u>EQ 75MG BASE</u> | <u>A204165 001</u> | Sep 15, 2014 |
| <u>AB</u> | | | <u>EQ 300MG BASE</u> | <u>A204165 002</u> | Sep 15, 2014 |
| <u>AB</u> | | SUN PHARM INDUSTRIES | <u>EQ 75MG BASE</u> | <u>A078133 001</u> | Jun 10, 2013 |
| <u>AB</u> | | SUN PHARMA GLOBAL | <u>EQ 75MG BASE</u> | <u>A090494 001</u> | May 17, 2012 |
| <u>AB</u> | | TEVA | <u>EQ 75MG BASE</u> | <u>A076999 001</u> | May 17, 2012 |
| <u>AB</u> | | TEVA PHARMS | <u>EQ 300MG BASE</u> | <u>A090625 001</u> | May 17, 2012 |
| <u>AB</u> | | TORRENT PHARMS LTD | <u>EQ 75MG BASE</u> | <u>A090844 001</u> | May 17, 2012 |
| <u>AB</u> | | WOCKHARDT LTD | <u>EQ 75MG BASE</u> | <u>A202266 001</u> | Aug 14, 2012 |
| <u>AB</u> | | | <u>EQ 300MG BASE</u> | <u>A202266 002</u> | Nov 20, 2012 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>EQ 75MG BASE</u> | <u>A201686 001</u> | Oct 10, 2012 |

PLAVIX

| | | | | | |
|-----------|------------|-------------------|----------------------|--------------------|--------------|
| <u>AB</u> | <u>+</u> | SANOFI AVENTIS US | <u>EQ 75MG BASE</u> | <u>N020839 001</u> | Nov 17, 1997 |
| <u>AB</u> | <u>+</u> ! | | <u>EQ 300MG BASE</u> | <u>N020839 002</u> | Sep 20, 2007 |

CLORAZEPATE DIPOTASSIUM

TABLET;ORAL

CLORAZEPATE DIPOTASSIUM

| | | | | | |
|-----------|----------|------------|---------------|--------------------|--------------|
| <u>AB</u> | | MYLAN | <u>3.75MG</u> | <u>A071858 002</u> | Jul 17, 1987 |
| <u>AB</u> | | | <u>7.5MG</u> | <u>A071858 003</u> | Jul 17, 1987 |
| <u>AB</u> | <u>!</u> | | <u>15MG</u> | <u>A071858 001</u> | Jul 17, 1987 |
| <u>AB</u> | | TARO PHARM | <u>3.75MG</u> | <u>A075731 003</u> | Apr 27, 2000 |
| <u>AB</u> | | | <u>7.5MG</u> | <u>A075731 002</u> | Apr 27, 2000 |
| <u>AB</u> | | | <u>15MG</u> | <u>A075731 001</u> | Apr 27, 2000 |

GEN-XENE

| | | | | | |
|-----------|--|------|---------------|--------------------|--------------|
| <u>AB</u> | | ALRA | <u>3.75MG</u> | <u>A071787 001</u> | Apr 26, 1988 |
| <u>AB</u> | | | <u>7.5MG</u> | <u>A071788 001</u> | Apr 26, 1988 |
| <u>AB</u> | | | <u>15MG</u> | <u>A071789 001</u> | Apr 26, 1988 |

TRANXENE

| | | | | | |
|-----------|----------|----------------|--------------|--------------------|--|
| <u>AB</u> | <u>+</u> | RECORDATI RARE | <u>7.5MG</u> | <u>N017105 007</u> | |
|-----------|----------|----------------|--------------|--------------------|--|

PRESCRIPTION DRUG PRODUCT LIST

CLOTRIMAZOLE

CREAM; TOPICAL

CLOTRIMAZOLE

| | | | | | |
|-----------|-----------------|-----------|----------------|------------|--------------|
| <u>AB</u> | FOUGERA PHARMS | <u>1%</u> | <u>A078338</u> | <u>001</u> | Sep 02, 2008 |
| <u>AB</u> | GLENMARK PHARMS | <u>1%</u> | <u>A090219</u> | <u>001</u> | Aug 03, 2010 |
| <u>AB</u> | ! TARO | <u>1%</u> | <u>A072640</u> | <u>001</u> | Aug 31, 1993 |

SOLUTION; TOPICAL

CLOTRIMAZOLE

| | | | | | |
|-----------|--------|-----------|----------------|------------|--------------|
| <u>AT</u> | ! TARO | <u>1%</u> | <u>A074580</u> | <u>001</u> | Jul 29, 1996 |
| <u>AT</u> | TEVA | <u>1%</u> | <u>A073306</u> | <u>001</u> | Feb 28, 1995 |

TROCHE/LOZENGE; ORAL

CLOTRIMAZOLE

| | | | | | |
|-----------|---------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | PADDOCK LLC | <u>10MG</u> | <u>A076763</u> | <u>001</u> | Oct 28, 2005 |
| <u>AB</u> | ! WEST-WARD PHARMS INT | <u>10MG</u> | <u>A076387</u> | <u>001</u> | Jul 29, 2004 |

CLOZAPINE

SUSPENSION; ORAL

VERSACLOZ

| | | | | | |
|---|-----------------|---------|---------|-----|--------------|
| + | ! TASMAN PHARMA | 50MG/ML | N203479 | 001 | Feb 06, 2013 |
|---|-----------------|---------|---------|-----|--------------|

TABLET; ORAL

CLOZAPINE

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>25MG</u> | <u>A202873</u> | <u>001</u> | Nov 25, 2015 |
| <u>AB</u> | | <u>100MG</u> | <u>A202873</u> | <u>002</u> | Nov 25, 2015 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>25MG</u> | <u>A206433</u> | <u>001</u> | Nov 29, 2016 |
| <u>AB</u> | | <u>50MG</u> | <u>A206433</u> | <u>002</u> | Nov 29, 2016 |
| <u>AB</u> | | <u>100MG</u> | <u>A206433</u> | <u>003</u> | Nov 29, 2016 |
| <u>AB</u> | | <u>200MG</u> | <u>A206433</u> | <u>004</u> | Nov 29, 2016 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>25MG</u> | <u>A074949</u> | <u>001</u> | Nov 26, 1997 |
| <u>AB</u> | | <u>50MG</u> | <u>A074949</u> | <u>004</u> | Apr 25, 2005 |
| <u>AB</u> | | <u>50MG</u> | <u>A076809</u> | <u>003</u> | Dec 16, 2005 |
| <u>AB</u> | | <u>100MG</u> | <u>A074949</u> | <u>002</u> | Nov 26, 1997 |
| <u>AB</u> | | <u>100MG</u> | <u>A076809</u> | <u>002</u> | Dec 16, 2005 |
| <u>AB</u> | | <u>200MG</u> | <u>A076809</u> | <u>001</u> | Dec 16, 2005 |
| <u>AB</u> | MAYNE PHARMA | <u>25MG</u> | <u>A203807</u> | <u>001</u> | Sep 17, 2015 |
| <u>AB</u> | | <u>50MG</u> | <u>A203807</u> | <u>003</u> | Aug 22, 2017 |
| <u>AB</u> | | <u>100MG</u> | <u>A203807</u> | <u>002</u> | Sep 17, 2015 |
| <u>AB</u> | | <u>200MG</u> | <u>A203807</u> | <u>004</u> | Aug 22, 2017 |
| <u>AB</u> | MYLAN | <u>25MG</u> | <u>A075417</u> | <u>001</u> | May 27, 1999 |
| <u>AB</u> | | <u>50MG</u> | <u>A075417</u> | <u>004</u> | Apr 15, 2010 |
| <u>AB</u> | | <u>100MG</u> | <u>A075417</u> | <u>002</u> | May 27, 1999 |
| <u>AB</u> | | <u>200MG</u> | <u>A075417</u> | <u>005</u> | Apr 15, 2010 |
| <u>AB</u> | SUN PHARM INDS INC | <u>25MG</u> | <u>A075713</u> | <u>001</u> | Nov 15, 2002 |
| <u>AB</u> | | <u>50MG</u> | <u>A075713</u> | <u>003</u> | Aug 19, 2005 |
| <u>AB</u> | | <u>100MG</u> | <u>A075713</u> | <u>002</u> | Nov 15, 2002 |
| <u>AB</u> | | <u>200MG</u> | <u>A075713</u> | <u>004</u> | Nov 07, 2017 |

CLOZARIL

| | | | | | |
|-----------|-----------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + HERITAGE LIFE | <u>25MG</u> | <u>N019758</u> | <u>001</u> | Sep 26, 1989 |
| <u>AB</u> | + | <u>100MG</u> | <u>N019758</u> | <u>002</u> | Sep 26, 1989 |

CLOZAPINE

| | | | | | |
|--|-------------------------|--------|---------|-----|--------------|
| | IVAX SUB TEVA PHARMS | 12.5MG | A074949 | 003 | Jul 31, 2003 |
|--|-------------------------|--------|---------|-----|--------------|

TABLET, ORALLY DISINTEGRATING; ORAL

CLOZAPINE

| | | | | | |
|-----------|------------------|---------------|----------------|------------|--------------|
| <u>AB</u> | BARR LABS INC | <u>12.5MG</u> | <u>A090308</u> | <u>003</u> | Apr 09, 2018 |
| <u>AB</u> | | <u>25MG</u> | <u>A090308</u> | <u>001</u> | Nov 25, 2015 |
| <u>AB</u> | | <u>100MG</u> | <u>A090308</u> | <u>002</u> | Nov 25, 2015 |
| <u>AB</u> | MYLAN PHARMS INC | <u>25MG</u> | <u>A201824</u> | <u>002</u> | Sep 15, 2015 |
| <u>AB</u> | | <u>100MG</u> | <u>A201824</u> | <u>003</u> | Sep 15, 2015 |
| <u>AB</u> | TEVA PHARMS USA | <u>150MG</u> | <u>A203039</u> | <u>001</u> | Nov 25, 2015 |
| <u>AB</u> | | <u>200MG</u> | <u>A203039</u> | <u>002</u> | Nov 25, 2015 |

FAZACLO ODT

| | | | | | |
|-----------|-------------------|---------------|----------------|------------|--------------|
| <u>AB</u> | + JAZZ PHARMS III | <u>12.5MG</u> | <u>N021590</u> | <u>004</u> | May 30, 2007 |
| <u>AB</u> | + | <u>25MG</u> | <u>N021590</u> | <u>001</u> | Feb 10, 2004 |
| <u>AB</u> | + | <u>100MG</u> | <u>N021590</u> | <u>002</u> | Feb 10, 2004 |
| <u>AB</u> | + | <u>150MG</u> | <u>N021590</u> | <u>005</u> | Jul 09, 2010 |
| <u>AB</u> | + | <u>200MG</u> | <u>N021590</u> | <u>006</u> | Jul 09, 2010 |

PRESCRIPTION DRUG PRODUCT LIST

COBICISTAT

TABLET;ORAL

TYBOST

+! GILEAD SCIENCES INC 150MG N203094 001 Sep 24, 2014

COBICISTAT; DARUNAVIR ETHANOLATE

TABLET;ORAL

PREZCOBIX

+! JANSSEN PRODS 150MG;EQ 800MG BASE N205395 001 Jan 29, 2015

COBICISTAT; DARUNAVIR ETHANOLATE; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

SYMITUZA

+! JANSSEN PRODS 150MG;EQ 800MG BASE;200MG;EQ 10MG BASE N210455 001 Jul 17, 2018

COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

GENVOYA

+! GILEAD SCIENCES INC 150MG;150MG;200MG;EQ 10MG BASE N207561 001 Nov 05, 2015

COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

STRIBILD

+! GILEAD SCIENCES INC 150MG;150MG;200MG;300MG N203100 001 Aug 27, 2012

COBIMETINIB FUMARATE

TABLET;ORAL

COTELLIC

+! GENENTECH INC EQ 20MG BASE N206192 001 Nov 10, 2015

COCAINE HYDROCHLORIDE

SOLUTION;NASAL

GOPRELTO

+! GENUS LIFESCIENCES 4% N209963 001 Dec 14, 2017

CODEINE PHOSPHATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE**AA** VINTAGE 10MG/5ML; 5MG/5ML; 6.25MG/5ML **A040660 001** Dec 07, 2006PROMETH VC W/ CODEINE**AA** ! ACTAVIS MID 10MG/5ML; 5MG/5ML; 6.25MG/5ML **A088764 001** Oct 31, 1984

ATLANTIC

PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE**AA** HI-TECH PHARMA CO 10MG/5ML; 5MG/5ML; 6.25MG/5ML **A040674 001** Dec 23, 2014PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE**AA** AMNEAL PHARMS 10MG/5ML; 5MG/5ML; 6.25MG/5ML **A200963 001** Aug 26, 2015CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE**AA** ! ACTAVIS MID 10MG/5ML; 6.25MG/5ML **A088763 001** Oct 31, 1984

ATLANTIC

AA AMNEAL PHARMS 10MG/5ML; 6.25MG/5ML **A200894 001** Apr 24, 2013**AA** HI TECH PHARMA 10MG/5ML; 6.25MG/5ML **A040151 001** Aug 26, 1997**AA** NOSTRUM LABS INC 10MG/5ML; 6.25MG/5ML **A090180 001** Mar 17, 2010**AA** TRIS PHARMA INC 10MG/5ML; 6.25MG/5ML **A200386 001** Jun 29, 2012**AA** WOCKHARDT BIO AG 10MG/5ML; 6.25MG/5ML **A088875 001** Dec 17, 1984PROMETHAZINE WITH CODEINE**AA** VINTAGE 10MG/5ML; 6.25MG/5ML **A040650 001** Jan 31, 2006CODEINE PHOSPHATE; PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP;ORAL

TRIACIN-C

! STI PHARMA LLC 10MG/5ML; 30MG/5ML; 1.25MG/5ML A088704 001 Mar 22, 1985

CODEINE SULFATE

TABLET;ORAL

CODEINE SULFATE**AB** LANNETT CO INC 15MG **A203046 001** Jun 13, 2014**AB** 30MG **A203046 002** Jun 13, 2014**AB** 60MG **A203046 003** Jun 13, 2014**AB** + WEST-WARD PHARMS 15MG **N022402 001** Jul 16, 2009

INT

AB + 30MG **N022402 002** Jul 16, 2009**AB** +! 60MG **N022402 003** Jul 16, 2009

PRESCRIPTION DRUG PRODUCT LIST

COLCHICINE

CAPSULE; ORAL

COLCHICINE

| | | | | |
|-----------|---------------|--------------|--------------------|--------------|
| AB | PAR PHARM INC | 0.6MG | A208678 001 | Nov 29, 2018 |
|-----------|---------------|--------------|--------------------|--------------|

MITIGARE

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| AB | +! HIKMA INTL PHARMS | 0.6MG | N204820 001 | Sep 26, 2014 |
|-----------|----------------------|--------------|--------------------|--------------|

TABLET; ORAL

COLCRYS

| | | | | |
|--|----------------------|-------|-------------|--------------|
| | +! TAKEDA PHARMS USA | 0.6MG | N022352 001 | Jul 29, 2009 |
|--|----------------------|-------|-------------|--------------|

COLCHICINE; PROBENECID

TABLET; ORAL

COL-PROBENECID

| | | | | |
|-----------|---------------|--------------------|--------------------|--|
| AB | ! WATSON LABS | 0.5MG;500MG | A084279 001 | |
|-----------|---------------|--------------------|--------------------|--|

PROBENECID AND COLCHICINE

| | | | | |
|-----------|-------------|--------------------|--------------------|--------------|
| AB | NOVAST LABS | 0.5MG;500MG | A040618 001 | May 13, 2008 |
|-----------|-------------|--------------------|--------------------|--------------|

COLESEVELAM HYDROCHLORIDE

FOR SUSPENSION; ORAL

COLESEVELAM HYDROCHLORIDE

| | | | | |
|-----------|---------------------|-----------------------|--------------------|--------------|
| AB | GLENMARK PHARMS LTD | 1.875GM/PACKET | A202190 001 | Jul 16, 2018 |
|-----------|---------------------|-----------------------|--------------------|--------------|

| | | | | |
|-----------|--|----------------------|--------------------|--------------|
| AB | | 3.75GM/PACKET | A202190 002 | Jul 16, 2018 |
|-----------|--|----------------------|--------------------|--------------|

WELCHOL

| | | | | |
|-----------|------------------|-----------------------|--------------------|--------------|
| AB | + DAIICHI SANKYO | 1.875GM/PACKET | N022362 001 | Oct 02, 2009 |
|-----------|------------------|-----------------------|--------------------|--------------|

| | | | | |
|-----------|----|----------------------|--------------------|--------------|
| AB | +! | 3.75GM/PACKET | N022362 002 | Oct 02, 2009 |
|-----------|----|----------------------|--------------------|--------------|

TABLET; ORAL

COLESEVELAM HYDROCHLORIDE

| | | | | |
|-----------|----------------|--------------|--------------------|--------------|
| AB | ALKEM LABS LTD | 625MG | A209038 001 | Oct 05, 2018 |
|-----------|----------------|--------------|--------------------|--------------|

| | | | | |
|-----------|--------------------|--------------|--------------------|--------------|
| AB | DR REDDYS LABS LTD | 625MG | A210889 001 | Oct 05, 2018 |
|-----------|--------------------|--------------|--------------------|--------------|

| | | | | |
|-----------|---------------------|--------------|--------------------|--------------|
| AB | GLENMARK PHARMS LTD | 625MG | A203480 001 | May 18, 2018 |
|-----------|---------------------|--------------|--------------------|--------------|

| | | | | |
|-----------|----------------|--------------|--------------------|--------------|
| AB | IMPAX LABS INC | 625MG | A091600 001 | May 16, 2018 |
|-----------|----------------|--------------|--------------------|--------------|

WELCHOL

| | | | | |
|-----------|-------------------|--------------|--------------------|--------------|
| AB | +! DAIICHI SANKYO | 625MG | N021176 001 | May 26, 2000 |
|-----------|-------------------|--------------|--------------------|--------------|

COLESTIPOL HYDROCHLORIDE

GRANULE; ORAL

COLESTID

| | | | | |
|-----------|--------------------|---------------------|--------------------|--------------|
| AB | + PHARMACIA UPJOHN | 5GM/SCOOPFUL | N017563 003 | Sep 22, 1995 |
|-----------|--------------------|---------------------|--------------------|--------------|

| | | | | |
|-----------|----|-------------------|--------------------|--------------|
| AB | +! | 5GM/PACKET | N017563 004 | Sep 22, 1995 |
|-----------|----|-------------------|--------------------|--------------|

COLESTIPOL HYDROCHLORIDE

| | | | | |
|-----------|------------|---------------------|--------------------|--------------|
| AB | IMPAX LABS | 5GM/SCOOPFUL | A077277 001 | May 02, 2006 |
|-----------|------------|---------------------|--------------------|--------------|

| | | | | |
|-----------|--|-------------------|--------------------|--------------|
| AB | | 5GM/PACKET | A077277 002 | May 02, 2006 |
|-----------|--|-------------------|--------------------|--------------|

FLAVORED COLESTID

| | | | | |
|--|--------------------|------------|-------------|--|
| | + PHARMACIA UPJOHN | 5GM/PACKET | N017563 001 | |
|--|--------------------|------------|-------------|--|

| | | | | |
|--|---|--------------|-------------|--|
| | + | 5GM/SCOOPFUL | N017563 002 | |
|--|---|--------------|-------------|--|

TABLET; ORAL

COLESTID

| | | | | |
|-----------|---------------------|------------|--------------------|--------------|
| AB | +! PHARMACIA UPJOHN | 1GM | N020222 001 | Jul 19, 1994 |
|-----------|---------------------|------------|--------------------|--------------|

COLESTIPOL HYDROCHLORIDE

| | | | | |
|-----------|------------|------------|--------------------|--------------|
| AB | IMPAX LABS | 1GM | A077510 001 | Oct 24, 2006 |
|-----------|------------|------------|--------------------|--------------|

COLISTIMETHATE SODIUM

INJECTABLE; INJECTION

COLISTIMETHATE SODIUM

| | | | | |
|-----------|-------------------|---------------------------|--------------------|--------------|
| AP | EMCURE PHARMS LTD | EQ 150MG BASE/VIAL | A202359 001 | Sep 28, 2012 |
|-----------|-------------------|---------------------------|--------------------|--------------|

| | | | | |
|-----------|--------------------|---------------------------|--------------------|--------------|
| AP | FRESENIUS KABI USA | EQ 150MG BASE/VIAL | A065364 001 | Apr 17, 2008 |
|-----------|--------------------|---------------------------|--------------------|--------------|

| | | | | |
|-----------|--------------|---------------------------|--------------------|--------------|
| AP | NEXUS PHARMS | EQ 150MG BASE/VIAL | A065177 001 | Mar 19, 2004 |
|-----------|--------------|---------------------------|--------------------|--------------|

| | | | | |
|-----------|---------------|---------------------------|--------------------|--------------|
| AP | SAGENT PHARMS | EQ 150MG BASE/VIAL | A201365 001 | Feb 19, 2014 |
|-----------|---------------|---------------------------|--------------------|--------------|

| | | | | |
|-----------|--------------|---------------------------|--------------------|--------------|
| AP | X GEN PHARMS | EQ 150MG BASE/VIAL | A064216 001 | Feb 26, 1999 |
|-----------|--------------|---------------------------|--------------------|--------------|

| | | | | |
|-----------|-------------------|---------------------------|--------------------|--------------|
| AP | XELLIA PHARMS APS | EQ 150MG BASE/VIAL | A205356 001 | May 29, 2015 |
|-----------|-------------------|---------------------------|--------------------|--------------|

COLY-MYCIN M

| | | | | |
|-----------|-------------------------|---------------------------|--------------------|--|
| AP | +! PAR STERILE PRODUCTS | EQ 150MG BASE/VIAL | N050108 002 | |
|-----------|-------------------------|---------------------------|--------------------|--|

COLISTIN SULFATE; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; THONZONIUM BROMIDE

SUSPENSION/DROPS; OTIC

COLY-MYCIN S

| | | | | |
|--|--------------------|--------------------------------------------------|-------------|--|
| | +! ENDO PHARMS INC | EQ 3MG BASE/ML;10MG/ML;EQ 3.3MG BASE/ML;0.5MG/ML | N050356 001 | |
|--|--------------------|--------------------------------------------------|-------------|--|

PRESCRIPTION DRUG PRODUCT LIST

CONIVAPTAN HYDROCHLORIDE

INJECTABLE; INTRAVENOUS

VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER

+! CUMBERLAND PHARMS 20MG/100ML (0.2MG/ML)

N021697 002 Oct 08, 2008

COPANLISIB DIHYDROCHLORIDE

POWDER; INTRAVENOUS

ALIQOPA

+! BAYER HEALTHCARE 60MG/VIAL

N209936 001 Sep 14, 2017

COPPER

INTRAUTERINE DEVICE; INTRAUTERINE

PARAGARD T 380A

+! COOPERSURGICAL 309MG/COPPER

N018680 001 Nov 15, 1984

CORTICORELIN OVINE TRIFLUORATE

INJECTABLE; INJECTION

ACTHREL

+! FERRING EQ 0.1MG BASE/VIAL

N020162 001 May 23, 1996

CORTICOTROPIN

INJECTABLE; INJECTION

H.P. ACTHAR GEL

+! MALLINCKRODT ARD 80 UNITS/ML

N008372 008

CORTISONE ACETATE

TABLET; ORAL

CORTISONE ACETATE

! HIKMA INTL PHARMS 25MG

A080776 002

COSYNTROPIN

INJECTABLE; INJECTION

CORTROSYN**AP** +! AMPHASTAR PHARMS **0.25MG/VIAL****N016750 001**

INC

COSYNTROPIN**AP** MYLAN INSTITUTIONAL **0.25MG/VIAL****A090574 001** Dec 17, 2009**AP** SANDOZ **0.25MG/VIAL****A202147 001** Jun 29, 2012CRISABOROLE

OINTMENT; TOPICAL

EUCRISA

+! ANACOR PHARMS INC 2%

N207695 001 Dec 14, 2016

CRIZOTINIB

CAPSULE; ORAL

XALKORI

+ PF PRISM CV 200MG

N202570 001 Aug 26, 2011

+! 250MG

N202570 002 Aug 26, 2011

CROFELEMER

TABLET, DELAYED RELEASE; ORAL

MYTESI

+! NAPO PHARMS INC 125MG

N202292 001 Dec 31, 2012

CROMOLYN SODIUM

CONCENTRATE; ORAL

CROMOLYN SODIUM**AA** AILEX PHARMS LLC **100MG/5ML****A209264 001** Oct 16, 2017**AA** MICRO LABS LTD **100MG/5ML****A202745 001** Apr 04, 2013

INDIA

AA RISING PHARMS **100MG/5ML****A202583 001** Oct 27, 2011GASTROCROM**AA** +! MYLAN SPECIALITY LP **100MG/5ML****N020479 001** Feb 29, 1996

SOLUTION; INHALATION

CROMOLYN SODIUM**AN** AILEX PHARMS LLC **10MG/ML****A209453 001** Oct 16, 2017**AN** MYLAN SPECIALITY LP **10MG/ML****A074209 001** Apr 26, 1994**AN** ! TEVA PHARMS **10MG/ML****A075271 001** Jan 18, 2000**AN** WOCKHARDT BIO AG **10MG/ML****A075346 001** Oct 25, 1999

SOLUTION/DROPS; OPHTHALMIC

CROMOLYN SODIUM**AT** ! AKORN **4%****A074706 001** Apr 29, 1998**AT** SANDOZ INC **4%****A075282 001** Jun 16, 1999

PRESCRIPTION DRUG PRODUCT LIST

CROTAMITON

CREAM; TOPICAL

EURAX

+! SUN PHARM INDS INC 10%

N006927 001

LOTION; TOPICAL

CROTAN**AT** MARNEL PHARMS 10%**A087204 001**EURAX**AT** +! SUN PHARM INDS INC 10%**N009112 003**CUPRIC CHLORIDE

INJECTABLE; INJECTION

CUPRIC CHLORIDE IN PLASTIC CONTAINER

+! HOSPIRA EQ 0.4MG COPPER/ML

N018960 001 Jun 26, 1986

CYANOCOBALAMIN

INJECTABLE; INJECTION

CYANOCOBALAMIN**AP** +! LUITPOLD 1MG/ML**A080737 001****AP** MYLAN LABS LTD 1MG/ML**A204829 001** Jun 05, 2017**AP** SOMERSET THERAPS
LLC 1MG/ML**A206503 001** Dec 11, 2015**AP** 1MG/ML**A209429 001** Dec 18, 2018**AP** VITRUVIAS THERAP 1MG/ML**A209255 001** Dec 18, 2018**AP** WEST-WARD PHARMS
INT 1MG/ML**A080515 002**VIBISONE**AP** ! FRESENIUS KABI USA 1MG/ML**A080557 003**

SPRAY, METERED; NASAL

NASCOBAL

+! ENDO PHARMS INC 0.5MG/SPRAY

N021642 001 Jan 31, 2005

CYCLOBENZAPRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

AMRIX**AB** + TEVA PHARMS INTL 15MG**N021777 001** Feb 01, 2007**AB** +! 30MG**N021777 002** Feb 01, 2007CYCLOBENZAPRINE HYDROCHLORIDE**AB** APOTEX INC 15MG**A206703 001** Jul 24, 2018**AB** 30MG**A206703 002** Jul 24, 2018

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE**AB** ACTAVIS LABS FL INC 5MG **A071611 002** Feb 03, 2006**AB** 7.5MG **A071611 003** Feb 03, 2006**AB** 10MG **A071611 001** May 03, 1989**AB** AUROBINDO PHARMA 5MG **A078643 001** Sep 26, 2008**AB** 10MG **A078643 002** Sep 26, 2008**AB** FRONTIDA BIOPHARM 5MG **A073541 002** Apr 06, 2006**AB** 10MG **A073541 001** May 23, 1995**AB** INVAGEN PHARMS 5MG **A090478 001** Jul 23, 2010**AB** 10MG **A090478 002** Jul 23, 2010**AB** JUBILANT CADISTA 5MG **A077563 001** Apr 19, 2006**AB** 7.5MG **A077563 003** Aug 25, 2017**AB** 10MG **A077563 002** Apr 19, 2006**AB** KVK TECH 5MG **A078048 001** Feb 28, 2011**AB** 10MG **A078048 002** Feb 28, 2011**AB** MYLAN PHARMS INC 5MG **A073144 002** Feb 03, 2006**AB** 7.5MG **A073144 003** Mar 25, 2013**AB** ! 10MG **A073144 001** May 30, 1991**AB** ORIT LABS LLC 5MG **A078218 002** Jun 19, 2015**AB** 10MG **A078218 001** Apr 18, 2008**AB** OXFORD PHARMS 5MG **A077209 002** Feb 03, 2006**AB** 10MG **A077209 001** Oct 04, 2005**AB** PLIVA 10MG **A074421 001** Sep 29, 1995**AB** PRINSTON INC 5MG **A077797 001** Feb 28, 2007**AB** 10MG **A077797 002** Feb 28, 2007**AB** RUBICON RES PVT LTD 5MG **A208170 001** May 31, 2017**AB** 7.5MG **A208170 002** May 31, 2017**AB** 10MG **A208170 003** May 31, 2017**AB** SUN PHARM INDS LTD 5MG **A078722 001** May 12, 2008**AB** 7.5MG **A078722 002** May 12, 2008**AB** 10MG **A078722 003** May 12, 2008

PRESCRIPTION DRUG PRODUCT LIST

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

AKPENTOLATE

| | | | | | |
|-----------|-------|-----------|----------------|------------|--------------|
| <u>AT</u> | AKORN | <u>1%</u> | <u>A040164</u> | <u>001</u> | Jan 13, 1997 |
|-----------|-------|-----------|----------------|------------|--------------|

CYCLOGYL

| | | | | | |
|-----------|---|----------------|-------------|----------------|------------|
| <u>AT</u> | ! | ALCON LABS INC | <u>0.5%</u> | <u>A084109</u> | <u>001</u> |
|-----------|---|----------------|-------------|----------------|------------|

| | | | | | |
|-----------|---|--|-----------|----------------|------------|
| <u>AT</u> | ! | | <u>1%</u> | <u>A084110</u> | <u>001</u> |
|-----------|---|--|-----------|----------------|------------|

CYCLOPENTOLATE HYDROCHLORIDE

| | | | | | | |
|-----------|--|-----------|-------------|----------------|------------|--------------|
| <u>AT</u> | | AKORN INC | <u>0.5%</u> | <u>A205937</u> | <u>001</u> | Dec 09, 2015 |
|-----------|--|-----------|-------------|----------------|------------|--------------|

PENTOLAIR

| | | | | | | |
|-----------|--|-----------------|-----------|----------------|------------|--------------|
| <u>AT</u> | | BAUSCH AND LOMB | <u>1%</u> | <u>A040075</u> | <u>001</u> | Apr 29, 1994 |
|-----------|--|-----------------|-----------|----------------|------------|--------------|

CYCLOGYL

| | | | | | |
|---|----------------|-----------|---------|-----|--|
| ! | ALCON LABS INC | <u>2%</u> | A084108 | 001 | |
|---|----------------|-----------|---------|-----|--|

CYCLOPENTOLATE HYDROCHLORIDE; PHENYLEPHRINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

CYCLOMYDRIL

| | | | | | |
|---|----------------|----------------|---------|-----|--|
| ! | ALCON LABS INC | <u>0.2%;1%</u> | A084300 | 001 | |
|---|----------------|----------------|---------|-----|--|

CYCLOPHOSPHAMIDE

CAPSULE;ORAL

CYCLOPHOSPHAMIDE

| | | | | | | |
|-----------|--|---------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | AMERIGEN PHARMS LTD | <u>25MG</u> | <u>A207014</u> | <u>001</u> | Mar 19, 2018 |
|-----------|--|---------------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>50MG</u> | <u>A207014</u> | <u>002</u> | Mar 19, 2018 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|----------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | STI PHARMA LLC | <u>25MG</u> | <u>A209872</u> | <u>001</u> | May 07, 2018 |
|-----------|--|----------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>50MG</u> | <u>A209872</u> | <u>002</u> | May 07, 2018 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | + | WEST-WARD PHARMS | <u>25MG</u> | <u>N203856</u> | <u>001</u> | Sep 16, 2013 |
|-----------|---|------------------|-------------|----------------|------------|--------------|

INT

| | | | | | | |
|-----------|---|---|-------------|----------------|------------|--------------|
| <u>AB</u> | + | ! | <u>50MG</u> | <u>N203856</u> | <u>002</u> | Sep 16, 2013 |
|-----------|---|---|-------------|----------------|------------|--------------|

INJECTABLE;INJECTION

CYCLOPHOSPHAMIDE

| | | | | | | |
|-----------|--|------------------|-------------------|----------------|------------|--------------|
| <u>AP</u> | | AMNEAL PHARMS CO | <u>500MG/VIAL</u> | <u>A210046</u> | <u>001</u> | May 25, 2018 |
|-----------|--|------------------|-------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-----------------|----------------|------------|--------------|
| <u>AP</u> | | | <u>1GM/VIAL</u> | <u>A210046</u> | <u>002</u> | May 25, 2018 |
|-----------|--|--|-----------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-----------------|----------------|------------|--------------|
| <u>AP</u> | | | <u>2GM/VIAL</u> | <u>A210046</u> | <u>003</u> | May 25, 2018 |
|-----------|--|--|-----------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|-----------------|-------------------|----------------|------------|--------------|
| <u>AP</u> | ! | BAXTER HLTHCARE | <u>500MG/VIAL</u> | <u>A040745</u> | <u>001</u> | May 21, 2008 |
|-----------|---|-----------------|-------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|--|-----------------|----------------|------------|--------------|
| <u>AP</u> | ! | | <u>1GM/VIAL</u> | <u>A040745</u> | <u>002</u> | May 21, 2008 |
|-----------|---|--|-----------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|--|-----------------|----------------|------------|--------------|
| <u>AP</u> | ! | | <u>2GM/VIAL</u> | <u>A040745</u> | <u>003</u> | May 21, 2008 |
|-----------|---|--|-----------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|---------------------|-------------------|----------------|------------|--------------|
| <u>AP</u> | | JIANGSU HENGRUI MED | <u>500MG/VIAL</u> | <u>A204555</u> | <u>001</u> | Oct 31, 2014 |
|-----------|--|---------------------|-------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-----------------|----------------|------------|--------------|
| <u>AP</u> | | | <u>1GM/VIAL</u> | <u>A204555</u> | <u>002</u> | Oct 31, 2014 |
|-----------|--|--|-----------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-----------------|----------------|------------|--------------|
| <u>AP</u> | | | <u>2GM/VIAL</u> | <u>A204555</u> | <u>003</u> | Oct 31, 2014 |
|-----------|--|--|-----------------|----------------|------------|--------------|

CYCLOSERINE

CAPSULE;ORAL

SEROMYCIN

| | | | | | |
|---|------------|--------------|---------|-----|--|
| ! | PURDUE GMP | <u>250MG</u> | A060593 | 001 | |
|---|------------|--------------|---------|-----|--|

CYCLOSPORINE

CAPSULE;ORAL

CYCLOSPORINE

| | | | | | | |
|------------|--|---------------|-------------|----------------|------------|--------------|
| <u>AB1</u> | | IVAX SUB TEVA | <u>25MG</u> | <u>A065110</u> | <u>003</u> | Mar 29, 2005 |
|------------|--|---------------|-------------|----------------|------------|--------------|

PHARMS

| | | | | | | |
|------------|--|--|--------------|----------------|------------|--------------|
| <u>AB1</u> | | | <u>100MG</u> | <u>A065110</u> | <u>002</u> | Mar 29, 2005 |
|------------|--|--|--------------|----------------|------------|--------------|

| | | | | | | |
|------------|--|--------------|-------------|----------------|------------|--------------|
| <u>AB1</u> | | MAYNE PHARMA | <u>25MG</u> | <u>A065044</u> | <u>002</u> | Dec 20, 2000 |
|------------|--|--------------|-------------|----------------|------------|--------------|

| | | | | | | |
|------------|--|--|--------------|----------------|------------|--------------|
| <u>AB1</u> | | | <u>100MG</u> | <u>A065044</u> | <u>001</u> | Dec 20, 2000 |
|------------|--|--|--------------|----------------|------------|--------------|

| | | | | | | |
|------------|--|--------|-------------|----------------|------------|--------------|
| <u>AB1</u> | | SANDOZ | <u>25MG</u> | <u>A065017</u> | <u>002</u> | Jan 13, 2000 |
|------------|--|--------|-------------|----------------|------------|--------------|

| | | | | | | |
|------------|--|--|--------------|----------------|------------|--------------|
| <u>AB1</u> | | | <u>100MG</u> | <u>A065017</u> | <u>001</u> | Jan 13, 2000 |
|------------|--|--|--------------|----------------|------------|--------------|

GENGRAF

| | | | | | | |
|------------|--|--------|-------------|----------------|------------|--------------|
| <u>AB1</u> | | ABBVIE | <u>25MG</u> | <u>A065003</u> | <u>001</u> | May 12, 2000 |
|------------|--|--------|-------------|----------------|------------|--------------|

| | | | | | | |
|------------|--|--|--------------|----------------|------------|--------------|
| <u>AB1</u> | | | <u>100MG</u> | <u>A065003</u> | <u>003</u> | May 12, 2000 |
|------------|--|--|--------------|----------------|------------|--------------|

NEORAL

| | | | | | | |
|------------|---|----------|-------------|----------------|------------|--------------|
| <u>AB1</u> | + | NOVARTIS | <u>25MG</u> | <u>N050715</u> | <u>001</u> | Jul 14, 1995 |
|------------|---|----------|-------------|----------------|------------|--------------|

| | | | | | | |
|------------|---|---|--------------|----------------|------------|--------------|
| <u>AB1</u> | + | ! | <u>100MG</u> | <u>N050715</u> | <u>002</u> | Jul 14, 1995 |
|------------|---|---|--------------|----------------|------------|--------------|

CYCLOSPORINE

| | | | | | | |
|------------|--|--------|-------------|----------------|------------|--------------|
| <u>AB2</u> | | APOTEX | <u>25MG</u> | <u>A065040</u> | <u>001</u> | May 09, 2002 |
|------------|--|--------|-------------|----------------|------------|--------------|

| | | | | | | |
|------------|--|--|--------------|----------------|------------|--------------|
| <u>AB2</u> | | | <u>100MG</u> | <u>A065040</u> | <u>002</u> | May 09, 2002 |
|------------|--|--|--------------|----------------|------------|--------------|

SANDIMMUNE

| | | | | | | |
|------------|---|----------|-------------|----------------|------------|--------------|
| <u>AB2</u> | + | NOVARTIS | <u>25MG</u> | <u>N050625</u> | <u>001</u> | Mar 02, 1990 |
|------------|---|----------|-------------|----------------|------------|--------------|

| | | | | | | |
|------------|---|---|--------------|----------------|------------|--------------|
| <u>AB2</u> | + | ! | <u>100MG</u> | <u>N050625</u> | <u>002</u> | Mar 02, 1990 |
|------------|---|---|--------------|----------------|------------|--------------|

| | | | | | | |
|----|---|--|-------------|---------|-----|--------------|
| BX | + | | <u>50MG</u> | N050625 | 003 | Nov 23, 1992 |
|----|---|--|-------------|---------|-----|--------------|

CYCLOSPORINE

| | | | | | | |
|--|--|---------------|-------------|---------|-----|--------------|
| | | IVAX SUB TEVA | <u>50MG</u> | A065110 | 001 | Mar 29, 2005 |
| | | PHARMS | | | | |

PRESCRIPTION DRUG PRODUCT LIST

CYCLOSPORINE

EMULSION;OPHTHALMIC

RESTASIS

+! ALLERGAN 0.05% N050790 001 Dec 23, 2002

RESTASIS MULTIDOSE

+! ALLERGAN 0.05% N050790 002 Oct 27, 2016

INJECTABLE; INJECTION

CYCLOSPORINE**AP** LUITPOLD **50MG/ML** **A065151 001** Oct 07, 2003**AP** WEST-WARD PHARMS **50MG/ML** **A065004 001** Oct 29, 1999

INT

SANDIMMUNE**AP** +! NOVARTIS **50MG/ML** **N050573 001** Nov 14, 1983

SOLUTION;OPHTHALMIC

CEQUA

+! SUN PHARMA GLOBAL 0.09% N210913 001 Aug 14, 2018

SOLUTION;ORAL

CYCLOSPORINE**AB1** ABBVIE **100MG/ML** **A065025 001** Mar 03, 2000**AB1** IVAX SUB TEVA **100MG/ML** **A065078 001** Mar 25, 2005

PHARMS

AB1 MAYNE PHARMA **100MG/ML** **A065054 001** Dec 18, 2001NEORAL**AB1** +! NOVARTIS **100MG/ML** **N050716 001** Jul 14, 1995CYCLOSPORINE**AB2** WOCKHARDT BIO AG **100MG/ML** **A065133 001** Sep 17, 2004SANDIMMUNE**AB2** +! NOVARTIS **100MG/ML** **N050574 001** Nov 14, 1983CYPROHEPTADINE HYDROCHLORIDE

SYRUP;ORAL

CYPROHEPTADINE HYDROCHLORIDE**AA** BIO-PHARM INC **2MG/5ML** **A204823 001** Dec 27, 2016**AA** INVATECH PHARMA **2MG/5ML** **A209108 001** Oct 16, 2018**AA** LANNETT CO INC **2MG/5ML** **A203191 001** Jul 13, 2017**AA** ! LYNE **2MG/5ML** **A040668 001** Jun 28, 2006**AA** PHARM ASSOC **2MG/5ML** **A091295 001** Mar 28, 2013

TABLET;ORAL

CYPROHEPTADINE HYDROCHLORIDE**AA** APEX PHARMS INC **4MG** **A207783 001** Dec 29, 2016**AA** APNAR PHARMA LP **4MG** **A207555 001** Jan 31, 2017**AA** BOSCOGEN **4MG** **A040644 001** May 30, 2006**AA** ! IVAX SUB TEVA **4MG** **A087056 001**

PHARMS

AA MOUNTAIN **4MG** **A040537 001** Sep 30, 2003**AA** NOVAST LABS **4MG** **A205087 001** Sep 23, 2015**AA** PAR PHARM **4MG** **A087129 001****AA** STRIDES PHARMA **4MG** **A209172 001** Apr 11, 2018**AA** TWI PHARMS **4MG** **A206553 001** Nov 29, 2016**AA** ZYDUS PHARMS USA **4MG** **A208938 001** May 19, 2017

INC

CYSTEAMINE BITARTRATE

CAPSULE;ORAL

CYSTAGON

+ MYLAN EQ 50MG BASE N020392 001 Aug 15, 1994

+! EQ 150MG BASE N020392 002 Aug 15, 1994

CAPSULE, DELAYED RELEASE;ORAL

PROCSBI

+ HORIZON PHARMA USA EQ 25MG BASE N203389 001 Apr 30, 2013

+! EQ 75MG BASE N203389 002 Apr 30, 2013

CYSTEAMINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

CYSTARAN

+! LEADIANT BIOSCI INC EQ 0.44% BASE N200740 001 Oct 02, 2012

CYTARABINE

INJECTABLE; INJECTION

CYTARABINE**AP** ! FRESENIUS KABI USA **100MG/ML** **A076512 001** Jan 15, 2004**AP** HONG KONG **20MG/ML** **A206190 001** Nov 09, 2017**AP** **100MG/ML** **A205696 001** Jul 17, 2018**AP** ! HOSPIRA **20MG/ML** **A071868 001** Jun 04, 1990**AP** ! **20MG/ML** **A072168 001** Aug 31, 1990

PRESCRIPTION DRUG PRODUCT LIST

CYTARABINE

INJECTABLE; INJECTION

CYTARABINE

| | | | | | | |
|-----------|---|------------------|-----------------|----------------|------------|--------------|
| AP | ! | | <u>20MG/ML</u> | <u>A072945</u> | <u>001</u> | Feb 28, 1994 |
| AP | | | <u>100MG/ML</u> | <u>A075383</u> | <u>001</u> | Nov 22, 1999 |
| AP | | MYLAN LABS LTD | <u>20MG/ML</u> | <u>A200914</u> | <u>001</u> | Dec 13, 2011 |
| AP | | | <u>20MG/ML</u> | <u>A200915</u> | <u>001</u> | Dec 13, 2011 |
| AP | | | <u>20MG/ML</u> | <u>A200916</u> | <u>001</u> | Dec 13, 2011 |
| AP | | | <u>100MG/ML</u> | <u>A201784</u> | <u>001</u> | Jan 30, 2012 |
| | | WEST-WARD PHARMS | 100MG/VIAL | A071471 | 001 | Aug 02, 1989 |
| | | INT | | | | |
| | ! | | 500MG/VIAL | A071472 | 001 | Aug 02, 1989 |
| | ! | | 1GM/VIAL | A074245 | 001 | Aug 31, 1994 |
| | ! | | 2GM/VIAL | A074245 | 002 | Aug 31, 1994 |

CYTARABINE; DAUNORUBICIN

POWDER; INTRAVENOUS

VYXEOS

| | | | | | | |
|---|---|----------------|------------|---------|-----|--------------|
| + | ! | CELATOR PHARMS | 100MG;44MG | N209401 | 001 | Aug 03, 2017 |
|---|---|----------------|------------|---------|-----|--------------|

DABIGATRAN ETEXILATE MESYLATE

CAPSULE; ORAL

PRADAXA

| | | | | | | |
|---|---|------------|---------------|---------|-----|--------------|
| + | | BOEHRINGER | EQ 75MG BASE | N022512 | 001 | Oct 19, 2010 |
| | | INGELHEIM | | | | |
| + | | | EQ 110MG BASE | N022512 | 003 | Nov 20, 2015 |
| + | ! | | EQ 150MG BASE | N022512 | 002 | Oct 19, 2010 |

DABRAFENIB MESYLATE

CAPSULE; ORAL

TAFINLAR

| | | | | | | |
|---|---|-----------------|--------------|---------|-----|--------------|
| + | | NOVARTIS PHARMS | EQ 50MG BASE | N202806 | 001 | May 29, 2013 |
| | | CORP | | | | |
| + | ! | | EQ 75MG BASE | N202806 | 002 | May 29, 2013 |

DACARBAZINE

INJECTABLE; INJECTION

DACARBAZINE

| | | | | | | |
|-----------|---|--------------------|-------------------|----------------|------------|--------------|
| AP | ! | FRESENIUS KABI USA | <u>200MG/VIAL</u> | <u>A075371</u> | <u>002</u> | Aug 27, 1999 |
| AP | | HOSPIRA | <u>200MG/VIAL</u> | <u>A075940</u> | <u>001</u> | Oct 18, 2001 |
| AP | | TEVA PHARMS USA | <u>200MG/VIAL</u> | <u>A075259</u> | <u>002</u> | Aug 27, 1998 |
| AP | ! | | <u>500MG/VIAL</u> | <u>A075259</u> | <u>001</u> | Sep 22, 2000 |
| AP | | WEST-WARD PHARMS | <u>200MG/VIAL</u> | <u>A075812</u> | <u>001</u> | Jun 15, 2001 |
| | | INT | | | | |
| AP | | | <u>500MG/VIAL</u> | <u>A075812</u> | <u>002</u> | Oct 31, 2002 |
| | ! | FRESENIUS KABI USA | 100MG/VIAL | A075371 | 001 | Aug 27, 1999 |

DACLATASVIR DIHYDROCHLORIDE

TABLET; ORAL

DAKLINZA

| | | | | | | |
|---|---|---------------|--------------|---------|-----|--------------|
| + | | BRISTOL-MYERS | EQ 30MG BASE | N206843 | 001 | Jul 24, 2015 |
| | | SQUIBB | | | | |
| + | | | EQ 60MG BASE | N206843 | 002 | Jul 24, 2015 |
| + | ! | | EQ 90MG BASE | N206843 | 003 | Apr 13, 2016 |

DACOMITINIB

TABLET; ORAL

VIZIMPRO

| | | | | | | |
|---|---|------------|------|---------|-----|--------------|
| + | | PFIZER INC | 15MG | N211288 | 001 | Sep 27, 2018 |
| + | | | 30MG | N211288 | 002 | Sep 27, 2018 |
| + | ! | | 45MG | N211288 | 003 | Sep 27, 2018 |

DACTINOMYCIN

INJECTABLE; INJECTION

COSMEGEN

| | | | | | | |
|-----------|---|---|----------------|-------------------|----------------|--------------|
| AP | + | ! | RECORDATI RARE | <u>0.5MG/VIAL</u> | <u>N050682</u> | <u>001</u> |
| AP | | | | <u>0.5MG/VIAL</u> | <u>A202562</u> | <u>001</u> |
| | | | LUITPOLD | | | Aug 23, 2013 |
| AP | | | | <u>0.5MG/VIAL</u> | <u>A203385</u> | <u>001</u> |
| | | | MYLAN LABS LTD | | | Nov 09, 2017 |

DALBAVANCIN HYDROCHLORIDE

POWDER; INTRAVENOUS

DALVANCE

| | | | | | | |
|---|---|--------------------|--------------------|---------|-----|--------------|
| + | ! | ALLERGAN SALES LLC | EQ 500MG BASE/VIAL | N021883 | 001 | May 23, 2014 |
|---|---|--------------------|--------------------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

DALFAMPRIDINE

TABLET, EXTENDED RELEASE;ORAL

AMPYRA

| | | | | | | |
|-----------|------------|--------|-------------|----------------|------------|--------------|
| AB | + ! | ACORDA | 10MG | N022250 | 001 | Jan 22, 2010 |
|-----------|------------|--------|-------------|----------------|------------|--------------|

DALFAMPRIDINE

| | | | | | | |
|-----------|--|----------------------|-------------|----------------|------------|--------------|
| AB | | ACCORD HLTHCARE | 10MG | A206863 | 001 | Jul 11, 2018 |
| AB | | ACTAVIS LABS FL INC | 10MG | A206836 | 001 | Jan 23, 2017 |
| AB | | ALKEM LABS LTD | 10MG | A206765 | 001 | Jul 30, 2018 |
| AB | | AUROBINDO PHARMA LTD | 10MG | A206811 | 001 | Jan 23, 2017 |
| AB | | WEST-WARD PHARMS INT | 10MG | A206646 | 001 | Oct 24, 2018 |

DALFOPRISTIN; QUINUPRISTININJECTABLE; INTRAVENOUS
SYNERCID

| | | | | | |
|------------|-------------|-----------------------|---------|-----|--------------|
| + ! | KING PHARMS | 350MG/VIAL;150MG/VIAL | N050748 | 001 | Sep 21, 1999 |
|------------|-------------|-----------------------|---------|-----|--------------|

DALTEPARIN SODIUMINJECTABLE; SUBCUTANEOUS
FRAGMIN

| | | | | | |
|------------|------------|-------------------------------|---------|-----|--------------|
| + | PFIZER INC | 2,500IU/0.2ML (12,500IU/ML) | N020287 | 001 | Dec 22, 1994 |
| + | | 5,000IU/0.2ML (25,000IU/ML) | N020287 | 003 | Mar 18, 1996 |
| + | | 7,500IU/0.3ML (25,000IU/ML) | N020287 | 005 | Apr 04, 2002 |
| + | | 10,000IU/ML (10,000IU/ML) | N020287 | 004 | Jan 30, 1998 |
| + | | 12,500IU/0.5ML (25,000IU/ML) | N020287 | 009 | May 01, 2007 |
| + | | 15,000IU/0.6ML (25,000IU/ML) | N020287 | 010 | May 01, 2007 |
| + | | 18,000IU/0.72ML (25,000IU/ML) | N020287 | 011 | May 01, 2007 |
| + ! | | 95,000IU/3.8ML (25,000IU/ML) | N020287 | 006 | Apr 04, 2002 |

DANAZOL

CAPSULE; ORAL

DANAZOL

| | | | | | | |
|-----------|----------|----------------|--------------|----------------|------------|--------------|
| AB | | BARR | 50MG | A074582 | 003 | May 29, 1998 |
| AB | | | 100MG | A074582 | 002 | May 29, 1998 |
| AB | ! | | 200MG | A074582 | 001 | Aug 09, 1996 |
| AB | | LANNETT CO INC | 50MG | A077246 | 002 | Apr 19, 2007 |
| AB | | | 100MG | A077246 | 003 | Apr 19, 2007 |
| AB | | | 200MG | A077246 | 001 | Sep 28, 2005 |

DANTROLENE SODIUM

CAPSULE; ORAL

DANTRIUUM

| | | | | | | |
|-----------|------------|----------------------|--------------|----------------|------------|--|
| AB | + | PAR STERILE PRODUCTS | 25MG | N017443 | 001 | |
| AB | + | | 50MG | N017443 | 003 | |
| AB | + ! | | 100MG | N017443 | 002 | |

DANTROLENE SODIUM

| | | | | | | |
|-----------|--|----------------|--------------|----------------|------------|--------------|
| AB | | ELITE LABS INC | 25MG | A076686 | 001 | Oct 24, 2005 |
| AB | | | 50MG | A076686 | 002 | Oct 24, 2005 |
| AB | | | 100MG | A076686 | 003 | Oct 24, 2005 |
| AB | | IMPAX LABS | 25MG | A076856 | 001 | Mar 01, 2005 |
| AB | | | 50MG | A076856 | 002 | Mar 01, 2005 |
| AB | | | 100MG | A076856 | 003 | Mar 01, 2005 |

FOR SUSPENSION; INTRAVENOUS

RYANODEX

| | | | | | |
|------------|--------------|------------|---------|-----|--------------|
| + ! | EAGLE PHARMS | 250MG/VIAL | N205579 | 001 | Jul 22, 2014 |
|------------|--------------|------------|---------|-----|--------------|

INJECTABLE; INJECTION

DANTRIUUM

| | | | | | | |
|-----------|------------|----------------------|------------------|----------------|------------|--|
| AP | + ! | PAR STERILE PRODUCTS | 20MG/VIAL | N018264 | 001 | |
|-----------|------------|----------------------|------------------|----------------|------------|--|

DANTROLENE SODIUM

| | | | | | | |
|-----------|--|---------------------|------------------|----------------|------------|--------------|
| AP | | HIKMA PHARMS | 20MG/VIAL | A204762 | 001 | Jun 19, 2017 |
| AP | | MYLAN INSTITUTIONAL | 20MG/VIAL | A205239 | 001 | Feb 18, 2016 |

REVONTO

| | | | | | | |
|-----------|--|--------------|------------------|----------------|------------|--------------|
| AP | | US WORLDMEDS | 20MG/VIAL | A078378 | 001 | Jul 24, 2007 |
|-----------|--|--------------|------------------|----------------|------------|--------------|

DAPAGLIFLOZIN

TABLET; ORAL

FARXIGA

| | | | | | |
|------------|----------------|------|---------|-----|--------------|
| + | ASTRAZENECA AB | 5MG | N202293 | 001 | Jan 08, 2014 |
| + ! | | 10MG | N202293 | 002 | Jan 08, 2014 |

PRESCRIPTION DRUG PRODUCT LIST

DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL
XIGDUO XR

| | | | | | |
|---|----------------|-------------|---------|-----|--------------|
| + | ASTRAZENECA AB | 2.5MG; 1GM | N205649 | 005 | Jul 28, 2017 |
| + | | 5MG; 500MG | N205649 | 001 | Oct 29, 2014 |
| + | | 5MG; 1GM | N205649 | 002 | Oct 29, 2014 |
| + | | 10MG; 500MG | N205649 | 003 | Oct 29, 2014 |
| + | ! | 10MG; 1GM | N205649 | 004 | Oct 29, 2014 |

DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE

TABLET; ORAL
QTERN

| | | | | | |
|---|----------------|-------------------|---------|-----|--------------|
| + | ASTRAZENECA AB | 10MG; EQ 5MG BASE | N209091 | 001 | Feb 27, 2017 |
|---|----------------|-------------------|---------|-----|--------------|

DAPSONE

GEL; TOPICAL

ACZONE

| | | | | | | |
|-----------|---|----------|-----------|----------------|------------|--------------|
| AB | + | ALLERGAN | 5% | N021794 | 001 | Jul 07, 2005 |
|-----------|---|----------|-----------|----------------|------------|--------------|

DAPSONE

| | | | | | | |
|-----------|--|------|-----------|----------------|------------|--------------|
| AB | | TARO | 5% | A209506 | 001 | Oct 16, 2017 |
|-----------|--|------|-----------|----------------|------------|--------------|

ACZONE

| | | | | | |
|---|-----------------|------|---------|-----|--------------|
| + | AQUA PHARMS LLC | 7.5% | N207154 | 001 | Feb 24, 2016 |
|---|-----------------|------|---------|-----|--------------|

TABLET; ORAL

DAPSONE

| | | | | | | |
|-----------|---|------------------|--------------|----------------|------------|--------------|
| AB | | ACTAVIS LLC | 25MG | A204380 | 001 | Mar 23, 2017 |
| AB | | | 100MG | A204380 | 002 | Mar 23, 2017 |
| AB | | ALVOGEN | 25MG | A205429 | 001 | Jan 07, 2016 |
| AB | | | 100MG | A205429 | 002 | Jan 07, 2016 |
| AB | | JACOBUS | 25MG | A086841 | 001 | |
| AB | ! | | 100MG | A086842 | 001 | |
| AB | | NOSTRUM LABS INC | 25MG | A203887 | 001 | May 06, 2016 |
| AB | | | 100MG | A203887 | 002 | May 06, 2016 |
| AB | | NOVITIUM PHARMA | 25MG | A206505 | 001 | Dec 01, 2016 |
| AB | | | 100MG | A206505 | 002 | Dec 01, 2016 |
| AB | | VIRTUS PHARMS | 25MG | A204074 | 001 | May 10, 2016 |
| AB | | | 100MG | A204074 | 002 | May 10, 2016 |

DAPTOMYCIN

POWDER; INTRAVENOUS

CUBICIN

| | | | | | | |
|-----------|---|-------------------|-------------------|----------------|------------|--------------|
| AP | + | CUBIST PHARMS LLC | 500MG/VIAL | N021572 | 002 | Sep 12, 2003 |
|-----------|---|-------------------|-------------------|----------------|------------|--------------|

DAPTOMYCIN

| | | | | | | |
|-----------|--|--------------------|-------------------|----------------|------------|--------------|
| AP | | CRANE PHARMS LLC | 500MG/VIAL | A206005 | 001 | Jun 15, 2016 |
| AP | | FRESENIUS KABI USA | 500MG/VIAL | A206077 | 001 | Apr 11, 2018 |
| AP | | HOSPIRA INC | 500MG/VIAL | A202857 | 001 | Sep 12, 2014 |
| AP | | MYLAN LABS LTD | 500MG/VIAL | A205037 | 001 | Jun 05, 2018 |
| AP | | TEVA PHARMS USA | 500MG/VIAL | A091039 | 001 | Mar 25, 2016 |

CUBICIN RF

| | | | | | |
|---|-------------------|------------|---------|-----|--------------|
| + | CUBIST PHARMS LLC | 500MG/VIAL | N021572 | 003 | Jul 06, 2016 |
|---|-------------------|------------|---------|-----|--------------|

POWDER; IV (INFUSION)

DAPTOMYCIN

| | | | | | |
|---|-------------------|------------|---------|-----|--------------|
| + | SAGENT PHARMS | 350MG/VIAL | N208385 | 001 | Sep 12, 2017 |
| + | XELLIA PHARMS APS | 350MG/VIAL | N209949 | 001 | Oct 20, 2017 |

DARIFENACIN HYDROBROMIDE

TABLET, EXTENDED RELEASE; ORAL

DARIFENACIN

| | | | | | | |
|-----------|--|---------------------|----------------------|----------------|------------|--------------|
| AB | | MACLEODS PHARMS LTD | EQ 7.5MG BASE | A207302 | 001 | Jul 28, 2017 |
| AB | | | EQ 15MG BASE | A207302 | 002 | Jul 28, 2017 |

DARIFENACIN HYDROBROMIDE

| | | | | | | |
|-----------|--|----------------------|----------------------|----------------|------------|--------------|
| AB | | ALEMBIC PHARMS LTD | EQ 7.5MG BASE | A207681 | 001 | Dec 08, 2017 |
| AB | | | EQ 15MG BASE | A207681 | 002 | Dec 08, 2017 |
| AB | | ANCHEN PHARMS | EQ 7.5MG BASE | A091190 | 001 | Mar 13, 2015 |
| AB | | | EQ 15MG BASE | A091190 | 002 | Mar 13, 2015 |
| AB | | AUROBINDO PHARMA LTD | EQ 7.5MG BASE | A206743 | 001 | Sep 19, 2016 |
| AB | | | EQ 15MG BASE | A206743 | 002 | Sep 19, 2016 |
| AB | | CIPLA | EQ 7.5MG BASE | A207664 | 001 | Sep 01, 2016 |
| AB | | | EQ 15MG BASE | A207664 | 002 | Sep 01, 2016 |
| AB | | JUBILANT GENERICS | EQ 7.5MG BASE | A205550 | 001 | Oct 12, 2016 |
| AB | | | EQ 15MG BASE | A205550 | 002 | Oct 12, 2016 |
| AB | | TORRENT PHARMS LTD | EQ 7.5MG BASE | A205209 | 001 | Nov 17, 2016 |
| AB | | | EQ 15MG BASE | A205209 | 002 | Nov 17, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

DARIFENACIN HYDROBROMIDE

TABLET, EXTENDED RELEASE;ORAL

ENABLEX

| | | | | | |
|-----------|---|------|----------------------|--------------------|--------------|
| AB | + | APIL | EQ 7.5MG BASE | N021513 001 | Dec 22, 2004 |
| AB | + | ! | EQ 15MG BASE | N021513 002 | Dec 22, 2004 |

DARUNAVIR ETHANOLATE

SUSPENSION;ORAL

PREZISTA

| | | | | |
|---|---------------|------------------|-------------|--------------|
| + | JANSSEN PRODS | EQ 100MG BASE/ML | N202895 001 | Dec 16, 2011 |
|---|---------------|------------------|-------------|--------------|

TABLET;ORAL

DARUNAVIR ETHANOLATE

| | | | | | |
|-----------|--|-----------------|----------------------|--------------------|--------------|
| AB | | TEVA PHARMS USA | EQ 600MG BASE | A202118 001 | Nov 21, 2017 |
|-----------|--|-----------------|----------------------|--------------------|--------------|

PREZISTA

| | | | | | |
|-----------|---|---------------|----------------------|--------------------|--------------|
| AB | + | JANSSEN PRODS | EQ 600MG BASE | N021976 002 | Feb 25, 2008 |
| | + | | EQ 75MG BASE | N021976 004 | Dec 18, 2008 |
| | + | | EQ 150MG BASE | N021976 005 | Dec 18, 2008 |
| | + | ! | EQ 800MG BASE | N021976 006 | Nov 09, 2012 |

DASABUVIR SODIUM ; OMBITASVIR; PARITAPREVIR; RITONAVIR

TABLET, TABLET;ORAL

VIEKIRA PAK (COPACKAGED)

| | | | | |
|---|------------|----------------------------------------------------|-------------|--------------|
| + | ABBVIE INC | EQ 250MG BASE,N/A,N/A,N/A; N/A,12.5MG,75MG,50MG | N206619 001 | Dec 19, 2014 |
|---|------------|----------------------------------------------------|-------------|--------------|

DASABUVIR SODIUM; OMBITASVIR; PARITAPREVIR; RITONAVIR

TABLET, EXTENDED RELEASE;ORAL

VIEKIRA XR

| | | | | |
|---|------------|-----------------------------------|-------------|--------------|
| + | ABBVIE INC | EQ 200MG BASE;8.33MG;50MG;33.33MG | N208624 001 | Jul 22, 2016 |
|---|------------|-----------------------------------|-------------|--------------|

DASATINIB

TABLET;ORAL

SPRYCEL

| | | | | |
|---|-------------------------|-------|-------------|--------------|
| + | BRISTOL MYERS SQUIBB | 20MG | N021986 001 | Jun 28, 2006 |
| + | | 50MG | N021986 002 | Jun 28, 2006 |
| + | | 70MG | N021986 003 | Jun 28, 2006 |
| + | | 80MG | N021986 005 | Oct 28, 2010 |
| + | ! | 100MG | N021986 004 | May 30, 2008 |
| + | | 140MG | N021986 006 | Oct 28, 2010 |

DAUNORUBICIN HYDROCHLORIDE

INJECTABLE;INJECTION

CERUBIDINE

| | | | | | |
|-----------|---|-------------------------|--------------------------|--------------------|--------------|
| AP | ! | WEST-WARD PHARMS INT | EQ 20MG BASE/VIAL | A064103 001 | Feb 03, 1995 |
|-----------|---|-------------------------|--------------------------|--------------------|--------------|

DAUNORUBICIN HYDROCHLORIDE

| | | | | | |
|-----------|---|-------------------------|--------------------------|--------------------|--------------|
| AP | | FRESENIUS KABI USA | EQ 20MG BASE/VIAL | A065000 001 | May 25, 1999 |
| AP | | TEVA PHARMS USA | EQ 5MG BASE/ML | A065035 001 | Jan 24, 2000 |
| AP | + | WEST-WARD PHARMS INT | EQ 5MG BASE/ML | N050731 001 | Jan 30, 1998 |
| | | FRESENIUS KABI USA | EQ 5MG BASE/VIAL | A065034 001 | Nov 20, 2001 |

DECITABINE

INJECTABLE;INTRAVENOUS

DACOGEN

| | | | | | |
|-----------|---|---------------------|------------------|--------------------|--------------|
| AP | + | OTSUKA PHARM CO LTD | 50MG/VIAL | N021790 001 | May 02, 2006 |
|-----------|---|---------------------|------------------|--------------------|--------------|

DECITABINE

| | | | | | |
|-----------|--|--------------------|------------------|--------------------|--------------|
| AP | | ACCORD HLTHCARE | 50MG/VIAL | A203475 001 | Feb 27, 2017 |
| AP | | CHEMI SPA | 50MG/VIAL | A206033 001 | Sep 22, 2017 |
| AP | | CIPLA | 50MG/VIAL | A208601 001 | Nov 16, 2017 |
| AP | | DR REDDYS LABS LTD | 50MG/VIAL | A203131 001 | Jul 11, 2013 |
| AP | | LUPIN LTD | 50MG/VIAL | A210756 001 | Nov 09, 2018 |
| AP | | PHARMASCIENCE INC | 50MG/VIAL | A204607 001 | May 31, 2017 |
| AP | | SAGENT PHARMS | 50MG/VIAL | A207100 001 | Mar 16, 2018 |
| AP | | SANDOZ INC | 50MG/VIAL | A202969 001 | Aug 28, 2014 |

POWDER;INTRAVENOUS

DECITABINE

| | | | | |
|---|-------------------|-----------|-------------|--------------|
| + | SUN PHARMA GLOBAL | 50MG/VIAL | N205582 001 | Jan 28, 2014 |
|---|-------------------|-----------|-------------|--------------|

DEFERASIROX

GRANULE;ORAL

JADENU SPRINKLE

| | | | | |
|---|-------------------------|-------|-------------|--------------|
| + | NOVARTIS PHARMS CORP | 90MG | N207968 001 | May 18, 2017 |
| + | | 180MG | N207968 002 | May 18, 2017 |
| + | ! | 360MG | N207968 003 | May 18, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

DEFERASIROX

TABLET; ORAL

JADENU

| | | | | |
|---|-------------------------|-------|-------------|--------------|
| + | NOVARTIS PHARMS CORP | 90MG | N206910 001 | Mar 30, 2015 |
| + | | 180MG | N206910 002 | Mar 30, 2015 |
| + | ! | 360MG | N206910 003 | Mar 30, 2015 |

TABLET, FOR SUSPENSION; ORAL

DEFERASIROX

| | | | | |
|-----------|-------------------|--------------|--------------------|--------------|
| AB | ACTAVIS ELIZABETH | 125MG | A203560 001 | Jan 26, 2016 |
| AB | | 250MG | A203560 002 | Jan 26, 2016 |
| AB | | 500MG | A203560 003 | Jan 26, 2016 |

EXJADE

| | | | | | |
|-----------|---|----------|--------------|--------------------|--------------|
| AB | + | NOVARTIS | 125MG | N021882 001 | Nov 02, 2005 |
| AB | + | | 250MG | N021882 002 | Nov 02, 2005 |
| AB | + | ! | 500MG | N021882 003 | Nov 02, 2005 |

DEFERIPRONE

SOLUTION; ORAL

FERRIPROX

| | | | | |
|---|--------------|----------|-------------|--------------|
| + | AOPHARMA INC | 80MG/ML | N208030 002 | Apr 20, 2018 |
| + | ! | 100MG/ML | N208030 001 | Sep 09, 2015 |

TABLET; ORAL

FERRIPROX

| | | | | |
|---|--------------|-------|-------------|--------------|
| + | AOPHARMA INC | 500MG | N021825 001 | Oct 14, 2011 |
|---|--------------|-------|-------------|--------------|

DEFEROXAMINE MESYLATE

INJECTABLE; INJECTION

DEFEROXAMINE MESYLATE

| | | | | |
|-----------|-------------------------|-------------------|--------------------|--------------|
| AP | FRESENIUS KABI USA | 500MG/VIAL | A078718 001 | Sep 15, 2009 |
| AP | | 2GM/VIAL | A078718 002 | Sep 15, 2009 |
| AP | GLAND PHARMA LTD | 500MG/VIAL | A207384 001 | Sep 29, 2017 |
| AP | | 2GM/VIAL | A207384 002 | Sep 29, 2017 |
| AP | HOSPIRA | 500MG/VIAL | A076019 001 | Mar 17, 2004 |
| AP | | 2GM/VIAL | A076019 002 | Mar 17, 2004 |
| AP | WEST-WARD PHARMS INT | 500MG/VIAL | A078086 001 | May 30, 2007 |
| AP | | 2GM/VIAL | A078086 002 | May 30, 2007 |

DESFERAL

| | | | | | |
|-----------|---|----------|-------------------|--------------------|--------------|
| AP | + | NOVARTIS | 500MG/VIAL | N016267 001 | |
| AP | + | ! | 2GM/VIAL | N016267 002 | May 25, 2000 |

DEFIBROTIDE SODIUM

SOLUTION; INTRAVENOUS

DEFITELIO

| | | | | |
|---|-----------------|-----------------------|-------------|--------------|
| + | JAZZ PHARMS INC | 200MG/2.5ML (80MG/ML) | N208114 001 | Mar 30, 2016 |
|---|-----------------|-----------------------|-------------|--------------|

DEFLAZACORT

SUSPENSION; ORAL

EMFLAZA

| | | | | |
|---|------------|------------|-------------|--------------|
| + | PTC THERAP | 22.75MG/ML | N208685 001 | Feb 09, 2017 |
|---|------------|------------|-------------|--------------|

TABLET; ORAL

EMFLAZA

| | | | | |
|---|------------|------|-------------|--------------|
| + | PTC THERAP | 6MG | N208684 001 | Feb 09, 2017 |
| + | | 18MG | N208684 002 | Feb 09, 2017 |
| + | | 30MG | N208684 003 | Feb 09, 2017 |
| + | ! | 36MG | N208684 004 | Feb 09, 2017 |

DEGARELIX ACETATE

POWDER; SUBCUTANEOUS

FIRMAGON

| | | | | |
|---|---------|--------------------|-------------|--------------|
| + | FERRING | EQ 80MG BASE/VIAL | N022201 001 | Dec 24, 2008 |
| + | ! | EQ 120MG BASE/VIAL | N022201 002 | Dec 24, 2008 |

DELAFLOXACIN MEGLUMINE

POWDER; INTRAVENOUS

BAXDELA

| | | | | |
|---|---------|--------------------|-------------|--------------|
| + | MELINTA | EQ 300MG BASE/VIAL | N208611 001 | Jun 19, 2017 |
|---|---------|--------------------|-------------|--------------|

TABLET; ORAL

BAXDELA

| | | | | |
|---|---------|---------------|-------------|--------------|
| + | MELINTA | EQ 450MG BASE | N208610 001 | Jun 19, 2017 |
|---|---------|---------------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

DELAVIRDINE MESYLATE

TABLET; ORAL

RESCRIPTOR

| | | |
|---|---------------|-------|
| + | VIIV HLTHCARE | 100MG |
| + | ! | 200MG |

| | | |
|---------|-----|--------------|
| N020705 | 001 | Apr 04, 1997 |
| N020705 | 002 | Jul 14, 1999 |

DEMECLOCYCLINE HYDROCHLORIDE

TABLET; ORAL

DEMECLOCYCLINE HYDROCHLORIDE

| | | | | | |
|-----------|-----------------|--------------|----------------|------------|--------------|
| <u>AB</u> | AKORN | <u>150MG</u> | <u>A065389</u> | <u>001</u> | Dec 01, 2008 |
| <u>AB</u> | | <u>300MG</u> | <u>A065389</u> | <u>002</u> | Dec 01, 2008 |
| <u>AB</u> | AMNEAL PHARM | <u>150MG</u> | <u>A065425</u> | <u>001</u> | Feb 27, 2008 |
| <u>AB</u> | ! | <u>300MG</u> | <u>A065425</u> | <u>002</u> | Feb 27, 2008 |
| <u>AB</u> | BARR | <u>150MG</u> | <u>A065171</u> | <u>001</u> | Dec 13, 2004 |
| <u>AB</u> | | <u>300MG</u> | <u>A065171</u> | <u>002</u> | Dec 13, 2004 |
| <u>AB</u> | EPIC PHARMA LLC | <u>150MG</u> | <u>A065447</u> | <u>001</u> | Aug 18, 2015 |
| <u>AB</u> | | <u>300MG</u> | <u>A065447</u> | <u>002</u> | Aug 18, 2015 |

DEOXYCHOLIC ACID

SOLUTION; SUBCUTANEOUS

KYBELLA

| | | |
|---|-------------------|--------------------|
| + | KYTHERA BIOPHARMS | 20MG/2ML (10MG/ML) |
|---|-------------------|--------------------|

| | | |
|---------|-----|--------------|
| N206333 | 001 | Apr 29, 2015 |
|---------|-----|--------------|

DESFLURANE

LIQUID; INHALATION

DESFLURANE

| | | | | | | | |
|-----------|------------------|-------------|------------------------|-------------|----------------|------------|--------------|
| <u>AN</u> | SHANGHAI HENGRUI | <u>100%</u> | <u>A208234</u> | <u>001</u> | Feb 26, 2018 | | |
| <u>AN</u> | + | ! | <u>BAXTER HLTHCARE</u> | <u>100%</u> | <u>N020118</u> | <u>001</u> | Sep 18, 1992 |

DESIPRAMINE HYDROCHLORIDE

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

| | | | | | | |
|-----------|---------------------|-------------------|----------------|----------------|--------------|--------------|
| <u>AB</u> | ACTAVIS TOTOWA | <u>10MG</u> | <u>A074430</u> | <u>001</u> | Feb 09, 1996 | |
| <u>AB</u> | | <u>25MG</u> | <u>A071601</u> | <u>001</u> | Jun 05, 1987 | |
| <u>AB</u> | | <u>50MG</u> | <u>A071588</u> | <u>001</u> | Jun 05, 1987 | |
| <u>AB</u> | | <u>75MG</u> | <u>A071602</u> | <u>001</u> | Oct 05, 1987 | |
| <u>AB</u> | | <u>100MG</u> | <u>A071766</u> | <u>001</u> | Oct 05, 1987 | |
| <u>AB</u> | | <u>150MG</u> | <u>A074430</u> | <u>002</u> | Feb 09, 1996 | |
| <u>AB</u> | AMNEAL PHARMS CO | <u>10MG</u> | <u>A208105</u> | <u>001</u> | Mar 17, 2016 | |
| <u>AB</u> | | <u>25MG</u> | <u>A208105</u> | <u>002</u> | Mar 17, 2016 | |
| <u>AB</u> | | <u>50MG</u> | <u>A208105</u> | <u>003</u> | Mar 17, 2016 | |
| <u>AB</u> | | <u>75MG</u> | <u>A208105</u> | <u>004</u> | Mar 17, 2016 | |
| <u>AB</u> | | <u>100MG</u> | <u>A208105</u> | <u>005</u> | Mar 17, 2016 | |
| <u>AB</u> | | <u>150MG</u> | <u>A208105</u> | <u>006</u> | Mar 17, 2016 | |
| <u>AB</u> | ANI PHARMS INC | <u>10MG</u> | <u>A205153</u> | <u>001</u> | Oct 28, 2016 | |
| <u>AB</u> | | <u>25MG</u> | <u>A205153</u> | <u>002</u> | Oct 28, 2016 | |
| <u>AB</u> | | <u>50MG</u> | <u>A205153</u> | <u>003</u> | Oct 28, 2016 | |
| <u>AB</u> | | <u>75MG</u> | <u>A205153</u> | <u>004</u> | Oct 28, 2016 | |
| <u>AB</u> | | <u>100MG</u> | <u>A205153</u> | <u>005</u> | Oct 28, 2016 | |
| <u>AB</u> | | <u>150MG</u> | <u>A205153</u> | <u>006</u> | Oct 28, 2016 | |
| <u>AB</u> | HERITAGE PHARMS INC | <u>10MG</u> | <u>A207433</u> | <u>001</u> | May 05, 2016 | |
| <u>AB</u> | | <u>25MG</u> | <u>A207433</u> | <u>002</u> | May 05, 2016 | |
| <u>AB</u> | | <u>50MG</u> | <u>A207433</u> | <u>003</u> | May 05, 2016 | |
| <u>AB</u> | | <u>75MG</u> | <u>A207433</u> | <u>004</u> | May 05, 2016 | |
| <u>AB</u> | | <u>100MG</u> | <u>A207433</u> | <u>005</u> | May 05, 2016 | |
| <u>AB</u> | | <u>150MG</u> | <u>A207433</u> | <u>006</u> | May 05, 2016 | |
| <u>AB</u> | INGENUS PHARMS LLC | <u>10MG</u> | <u>A204963</u> | <u>001</u> | Dec 26, 2017 | |
| <u>AB</u> | | <u>25MG</u> | <u>A204963</u> | <u>002</u> | Dec 26, 2017 | |
| <u>AB</u> | | <u>50MG</u> | <u>A204963</u> | <u>003</u> | Dec 26, 2017 | |
| <u>AB</u> | | <u>75MG</u> | <u>A204963</u> | <u>004</u> | Dec 26, 2017 | |
| <u>AB</u> | | <u>100MG</u> | <u>A204963</u> | <u>005</u> | Dec 26, 2017 | |
| <u>AB</u> | | <u>150MG</u> | <u>A204963</u> | <u>006</u> | Dec 26, 2017 | |
| <u>AB</u> | SANDOZ | <u>10MG</u> | <u>A072099</u> | <u>001</u> | May 24, 1988 | |
| <u>AB</u> | | <u>25MG</u> | <u>A072100</u> | <u>001</u> | May 24, 1988 | |
| <u>AB</u> | | <u>50MG</u> | <u>A072101</u> | <u>001</u> | May 24, 1988 | |
| <u>AB</u> | | <u>75MG</u> | <u>A072102</u> | <u>001</u> | Jun 20, 1988 | |
| <u>AB</u> | | <u>100MG</u> | <u>A072103</u> | <u>001</u> | Jun 20, 1988 | |
| <u>AB</u> | | <u>150MG</u> | <u>A072104</u> | <u>001</u> | Jun 20, 1988 | |
| | <u>NORPRAMIN</u> | | | | | |
| <u>AB</u> | + | US PHARM HOLDINGS | <u>10MG</u> | <u>N014399</u> | <u>007</u> | Feb 11, 1982 |
| <u>AB</u> | + | | <u>25MG</u> | <u>N014399</u> | <u>001</u> | |
| <u>AB</u> | + | | <u>50MG</u> | <u>N014399</u> | <u>003</u> | |
| <u>AB</u> | + | | <u>75MG</u> | <u>N014399</u> | <u>004</u> | |

PRESCRIPTION DRUG PRODUCT LIST

DESIPRAMINE HYDROCHLORIDE

TABLET; ORAL

NORPRAMIN

| | | | | | | |
|-----------|------------|--|--------------|----------------|------------|--|
| AB | + ! | | 100MG | N014399 | 005 | |
| AB | + | | 150MG | N014399 | 006 | |

DES Loratadine

SOLUTION; ORAL

CLARINEX

| | | | | | | |
|-----------------------|------------|---------------------|-----------------|----------------|------------|--------------|
| AA | + ! | MERCK SHARP DOHME | 0.5MG/ML | N021300 | 001 | Sep 01, 2004 |
| DES Loratadine | | | | | | |
| AA | | TARO PHARM | 0.5MG/ML | A202936 | 001 | May 26, 2016 |
| AA | | TARO PHARM INDS LTD | 0.5MG/ML | A202592 | 001 | Jun 30, 2015 |

TABLET; ORAL

CLARINEX

| | | | | | | |
|-----------------------|------------|--------------------|------------|----------------|------------|--------------|
| AB | + ! | MERCK SHARP DOHME | 5MG | N021165 | 001 | Dec 21, 2001 |
| DES Loratadine | | | | | | |
| AB | | BELCHER PHARMS | 5MG | A078355 | 001 | Apr 19, 2012 |
| AB | | DR REDDYS LABS LTD | 5MG | A078365 | 001 | Mar 08, 2011 |
| AB | | LUPIN PHARMS | 5MG | A078352 | 001 | Oct 25, 2010 |
| AB | | MYLAN PHARMS INC | 5MG | A078351 | 001 | Feb 10, 2012 |
| AB | | ORCHID HLTHCARE | 5MG | A078357 | 001 | Feb 19, 2010 |
| AB | | PERRIGO R AND D | 5MG | A078361 | 001 | Dec 22, 2011 |
| AB | | SANDOZ | 5MG | A078364 | 001 | Dec 03, 2010 |
| AB | | SUN PHARM INDS | 5MG | A078359 | 001 | Nov 16, 2010 |

TABLET, ORALLY DISINTEGRATING; ORAL

CLARINEX

| | | | | | | |
|-----------------------|------------|-------------------|--------------|----------------|------------|--------------|
| AB | + | MERCK SHARP DOHME | 2.5MG | N021312 | 002 | Jul 14, 2005 |
| AB | + ! | | 5MG | N021312 | 001 | Jun 26, 2002 |
| DES Loratadine | | | | | | |
| AB | | REDDYS | 2.5MG | A078367 | 001 | Jul 12, 2010 |
| AB | | | 5MG | A078367 | 002 | Jul 12, 2010 |

DES Loratadine; Pseudoephedrine Sulfate

TABLET, EXTENDED RELEASE; ORAL

CLARINEX D 24 HOUR

| | | | | | | |
|-----------------------------------------------------------|------------|--------------------|--------------------|----------------|------------|--------------|
| AB | + ! | MERCK SHARP DOHME | 5MG;240MG | N021605 | 001 | Mar 03, 2005 |
| DES Loratadine AND Pseudoephedrine Sulfate 24 HOUR | | | | | | |
| AB | | DR REDDYS LABS LTD | 5MG;240MG | A078366 | 001 | Apr 26, 2011 |
| CLARINEX-D 12 HOUR | | | | | | |
| | + ! | MERCK SHARP DOHME | 2.5MG;120MG | N021313 | 001 | Feb 01, 2006 |

Desmopressin Acetate

INJECTABLE; INJECTION

DDAVP

| | | | | | | |
|-----------------------------|------------|--------------------|-------------------|----------------|------------|--------------|
| AP | + ! | FERRING PHARMS INC | 0.004MG/ML | N018938 | 001 | Mar 30, 1984 |
| Desmopressin Acetate | | | | | | |
| AP | | SAGENT PHARMS | 0.004MG/ML | A204695 | 001 | Aug 22, 2017 |
| AP | | | 0.004MG/ML | A204751 | 001 | Aug 22, 2017 |
| AP | | SUN PHARM INDS LTD | 0.004MG/ML | A091280 | 001 | Jan 25, 2013 |

SOLUTION; NASAL

DDAVP

| | | | | | | |
|-----------------------------|------------|--------------------|--------------|----------------|------------|--------------|
| AB | + ! | FERRING PHARMS INC | 0.01% | N017922 | 001 | |
| Desmopressin Acetate | | | | | | |
| AB | | SUN PHARM INDS | 0.01% | A077212 | 001 | Apr 12, 2012 |

SPRAY, METERED; NASAL

DDAVP (NEEDS NO REFRIGERATION)

| | | | | | | |
|------------------------------------------------------|------------|----------------------|---------------------|----------------|------------|--------------|
| AB | + ! | FERRING PHARMS INC | 0.01MG/SPRAY | N017922 | 003 | Aug 07, 1996 |
| Desmopressin Acetate | | | | | | |
| AB | ! | BAUSCH AND LOMB | 0.01MG/SPRAY | A074830 | 001 | Jan 25, 1999 |
| Desmopressin Acetate (NEEDS NO REFRIGERATION) | | | | | | |
| AB | | APOTEX INC | 0.01MG/SPRAY | A076703 | 001 | Jan 27, 2005 |
| AB | | SUN PHARMA GLOBAL | 0.01MG/SPRAY | A078271 | 001 | Dec 23, 2013 |
| AB | | ZYDUS PHARMS USA INC | 0.01MG/SPRAY | A091345 | 001 | Oct 03, 2017 |

MINIRIN

| | | | | | | |
|----------------------------------|------------|--------------------|------------------------|----------------|------------|--------------|
| AB | + ! | FERRING | 0.01MG/SPRAY | N021333 | 001 | Sep 16, 2002 |
| NOCTIVA | | | | | | |
| | + | AVADEL SPECLT | 0.00083MG/SPRAY | N201656 | 001 | Mar 03, 2017 |
| | + ! | | 0.00166MG/SPRAY | N201656 | 002 | Mar 03, 2017 |
| STIMATE (NEEDS NO REFRIGERATION) | | | | | | |
| | + ! | FERRING PHARMS INC | 0.15MG/SPRAY | N020355 | 002 | Oct 24, 2007 |

PRESCRIPTION DRUG PRODUCT LIST

DESMOPRESSIN ACETATE

TABLET;ORAL

DDAVP

| | | | | | | |
|-----------|---|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | FERRING PHARMS INC | <u>0.1MG</u> | <u>N019955</u> | <u>001</u> | Sep 06, 1995 |
| <u>AB</u> | + | ! | <u>0.2MG</u> | <u>N019955</u> | <u>002</u> | Sep 06, 1995 |

DESMOPRESSIN ACETATE

| | | | | | | |
|-----------|--|---------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | ACTAVIS LABS FL INC | <u>0.1MG</u> | <u>A076470</u> | <u>001</u> | Jul 01, 2005 |
| <u>AB</u> | | | <u>0.2MG</u> | <u>A076470</u> | <u>002</u> | Jul 01, 2005 |
| <u>AB</u> | | APOTEX INC | <u>0.1MG</u> | <u>A077414</u> | <u>001</u> | Mar 07, 2006 |
| <u>AB</u> | | | <u>0.2MG</u> | <u>A077414</u> | <u>002</u> | Mar 07, 2006 |
| <u>AB</u> | | GLENMARK PHARMS LTD | <u>0.1MG</u> | <u>A201831</u> | <u>001</u> | May 28, 2015 |
| <u>AB</u> | | | <u>0.2MG</u> | <u>A201831</u> | <u>002</u> | May 28, 2015 |
| <u>AB</u> | | HERITAGE PHARMA | <u>0.1MG</u> | <u>A207880</u> | <u>001</u> | May 26, 2017 |
| <u>AB</u> | | | <u>0.2MG</u> | <u>A207880</u> | <u>002</u> | May 26, 2017 |
| <u>AB</u> | | IMPAX LABS INC | <u>0.1MG</u> | <u>A077122</u> | <u>001</u> | Jan 25, 2006 |
| <u>AB</u> | | | <u>0.2MG</u> | <u>A077122</u> | <u>002</u> | Jan 25, 2006 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>0.1MG</u> | <u>A200653</u> | <u>001</u> | Jun 27, 2014 |
| <u>AB</u> | | | <u>0.2MG</u> | <u>A200653</u> | <u>002</u> | Jun 27, 2014 |

TABLET;SUBLINGUAL

NOCDURNA

| | | | | | | |
|--|---|--------------------|----------|---------|-----|--------------|
| | + | FERRING PHARMS INC | 0.0277MG | N022517 | 001 | Jun 21, 2018 |
| | + | ! | 0.0553MG | N022517 | 002 | Jun 21, 2018 |

DESOGESTREL; ETHINYL ESTRADIOL

TABLET;ORAL-28

BEKYREE

| | | | | | | |
|-----------|--|-----------|---------------------------------|----------------|------------|--------------|
| <u>AB</u> | | LUPIN LTD | <u>0.15MG,N/A;0.02MG,0.01MG</u> | <u>A202226</u> | <u>001</u> | Aug 12, 2015 |
|-----------|--|-----------|---------------------------------|----------------|------------|--------------|

CYCLESSA

| | | | | | | | |
|-----------|---|---|-------------------------|-----------------------------------------------------|----------------|------------|--------------|
| <u>AB</u> | + | ! | <u>ASPEN GLOBAL INC</u> | <u>0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.025MG</u> | <u>N021090</u> | <u>001</u> | Dec 20, 2000 |
|-----------|---|---|-------------------------|-----------------------------------------------------|----------------|------------|--------------|

DESOGEN

| | | | | | | |
|-----------|--|-----------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | | ORGANON USA INC | <u>0.15MG;0.03MG</u> | <u>N020071</u> | <u>002</u> | Dec 10, 1992 |
|-----------|--|-----------------|----------------------|----------------|------------|--------------|

DESOGESTREL AND ETHINYL ESTRADIOL

| | | | | | | |
|-----------|---|----------------------|-----------------------------------------------------|----------------|------------|--------------|
| <u>AB</u> | | ACCORD HLTHCARE | <u>0.15MG,N/A;0.02MG,0.01MG</u> | <u>A209170</u> | <u>001</u> | Jun 05, 2017 |
| <u>AB</u> | | | <u>0.15MG;0.03MG</u> | <u>A207067</u> | <u>001</u> | Sep 13, 2018 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>0.15MG,N/A;0.02MG,0.01MG</u> | <u>A206853</u> | <u>001</u> | Mar 22, 2017 |
| <u>AB</u> | ! | DURAMED PHARMS BARR | <u>0.15MG;0.03MG</u> | <u>A075256</u> | <u>002</u> | Aug 12, 1999 |
| <u>AB</u> | | MAYNE PHARMA | <u>0.15MG,N/A;0.02MG,0.01MG</u> | <u>A076916</u> | <u>001</u> | Dec 29, 2008 |
| <u>AB</u> | | | <u>0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.025MG</u> | <u>A077182</u> | <u>001</u> | Jan 24, 2006 |
| <u>AB</u> | | MYLAN LABS LTD | <u>0.15MG,N/A;0.02MG,0.01MG</u> | <u>A202296</u> | <u>001</u> | Aug 30, 2013 |
| <u>AB</u> | | | <u>0.15MG;0.03MG</u> | <u>A202085</u> | <u>001</u> | May 20, 2015 |
| <u>AB</u> | | NOVAST LABS | <u>0.15MG;0.03MG</u> | <u>A091234</u> | <u>001</u> | Jul 12, 2013 |
| <u>AB</u> | | WATSON LABS | <u>0.15MG;0.03MG</u> | <u>A076915</u> | <u>001</u> | Jul 29, 2005 |
| | | <u>EMOQUETTE</u> | | | | |
| <u>AB</u> | | VINTAGE PHARMS LLC | <u>0.15MG;0.03MG</u> | <u>A076675</u> | <u>001</u> | Feb 25, 2011 |
| | | <u>ENSKYCE</u> | | | | |
| <u>AB</u> | | LUPIN LTD | <u>0.15MG;0.03MG</u> | <u>A201887</u> | <u>001</u> | Mar 07, 2013 |
| | | <u>ISIBLOOM</u> | | | | |
| <u>AB</u> | | LABS LEON FARMA | <u>0.15MG;0.03MG</u> | <u>A202789</u> | <u>001</u> | Aug 12, 2015 |
| | | <u>KALLIGA</u> | | | | |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>0.15MG;0.03MG</u> | <u>A207081</u> | <u>001</u> | May 17, 2017 |
| | | <u>KARIVA</u> | | | | |
| <u>AB</u> | ! | BARR | <u>0.15MG,N/A;0.02MG,0.01MG</u> | <u>A075863</u> | <u>001</u> | Apr 05, 2002 |
| | | <u>KIMIDESS</u> | | | | |
| <u>AB</u> | | VINTAGE PHARMS | <u>0.15MG,N/A;0.02MG,0.01MG</u> | <u>A076681</u> | <u>001</u> | Apr 30, 2015 |
| | | <u>PIMTREA</u> | | | | |
| <u>AB</u> | | NOVAST LABS | <u>0.15MG,N/A;0.02MG,0.01MG</u> | <u>A091247</u> | <u>001</u> | Aug 01, 2013 |
| | | <u>VELIVET</u> | | | | |
| <u>AB</u> | | DURAMED PHARMS BARR | <u>0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.025MG</u> | <u>A076455</u> | <u>001</u> | Feb 24, 2004 |
| | | <u>VIORELE</u> | | | | |
| <u>AB</u> | | GLENMARK GENERICS | <u>0.15MG,N/A;0.02MG,0.01MG</u> | <u>A091346</u> | <u>001</u> | Apr 02, 2012 |
| | | <u>VOLNEA</u> | | | | |
| <u>AB</u> | | LABS LEON FARMA | <u>0.15MG,N/A;0.02MG,0.01MG</u> | <u>A202689</u> | <u>001</u> | Sep 09, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

DESONIDE

AEROSOL, FOAM;TOPICAL

VERDESO

+! AQUA PHARMS 0.05% N021978 001 Sep 19, 2006

CREAM;TOPICAL

DESONIDE**AB** G AND W LABS INC 0.05% **A074027 001** Sep 28, 1992**AB** GLENMARK PHARMS 0.05% **A209729 001** Jul 24, 2017**AB** +! PERRIGO NEW YORK 0.05% **N017010 001****AB** TARO 0.05% **A073548 001** Jun 30, 1992DESOWEN**AB** GALDERMA LABS LP 0.05% **N019048 001** Dec 14, 1984

GEL;TOPICAL

DESONATE

+! LEO PHARMA AS 0.05% N021844 001 Oct 20, 2006

LOTION;TOPICAL

DESONIDE**AB** FOUGERA PHARMS 0.05% **A075860 001** Mar 19, 2002**AB** GLENMARK PHARMS 0.05% **A209494 001** Sep 26, 2017**AB** TARO PHARM 0.05% **A202161 001** Oct 31, 2014**AB** TELIGENT PHARMA INC 0.05% **A207855 001** Sep 28, 2017DESOWEN**AB** ! GALDERMA LABS LP 0.05% **A072354 001** Jan 24, 1992

OINTMENT;TOPICAL

DESONIDE**AB** FOUGERA PHARMS 0.05% **A075751 001** Mar 12, 2001**AB** GLENMARK PHARMS LTD 0.05% **A209996 001** Sep 15, 2017**AB** HI-TECH PHARMACAL 0.05% **A208836 001** Mar 27, 2017**AB** +! PERRIGO NEW YORK 0.05% **N017426 001****AB** TARO 0.05% **A074254 001** Aug 03, 1994DESOWEN**AB** GALDERMA LABS LP 0.05% **A071425 001** Jun 15, 1988DESOXIMETASONE

CREAM;TOPICAL

DESOXIMETASONE**AB** ACTAVIS MID 0.25% **A205082 001** Sep 04, 2015

ATLANTIC

AB AKORN 0.05% **A203787 001** Jan 06, 2017**AB** 0.25% **A203234 001** Jun 12, 2015**AB** FOUGERA PHARMS 0.25% **A078369 001** Jun 29, 2010**AB** LUPIN ATLANTIS 0.05% **A208163 001** Jan 10, 2017**AB** 0.25% **A208164 001** Jan 09, 2017**AB** PERRIGO NEW YORK 0.25% **A076510 001** Jul 01, 2003**AB** RICONPHARMA LLC 0.05% **A210980 001** Dec 21, 2018**AB** RISING PHARMS 0.25% **A205594 001** Jul 02, 2018**AB** ZYDUS PHARMS USA 0.25% **A205620 001** Sep 28, 2018

INC

TOPICORT**AB** ! TARO PHARM INDS LTD 0.05% **A073210 001** Nov 30, 1990**AB** ! 0.25% **A073193 001** Nov 30, 1990

GEL;TOPICAL

DESOXIMETASONE**AB** AKORN 0.05% **A090727 001** Mar 10, 2011**AB** PERRIGO NEW YORK 0.05% **A077552 001** Jan 09, 2006**AB** RISING PHARMS 0.05% **A204675 001** Aug 12, 2016TOPICORT**AB** ! TARO PHARM INDS LTD 0.05% **A074904 001** Jul 14, 1998

OINTMENT;TOPICAL

DESOXIMETASONE**AB** ACTAVIS MID 0.25% **A204965 001** Nov 07, 2016

ATLANTIC

AB AKORN 0.25% **A201005 001** Apr 24, 2014**AB** FOUGERA PHARMS 0.25% **A078657 001** Sep 28, 2012**AB** G AND W LABS INC 0.25% **A206740 001** Dec 23, 2016**AB** GLENMARK GENERICS 0.25% **A202838 001** Sep 20, 2013**AB** LUPIN ATLANTIS 0.05% **A208044 001** Dec 12, 2016**AB** 0.25% **A208104 001** Dec 01, 2016**AB** NOVEL LABS INC 0.25% **A206792 001** May 10, 2016**AB** PERRIGO ISRAEL 0.25% **A077770 001** Apr 20, 2015**AB** RISING PHARMS 0.25% **A204272 001** Nov 30, 2016**AB** TELIGENT PHARMA INC 0.05% **A209973 001** Oct 23, 2018**AB** 0.25% **A208101 001** Feb 25, 2016

PRESCRIPTION DRUG PRODUCT LIST

DESOXIMETASONE

OINTMENT; TOPICAL

DESOXIMETASONE

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| AB | ZYDUS PHARMS USA INC | 0.25% | A205206 001 | Sep 19, 2017 |
|-----------|-------------------------|--------------|--------------------|--------------|

TOPICORT

| | | | | | |
|-----------|------------|---------------------|--------------|--------------------|--------------|
| AB | + ! | TARO PHARM INDS LTD | 0.05% | N018594 001 | Jan 17, 1985 |
| AB | ! | | 0.25% | A074286 001 | Jun 07, 1996 |

SPRAY; TOPICAL

DESOXIMETASONE

| | | | | |
|-----------|----------------|--------------|--------------------|--------------|
| AT | LUPIN ATLANTIS | 0.25% | A208124 001 | Mar 16, 2018 |
| AT | PERRIGO ISRAEL | 0.25% | A206441 001 | Jan 20, 2017 |

TOPICORT

| | | | | | |
|-----------|------------|-------------|--------------|--------------------|--------------|
| AT | + ! | TARO PHARMS | 0.25% | N204141 001 | Apr 11, 2013 |
|-----------|------------|-------------|--------------|--------------------|--------------|

DESVENLAFAXINE

TABLET, EXTENDED RELEASE; ORAL

DESVENLAFAXINE

| | | | | | |
|----|------------|--------------------|-------|-------------|--------------|
| BC | + ! | ALEMBIC PHARMS LTD | 50MG | N204150 001 | Mar 04, 2013 |
| BC | + ! | | 100MG | N204150 002 | Mar 04, 2013 |

KHEDEZLA

| | | | | | |
|----|--|---------------------|-------|-------------|--------------|
| BC | | OSMOTICA PHARM CORP | 50MG | N204683 001 | Jul 10, 2013 |
| BC | | | 100MG | N204683 002 | Jul 10, 2013 |

DESVENLAFAXINE SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

DESVENLAFAXINE SUCCINATE

| | | | | | |
|-----------|------------|-------------------------|----------------------|--------------------|--------------|
| AB | | ACTAVIS LABS FL | EQ 25MG BASE | A204065 001 | Jul 29, 2016 |
| AB | | | EQ 50MG BASE | A204065 002 | Jul 29, 2016 |
| AB | | | EQ 100MG BASE | A204065 003 | Jul 29, 2016 |
| AB | | ALEMBIC PHARMS LTD | EQ 25MG BASE | A204003 003 | Sep 14, 2018 |
| AB | | | EQ 50MG BASE | A204003 001 | Jun 29, 2015 |
| AB | | | EQ 100MG BASE | A204003 002 | Jun 29, 2015 |
| AB | | CASI PHARMS INC | EQ 50MG BASE | A204028 001 | Jun 29, 2015 |
| AB | | | EQ 100MG BASE | A204028 002 | Jun 29, 2015 |
| AB | | LUPIN LTD | EQ 50MG BASE | A204172 001 | Jun 29, 2015 |
| AB | | | EQ 100MG BASE | A204172 002 | Jun 29, 2015 |
| AB | | MYLAN PHARMS INC | EQ 50MG BASE | A204095 001 | Jun 29, 2015 |
| AB | | | EQ 100MG BASE | A204095 002 | Jun 29, 2015 |
| AB | | WEST-WARD PHARMS INT | EQ 25MG BASE | A204082 002 | Aug 28, 2017 |
| AB | | | EQ 50MG BASE | A204082 001 | Feb 16, 2016 |
| AB | | | EQ 100MG BASE | A204083 001 | Feb 16, 2016 |
| AB | | YICHANG HUMANWELL | EQ 50MG BASE | A210014 001 | Oct 01, 2018 |
| AB | | | EQ 100MG BASE | A210014 002 | Oct 01, 2018 |
| AB | | ZYDUS PHARMS USA INC | EQ 50MG BASE | A204020 001 | Oct 11, 2017 |
| AB | | | EQ 100MG BASE | A204020 002 | Oct 11, 2017 |
| | | PRISTIO | | | |
| AB | + | PF PRISM CV | EQ 25MG BASE | N021992 003 | Aug 20, 2014 |
| AB | + ! | | EQ 50MG BASE | N021992 001 | Feb 29, 2008 |
| AB | + ! | | EQ 100MG BASE | N021992 002 | Feb 29, 2008 |

DEUTETRABENAZINE

TABLET; ORAL

AUSTEDO

| | | | | |
|------------|--------------------|------|-------------|--------------|
| + | TEVA BRANDED PHARM | 6MG | N208082 001 | Apr 03, 2017 |
| + | | 9MG | N208082 002 | Apr 03, 2017 |
| + ! | | 12MG | N208082 003 | Apr 03, 2017 |

DEXAMETHASONE

CONCENTRATE; ORAL

DEXAMETHASONE INTENSOL

| | | | | |
|----------|-------------------------|--------|-------------|--------------|
| ! | WEST-WARD PHARMS INT | 1MG/ML | A088252 001 | Sep 01, 1983 |
|----------|-------------------------|--------|-------------|--------------|

ELIXIR; ORAL

DEXAMETHASONE

| | | | | |
|-----------|------------------|------------------|--------------------|--------------------|
| AA | LANNETT CO INC | 0.5MG/5ML | A091188 001 | May 11, 2011 |
| AA | LYNE | 0.5MG/5ML | A090891 001 | Jul 12, 2011 |
| AA | ! | STI PHARMA LLC | 0.5MG/5ML | A084754 001 |
| AA | WOCKHARDT BIO AG | 0.5MG/5ML | A088254 001 | Jul 27, 1983 |

IMPLANT; INTRAVITREAL

OZURDEX

| | | | | |
|------------|----------|-------|-------------|--------------|
| + ! | ALLERGAN | 0.7MG | N022315 001 | Jun 17, 2009 |
|------------|----------|-------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

DEXAMETHASONE

INSERT;OPHTHALMIC

DEXTENZA

+! OCULAR THERAPEUTIX 0.4MG N208742 001 Nov 30, 2018

SOLUTION;ORAL

DEXAMETHASONE

! WEST-WARD PHARMS 0.5MG/5ML A088248 001 Sep 01, 1983
INT

SUSPENSION;INTRAOCULAR

DEXYCU KIT

+! EYEPOINT PHARMS 9% N208912 001 Feb 09, 2018

SUSPENSION/DROPS;OPHTHALMIC

MAXIDEX

+! NOVARTIS PHARMS 0.1% N013422 001
CORP

TABLET;ORAL

DEXAMETHASONEAB ECR 1.5MG A040700 001 Aug 15, 2008AB LARKEN LABS INC 1.5MG A201270 001 Jul 17, 2017AB WEST-WARD PHARMS 1.5MG A084610 001

INT

BP FERA PHARMS LLC 0.5MG A088481 002 Apr 28, 1983

BP 0.75MG A088481 003 Apr 28, 1983

BP 4MG A088481 004 Apr 28, 1983

BP 6MG A088481 001 Nov 28, 1983

BP WEST-WARD PHARMS 0.5MG A084611 001

INT

BP 0.75MG A084613 001

BP 1MG A088306 001 Sep 15, 1983

BP 2MG A087916 001 Aug 26, 1982

BP 4MG A084612 001

BP ! 6MG A088316 001 Sep 15, 1983

BP XSPIRE PHARMA 1.5MG A088237 001 Apr 28, 1983

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE;INJECTION

DEXAMETHASONE SODIUM PHOSPHATEAP AMNEAL PHARMS CO EQ 4MG PHOSPHATE/ML A208689 001 Aug 22, 2018AP AUROBINDO PHARMA EQ 4MG PHOSPHATE/ML A206781 001 Dec 01, 2015

LTD

AP FRESENIUS KABI USA EQ 4MG PHOSPHATE/ML A084916 001AP EQ 4MG PHOSPHATE/ML A203129 001 Sep 30, 2015AP ! EQ 10MG PHOSPHATE/ML A040572 001 Apr 22, 2005AP EQ 10MG PHOSPHATE/ML A209192 001 Jul 06, 2018AP ! LUITPOLD EQ 4MG PHOSPHATE/ML A087440 001 Jul 21, 1982AP MYLAN LABS LTD EQ 4MG PHOSPHATE/ML A040803 001 Aug 29, 2008AP EQ 10MG PHOSPHATE/ML A040802 001 Aug 29, 2008AP SOMERSET THERAPS EQ 4MG PHOSPHATE/ML A207521 001 Jun 08, 2018

LLC

AP WEST-WARD PHARMS EQ 4MG PHOSPHATE/ML A084282 001

INT

AP ! EQ 10MG PHOSPHATE/ML A087702 001 Sep 07, 1982DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREEAP AMNEAL PHARMS CO EQ 10MG PHOSPHATE/ML A208690 001 Aug 22, 2018AP ! FRESENIUS KABI USA EQ 10MG PHOSPHATE/ML A040491 001 Apr 11, 2003AP SOMERSET THERAPS EQ 10MG PHOSPHATE/ML A207442 001 Apr 19, 2018

LLC

SOLUTION/DROPS;OPHTHALMIC, OTIC

DEXAMETHASONE SODIUM PHOSPHATEAT BAUSCH AND LOMB EQ 0.1% PHOSPHATE A040069 001 Jul 26, 1996AT ! SANDOZ INC EQ 0.1% PHOSPHATE A088771 001 Jan 16, 1985DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

MAXITROLAT +! NOVARTIS PHARMS 0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM N050065 002

CORP

NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONEAT BAUSCH AND LOMB 0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM A064063 001 Jul 25, 1994AT PERRIGO CO 0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM A062938 001 Jul 31, 1989

TENNESSEE

SUSPENSION/DROPS;OPHTHALMIC

DEXASPORINAT BAUSCH AND LOMB 0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML A064135 001 Sep 13, 1995

PRESCRIPTION DRUG PRODUCT LIST

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SUSPENSION/DROPS;OPHTHALMIC

MAXITROL

| | | | | | |
|-----------|------------|-------------------------|----------------------------------------------|--------------------|--------------|
| <u>AT</u> | <u>+</u> ! | NOVARTIS PHARMS CORP | <u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u> | <u>N050023 002</u> | |
| <u>AT</u> | | SANDOZ INC | <u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u> | <u>A062341 001</u> | May 22, 1984 |

DEXAMETHASONE; TOBRAMYCIN

OINTMENT;OPHTHALMIC

TOBRADEX

| | | | | |
|------------|-------------------------|------------------|--------------------|--------------|
| <u>+</u> ! | NOVARTIS PHARMS CORP | <u>0.1%;0.3%</u> | <u>N050616 001</u> | Sep 28, 1988 |
|------------|-------------------------|------------------|--------------------|--------------|

SUSPENSION/DROPS;OPHTHALMIC

TOBRADEX

| | | | | | |
|-----------|------------|-------------------------|------------------|--------------------|--------------|
| <u>AB</u> | <u>+</u> ! | NOVARTIS PHARMS CORP | <u>0.1%;0.3%</u> | <u>N050592 001</u> | Aug 18, 1988 |
|-----------|------------|-------------------------|------------------|--------------------|--------------|

TOBRAMYCIN AND DEXAMETHASONE

| | | | | | |
|-----------|------------|-------------------------|-------------------|--------------------|--------------|
| <u>AB</u> | | BAUSCH AND LOMB | <u>0.1%;0.3%</u> | <u>A064134 001</u> | Oct 27, 1999 |
| | <u>+</u> ! | NOVARTIS PHARMS CORP | <u>0.05%;0.3%</u> | <u>N050818 001</u> | Feb 13, 2009 |

DEXCHLORPHENIRAMINE MALEATE

SYRUP;ORAL

DEXCHLORPHENIRAMINE MALEATE

| | | | | | |
|-----------|----------|------------------|----------------|--------------------|--------------|
| <u>AA</u> | <u>!</u> | WOCKHARDT BIO AG | <u>2MG/5ML</u> | <u>A088251 001</u> | Mar 23, 1984 |
|-----------|----------|------------------|----------------|--------------------|--------------|

POLMON

| | | | | | |
|-----------|--|---------------------|----------------|--------------------|--------------|
| <u>AA</u> | | CAPELLON PHARMS LLC | <u>2MG/5ML</u> | <u>A202520 001</u> | Jul 16, 2018 |
|-----------|--|---------------------|----------------|--------------------|--------------|

DEXLANSOPRAZOLE

CAPSULE, DELAYED RELEASE;ORAL

DEXILANT

| | | | | | |
|-----------|------------|-------------------|-------------|--------------------|--------------|
| <u>AB</u> | <u>+</u> ! | TAKEDA PHARMS USA | <u>60MG</u> | <u>N022287 002</u> | Jan 30, 2009 |
|-----------|------------|-------------------|-------------|--------------------|--------------|

DEXLANSOPRAZOLE

| | | | | | |
|-----------|----------|-------------------|-------------|--------------------|--------------|
| <u>AB</u> | | PAR PHARM INC | <u>60MG</u> | <u>A202294 001</u> | Apr 19, 2017 |
| | <u>+</u> | TAKEDA PHARMS USA | <u>30MG</u> | <u>N022287 001</u> | Jan 30, 2009 |

DEXMEDETOMIDINE HYDROCHLORIDE

INJECTABLE;INJECTION

DEXMEDETOMIDINE HYDROCHLORIDE

| | | | | | |
|-----------|------------|-------------------------|-----------------------------------------------|--------------------|--------------|
| <u>AP</u> | | ACCORD HLTHCARE | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>A204023 001</u> | Feb 09, 2016 |
| <u>AP</u> | | ACTAVIS INC | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>A204686 001</u> | Oct 17, 2016 |
| <u>AP</u> | | AKORN INC | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>A202585 001</u> | Nov 24, 2014 |
| <u>AP</u> | | AUROBINDO PHARMA LTD | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>A205867 001</u> | Mar 17, 2016 |
| <u>AP</u> | | BAXTER HLTHCARE CORP | <u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u> | <u>A208532 001</u> | Aug 21, 2018 |
| <u>AP</u> | | | <u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u> | <u>A208532 002</u> | Aug 21, 2018 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u> | <u>A208129 001</u> | Nov 29, 2018 |
| <u>AP</u> | | | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>A201072 001</u> | Sep 18, 2015 |
| <u>AP</u> | | | <u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u> | <u>A208129 002</u> | Nov 29, 2018 |
| <u>AP</u> | | | <u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u> | <u>A208129 003</u> | Nov 29, 2018 |
| <u>AP</u> | | JIANGSU HENGRUI MED | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>A209065 001</u> | Sep 19, 2017 |
| <u>AP</u> | | LUITPOLD | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>A203773 001</u> | May 12, 2017 |
| <u>AP</u> | | MYLAN INSTITUTIONAL | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>A202881 001</u> | Aug 18, 2014 |
| <u>AP</u> | | PAR STERILE PRODUCTS | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>A203972 001</u> | Aug 18, 2014 |
| <u>AP</u> | | SANDOZ INC | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>A091465 001</u> | Jun 14, 2016 |
| <u>AP</u> | | SUN PHARM INDS INC | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>A202126 001</u> | Aug 20, 2015 |
| <u>AP</u> | | TEVA PHARMS USA | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>A205272 001</u> | Nov 28, 2017 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>A205046 001</u> | Apr 26, 2017 |
| <u>AP</u> | | ZYDUS PHARMS USA INC | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>A206798 001</u> | Feb 27, 2018 |
| | | | | <u>N021038 004</u> | Nov 14, 2014 |
| <u>AP</u> | <u>+</u> ! | HOSPIRA | <u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u> | <u>N021038 001</u> | Dec 17, 1999 |
| <u>AP</u> | <u>+</u> ! | | <u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u> | <u>N021038 002</u> | Mar 13, 2013 |
| <u>AP</u> | <u>+</u> ! | | <u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u> | <u>N021038 003</u> | Mar 13, 2013 |

SOLUTION;INTRAVENOUS

DEXMEDETOMIDINE HYDROCHLORIDE

| | | | | |
|------------|-----------------|-----------------------------------------------|--------------------|--------------|
| <u>+</u> ! | HQ SPCLT PHARMA | <u>EQ 1MG BASE/10ML (EQ 100MCG BASE/ML)</u> | <u>N206628 002</u> | Oct 21, 2015 |
| <u>+</u> | | <u>EQ 200MG BASE/50ML (EQ 4MCG BASE/ML)</u> | <u>N206628 003</u> | Jun 22, 2018 |
| <u>+</u> | | <u>EQ 400MCG BASE/4ML (EQ 100MCG BASE/ML)</u> | <u>N206628 001</u> | Oct 21, 2015 |
| <u>+</u> | | <u>EQ 400MG BASE/100ML (EQ 4MCG BASE/ML)</u> | <u>N206628 004</u> | Jun 22, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

DEXMETHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

| | | | | |
|-----------------------------------------|--------------------------|--------------|--------------------|---------------------------------|
| <u>AB</u> | ADARE PHARMS INC | <u>5MG</u> | <u>A210279 001</u> | Oct 09, 2018 |
| <u>AB</u> | | <u>10MG</u> | <u>A210279 002</u> | Oct 09, 2018 |
| <u>AB</u> | | <u>15MG</u> | <u>A210279 003</u> | Oct 09, 2018 |
| <u>AB</u> | | <u>20MG</u> | <u>A210279 004</u> | Oct 09, 2018 |
| <u>AB</u> | | <u>25MG</u> | <u>A210279 005</u> | Oct 09, 2018 |
| <u>AB</u> | | <u>30MG</u> | <u>A210279 006</u> | Oct 09, 2018 |
| <u>AB</u> | | <u>35MG</u> | <u>A210279 007</u> | Oct 09, 2018 |
| <u>AB</u> | | <u>40MG</u> | <u>A210279 008</u> | Oct 09, 2018 |
| <u>AB</u> | IMPAX LABS INC | <u>5MG</u> | <u>A079108 001</u> | Aug 05, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A079108 002</u> | Aug 05, 2015 |
| <u>AB</u> | | <u>15MG</u> | <u>A079108 003</u> | May 19, 2014 |
| <u>AB</u> | | <u>20MG</u> | <u>A079108 004</u> | Dec 21, 2015 |
| <u>AB</u> | | <u>25MG</u> | <u>A203614 001</u> | Jul 05, 2017 |
| <u>AB</u> | | <u>30MG</u> | <u>A079108 005</u> | Nov 21, 2013 |
| <u>AB</u> | | <u>35MG</u> | <u>A203614 002</u> | Jul 05, 2017 |
| <u>AB</u> | INTELLIPHARMACEUTIC S | <u>15MG</u> | <u>A078992 003</u> | Nov 18, 2013 |
| <u>AB</u> | | <u>30MG</u> | <u>A078992 004</u> | Nov 18, 2013 |
| <u>AB</u> | MYLAN PHARMS INC | <u>5MG</u> | <u>A204266 001</u> | Aug 25, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A204266 002</u> | Aug 25, 2015 |
| <u>AB</u> | | <u>15MG</u> | <u>A204266 003</u> | Aug 25, 2015 |
| <u>AB</u> | | <u>20MG</u> | <u>A204266 004</u> | Dec 21, 2015 |
| <u>AB</u> | | <u>30MG</u> | <u>A202580 001</u> | Aug 28, 2013 |
| <u>AB</u> | | <u>40MG</u> | <u>A204266 007</u> | Aug 25, 2015 |
| <u>AB</u> | PAR PHARM INC | <u>5MG</u> | <u>A202842 001</u> | Nov 30, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A202842 002</u> | Nov 30, 2016 |
| <u>AB</u> | | <u>15MG</u> | <u>A202842 003</u> | Nov 30, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A202842 004</u> | Nov 30, 2016 |
| <u>AB</u> | | <u>25MG</u> | <u>A202842 005</u> | Nov 30, 2016 |
| <u>AB</u> | | <u>30MG</u> | <u>A202842 006</u> | Nov 30, 2016 |
| <u>AB</u> | | <u>35MG</u> | <u>A202842 007</u> | Nov 30, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A202842 008</u> | Nov 30, 2016 |
| <u>AB</u> | TEVA PHARMS USA | <u>5MG</u> | <u>A078908 001</u> | Nov 19, 2013 |
| <u>AB</u> | | <u>10MG</u> | <u>A078908 002</u> | Nov 19, 2013 |
| <u>AB</u> | | <u>15MG</u> | <u>A078908 004</u> | May 19, 2014 |
| <u>AB</u> | | <u>20MG</u> | <u>A078908 003</u> | Nov 19, 2013 |
| <u>AB</u> | | <u>25MG</u> | <u>A202731 001</u> | Jul 05, 2017 |
| <u>AB</u> | | <u>30MG</u> | <u>A202731 003</u> | May 19, 2014 |
| <u>AB</u> | | <u>35MG</u> | <u>A202731 004</u> | Jul 05, 2017 |
| <u>AB</u> | | <u>40MG</u> | <u>A202731 002</u> | Nov 19, 2013 |
| <u>FOCALIN XR</u> | | | | |
| <u>AB</u> | + | NOVARTIS | <u>5MG</u> | <u>N021802 001</u> May 26, 2005 |
| <u>AB</u> | + | | <u>10MG</u> | <u>N021802 002</u> May 26, 2005 |
| <u>AB</u> | + | | <u>15MG</u> | <u>N021802 004</u> Aug 01, 2006 |
| <u>AB</u> | + | | <u>20MG</u> | <u>N021802 003</u> May 26, 2005 |
| <u>AB</u> | + | | <u>25MG</u> | <u>N021802 008</u> Apr 21, 2011 |
| <u>AB</u> | + | | <u>30MG</u> | <u>N021802 005</u> Oct 23, 2009 |
| <u>AB</u> | + | | <u>35MG</u> | <u>N021802 007</u> Apr 21, 2011 |
| <u>AB</u> | + | | <u>40MG</u> | <u>N021802 006</u> Aug 11, 2010 |
| TABLET;ORAL | | | | |
| <u>DEXMETHYLPHENIDATE HYDROCHLORIDE</u> | | | | |
| <u>AB</u> | ABHAI INC | <u>2.5MG</u> | <u>A206931 001</u> | Dec 04, 2015 |
| <u>AB</u> | | <u>5MG</u> | <u>A206931 002</u> | Dec 04, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A206931 003</u> | Dec 04, 2015 |
| <u>AB</u> | CEDIPROF INC | <u>5MG</u> | <u>A209211 001</u> | Sep 19, 2018 |
| <u>AB</u> | | <u>10MG</u> | <u>A209211 002</u> | Sep 19, 2018 |
| <u>AB</u> | LANNETT CO INC | <u>2.5MG</u> | <u>A209468 001</u> | Sep 25, 2017 |
| <u>AB</u> | | <u>5MG</u> | <u>A209468 002</u> | Sep 25, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A209468 003</u> | Sep 25, 2017 |
| <u>AB</u> | NOVEL LABS INC | <u>2.5MG</u> | <u>A204534 001</u> | Dec 04, 2015 |
| <u>AB</u> | | <u>5MG</u> | <u>A204534 002</u> | Dec 04, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A204534 003</u> | Dec 04, 2015 |
| <u>AB</u> | RHODES PHARMS | <u>2.5MG</u> | <u>A208756 001</u> | Nov 20, 2017 |
| <u>AB</u> | | <u>5MG</u> | <u>A208756 002</u> | Nov 20, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A208756 003</u> | Nov 20, 2017 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>2.5MG</u> | <u>A201231 001</u> | Sep 24, 2015 |
| <u>AB</u> | | <u>5MG</u> | <u>A201231 002</u> | Sep 24, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A201231 003</u> | Sep 24, 2015 |
| <u>AB</u> | TEVA PHARMS | <u>2.5MG</u> | <u>A077107 003</u> | Jan 29, 2007 |

PRESCRIPTION DRUG PRODUCT LIST

DEXMETHYLPHENIDATE HYDROCHLORIDE

TABLET; ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

| | | | | |
|-----------|-----------------|--------------|--------------------|--------------|
| <u>AB</u> | | <u>5MG</u> | <u>A077107 001</u> | Jan 29, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A077107 002</u> | Jan 29, 2007 |
| <u>AB</u> | TRIS PHARMA INC | <u>2.5MG</u> | <u>A207901 001</u> | Aug 26, 2016 |
| <u>AB</u> | | <u>5MG</u> | <u>A207901 002</u> | Aug 26, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A207901 003</u> | Aug 26, 2016 |

FOCALIN

| | | | | | |
|-----------|---|----------|--------------|--------------------|--------------|
| <u>AB</u> | + | NOVARTIS | <u>2.5MG</u> | <u>N021278 001</u> | Nov 13, 2001 |
| <u>AB</u> | + | | <u>5MG</u> | <u>N021278 002</u> | Nov 13, 2001 |
| <u>AB</u> | + | ! | <u>10MG</u> | <u>N021278 003</u> | Nov 13, 2001 |

DEXRAZOXANE HYDROCHLORIDE

INJECTABLE; INJECTION

DEXRAZOXANE HYDROCHLORIDE

| | | | | | | | |
|-----------|---|-------------------------|---------------------------|---------------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | | GLAND PHARMA LTD | <u>EQ 500MG BASE/VIAL</u> | <u>A207321 001</u> | Nov 28, 2016 | | |
| <u>AP</u> | | MYLAN INSTITUTIONAL | <u>EQ 250MG BASE/VIAL</u> | <u>A200752 001</u> | Oct 19, 2011 | | |
| <u>AP</u> | | | <u>EQ 500MG BASE/VIAL</u> | <u>A200752 002</u> | Oct 19, 2011 | | |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>EQ 250MG BASE/VIAL</u> | <u>A076068 001</u> | Sep 28, 2004 | | |
| <u>AP</u> | | | <u>EQ 500MG BASE/VIAL</u> | <u>A076068 002</u> | Sep 28, 2004 | | |
| <u>AP</u> | + | ! | PHARMACIA AND UPJOHN | <u>EQ 250MG BASE/VIAL</u> | <u>N020212 001</u> | May 26, 1995 | |
| <u>AP</u> | + | ! | | <u>EQ 500MG BASE/VIAL</u> | <u>N020212 002</u> | May 26, 1995 | |
| | | TOTECT | | | | | |
| | | + | ! | CLINIGEN HLTHCARE | <u>EQ 500MG BASE/VIAL</u> | <u>N022025 001</u> | Sep 06, 2007 |

DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

DEXEDRINE

| | | | | | |
|-----------|---|----------------|-------------|--------------------|--|
| <u>AB</u> | + | IMPAX LABS INC | <u>5MG</u> | <u>N017078 001</u> | |
| <u>AB</u> | + | | <u>10MG</u> | <u>N017078 002</u> | |
| <u>AB</u> | + | ! | <u>15MG</u> | <u>N017078 003</u> | |

DEXTROAMPHETAMINE SULFATE

| | | | | | |
|-----------|--|-------------------|-------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS ELIZABETH | <u>5MG</u> | <u>A203901 001</u> | Nov 30, 2012 |
| <u>AB</u> | | | <u>10MG</u> | <u>A203901 002</u> | Nov 30, 2012 |
| <u>AB</u> | | | <u>15MG</u> | <u>A203901 003</u> | Nov 30, 2012 |
| <u>AB</u> | | MAYNE PHARMA | <u>5MG</u> | <u>A076137 001</u> | Jan 18, 2002 |
| <u>AB</u> | | | <u>10MG</u> | <u>A076137 002</u> | Jan 18, 2002 |
| <u>AB</u> | | | <u>15MG</u> | <u>A076137 003</u> | Jan 18, 2002 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>5MG</u> | <u>A206735 001</u> | Jan 27, 2016 |
| <u>AB</u> | | | <u>10MG</u> | <u>A206735 002</u> | Jan 27, 2016 |
| <u>AB</u> | | | <u>15MG</u> | <u>A206735 003</u> | Jan 27, 2016 |
| <u>AB</u> | | NESHER PHARMS | <u>5MG</u> | <u>A209111 001</u> | Jun 27, 2017 |
| <u>AB</u> | | | <u>10MG</u> | <u>A209111 002</u> | Jun 27, 2017 |
| <u>AB</u> | | | <u>15MG</u> | <u>A209111 003</u> | Jun 27, 2017 |
| <u>AB</u> | | SPECGX LLC | <u>5MG</u> | <u>A076353 001</u> | May 06, 2003 |
| <u>AB</u> | | | <u>10MG</u> | <u>A076353 002</u> | May 06, 2003 |
| <u>AB</u> | | | <u>15MG</u> | <u>A076353 003</u> | May 06, 2003 |
| <u>AB</u> | | VINTAGE PHARMS | <u>5MG</u> | <u>A205673 001</u> | Oct 31, 2017 |
| <u>AB</u> | | | <u>10MG</u> | <u>A205673 002</u> | Oct 31, 2017 |
| <u>AB</u> | | | <u>15MG</u> | <u>A205673 003</u> | Oct 31, 2017 |

SOLUTION; ORAL

DEXTROAMPHETAMINE SULFATE

| | | | | | |
|-----------|---|-----------------|----------------|--------------------|--------------|
| <u>AA</u> | ! | OUTLOOK PHARMS | <u>5MG/5ML</u> | <u>A040776 001</u> | Jan 29, 2008 |
| <u>AA</u> | | TRIS PHARMA INC | <u>5MG/5ML</u> | <u>A203644 001</u> | May 29, 2013 |

TABLET; ORAL

DEXTROAMPHETAMINE SULFATE

| | | | | | |
|-----------|---|---------------------|-------------|--------------------|--------------|
| <u>AA</u> | | ARBOR PHARMS LLC | <u>5MG</u> | <u>A090533 002</u> | Oct 25, 2011 |
| <u>AA</u> | | | <u>10MG</u> | <u>A090533 004</u> | Oct 25, 2011 |
| <u>AA</u> | | AUROLIFE PHARMA LLC | <u>5MG</u> | <u>A202893 001</u> | Jul 31, 2013 |
| <u>AA</u> | | | <u>10MG</u> | <u>A202893 002</u> | Jul 31, 2013 |
| <u>AA</u> | | AVANTHI INC | <u>5MG</u> | <u>A203548 001</u> | Nov 23, 2015 |
| <u>AA</u> | | | <u>10MG</u> | <u>A203548 002</u> | Nov 23, 2015 |
| <u>AA</u> | | BARR | <u>5MG</u> | <u>A040361 001</u> | Jan 31, 2001 |
| <u>AA</u> | ! | | <u>10MG</u> | <u>A040361 002</u> | Jan 31, 2001 |
| <u>AA</u> | | NESHER PHARMS | <u>5MG</u> | <u>A206588 001</u> | Mar 28, 2016 |
| <u>AA</u> | | | <u>10MG</u> | <u>A206588 002</u> | Mar 28, 2016 |
| <u>AA</u> | | NOVEL LABS INC | <u>5MG</u> | <u>A204330 001</u> | Mar 16, 2016 |
| <u>AA</u> | | | <u>10MG</u> | <u>A204330 002</u> | Mar 16, 2016 |
| <u>AA</u> | | NUVO PHARM | <u>5MG</u> | <u>A210059 001</u> | Oct 18, 2017 |
| <u>AA</u> | | | <u>10MG</u> | <u>A210059 002</u> | Oct 18, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMPHETAMINE SULFATE

| | | | | | |
|-----------|------------------|-------------|----------------|------------|--------------|
| <u>AA</u> | SPECGX LLC | <u>5MG</u> | <u>A040436</u> | <u>001</u> | Jan 29, 2002 |
| <u>AA</u> | | <u>10MG</u> | <u>A040436</u> | <u>002</u> | Jan 29, 2002 |
| | ARBOR PHARMS LLC | 2.5MG | A090533 | 001 | Oct 25, 2011 |
| | | 7.5MG | A090533 | 003 | Oct 25, 2011 |
| | | 15MG | A090533 | 005 | Oct 25, 2011 |
| | | 20MG | A090533 | 006 | Oct 25, 2011 |
| | | 30MG | A090533 | 007 | Oct 25, 2011 |

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETHAZINE DM

| | | | | | | |
|-----------|---|---------|-----------------------------|----------------|------------|--------------|
| <u>AA</u> | ! | VINTAGE | <u>15MG/5ML; 6.25MG/5ML</u> | <u>A040649</u> | <u>001</u> | Feb 14, 2006 |
|-----------|---|---------|-----------------------------|----------------|------------|--------------|

PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

| | | | | | | |
|-----------|--|----------------|-----------------------------|----------------|------------|--------------|
| <u>AA</u> | | HI TECH PHARMA | <u>15MG/5ML; 6.25MG/5ML</u> | <u>A040027</u> | <u>001</u> | Jul 31, 1996 |
|-----------|--|----------------|-----------------------------|----------------|------------|--------------|

PROMETHAZINE W/ DEXTROMETHORPHAN

| | | | | | | |
|-----------|--|------------------|-----------------------------|----------------|------------|--------------|
| <u>AA</u> | | WOCKHARDT BIO AG | <u>15MG/5ML; 6.25MG/5ML</u> | <u>A088864</u> | <u>001</u> | Jan 04, 1985 |
|-----------|--|------------------|-----------------------------|----------------|------------|--------------|

DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE

CAPSULE; ORAL

DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE

| | | | | | | |
|-----------|--|-------------------|-------------------|----------------|------------|--------------|
| <u>AB</u> | | ACTAVIS ELIZABETH | <u>20MG; 10MG</u> | <u>A202934</u> | <u>001</u> | Oct 10, 2017 |
|-----------|--|-------------------|-------------------|----------------|------------|--------------|

NUDEXTA

| | | | | | | |
|-----------|---|---------------|-------------------|----------------|------------|--------------|
| <u>AB</u> | + | AVANIR PHARMS | <u>20MG; 10MG</u> | <u>N021879</u> | <u>001</u> | Oct 29, 2010 |
|-----------|---|---------------|-------------------|----------------|------------|--------------|

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 10% IN PLASTIC CONTAINER

| | | | | | | |
|-----------|---|---------|-------------------|----------------|------------|--------------|
| <u>AP</u> | + | B BRAUN | <u>10GM/100ML</u> | <u>N019626</u> | <u>004</u> | Feb 02, 1988 |
|-----------|---|---------|-------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|-----------------|-------------------|----------------|------------|--|
| <u>AP</u> | + | BAXTER HLTHCARE | <u>10GM/100ML</u> | <u>N016694</u> | <u>001</u> | |
|-----------|---|-----------------|-------------------|----------------|------------|--|

| | | | | | | |
|-----------|--|--------------------|-------------------|----------------|------------|--------------|
| <u>AP</u> | | FRESENIUS KABI USA | <u>10GM/100ML</u> | <u>A209448</u> | <u>001</u> | Jul 16, 2018 |
|-----------|--|--------------------|-------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|-----------------|-------------------|----------------|------------|--|
| <u>AP</u> | + | ICU MEDICAL INC | <u>10GM/100ML</u> | <u>N018080</u> | <u>001</u> | |
|-----------|---|-----------------|-------------------|----------------|------------|--|

DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | | |
|-----------|---|---------|----------------|----------------|------------|--|
| <u>AP</u> | + | B BRAUN | <u>50MG/ML</u> | <u>N016730</u> | <u>002</u> | |
|-----------|---|---------|----------------|----------------|------------|--|

| | | | | | | |
|-----------|---|--|------------------|----------------|------------|--|
| <u>AP</u> | + | | <u>5GM/100ML</u> | <u>N016730</u> | <u>001</u> | |
|-----------|---|--|------------------|----------------|------------|--|

| | | | | | | |
|-----------|---|--|------------------|----------------|------------|--------------|
| <u>AP</u> | + | | <u>5GM/100ML</u> | <u>N019626</u> | <u>002</u> | Feb 02, 1988 |
|-----------|---|--|------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|-----------------|----------------|----------------|------------|--------------|
| <u>AP</u> | + | BAXTER HLTHCARE | <u>50MG/ML</u> | <u>N016673</u> | <u>003</u> | Oct 30, 1985 |
|-----------|---|-----------------|----------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|--|----------------|----------------|------------|--------------|
| <u>AP</u> | + | | <u>50MG/ML</u> | <u>N020179</u> | <u>002</u> | Dec 07, 1992 |
|-----------|---|--|----------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|--|------------------|----------------|------------|--|
| <u>AP</u> | + | | <u>5GM/100ML</u> | <u>N016673</u> | <u>001</u> | |
|-----------|---|--|------------------|----------------|------------|--|

| | | | | | | |
|-----------|---|--|------------------|----------------|------------|--------------|
| <u>AP</u> | + | | <u>5GM/100ML</u> | <u>N020179</u> | <u>001</u> | Dec 07, 1992 |
|-----------|---|--|------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--------------------|----------------|----------------|------------|--------------|
| <u>AP</u> | | FRESENIUS KABI USA | <u>50MG/ML</u> | <u>A207449</u> | <u>001</u> | Oct 21, 2016 |
|-----------|--|--------------------|----------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|---------|------------------|----------------|------------|--------------|
| <u>AP</u> | + | HOSPIRA | <u>5GM/100ML</u> | <u>N019466</u> | <u>001</u> | Jul 15, 1985 |
|-----------|---|---------|------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|--|------------------|----------------|------------|--------------|
| <u>AP</u> | + | | <u>5GM/100ML</u> | <u>N019479</u> | <u>001</u> | Sep 17, 1985 |
|-----------|---|--|------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|-----------------|----------------|----------------|------------|--|
| <u>AP</u> | + | ICU MEDICAL INC | <u>50MG/ML</u> | <u>N016367</u> | <u>002</u> | |
|-----------|---|-----------------|----------------|----------------|------------|--|

DEXTROSE 50% IN PLASTIC CONTAINER

| | | | | | | |
|-----------|---|-----------------|-------------------|----------------|------------|--------------|
| <u>AP</u> | + | BAXTER HLTHCARE | <u>50GM/100ML</u> | <u>N020047</u> | <u>001</u> | Jul 02, 1991 |
|-----------|---|-----------------|-------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|-----------------|-------------------|----------------|------------|--------------|
| <u>AP</u> | + | ICU MEDICAL INC | <u>50GM/100ML</u> | <u>N018563</u> | <u>001</u> | Mar 23, 1982 |
|-----------|---|-----------------|-------------------|----------------|------------|--------------|

DEXTROSE 70% IN PLASTIC CONTAINER

| | | | | | | |
|-----------|---|---------|-------------------|----------------|------------|--------------|
| <u>AP</u> | + | B BRAUN | <u>70GM/100ML</u> | <u>N019626</u> | <u>005</u> | Feb 18, 2015 |
|-----------|---|---------|-------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|-----------------|-------------------|----------------|------------|--------------|
| <u>AP</u> | + | BAXTER HLTHCARE | <u>70GM/100ML</u> | <u>N017521</u> | <u>006</u> | Mar 26, 1982 |
|-----------|---|-----------------|-------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|--|-------------------|----------------|------------|--------------|
| <u>AP</u> | + | | <u>70GM/100ML</u> | <u>N020047</u> | <u>003</u> | Jul 02, 1991 |
|-----------|---|--|-------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|-----------------|-------------------|----------------|------------|--------------|
| <u>AP</u> | + | ICU MEDICAL INC | <u>70GM/100ML</u> | <u>N018561</u> | <u>001</u> | Mar 23, 1982 |
|-----------|---|-----------------|-------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|--|-------------------|----------------|------------|--------------|
| <u>AP</u> | + | | <u>70GM/100ML</u> | <u>N019893</u> | <u>001</u> | Dec 26, 1989 |
|-----------|---|--|-------------------|----------------|------------|--------------|

DEXTROSE 20% IN PLASTIC CONTAINER

| | | | | | | |
|--|---|-----------------|------------|---------|-----|--------------|
| | + | ICU MEDICAL INC | 20GM/100ML | N018564 | 001 | Mar 23, 1982 |
|--|---|-----------------|------------|---------|-----|--------------|

DEXTROSE 25%

| | | | | | | |
|--|---|---------|----------|---------|-----|--------------|
| | + | HOSPIRA | 250MG/ML | N019445 | 002 | Nov 23, 1998 |
|--|---|---------|----------|---------|-----|--------------|

DEXTROSE 30% IN PLASTIC CONTAINER

| | | | | | | |
|--|---|-----------------|------------|---------|-----|--------------|
| | + | ICU MEDICAL INC | 30GM/100ML | N019345 | 001 | Jan 26, 1985 |
|--|---|-----------------|------------|---------|-----|--------------|

DEXTROSE 40% IN PLASTIC CONTAINER

| | | | | | | |
|--|---|-----------------|------------|---------|-----|--------------|
| | + | ICU MEDICAL INC | 40GM/100ML | N018562 | 001 | Mar 23, 1982 |
|--|---|-----------------|------------|---------|-----|--------------|

DEXTROSE 50%

| | | | | | | |
|--|---|---------|----------|---------|-----|--------------|
| | + | HOSPIRA | 500MG/ML | N019445 | 003 | Sep 03, 2014 |
|--|---|---------|----------|---------|-----|--------------|

DEXTROSE 50% IN PLASTIC CONTAINER

| | | | | | | |
|--|---|---------|----------|---------|-----|--------------|
| | + | HOSPIRA | 500MG/ML | N019445 | 001 | Jun 03, 1986 |
|--|---|---------|----------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | |
|-----------------|-------------------------------------------------|---------|-----|
| ICU MEDICAL INC | 5GM/100ML; 21MG/100ML; 128MG/100ML; 234MG/100ML | N017610 | 001 |
|-----------------|-------------------------------------------------|---------|-----|

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE

INJECTABLE; INJECTION

ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|---------|-------------------------------------------------------------|---------|-----|--------------|
| B BRAUN | 5GM/100ML; 31MG/100ML; 130MG/100ML; 26MG/100ML; 320MG/100ML | N019873 | 001 | Jun 10, 1993 |
|---------|-------------------------------------------------------------|---------|-----|--------------|

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 5% AND ELECTROLYTE NO. 48 IN PLASTIC CONTAINER

| | | | |
|-----------------|-------------------------------------------------------------------------|---------|-----|
| BAXTER HLTHCARE | 5GM/100ML; 31MG/100ML; 141MG/100ML; 20MG/100ML; 12MG/100ML; 260MG/100ML | N017484 | 001 |
|-----------------|-------------------------------------------------------------------------|---------|-----|

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM LACTATE; SODIUM PHOSPHATE, MONOBASIC ANHYDROUS

INJECTABLE; INJECTION

IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|-----------------|-------------------------------------------------------------------------|---------|-----|--------------|
| ICU MEDICAL INC | 5GM/100ML; 30MG/100ML; 141MG/100ML; 15MG/100ML; 260MG/100ML; 25MG/100ML | N019513 | 001 | May 08, 1986 |
|-----------------|-------------------------------------------------------------------------|---------|-----|--------------|

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | |
|-----------------|--------------------------------------------------------------------------|---------|-----|
| ICU MEDICAL INC | 5GM/100ML; 30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML | N017609 | 001 |
|-----------------|--------------------------------------------------------------------------|---------|-----|

DEXTROSE; POTASSIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER

| | | | | | |
|----|---|-----------------|------------------------|---------|-----|
| AP | + | BAXTER HLTHCARE | 5GM/100ML; 150MG/100ML | N017634 | 001 |
|----|---|-----------------|------------------------|---------|-----|

DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER

| | | | | | |
|----|---|-----------------|------------------------|---------|-----|
| AP | + | BAXTER HLTHCARE | 5GM/100ML; 224MG/100ML | N017634 | 003 |
|----|---|-----------------|------------------------|---------|-----|

DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

| | | | | | |
|----|---|-----------------|------------------------|---------|-----|
| AP | + | BAXTER HLTHCARE | 5GM/100ML; 300MG/100ML | N017634 | 002 |
|----|---|-----------------|------------------------|---------|-----|

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | | |
|----|--|---------|------------------------|---------|-----|--------------|
| AP | | B BRAUN | 5GM/100ML; 150MG/100ML | N019699 | 004 | Sep 29, 1989 |
|----|--|---------|------------------------|---------|-----|--------------|

POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | | |
|----|--|---------|------------------------|---------|-----|--------------|
| AP | | B BRAUN | 5GM/100ML; 300MG/100ML | N019699 | 006 | Sep 29, 1989 |
|----|--|---------|------------------------|---------|-----|--------------|

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | |
|----|--|-----------------|------------------------|---------|-----|
| AP | | ICU MEDICAL INC | 5GM/100ML; 224MG/100ML | N018371 | 003 |
|----|--|-----------------|------------------------|---------|-----|

DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER

| | | | | |
|---|-----------------|-----------------------|---------|-----|
| + | BAXTER HLTHCARE | 5GM/100ML; 75MG/100ML | N017634 | 004 |
|---|-----------------|-----------------------|---------|-----|

POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | |
|-----------------|------------------------|---------|-----|
| ICU MEDICAL INC | 5GM/100ML; 149MG/100ML | N018371 | 001 |
|-----------------|------------------------|---------|-----|

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | |
|-----------------|------------------------|---------|-----|
| ICU MEDICAL INC | 5GM/100ML; 298MG/100ML | N018371 | 002 |
|-----------------|------------------------|---------|-----|

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ

| | | | | | | |
|----|--|-----------------|------------------------------------|---------|-----|--------------|
| AP | | BAXTER HLTHCARE | 5GM/100ML; 75MG/100ML; 200MG/100ML | N018037 | 006 | Apr 13, 1982 |
|----|--|-----------------|------------------------------------|---------|-----|--------------|

| | | | | | | |
|----|--|-----------------|-------------------------------------|---------|-----|--------------|
| AP | | BAXTER HLTHCARE | 5GM/100ML; 150MG/100ML; 200MG/100ML | N018037 | 007 | Apr 13, 1982 |
|----|--|-----------------|-------------------------------------|---------|-----|--------------|

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K)

| | | | | | |
|----|--|-----------------|-------------------------------------|---------|-----|
| AP | | BAXTER HLTHCARE | 5GM/100ML; 224MG/100ML; 200MG/100ML | N018037 | 004 |
|----|--|-----------------|-------------------------------------|---------|-----|

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ

| | | | | | | |
|----|--|-----------------|-------------------------------------|---------|-----|--------------|
| AP | | BAXTER HLTHCARE | 5GM/100ML; 150MG/100ML; 200MG/100ML | N018037 | 008 | Apr 13, 1982 |
|----|--|-----------------|-------------------------------------|---------|-----|--------------|

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K)

| | | | | | |
|----|--|-----------------|-------------------------------------|---------|-----|
| AP | | BAXTER HLTHCARE | 5GM/100ML; 300MG/100ML; 200MG/100ML | N018037 | 001 |
|----|--|-----------------|-------------------------------------|---------|-----|

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ

| | | | | | | |
|----|--|-----------------|-------------------------------------|---------|-----|--------------|
| AP | | BAXTER HLTHCARE | 5GM/100ML; 224MG/100ML; 200MG/100ML | N018037 | 005 | Apr 13, 1982 |
|----|--|-----------------|-------------------------------------|---------|-----|--------------|

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ

| | | | | | | |
|----|--|-----------------|-------------------------------------|---------|-----|--------------|
| AP | | BAXTER HLTHCARE | 5GM/100ML; 300MG/100ML; 200MG/100ML | N018037 | 009 | Apr 13, 1982 |
|----|--|-----------------|-------------------------------------|---------|-----|--------------|

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ

| | | | | | |
|----|--|-----------------|------------------------------------|---------|-----|
| AP | | BAXTER HLTHCARE | 5GM/100ML; 75MG/100ML; 200MG/100ML | N018037 | 002 |
|----|--|-----------------|------------------------------------|---------|-----|

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K)

| | | | | | |
|----|--|-----------------|-------------------------------------|---------|-----|
| AP | | BAXTER HLTHCARE | 5GM/100ML; 150MG/100ML; 200MG/100ML | N018037 | 003 |
|----|--|-----------------|-------------------------------------|---------|-----|

DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER

| | | | | | | |
|----|--|-----------------|------------------------------------|---------|-----|--------------|
| AP | | BAXTER HLTHCARE | 5GM/100ML; 75MG/100ML; 330MG/100ML | N018629 | 005 | Mar 23, 1982 |
|----|--|-----------------|------------------------------------|---------|-----|--------------|

| | | | | | | |
|----|--|-----------------|-------------------------------------|---------|-----|--------------|
| AP | | BAXTER HLTHCARE | 5GM/100ML; 150MG/100ML; 330MG/100ML | N018629 | 002 | Mar 23, 1982 |
|----|--|-----------------|-------------------------------------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDEINJECTABLE; INJECTION

| <u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER</u> | | | |
|-------------------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------|-------------------------------------------|---------------------------------|
| <u>AP</u> | <u>BAXTER HLTHCARE</u> | <u>5GM/100ML;224MG/100ML;330MG/100ML</u> | <u>N018629 003</u> Mar 23, 1982 |
| <u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>BAXTER HLTHCARE</u> | <u>5GM/100ML;150MG/100ML;330MG/100ML</u> | <u>N018629 004</u> Mar 23, 1982 |
| <u>AP</u> | | <u>5GM/100ML;300MG/100ML;330MG/100ML</u> | <u>N018629 006</u> Mar 23, 1982 |
| <u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>BAXTER HLTHCARE</u> | <u>5GM/100ML;224MG/100ML;330MG/100ML</u> | <u>N018629 007</u> Mar 23, 1982 |
| <u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>BAXTER HLTHCARE</u> | <u>5GM/100ML;300MG/100ML;330MG/100ML</u> | <u>N018629 008</u> Mar 23, 1982 |
| <u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>BAXTER HLTHCARE</u> | <u>5GM/100ML;75MG/100ML;330MG/100ML</u> | <u>N018629 001</u> Mar 23, 1982 |
| <u>DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>BAXTER HLTHCARE</u> | <u>5GM/100ML;300MG/100ML;450MG/100ML</u> | <u>N018008 010</u> |
| <u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>B BRAUN</u> | <u>5GM/100ML;75MG/100ML;200MG/100ML</u> | <u>N019630 008</u> Feb 17, 1988 |
| <u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>B BRAUN</u> | <u>5GM/100ML;75MG/100ML;330MG/100ML</u> | <u>N019630 014</u> Feb 17, 1988 |
| <u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>B BRAUN</u> | <u>5GM/100ML;75MG/100ML;450MG/100ML</u> | <u>N019630 020</u> Feb 17, 1988 |
| <u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>B BRAUN</u> | <u>5GM/100ML;75MG/100ML;900MG/100ML</u> | <u>N019630 026</u> Feb 17, 1988 |
| <u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>B BRAUN</u> | <u>5GM/100ML;150MG/100ML;200MG/100ML</u> | <u>N019630 010</u> Feb 17, 1988 |
| <u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>B BRAUN</u> | <u>5GM/100ML;150MG/100ML;330MG/100ML</u> | <u>N019630 016</u> Feb 17, 1988 |
| <u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>B BRAUN</u> | <u>5GM/100ML;150MG/100ML;450MG/100ML</u> | <u>N019630 022</u> Feb 17, 1988 |
| <u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>B BRAUN</u> | <u>5GM/100ML;150MG/100ML;900MG/100ML</u> | <u>N019630 028</u> Feb 17, 1988 |
| <u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>B BRAUN</u> | <u>5GM/100ML;300MG/100ML;200MG/100ML</u> | <u>N019630 012</u> Feb 17, 1988 |
| <u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>B BRAUN</u> | <u>5GM/100ML;300MG/100ML;330MG/100ML</u> | <u>N019630 018</u> Feb 17, 1988 |
| <u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>B BRAUN</u> | <u>5GM/100ML;300MG/100ML;450MG/100ML</u> | <u>N019630 024</u> Feb 17, 1988 |
| <u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>B BRAUN</u> | <u>5GM/100ML;300MG/100ML;900MG/100ML</u> | <u>N019630 030</u> Feb 17, 1988 |
| <u>POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>BAXTER HLTHCARE</u> | <u>5GM/100ML;75MG/100ML;450MG/100ML</u> | <u>N018008 005</u> Apr 28, 1982 |
| <u>AP</u> | | <u>5GM/100ML;150MG/100ML;450MG/100ML</u> | <u>N018008 006</u> Apr 28, 1982 |
| <u>AP</u> | <u>+!</u> <u>ICU MEDICAL INC</u> | <u>5GM/100ML;74.5MG/100ML;450MG/100ML</u> | <u>N018362 009</u> Jul 05, 1983 |
| <u>AP</u> | <u>+!</u> | <u>5GM/100ML;149MG/100ML;450MG/100ML</u> | <u>N018362 005</u> Mar 28, 1988 |
| <u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>BAXTER HLTHCARE</u> | <u>5GM/100ML;150MG/100ML;450MG/100ML</u> | <u>N018008 007</u> Apr 28, 1982 |
| <u>AP</u> | <u>+!</u> <u>ICU MEDICAL INC</u> | <u>5GM/100ML;149MG/100ML;450MG/100ML</u> | <u>N018362 010</u> Jul 05, 1983 |
| <u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>BAXTER HLTHCARE</u> | <u>5GM/100ML;150MG/100ML;900MG/100ML</u> | <u>N019308 005</u> Apr 05, 1985 |
| <u>AP</u> | <u>+!</u> <u>ICU MEDICAL INC</u> | <u>5GM/100ML;149MG/100ML;900MG/100ML</u> | <u>N019691 005</u> Mar 24, 1988 |
| <u>POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>BAXTER HLTHCARE</u> | <u>5GM/100ML;224MG/100ML;450MG/100ML</u> | <u>N018008 008</u> Apr 28, 1982 |
| <u>AP</u> | <u>+!</u> <u>ICU MEDICAL INC</u> | <u>5GM/100ML;224MG/100ML;450MG/100ML</u> | <u>N018362 002</u> |
| <u>POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>BAXTER HLTHCARE</u> | <u>5GM/100ML;300MG/100ML;450MG/100ML</u> | <u>N018008 009</u> Apr 28, 1982 |
| <u>AP</u> | <u>+!</u> <u>ICU MEDICAL INC</u> | <u>5GM/100ML;298MG/100ML;450MG/100ML</u> | <u>N018362 003</u> |
| <u>POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u> | | | |
| <u>AP</u> | <u>BAXTER HLTHCARE</u> | <u>5GM/100ML;300MG/100ML;900MG/100ML</u> | <u>N019308 007</u> Apr 05, 1985 |
| <u>AP</u> | <u>+!</u> <u>ICU MEDICAL INC</u> | <u>5GM/100ML;298MG/100ML;900MG/100ML</u> | <u>N019691 009</u> Mar 24, 1988 |
| | <u>POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u> | | |
| | <u>B BRAUN</u> | <u>10GM/100ML;37MG/100ML;200MG/100ML</u> | <u>N019630 031</u> Feb 17, 1988 |
| | <u>POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u> | | |
| | <u>B BRAUN</u> | <u>10GM/100ML;37MG/100ML;450MG/100ML</u> | <u>N019630 037</u> Feb 17, 1988 |
| | <u>POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u> | | |
| | <u>B BRAUN</u> | <u>10GM/100ML;37MG/100ML;900MG/100ML</u> | <u>N019630 043</u> Feb 17, 1988 |
| | <u>POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER</u> | | |
| | <u>B BRAUN</u> | <u>5GM/100ML;37MG/100ML;110MG/100ML</u> | <u>N019630 001</u> Feb 17, 1988 |
| | <u>POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u> | | |
| | <u>B BRAUN</u> | <u>5GM/100ML;37MG/100ML;200MG/100ML</u> | <u>N019630 007</u> Feb 17, 1988 |
| | <u>POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u> | | |
| | <u>B BRAUN</u> | <u>5GM/100ML;37MG/100ML;330MG/100ML</u> | <u>N019630 013</u> Feb 17, 1988 |

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | | | | |
|--------------------------|-----------------|----------------------------------------|----|-------------------|--------------------------|
| POTASSIUM CHLORIDE 10MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.225% | IN | PLASTIC CONTAINER | |
| + | ICU MEDICAL INC | 5GM/100ML;74.5MG/100ML;225MG/100ML | | | N018365 002 Jul 05, 1983 |
| + | | 5GM/100ML;149MG/100ML;225MG/100ML | | | N018365 006 Mar 28, 1988 |
| POTASSIUM CHLORIDE 10MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.3% | IN | PLASTIC CONTAINER | |
| | ICU MEDICAL INC | 5GM/100ML;74.5MG/100ML;300MG/100ML | | | N018876 001 Jan 17, 1986 |
| | | 5GM/100ML;149MG/100ML;300MG/100ML | | | N018876 006 Mar 28, 1988 |
| POTASSIUM CHLORIDE 10MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.9% | IN | PLASTIC CONTAINER | |
| | BAXTER HLTHCARE | 5GM/100ML;75MG/100ML;900MG/100ML | | | N019308 004 Apr 05, 1985 |
| | | 5GM/100ML;150MG/100ML;900MG/100ML | | | N019308 002 Apr 05, 1985 |
| POTASSIUM CHLORIDE 15MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.225% | IN | PLASTIC CONTAINER | |
| | ICU MEDICAL INC | 5GM/100ML;224MG/100ML;225MG/100ML | | | N018365 008 Mar 28, 1988 |
| POTASSIUM CHLORIDE 15MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.3% | IN | PLASTIC CONTAINER | |
| | ICU MEDICAL INC | 5GM/100ML;224MG/100ML;300MG/100ML | | | N018876 007 Mar 28, 1988 |
| POTASSIUM CHLORIDE 20MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.225% | IN | PLASTIC CONTAINER | |
| + | ICU MEDICAL INC | 5GM/100ML;149MG/100ML;225MG/100ML | | | N018365 001 |
| + | | 5GM/100ML;298MG/100ML;225MG/100ML | | | N018365 009 Mar 28, 1988 |
| POTASSIUM CHLORIDE 20MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.3% | IN | PLASTIC CONTAINER | |
| | ICU MEDICAL INC | 5GM/100ML;298MG/100ML;300MG/100ML | | | N018876 008 Mar 28, 1988 |
| POTASSIUM CHLORIDE 20MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.9% | IN | PLASTIC CONTAINER | |
| | BAXTER HLTHCARE | 5GM/100ML;300MG/100ML;900MG/100ML | | | N019308 003 Apr 05, 1985 |
| POTASSIUM CHLORIDE 20MEQ | IN | DEXTROSE 5% IN SODIUM CHLORIDE 0.3% | IN | PLASTIC CONTAINER | |
| | ICU MEDICAL INC | 5GM/100ML;149MG/100ML;300MG/100ML | | | N018876 002 Jan 17, 1986 |
| POTASSIUM CHLORIDE 30MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.225% | IN | PLASTIC CONTAINER | |
| | ICU MEDICAL INC | 5GM/100ML;224MG/100ML;225MG/100ML | | | N018365 003 Jul 05, 1983 |
| POTASSIUM CHLORIDE 30MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.3% | IN | PLASTIC CONTAINER | |
| | ICU MEDICAL INC | 5GM/100ML;224MG/100ML;300MG/100ML | | | N018876 003 Jan 17, 1986 |
| POTASSIUM CHLORIDE 30MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.9% | IN | PLASTIC CONTAINER | |
| | BAXTER HLTHCARE | 5GM/100ML;224MG/100ML;900MG/100ML | | | N019308 006 Apr 05, 1985 |
| POTASSIUM CHLORIDE 40MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.225% | IN | PLASTIC CONTAINER | |
| | ICU MEDICAL INC | 5GM/100ML;298MG/100ML;225MG/100ML | | | N018365 004 Jul 05, 1983 |
| POTASSIUM CHLORIDE 40MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.3% | IN | PLASTIC CONTAINER | |
| | ICU MEDICAL INC | 5GM/100ML;298MG/100ML;300MG/100ML | | | N018876 004 Mar 28, 1988 |
| POTASSIUM CHLORIDE 5MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.225% | IN | PLASTIC CONTAINER | |
| | ICU MEDICAL INC | 5GM/100ML;74.5MG/100ML;225MG/100ML | | | N018365 005 Mar 28, 1988 |
| | | 5GM/100ML;149MG/100ML;225MG/100ML | | | N018365 007 Mar 28, 1988 |
| POTASSIUM CHLORIDE 5MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.3% | IN | PLASTIC CONTAINER | |
| | ICU MEDICAL INC | 5GM/100ML;74.5MG/100ML;300MG/100ML | | | N018876 005 Mar 28, 1988 |
| | | 5GM/100ML;149MG/100ML;300MG/100ML | | | N018876 009 Mar 28, 1988 |
| POTASSIUM CHLORIDE 5MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.45% | IN | PLASTIC CONTAINER | |
| | BAXTER HLTHCARE | 5GM/100ML;150MG/100ML;450MG/100ML | | | N018008 004 |
| POTASSIUM CHLORIDE 5MEQ | IN | DEXTROSE 5% AND SODIUM CHLORIDE 0.9% | IN | PLASTIC CONTAINER | |
| | BAXTER HLTHCARE | 5GM/100ML;150MG/100ML;900MG/100ML | | | N019308 001 Apr 05, 1985 |

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

| | | | | | |
|--------------------------------------------------------------------|-----------------|--------------------------------|--------------------------------|--|---------------------------------|
| AP | B BRAUN | <u>2.5GM/100ML;450MG/100ML</u> | | | N019631 004 Feb 24, 1988 |
| AP | + | <u>BAXTER HLTHCARE</u> | <u>2.5GM/100ML;450MG/100ML</u> | | N016697 001 |
| <u>DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u> | | | | | |
| AP | B BRAUN | <u>5GM/100ML;200MG/100ML</u> | | | N019631 007 Feb 24, 1988 |
| <u>DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u> | | | | | |
| AP | B BRAUN | <u>5GM/100ML;330MG/100ML</u> | | | N019631 008 Feb 24, 1988 |
| <u>DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u> | | | | | |
| AP | B BRAUN | <u>5GM/100ML;450MG/100ML</u> | | | N019631 009 Feb 24, 1988 |
| AP | + | <u>ICU MEDICAL INC</u> | <u>5GM/100ML;450MG/100ML</u> | | N017607 001 |
| <u>DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u> | | | | | |
| AP | B BRAUN | <u>5GM/100ML;900MG/100ML</u> | | | N019631 010 Feb 24, 1988 |
| AP | + | <u>ICU MEDICAL INC</u> | <u>5GM/100ML;900MG/100ML</u> | | N017585 001 |
| <u>DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u> | | | | | |
| AP | BAXTER HLTHCARE | <u>5GM/100ML;200MG/100ML</u> | | | N016689 001 |
| <u>DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u> | | | | | |
| AP | BAXTER HLTHCARE | <u>5GM/100ML;330MG/100ML</u> | | | N016687 001 |
| <u>DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u> | | | | | |
| AP | BAXTER HLTHCARE | <u>5GM/100ML;450MG/100ML</u> | | | N016683 001 |
| <u>DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u> | | | | | |
| AP | BAXTER HLTHCARE | <u>5GM/100ML;900MG/100ML</u> | | | N016678 001 |
| <u>DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER</u> | | | | | |
| | B BRAUN | 10GM/100ML;110MG/100ML | | | N019631 011 Feb 24, 1988 |
| <u>DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u> | | | | | |
| | B BRAUN | 10GM/100ML;200MG/100ML | | | N019631 012 Feb 24, 1988 |

PRESCRIPTION DRUG PRODUCT LIST

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | |
|---------------------------------------------------------------------------------------------------------|-------------|--------------|
| DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER B BRAUN 10GM/100ML;330MG/100ML | N019631 013 | Feb 24, 1988 |
| DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER B BRAUN 10GM/100ML;450MG/100ML | N019631 014 | Feb 24, 1988 |
| DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER B BRAUN 10GM/100ML;900MG/100ML | N019631 015 | Feb 24, 1988 |
| DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER B BRAUN 2.5GM/100ML;110MG/100ML | N019631 001 | Feb 24, 1988 |
| DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER B BRAUN 2.5GM/100ML;200MG/100ML | N019631 002 | Feb 24, 1988 |
| DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER B BRAUN 2.5GM/100ML;330MG/100ML | N019631 003 | Feb 24, 1988 |
| DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER B BRAUN 2.5GM/100ML;900MG/100ML | N019631 005 | Feb 24, 1988 |
| DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER B BRAUN 3.3GM/100ML;300MG/100ML | N019631 016 | Jan 19, 1990 |
| DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER B BRAUN 5GM/100ML;110MG/100ML | N019631 006 | Feb 24, 1988 |
| DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER +! ICU MEDICAL INC 5GM/100ML;225MG/100ML | N017606 001 | |
| DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER +! ICU MEDICAL INC 5GM/100ML;300MG/100ML | N017799 001 | |

DIATRIZOATE MEGLUMINE

SOLUTION; URETHRAL

| | | |
|------------------------------------|-------------|--------------|
| CYSTOGRAFIN + BRACCO 30% | N010040 018 | |
| CYSTOGRAFIN DILUTE + BRACCO 18% | N010040 022 | Nov 09, 1982 |

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

SOLUTION; ORAL, RECTAL

GASTROGRAFIN

| | | |
|------------------------------------------------------------|--------------------|--|
| AA +! BRACCO 66%;10% | N011245 003 | |
| AA <u>MD-GASTROVIEW</u> LIEBEL-FLARSHEIM 66%;10% | A087388 001 | |

DIAZEPAM

CONCENTRATE; ORAL

DIAZEPAM

| | | |
|---------------------------------------------------------------------|--------------------|--------------|
| AA LANNETT CO INC 5MG/ML | A204433 001 | Apr 14, 2014 |
| AA ! <u>DIAZEPAM INTENSOL</u> WEST-WARD PHARMS INT 5MG/ML | A071415 001 | Apr 03, 1987 |

GEL; RECTAL

DIASTAT

| | | |
|-------------------------------------------------------------|-------------|--------------|
| + VALEANT PHARMS NORTH 2.5MG/0.5ML (5MG/ML) | N020648 001 | Jul 29, 1997 |
| DIASTAT ACUDIAL + VALEANT PHARMS NORTH 10MG/2ML (5MG/ML) | N020648 007 | Sep 15, 2005 |
| +! 20MG/4ML (5MG/ML) | N020648 006 | Sep 15, 2005 |

INJECTABLE; INJECTION

DIAZEPAM

| | | |
|-----------------------------|-------------|--------------|
| ! HOSPIRA 10MG/2ML (5MG/ML) | A072079 001 | Dec 20, 1988 |
| ! 50MG/10ML (5MG/ML) | A071583 001 | Oct 13, 1987 |

SOLUTION; ORAL

DIAZEPAM

| | | |
|------------------------------------------|--------------------|--------------|
| AA LANNETT CO INC 5MG/5ML | A206477 001 | Jun 24, 2016 |
| AA ! WEST-WARD PHARMS INT 5MG/5ML | A070928 001 | Apr 03, 1987 |

TABLET; ORAL

DIAZEPAM

| | | |
|------------------------------------|--------------------|--------------|
| AB BARR 2MG | A070152 001 | Nov 01, 1985 |
| AB 10MG | A070154 001 | Nov 01, 1985 |
| AB IVAX SUB TEVA PHARMS 2MG | A071307 001 | Dec 10, 1986 |
| AB 5MG | A071321 001 | Dec 10, 1986 |
| AB 10MG | A071322 001 | Dec 10, 1986 |
| AB MAYNE PHARMA 2MG | A071134 001 | Feb 03, 1987 |
| AB 5MG | A071135 001 | Feb 03, 1987 |
| AB 10MG | A071136 001 | Feb 03, 1987 |
| AB MYLAN 2MG | A070325 002 | Sep 04, 1985 |

PRESCRIPTION DRUG PRODUCT LIST

DIAZEPAM

TABLET; ORAL

DIAZEPAM

| | | | | |
|-----------|----------------|-------------|--------------------|--------------|
| <u>AB</u> | | <u>5MG</u> | <u>A070325 003</u> | Sep 04, 1985 |
| <u>AB</u> | | <u>10MG</u> | <u>A070325 001</u> | Sep 04, 1985 |
| <u>AB</u> | VINTAGE PHARMS | <u>2MG</u> | <u>A077749 001</u> | Mar 31, 2006 |
| <u>AB</u> | | <u>5MG</u> | <u>A077749 002</u> | Mar 31, 2006 |
| <u>AB</u> | | <u>10MG</u> | <u>A077749 003</u> | Mar 31, 2006 |

VALIUM

| | | | | |
|-----------|---------|-------------|--------------------|--|
| <u>AB</u> | + ROCHE | <u>2MG</u> | <u>N013263 002</u> | |
| <u>AB</u> | + | <u>5MG</u> | <u>N013263 004</u> | |
| <u>AB</u> | + | <u>10MG</u> | <u>N013263 006</u> | |

DIAZOXIDE

SUSPENSION; ORAL

PROGLYCEM

| | | | | |
|---|--------------------|---------|-------------|--|
| + | TEVA BRANDED PHARM | 50MG/ML | N017453 001 | |
|---|--------------------|---------|-------------|--|

DICHLORPHENAMIDE

TABLET; ORAL

KEVEYIS

| | | | | |
|---|-----------------|------|-------------|--------------|
| + | STRONGBRIDGE US | 50MG | N011366 002 | Aug 07, 2015 |
|---|-----------------|------|-------------|--------------|

DICLOFENAC

CAPSULE; ORAL

ZORVOLEX

| | | | | |
|---|------------------|------|-------------|--------------|
| + | IROKO PHARMS LLC | 18MG | N204592 001 | Oct 18, 2013 |
| + | | 35MG | N204592 002 | Oct 18, 2013 |

DICLOFENAC EPOLAMINE

PATCH; TOPICAL

FLECTOR

| | | | | |
|---|--------------|------|-------------|--------------|
| + | INST BIOCHEM | 1.3% | N021234 001 | Jan 31, 2007 |
|---|--------------|------|-------------|--------------|

SYSTEM; TOPICAL

LICART

| | | | | |
|---|---------------|------|-------------|--------------|
| + | IBSA INST BIO | 1.3% | N206976 001 | Dec 19, 2018 |
|---|---------------|------|-------------|--------------|

DICLOFENAC POTASSIUM

CAPSULE; ORAL

DICLOFENAC POTASSIUM

| | | | | |
|-----------|----------------|-------------|--------------------|--------------|
| <u>AB</u> | BIONPHARMA INC | <u>25MG</u> | <u>A204648 001</u> | Feb 23, 2016 |
|-----------|----------------|-------------|--------------------|--------------|

ZIPSOR

| | | | | | |
|-----------|---|----------|-------------|--------------------|--------------|
| <u>AB</u> | + | ASSERTIO | <u>25MG</u> | <u>N022202 001</u> | Jun 16, 2009 |
|-----------|---|----------|-------------|--------------------|--------------|

FOR SOLUTION; ORAL

CAMBIA

| | | | | | |
|-----------|---|----------|-------------|--------------------|--------------|
| <u>AB</u> | + | ASSERTIO | <u>50MG</u> | <u>N022165 001</u> | Jun 17, 2009 |
|-----------|---|----------|-------------|--------------------|--------------|

DICLOFENAC POTASSIUM

| | | | | |
|-----------|----------|-------------|--------------------|--------------|
| <u>AB</u> | PAR FORM | <u>50MG</u> | <u>A202964 001</u> | May 02, 2016 |
|-----------|----------|-------------|--------------------|--------------|

TABLET; ORAL

DICLOFENAC POTASSIUM

| | | | | |
|-----------|--------|-------------|--------------------|--------------|
| <u>AB</u> | APOTEX | <u>50MG</u> | <u>A076561 001</u> | Mar 18, 2004 |
|-----------|--------|-------------|--------------------|--------------|

| | | | | |
|-----------|-----------------|-------------|--------------------|--------------|
| <u>AB</u> | CASI PHARMS INC | <u>50MG</u> | <u>A075229 001</u> | Nov 20, 1998 |
|-----------|-----------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|---|-------|-------------|--------------------|--------------|
| <u>AB</u> | ! | MYLAN | <u>50MG</u> | <u>A075463 001</u> | Jul 26, 1999 |
|-----------|---|-------|-------------|--------------------|--------------|

| | | | | |
|-----------|------|-------------|--------------------|--------------|
| <u>AB</u> | TEVA | <u>50MG</u> | <u>A075219 001</u> | Aug 06, 1998 |
|-----------|------|-------------|--------------------|--------------|

DICLOFENAC SODIUM

GEL; TOPICAL

DICLOFENAC SODIUM

| | | | | |
|-----------|----------------------|-----------|--------------------|--------------|
| <u>AB</u> | ACTAVIS MID ATLANTIC | <u>3%</u> | <u>A206493 001</u> | Dec 02, 2015 |
|-----------|----------------------|-----------|--------------------|--------------|

| | | | | |
|-----------|---------------|-----------|--------------------|--------------|
| <u>AB</u> | AMNEAL PHARMS | <u>1%</u> | <u>A208077 001</u> | Mar 18, 2016 |
|-----------|---------------|-----------|--------------------|--------------|

| | | | | |
|-----------|-------|-----------|--------------------|--------------|
| <u>AB</u> | CIPLA | <u>1%</u> | <u>A209903 001</u> | Aug 03, 2018 |
|-----------|-------|-----------|--------------------|--------------|

| | | | | |
|-----------|---------------------|-----------|--------------------|--------------|
| <u>AB</u> | GLENMARK PHARMS LTD | <u>3%</u> | <u>A208301 001</u> | Sep 13, 2016 |
|-----------|---------------------|-----------|--------------------|--------------|

| | | | | |
|-----------|-------------------|-----------|--------------------|--------------|
| <u>AB</u> | HI-TECH PHARMACAL | <u>1%</u> | <u>A209484 001</u> | Nov 21, 2018 |
|-----------|-------------------|-----------|--------------------|--------------|

| | | | | |
|-----------|------------------|-----------|--------------------|--------------|
| <u>AB</u> | PERRIGO UK FINCO | <u>3%</u> | <u>A210893 001</u> | Jul 27, 2018 |
|-----------|------------------|-----------|--------------------|--------------|

| | | | | |
|-----------|------|-----------|--------------------|--------------|
| <u>AB</u> | TARO | <u>3%</u> | <u>A206298 001</u> | Apr 28, 2016 |
|-----------|------|-----------|--------------------|--------------|

| | | | | |
|-----------|--------|-----------|--------------------|--------------|
| <u>AB</u> | TOLMAR | <u>3%</u> | <u>A200936 001</u> | Oct 28, 2013 |
|-----------|--------|-----------|--------------------|--------------|

SOLARAZE

| | | | | | |
|-----------|---|----------------|-----------|--------------------|--------------|
| <u>AB</u> | + | FOUGERA PHARMS | <u>3%</u> | <u>N021005 001</u> | Oct 16, 2000 |
|-----------|---|----------------|-----------|--------------------|--------------|

VOLTAREN

| | | | | | |
|-----------|---|----------------------|-----------|--------------------|--------------|
| <u>AB</u> | + | GLAXOSMITHKLINE CONS | <u>1%</u> | <u>N022122 001</u> | Oct 17, 2007 |
|-----------|---|----------------------|-----------|--------------------|--------------|

SOLUTION; TOPICAL

DICLOFENAC SODIUM

| | | | | |
|-----------|---------------|-------------|--------------------|--------------|
| <u>AT</u> | AMNEAL PHARMS | <u>1.5%</u> | <u>A206116 001</u> | Sep 02, 2016 |
|-----------|---------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|---|------------|-------------|--------------------|--------------|
| <u>AT</u> | ! | APOTEX INC | <u>1.5%</u> | <u>A202027 001</u> | May 27, 2014 |
|-----------|---|------------|-------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

DICLOFENAC SODIUM

SOLUTION;TOPICAL

DICLOFENAC SODIUM

| | | | | | |
|-----------|-------------------------|-------------|----------------|------------|--------------|
| <u>AT</u> | LUPIN LTD | <u>1.5%</u> | <u>A204132</u> | <u>001</u> | Aug 20, 2015 |
| <u>AT</u> | NOVEL LABS INC | <u>1.5%</u> | <u>A205878</u> | <u>001</u> | Dec 09, 2015 |
| <u>AT</u> | RICONPHARMA LLC | <u>1.5%</u> | <u>A206715</u> | <u>001</u> | Aug 07, 2017 |
| <u>AT</u> | TARO | <u>1.5%</u> | <u>A203818</u> | <u>001</u> | Nov 26, 2014 |
| <u>AT</u> | TELIGENT PHARMA INC | <u>1.5%</u> | <u>A202769</u> | <u>001</u> | Jul 08, 2015 |
| <u>AT</u> | TWI PHARMS | <u>1.5%</u> | <u>A202393</u> | <u>001</u> | Nov 24, 2014 |
| <u>AT</u> | WATSON LABS INC | <u>1.5%</u> | <u>A202852</u> | <u>001</u> | Nov 24, 2014 |
| <u>AT</u> | ZYDUS PHARMS USA INC | <u>1.5%</u> | <u>A206411</u> | <u>001</u> | Apr 17, 2018 |

PENNSAID

+! HZNP

2%

N204623 001 Jan 16, 2014

SOLUTION/DROPS;OPHTHALMIC

DICLOFENAC SODIUM

| | | | | | |
|-----------|--------------------|-------------|----------------|------------|--------------|
| <u>AT</u> | AKORN | <u>0.1%</u> | <u>A077845</u> | <u>001</u> | Apr 17, 2008 |
| <u>AT</u> | ALTAIRE PHARMS INC | <u>0.1%</u> | <u>A203383</u> | <u>001</u> | Nov 16, 2015 |
| <u>AT</u> | BAUSCH AND LOMB | <u>0.1%</u> | <u>A078792</u> | <u>001</u> | Dec 28, 2007 |
| <u>AT</u> | RISING PHARMS | <u>0.1%</u> | <u>A078553</u> | <u>001</u> | Dec 28, 2007 |
| <u>AT</u> | SANDOZ INC | <u>0.1%</u> | <u>A078031</u> | <u>001</u> | Feb 06, 2008 |

VOLTAREN

| | | | | | |
|-----------|-------------|-------------|----------------|------------|--------------|
| <u>AT</u> | +! NOVARTIS | <u>0.1%</u> | <u>N020037</u> | <u>001</u> | Mar 28, 1991 |
|-----------|-------------|-------------|----------------|------------|--------------|

TABLET, DELAYED RELEASE;ORAL

DICLOFENAC SODIUM

| | | | | | |
|-----------|-------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | ACTAVIS ELIZABETH | <u>50MG</u> | <u>A074514</u> | <u>001</u> | Mar 26, 1996 |
| <u>AB</u> | | <u>75MG</u> | <u>A074514</u> | <u>002</u> | Mar 26, 1996 |
| <u>AB</u> | CARLSBAD | <u>25MG</u> | <u>A075185</u> | <u>002</u> | Nov 13, 1998 |
| <u>AB</u> | | <u>50MG</u> | <u>A075185</u> | <u>003</u> | Nov 13, 1998 |
| <u>AB</u> | | <u>75MG</u> | <u>A075185</u> | <u>001</u> | Nov 13, 1998 |
| <u>AB</u> | ! CASI PHARMS INC | <u>25MG</u> | <u>A074376</u> | <u>001</u> | Sep 28, 1995 |
| <u>AB</u> | ! | <u>50MG</u> | <u>A074376</u> | <u>002</u> | Sep 28, 1995 |
| <u>AB</u> | ! | <u>75MG</u> | <u>A074394</u> | <u>001</u> | Nov 30, 1995 |
| <u>AB</u> | MYLAN PHARMS INC | <u>50MG</u> | <u>A075281</u> | <u>002</u> | Feb 12, 2002 |
| <u>AB</u> | | <u>75MG</u> | <u>A075281</u> | <u>003</u> | Feb 12, 2002 |
| <u>AB</u> | UNIQUE PHARM LABS | <u>25MG</u> | <u>A090066</u> | <u>001</u> | Dec 01, 2010 |
| <u>AB</u> | | <u>50MG</u> | <u>A090066</u> | <u>002</u> | Dec 01, 2010 |
| <u>AB</u> | | <u>75MG</u> | <u>A077863</u> | <u>003</u> | Jun 08, 2007 |

TABLET, EXTENDED RELEASE;ORAL

DICLOFENAC SODIUM

| | | | | | |
|-----------|--------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ! DEXCEL LTD | <u>100MG</u> | <u>A076201</u> | <u>001</u> | Nov 06, 2002 |
| <u>AB</u> | MYLAN | <u>100MG</u> | <u>A076152</u> | <u>001</u> | Dec 13, 2001 |
| <u>AB</u> | VPNA | <u>100MG</u> | <u>A075492</u> | <u>001</u> | Feb 11, 2000 |

DICLOFENAC SODIUM; MISOPROSTOL

TABLET, DELAYED RELEASE;ORAL

ARTHROTEC

| | | | | | |
|-----------|-----------------|-------------------|----------------|------------|--------------|
| <u>AB</u> | + GD SEARLE LLC | <u>50MG;0.2MG</u> | <u>N020607</u> | <u>001</u> | Dec 24, 1997 |
| <u>AB</u> | +! | <u>75MG;0.2MG</u> | <u>N020607</u> | <u>002</u> | Dec 24, 1997 |

DICLOFENAC SODIUM AND MISOPROSTOL

| | | | | | |
|-----------|---------------------|-------------------|----------------|------------|--------------|
| <u>AB</u> | ACTAVIS LABS FL INC | <u>50MG;0.2MG</u> | <u>A201089</u> | <u>001</u> | Jul 09, 2012 |
| <u>AB</u> | | <u>75MG;0.2MG</u> | <u>A201089</u> | <u>002</u> | Jul 09, 2012 |
| <u>AB</u> | AMNEAL PHARMS | <u>50MG;0.2MG</u> | <u>A203995</u> | <u>001</u> | Nov 25, 2016 |
| <u>AB</u> | | <u>75MG;0.2MG</u> | <u>A203995</u> | <u>002</u> | Nov 25, 2016 |
| <u>AB</u> | EXELA HOLDINGS | <u>50MG;0.2MG</u> | <u>A200540</u> | <u>001</u> | Mar 14, 2014 |
| <u>AB</u> | | <u>75MG;0.2MG</u> | <u>A200540</u> | <u>002</u> | Mar 14, 2014 |
| <u>AB</u> | SANDOZ | <u>50MG;0.2MG</u> | <u>A200158</u> | <u>001</u> | May 09, 2013 |
| <u>AB</u> | | <u>75MG;0.2MG</u> | <u>A200158</u> | <u>002</u> | May 09, 2013 |

DICLOXACILLIN SODIUM

CAPSULE;ORAL

DICLOXACILLIN SODIUM

| | | | | | |
|-----------|--------|----------------------|----------------|------------|--------------|
| <u>AB</u> | SANDOZ | <u>EQ 250MG BASE</u> | <u>A061454</u> | <u>001</u> | |
| <u>AB</u> | ! | <u>EQ 500MG BASE</u> | <u>A061454</u> | <u>003</u> | |
| <u>AB</u> | TEVA | <u>EQ 250MG BASE</u> | <u>A062286</u> | <u>001</u> | Jun 03, 1982 |
| <u>AB</u> | | <u>EQ 500MG BASE</u> | <u>A062286</u> | <u>002</u> | Jun 03, 1982 |
| | SANDOZ | <u>EQ 125MG BASE</u> | <u>A061454</u> | <u>002</u> | |

PRESCRIPTION DRUG PRODUCT LIST

DICYCLOMINE HYDROCHLORIDE

CAPSULE; ORAL

BENTYL

| | | | | | | |
|-----------|------------|--------------------|-------------|----------------|------------|--------------|
| AB | + ! | ALLERGAN SALES LLC | 10MG | N007409 | 003 | Oct 15, 1984 |
|-----------|------------|--------------------|-------------|----------------|------------|--------------|

DICYCLOMINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|-------------|-------------|----------------|------------|--------------|
| AB | | LANNETT | 10MG | A084285 | 001 | |
| AB | | MYLAN | 10MG | A040319 | 001 | Sep 07, 1999 |
| AB | | WATSON LABS | 10MG | A085082 | 001 | Jun 19, 1986 |
| AB | | WEST WARD | 10MG | A040204 | 001 | Feb 28, 1997 |

INJECTABLE; INJECTION

BENTYL

| | | | | | | |
|-----------|------------|--------------------|----------------|----------------|------------|--------------|
| AP | + ! | ALLERGAN SALES LLC | 10MG/ML | N008370 | 001 | Oct 15, 1984 |
|-----------|------------|--------------------|----------------|----------------|------------|--------------|

BENTYL PRESERVATIVE FREE

| | | | | | | |
|-----------|------------|--------------------|----------------|----------------|------------|--------------|
| AP | + ! | ALLERGAN SALES LLC | 10MG/ML | N008370 | 002 | Oct 15, 1984 |
|-----------|------------|--------------------|----------------|----------------|------------|--------------|

DICYCLOMINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|---------------------|----------------|----------------|------------|--------------|
| AP | | AKORN INC | 10MG/ML | A207084 | 001 | May 04, 2018 |
| AP | | APC PHARMS LLC | 10MG/ML | A210979 | 001 | Jul 02, 2018 |
| AP | | LUITPOLD | 10MG/ML | A208353 | 001 | Feb 17, 2017 |
| AP | | RENAISSANCE SSA LLC | 10MG/ML | A207076 | 001 | Nov 02, 2018 |

DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE)

| | | | | | | |
|-----------|--|-------------------------|----------------|----------------|------------|--------------|
| AP | | WEST-WARD PHARMS INT | 10MG/ML | A040465 | 001 | Jun 30, 2003 |
|-----------|--|-------------------------|----------------|----------------|------------|--------------|

SYRUP; ORAL

DICYCLOMINE HYDROCHLORIDE

| | | | | | |
|---|----------|-----------------|----------------|------------|--------------|
| ! | GENERICS | 10MG/5ML | A040169 | 001 | Mar 24, 2005 |
|---|----------|-----------------|----------------|------------|--------------|

TABLET; ORAL

BENTYL

| | | | | | | |
|-----------|------------|--------------------|-------------|----------------|------------|--------------|
| AB | + ! | ALLERGAN SALES LLC | 20MG | N007409 | 001 | Oct 15, 1984 |
|-----------|------------|--------------------|-------------|----------------|------------|--------------|

DICYCLOMINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|--------------|-------------|----------------|------------|--------------|
| AB | | HIKMA PHARMS | 20MG | A040161 | 001 | Oct 01, 1996 |
| AB | | LANNETT | 20MG | A040230 | 001 | Feb 26, 1999 |
| AB | | MYLAN | 20MG | A040317 | 001 | Sep 07, 1999 |
| AB | | WATSON LABS | 20MG | A085223 | 001 | Jul 30, 1986 |

DIDANOSINE

CAPSULE, DELAYED REL PELLETS; ORAL

DIDANOSINE

| | | | | | | |
|-----------|--|------------------|--------------|----------------|------------|--------------|
| AB | | AUROBINDO PHARMA | 125MG | A090094 | 001 | Sep 24, 2008 |
| AB | | | 200MG | A090094 | 002 | Sep 24, 2008 |
| AB | | | 250MG | A090094 | 003 | Sep 24, 2008 |
| AB | | | 400MG | A090094 | 004 | Sep 24, 2008 |

VIDEX EC

| | | | | | | |
|-----------|------------|-------------------------|--------------|----------------|------------|--------------|
| AB | + | BRISTOL MYERS SQUIBB | 125MG | N021183 | 001 | Oct 31, 2000 |
| AB | + | | 200MG | N021183 | 002 | Oct 31, 2000 |
| AB | + | | 250MG | N021183 | 003 | Oct 31, 2000 |
| AB | + ! | | 400MG | N021183 | 004 | Oct 31, 2000 |

FOR SOLUTION; ORAL

VIDEX

| | | | | | | |
|---|---|-------------------------|----------------|----------------|------------|--------------|
| + | ! | BRISTOL-MYERS SQUIBB | 10MG/ML | N020156 | 001 | Oct 09, 1991 |
|---|---|-------------------------|----------------|----------------|------------|--------------|

DIENOGEST; ESTRADIOL VALERATE

TABLET; ORAL

NATAZIA

| | | | | | | |
|---|---|----------------|---------------------------------------------------------|----------------|------------|--------------|
| + | ! | BAYER HLTHCARE | N/A, 2MG, 3MG, N/A, N/A; 3MG, 2MG, 2MG, 1MG, N/A | N022252 | 001 | May 06, 2010 |
|---|---|----------------|---------------------------------------------------------|----------------|------------|--------------|

DIETHYLPROPION HYDROCHLORIDE

TABLET; ORAL

DIETHYLPROPION HYDROCHLORIDE

| | | | | | | |
|-----------|--|----------------|-------------|----------------|------------|--------------|
| AA | | AVANTHI INC | 25MG | A201212 | 001 | Dec 22, 2010 |
| AA | | LANNETT CO INC | 25MG | A200177 | 001 | Jul 18, 2011 |

TENUATE

| | | | | | | |
|-----------|------------|--------------------|-------------|----------------|------------|--|
| AA | + ! | TEVA BRANDED PHARM | 25MG | N011722 | 002 | |
|-----------|------------|--------------------|-------------|----------------|------------|--|

TABLET, EXTENDED RELEASE; ORAL

DIETHYLPROPION HYDROCHLORIDE

| | | | | | | |
|-----------|--|----------------|-------------|----------------|------------|--------------|
| AB | | LANNETT CO INC | 75MG | A091680 | 001 | Oct 24, 2011 |
|-----------|--|----------------|-------------|----------------|------------|--------------|

TENUATE DOSPAN

| | | | | | | |
|-----------|------------|--------------------|-------------|----------------|------------|--|
| AB | + ! | TEVA BRANDED PHARM | 75MG | N012546 | 001 | |
|-----------|------------|--------------------|-------------|----------------|------------|--|

PRESCRIPTION DRUG PRODUCT LIST

DIFLORASONE DIACETATE

CREAM; TOPICAL

DIFLORASONE DIACETATE

| | | | | | | |
|----|---|----------------|-------|---------|-----|--------------|
| BX | ! | FOUGERA PHARMS | 0.05% | A076263 | 001 | Dec 20, 2002 |
| BX | ! | TARO | 0.05% | A075508 | 001 | Apr 24, 2000 |

OINTMENT; TOPICAL

DIFLORASONE DIACETATE

| | | | | | | |
|-----------|---|---------------------|--------------|-----------------------|-------------------|--------------|
| AB | | AKORN | <u>0.05%</u> | <u>A206572</u> | <u>001</u> | Jul 24, 2015 |
| AB | | FOUGERA PHARMS | <u>0.05%</u> | <u>A075374</u> | <u>001</u> | Apr 27, 1999 |
| AB | | RISING PHARMS | <u>0.05%</u> | <u>A207440</u> | <u>001</u> | Feb 27, 2017 |
| AB | ! | TARO | <u>0.05%</u> | <u>A075331</u> | <u>001</u> | May 14, 1999 |
| AB | | TELIGENT PHARMA INC | <u>0.05%</u> | <u>A210753</u> | <u>001</u> | Jun 12, 2018 |

DIFLUNISAL

TABLET; ORAL

DIFLUNISAL

| | | | | | | |
|-----------|---|-------------------------|--------------|-----------------------|-------------------|--------------|
| AB | | HERITAGE PHARMA | <u>500MG</u> | <u>A202845</u> | <u>001</u> | Mar 08, 2012 |
| AB | ! | TEVA | <u>500MG</u> | <u>A073673</u> | <u>001</u> | Jul 31, 1992 |
| AB | | ZYDUS PHARMS USA INC | <u>500MG</u> | <u>A203547</u> | <u>001</u> | Jun 16, 2017 |

DIFLUPREDNATE

EMULSION; OPHTHALMIC

DUREZOL

| | | | | | | |
|---|---|-------------------------|-------|---------|-----|--------------|
| + | ! | NOVARTIS PHARMS CORP | 0.05% | N022212 | 001 | Jun 23, 2008 |
|---|---|-------------------------|-------|---------|-----|--------------|

DIGOXIN

ELIXIR; ORAL

DIGOXIN

| | | | | | | |
|---|---|-------------------------|-----------|---------|-----|--------------|
| + | ! | WEST-WARD PHARMS INT | 0.05MG/ML | N021648 | 001 | Aug 26, 2004 |
|---|---|-------------------------|-----------|---------|-----|--------------|

INJECTABLE; INJECTION

DIGOXIN

| | | | | | | |
|-----------|--|-------------------------|------------------|-----------------------|-------------------|--------------|
| AP | | SANDOZ INC | <u>0.25MG/ML</u> | <u>A040481</u> | <u>001</u> | Aug 21, 2003 |
| AP | | WEST-WARD PHARMS INT | <u>0.25MG/ML</u> | <u>A083391</u> | <u>001</u> | |

LANOXIN

| | | | | | | |
|-----------|---|-------------------|------------------|-----------------------|-------------------|--|
| AP | + | COVIS PHARMA BV | <u>0.25MG/ML</u> | <u>N009330</u> | <u>002</u> | |
| | | LANOXIN PEDIATRIC | | | | |
| | + | COVIS PHARMA BV | 0.1MG/ML | N009330 | 004 | |

TABLET; ORAL

DIGOXIN

| | | | | | | |
|-----------|--|--------------------|----------------|-----------------------|-------------------|--------------|
| AB | | HIKMA INTL PHARMS | <u>0.125MG</u> | <u>A077002</u> | <u>002</u> | Oct 30, 2007 |
| AB | | | <u>0.25MG</u> | <u>A077002</u> | <u>001</u> | Oct 30, 2007 |
| AB | | IMPAX LABS | <u>0.125MG</u> | <u>A078556</u> | <u>001</u> | Jul 20, 2009 |
| AB | | | <u>0.25MG</u> | <u>A078556</u> | <u>002</u> | Jul 20, 2009 |
| AB | | MYLAN PHARMS INC | <u>0.125MG</u> | <u>A040282</u> | <u>001</u> | Dec 23, 1999 |
| AB | | | <u>0.25MG</u> | <u>A040282</u> | <u>002</u> | Dec 23, 1999 |
| AB | | STEVENS J | <u>0.125MG</u> | <u>A076268</u> | <u>001</u> | Jul 26, 2002 |
| AB | | | <u>0.25MG</u> | <u>A076268</u> | <u>002</u> | Jul 26, 2002 |
| AB | | SUN PHARM INDS INC | <u>0.125MG</u> | <u>A076363</u> | <u>001</u> | Jan 31, 2003 |
| AB | | | <u>0.25MG</u> | <u>A076363</u> | <u>002</u> | Jan 31, 2003 |

LANOXIN

| | | | | | | |
|-----------|---|-------------------------|----------------|-----------------------|-------------------|--------------|
| AB | + | CONCORDIA PHARMS INC | <u>0.125MG</u> | <u>N020405</u> | <u>002</u> | Sep 30, 1997 |
| AB | + | | <u>0.25MG</u> | <u>N020405</u> | <u>004</u> | Sep 30, 1997 |
| | + | | 0.0625MG | N020405 | 001 | Sep 30, 1997 |

DIHYDROERGOTAMINE MESYLATE

INJECTABLE; INJECTION

D.H.E. 45

| | | | | | | |
|-----------|---|---------|---------------|-----------------------|-------------------|--|
| AP | + | VALEANT | <u>1MG/ML</u> | <u>N005929</u> | <u>001</u> | |
|-----------|---|---------|---------------|-----------------------|-------------------|--|

DIHYDROERGOTAMINE MESYLATE

| | | | | | | |
|-----------|--|-------------------------|---------------|-----------------------|-------------------|--------------|
| AP | | HIKMA PHARMS | <u>1MG/ML</u> | <u>A206621</u> | <u>001</u> | Sep 15, 2017 |
| AP | | PADDOCK LLC | <u>1MG/ML</u> | <u>A040475</u> | <u>001</u> | Apr 28, 2003 |
| AP | | SAGENT PHARMS | <u>1MG/ML</u> | <u>A207264</u> | <u>001</u> | Jul 11, 2018 |
| AP | | WEST-WARD PHARMS INT | <u>1MG/ML</u> | <u>A040453</u> | <u>001</u> | Jun 09, 2003 |

SPRAY, METERED; NASAL

MIGRANAL

| | | | | | | |
|---|---|---------|-----------|---------|-----|--------------|
| + | ! | VALEANT | 0.5MG/INH | N020148 | 001 | Dec 08, 1997 |
|---|---|---------|-----------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

DILTIAZEM HYDROCHLORIDE

| | | | | |
|--------------------------------|-------------------------|--------------|--------------------|--------------|
| <u>AB2</u> | APOTEX | <u>120MG</u> | <u>A074943 003</u> | Dec 19, 2000 |
| <u>AB2</u> | | <u>180MG</u> | <u>A074943 002</u> | Dec 19, 2000 |
| <u>AB2</u> | | <u>240MG</u> | <u>A074943 001</u> | Aug 06, 1998 |
| <u>AB2</u> | MYLAN | <u>120MG</u> | <u>A075124 002</u> | Mar 18, 1998 |
| <u>AB2</u> | | <u>180MG</u> | <u>A075124 003</u> | Mar 18, 1998 |
| <u>AB2</u> | ! | <u>240MG</u> | <u>A075124 001</u> | Mar 18, 1998 |
| <u>CARDIZEM CD</u> | | | | |
| <u>AB3</u> | + VALEANT INTL | <u>120MG</u> | <u>N020062 001</u> | Aug 10, 1992 |
| <u>AB3</u> | + | <u>180MG</u> | <u>N020062 002</u> | Dec 27, 1991 |
| <u>AB3</u> | + | <u>240MG</u> | <u>N020062 003</u> | Dec 27, 1991 |
| <u>AB3</u> | + | <u>300MG</u> | <u>N020062 004</u> | Dec 27, 1991 |
| <u>AB3</u> | +! | <u>360MG</u> | <u>N020062 005</u> | Aug 24, 1999 |
| <u>CARTIA XT</u> | | | | |
| <u>AB3</u> | ACTAVIS LABS FL INC | <u>120MG</u> | <u>A074752 002</u> | Jul 09, 1998 |
| <u>AB3</u> | | <u>180MG</u> | <u>A074752 001</u> | Jul 09, 1998 |
| <u>AB3</u> | | <u>240MG</u> | <u>A074752 003</u> | Jul 09, 1998 |
| <u>AB3</u> | | <u>300MG</u> | <u>A074752 004</u> | Jul 09, 1998 |
| <u>DILTIAZEM HYDROCHLORIDE</u> | | | | |
| <u>AB3</u> | ACTAVIS ELIZABETH | <u>360MG</u> | <u>A202463 001</u> | Dec 07, 2012 |
| <u>AB3</u> | PAR PHARM | <u>120MG</u> | <u>A074984 001</u> | Dec 20, 1999 |
| <u>AB3</u> | | <u>180MG</u> | <u>A074984 002</u> | Dec 20, 1999 |
| <u>AB3</u> | | <u>240MG</u> | <u>A074984 003</u> | Dec 20, 1999 |
| <u>AB3</u> | | <u>300MG</u> | <u>A074984 004</u> | Dec 20, 1999 |
| <u>AB3</u> | | <u>360MG</u> | <u>A209766 001</u> | May 30, 2018 |
| <u>AB3</u> | SUN PHARM INDS LTD | <u>120MG</u> | <u>A203023 001</u> | Jun 08, 2017 |
| <u>AB3</u> | | <u>180MG</u> | <u>A203023 002</u> | Jun 08, 2017 |
| <u>AB3</u> | | <u>240MG</u> | <u>A203023 003</u> | Jun 08, 2017 |
| <u>AB3</u> | | <u>300MG</u> | <u>A203023 004</u> | Jun 08, 2017 |
| <u>AB3</u> | | <u>360MG</u> | <u>A203023 005</u> | Jun 08, 2017 |
| <u>AB3</u> | SUN PHARMA GLOBAL | <u>120MG</u> | <u>A090492 001</u> | Oct 28, 2011 |
| <u>AB3</u> | | <u>180MG</u> | <u>A090492 002</u> | Oct 28, 2011 |
| <u>AB3</u> | | <u>240MG</u> | <u>A090492 003</u> | Oct 28, 2011 |
| <u>AB3</u> | | <u>300MG</u> | <u>A090492 004</u> | Oct 28, 2011 |
| <u>AB3</u> | | <u>360MG</u> | <u>A090492 005</u> | Oct 28, 2011 |
| <u>AB3</u> | TWI PHARMS | <u>120MG</u> | <u>A205231 001</u> | Aug 30, 2018 |
| <u>AB3</u> | | <u>180MG</u> | <u>A205231 002</u> | Aug 30, 2018 |
| <u>AB3</u> | | <u>240MG</u> | <u>A205231 003</u> | Aug 30, 2018 |
| <u>AB3</u> | | <u>300MG</u> | <u>A205231 004</u> | Aug 30, 2018 |
| <u>AB3</u> | | <u>360MG</u> | <u>A205231 005</u> | Aug 30, 2018 |
| <u>AB3</u> | VALEANT PHARMS NORTH | <u>120MG</u> | <u>A075116 001</u> | Dec 23, 1999 |
| <u>AB3</u> | | <u>180MG</u> | <u>A075116 002</u> | Dec 23, 1999 |
| <u>AB3</u> | | <u>240MG</u> | <u>A075116 003</u> | Dec 23, 1999 |
| <u>AB3</u> | | <u>300MG</u> | <u>A075116 004</u> | Dec 23, 1999 |
| <u>AB3</u> | ZYDUS PHARMS USA INC | <u>120MG</u> | <u>A206534 001</u> | Aug 08, 2017 |
| <u>AB3</u> | | <u>180MG</u> | <u>A206534 002</u> | Aug 08, 2017 |
| <u>AB3</u> | | <u>240MG</u> | <u>A206534 003</u> | Aug 08, 2017 |
| <u>AB3</u> | | <u>300MG</u> | <u>A206534 004</u> | Aug 08, 2017 |
| <u>AB3</u> | | <u>360MG</u> | <u>A206534 005</u> | Aug 08, 2017 |
| <u>AB4</u> | SANDOZ | <u>120MG</u> | <u>A091022 001</u> | Sep 28, 2012 |
| <u>AB4</u> | | <u>180MG</u> | <u>A091022 002</u> | Sep 28, 2012 |
| <u>AB4</u> | | <u>240MG</u> | <u>A091022 003</u> | Sep 28, 2012 |
| <u>AB4</u> | | <u>300MG</u> | <u>A091022 004</u> | Sep 28, 2012 |
| <u>AB4</u> | | <u>360MG</u> | <u>A091022 005</u> | Sep 28, 2012 |
| <u>AB4</u> | | <u>420MG</u> | <u>A091022 006</u> | Sep 28, 2012 |
| <u>AB4</u> | SUN PHARMA GLOBAL | <u>120MG</u> | <u>A090421 001</u> | Nov 15, 2010 |
| <u>AB4</u> | | <u>180MG</u> | <u>A090421 002</u> | Nov 15, 2010 |
| <u>AB4</u> | | <u>240MG</u> | <u>A090421 003</u> | Nov 15, 2010 |
| <u>AB4</u> | | <u>300MG</u> | <u>A090421 004</u> | Nov 15, 2010 |
| <u>AB4</u> | | <u>360MG</u> | <u>A090421 005</u> | Nov 15, 2010 |
| <u>AB4</u> | ZYDUS PHARMS USA INC | <u>120MG</u> | <u>A206641 001</u> | Aug 11, 2017 |
| <u>AB4</u> | | <u>180MG</u> | <u>A206641 002</u> | Aug 11, 2017 |
| <u>AB4</u> | | <u>240MG</u> | <u>A206641 003</u> | Aug 11, 2017 |
| <u>AB4</u> | | <u>300MG</u> | <u>A206641 004</u> | Aug 11, 2017 |
| <u>AB4</u> | | <u>360MG</u> | <u>A206641 005</u> | Aug 11, 2017 |
| <u>AB4</u> | | <u>420MG</u> | <u>A206641 006</u> | Aug 11, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

DILTZAC

| | | | | |
|------------|------------|--------------|--------------------|--------------|
| <u>AB4</u> | APOTEX INC | <u>120MG</u> | <u>A076395 001</u> | Feb 01, 2006 |
| <u>AB4</u> | | <u>180MG</u> | <u>A076395 002</u> | Feb 01, 2006 |
| <u>AB4</u> | | <u>240MG</u> | <u>A076395 003</u> | Feb 01, 2006 |
| <u>AB4</u> | | <u>300MG</u> | <u>A076395 004</u> | Feb 01, 2006 |
| <u>AB4</u> | | <u>360MG</u> | <u>A076395 005</u> | Feb 01, 2006 |

TAZTIA XT

| | | | | |
|------------|---------------------|--------------|--------------------|--------------|
| <u>AB4</u> | ACTAVIS LABS FL INC | <u>120MG</u> | <u>A075401 001</u> | Apr 10, 2003 |
| <u>AB4</u> | | <u>180MG</u> | <u>A075401 002</u> | Apr 10, 2003 |
| <u>AB4</u> | | <u>240MG</u> | <u>A075401 003</u> | Apr 10, 2003 |
| <u>AB4</u> | | <u>300MG</u> | <u>A075401 004</u> | Apr 10, 2003 |
| <u>AB4</u> | | <u>360MG</u> | <u>A075401 005</u> | Apr 10, 2003 |

TIAZAC

| | | | | |
|------------|---------------------------|--------------|--------------------|--------------|
| <u>AB4</u> | + VALEANT PHARMS NORTH | <u>120MG</u> | <u>N020401 001</u> | Sep 11, 1995 |
| <u>AB4</u> | + | <u>180MG</u> | <u>N020401 002</u> | Sep 11, 1995 |
| <u>AB4</u> | + | <u>240MG</u> | <u>N020401 003</u> | Sep 11, 1995 |
| <u>AB4</u> | + | <u>300MG</u> | <u>N020401 004</u> | Sep 11, 1995 |
| <u>AB4</u> | + | <u>360MG</u> | <u>N020401 005</u> | Sep 11, 1995 |
| <u>AB4</u> | + | <u>420MG</u> | <u>N020401 006</u> | Oct 16, 1998 |

DILTIAZEM HYDROCHLORIDE

| | | | | |
|----|---------|-------|-------------|--------------|
| BC | ! MYLAN | 120MG | A074910 003 | May 02, 1997 |
| | | 60MG | A074910 001 | May 02, 1997 |
| | | 90MG | A074910 002 | May 02, 1997 |

INJECTABLE; INJECTION

DILTIAZEM HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|---------------|--------------------|--------------|
| <u>AP</u> | AKORN INC | <u>5MG/ML</u> | <u>A075086 001</u> | Apr 09, 1998 |
| <u>AP</u> | ! ATHENEX INC | <u>5MG/ML</u> | <u>A074617 001</u> | Feb 28, 1996 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>5MG/ML</u> | <u>A202651 001</u> | Aug 09, 2012 |
| <u>AP</u> | HOSPIRA | <u>5MG/ML</u> | <u>A074941 001</u> | Apr 15, 1998 |
| <u>AP</u> | INTL MEDICATION | <u>5MG/ML</u> | <u>A075749 001</u> | Nov 21, 2001 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>5MG/ML</u> | <u>A078538 001</u> | Dec 17, 2008 |
| | ! HOSPIRA | 100MG/VIAL | A075853 001 | Dec 17, 2002 |

TABLET; ORAL

CARDIZEM

| | | | | |
|-----------|----------------|--------------|--------------------|--------------|
| <u>AB</u> | + VALEANT INTL | <u>30MG</u> | <u>N018602 001</u> | Nov 05, 1982 |
| <u>AB</u> | + | <u>60MG</u> | <u>N018602 002</u> | Nov 05, 1982 |
| <u>AB</u> | + | <u>90MG</u> | <u>N018602 003</u> | Dec 08, 1986 |
| <u>AB</u> | + | <u>120MG</u> | <u>N018602 004</u> | Dec 08, 1986 |

DILTIAZEM HYDROCHLORIDE

| | | | | |
|-----------|-------|--------------|--------------------|--------------|
| <u>AB</u> | MYLAN | <u>30MG</u> | <u>A072838 004</u> | Nov 05, 1992 |
| <u>AB</u> | | <u>60MG</u> | <u>A072838 003</u> | Nov 05, 1992 |
| <u>AB</u> | | <u>90MG</u> | <u>A072838 002</u> | Nov 05, 1992 |
| <u>AB</u> | | <u>120MG</u> | <u>A072838 001</u> | Nov 05, 1992 |
| <u>AB</u> | TEVA | <u>30MG</u> | <u>A074185 001</u> | May 31, 1995 |
| <u>AB</u> | | <u>60MG</u> | <u>A074185 002</u> | May 31, 1995 |
| <u>AB</u> | | <u>90MG</u> | <u>A074185 003</u> | May 31, 1995 |
| <u>AB</u> | | <u>120MG</u> | <u>A074185 004</u> | May 31, 1995 |

TABLET, EXTENDED RELEASE;ORAL

CARDIZEM LA

| | | | | |
|-----------|----------------|--------------|--------------------|--------------|
| <u>AB</u> | + VALEANT INTL | <u>120MG</u> | <u>N021392 001</u> | Feb 06, 2003 |
| <u>AB</u> | + | <u>180MG</u> | <u>N021392 002</u> | Feb 06, 2003 |
| <u>AB</u> | + | <u>240MG</u> | <u>N021392 003</u> | Feb 06, 2003 |
| <u>AB</u> | + | <u>300MG</u> | <u>N021392 004</u> | Feb 06, 2003 |
| <u>AB</u> | + | <u>360MG</u> | <u>N021392 005</u> | Feb 06, 2003 |
| <u>AB</u> | + | <u>420MG</u> | <u>N021392 006</u> | Feb 06, 2003 |

DILTIAZEM HYDROCHLORIDE

| | | | | |
|-----------|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS LABS FL INC | <u>120MG</u> | <u>A077686 006</u> | Mar 15, 2010 |
| <u>AB</u> | | <u>180MG</u> | <u>A077686 005</u> | Mar 15, 2010 |
| <u>AB</u> | | <u>240MG</u> | <u>A077686 004</u> | Mar 15, 2010 |
| <u>AB</u> | | <u>300MG</u> | <u>A077686 003</u> | Mar 15, 2010 |
| <u>AB</u> | | <u>360MG</u> | <u>A077686 002</u> | Mar 15, 2010 |
| <u>AB</u> | | <u>420MG</u> | <u>A077686 001</u> | Mar 15, 2010 |

PRESCRIPTION DRUG PRODUCT LIST

DIMENHYDRINATE

INJECTABLE; INJECTION

DIMENHYDRINATE

! FRESenius KABI USA 50MG/ML

A040519 001 Jun 23, 2004

DIMERCAPROL

INJECTABLE; INJECTION

BAL

+! AKORN 10%

N005939 001

DIMETHYL FUMARATE

CAPSULE, DELAYED RELEASE; ORAL

TECFIDERA

+ BIOGEN IDEC INC 120MG

N204063 001 Mar 27, 2013

+! 240MG

N204063 002 Mar 27, 2013

DIMETHYL SULFOXIDE

SOLUTION; INTRAVESICAL

DIMETHYL SULFOXIDE**AT** MYLAN INSTITUTIONAL **50%****A076185 001** Nov 29, 2002**RIMSO-50****AT** +! MYLAN INSTITUTIONAL **50%****N017788 001**DINOPROSTONE

GEL; ENDOCERVICAL

PREPIDIL

+! PHARMACIA AND 0.5MG/3GM

N019617 001 Dec 09, 1992

UPJOHN

INSERT, EXTENDED RELEASE; VAGINAL

CERVIDIL

+! FERRING PHARMS INC 10MG

N020411 001 Mar 30, 1995

SUPPOSITORY; VAGINAL

PROSTIN E2

+! PHARMACIA AND 20MG

N017810 001

UPJOHN

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

! BARR 50MG

A080738 001

ELIXIR; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

! PHARM ASSOC 12.5MG/5ML

A087513 001 Feb 10, 1982

INJECTABLE; INJECTION

DIPHENHYDRAMINE HYDROCHLORIDE**AP** APP PHARMS **50MG/ML****A040466 001** May 28, 2002**AP** HOSPIRA **50MG/ML****A040140 001** Nov 20, 1998**AP** MICRO LABS **50MG/ML****A205723 001** Aug 22, 2018**AP** MYLAN INSTITUTIONAL **50MG/ML****A040498 001** Jul 12, 2005**AP** ! WEST-WARD PHARMS **50MG/ML****A080817 002**

INT

DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE**AP** FRESenius KABI USA **50MG/ML****A091526 001** Mar 26, 2013DIPYRIDAMOLE

INJECTABLE; INJECTION

DIPYRIDAMOLE**AP** ! ATHENEX INC **5MG/ML****A074939 001** Apr 13, 1998**AP** FRESenius KABI USA **5MG/ML****A074956 001** Sep 30, 1998**AP** WEST-WARD PHARMS **5MG/ML****A074521 001** Oct 18, 1996

INT

TABLET; ORAL

DIPYRIDAMOLE**AB** BARR **25MG****A087184 001** Oct 03, 1990**AB** **50MG****A087716 001** Oct 03, 1990**AB** **75MG****A087717 001** Oct 03, 1990**AB** IMPAX LABS **25MG****A040782 001** Jul 18, 2007**AB** **50MG****A040782 002** Jul 18, 2007**AB** **75MG****A040782 003** Jul 18, 2007**AB** MURTY PHARMS **25MG****A040733 001** Feb 13, 2007**AB** **50MG****A040733 002** Feb 13, 2007**AB** **75MG****A040733 003** Feb 13, 2007**AB** ZYDUS PHARMS USA **25MG****A040874 001** Jan 28, 2008

INC

AB **50MG****A040874 002** Jan 28, 2008**AB** **75MG****A040874 003** Jan 28, 2008

PRESCRIPTION DRUG PRODUCT LIST

DIPYRIDAMOLE

TABLET; ORAL

PERSANTINE

| | | | | | |
|-----------|---|-------------------------|-------------|--------------------|--------------|
| <u>AB</u> | + | BOEHRINGER INGELHEIM | <u>25MG</u> | <u>N012836 003</u> | Dec 22, 1986 |
| <u>AB</u> | + | | <u>50MG</u> | <u>N012836 004</u> | Feb 06, 1987 |
| <u>AB</u> | + | ! | <u>75MG</u> | <u>N012836 005</u> | Feb 06, 1987 |

DISOPYRAMIDE PHOSPHATE

CAPSULE; ORAL

DISOPYRAMIDE PHOSPHATE

| | | | | | |
|-----------|--|--------------|----------------------|--------------------|--------------|
| <u>AB</u> | | MAYNE PHARMA | <u>EQ 100MG BASE</u> | <u>A070173 001</u> | May 31, 1985 |
| <u>AB</u> | | | <u>EQ 150MG BASE</u> | <u>A070173 002</u> | May 31, 1985 |
| <u>AB</u> | | TEVA | <u>EQ 100MG BASE</u> | <u>A070101 001</u> | Feb 22, 1985 |
| <u>AB</u> | | | <u>EQ 150MG BASE</u> | <u>A070102 001</u> | Feb 22, 1985 |

NORPACE

| | | | | | |
|-----------|---|---------------|----------------------|--------------------|--|
| <u>AB</u> | + | GD SEARLE LLC | <u>EQ 100MG BASE</u> | <u>N017447 001</u> | |
| <u>AB</u> | + | ! | <u>EQ 150MG BASE</u> | <u>N017447 002</u> | |

CAPSULE, EXTENDED RELEASE; ORAL

NORPACE CR

| | | | | | |
|--|---|---------------|----------------------|--------------------|--------------|
| | + | GD SEARLE LLC | <u>EQ 100MG BASE</u> | <u>N018655 001</u> | Jul 20, 1982 |
| | + | ! | <u>EQ 150MG BASE</u> | <u>N018655 002</u> | Jul 20, 1982 |

DISULFIRAM

TABLET; ORAL

ANTABUSE

| | | | | | |
|-----------|--|----------------|--------------|--------------------|--------------|
| <u>AB</u> | | ODYSSEY PHARMS | <u>250MG</u> | <u>A088482 001</u> | Dec 08, 1983 |
| <u>AB</u> | | ! | <u>500MG</u> | <u>A088483 001</u> | Dec 08, 1983 |

DISULFIRAM

| | | | | | |
|-----------|--|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | | ALVOGEN MALTA | <u>250MG</u> | <u>A091681 001</u> | Aug 08, 2013 |
| <u>AB</u> | | CHARTWELL MOLECULES | <u>250MG</u> | <u>A091563 001</u> | Dec 31, 2012 |
| <u>AB</u> | | | <u>500MG</u> | <u>A091563 002</u> | Dec 31, 2012 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>250MG</u> | <u>A203916 001</u> | Mar 04, 2015 |
| <u>AB</u> | | | <u>500MG</u> | <u>A203916 002</u> | Mar 04, 2015 |
| <u>AB</u> | | SIGMAPHARM LABS LLC | <u>250MG</u> | <u>A091619 001</u> | Mar 28, 2011 |
| <u>AB</u> | | | <u>500MG</u> | <u>A091619 002</u> | Mar 28, 2011 |
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>250MG</u> | <u>A202652 001</u> | Feb 05, 2014 |
| <u>AB</u> | | | <u>500MG</u> | <u>A202652 002</u> | Feb 05, 2014 |

DIVALPROEX SODIUM

CAPSULE, DELAYED REL PELLETS; ORAL

DEPAKOTE

| | | | | | |
|-----------|---|--------|-------------------------------|--------------------|--------------|
| <u>AB</u> | + | ABEVIE | <u>EQ 125MG VALPROIC ACID</u> | <u>N019680 001</u> | Sep 12, 1989 |
|-----------|---|--------|-------------------------------|--------------------|--------------|

DIVALPROEX SODIUM

| | | | | | |
|-----------|--|-------------------------|-------------------------------|--------------------|--------------|
| <u>AB</u> | | DR REDDYS LABS LTD | <u>EQ 125MG VALPROIC ACID</u> | <u>A078979 001</u> | Jan 23, 2009 |
| <u>AB</u> | | MYLAN | <u>EQ 125MG VALPROIC ACID</u> | <u>A090407 001</u> | Mar 28, 2011 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>EQ 125MG VALPROIC ACID</u> | <u>A078919 001</u> | Jan 27, 2009 |

TABLET, DELAYED RELEASE; ORAL

DEPAKOTE

| | | | | | |
|-----------|---|--------|-------------------------------|--------------------|--------------|
| <u>AB</u> | + | ABEVIE | <u>EQ 125MG VALPROIC ACID</u> | <u>N018723 003</u> | Oct 26, 1984 |
| <u>AB</u> | + | | <u>EQ 250MG VALPROIC ACID</u> | <u>N018723 001</u> | Mar 10, 1983 |
| <u>AB</u> | + | ! | <u>EQ 500MG VALPROIC ACID</u> | <u>N018723 002</u> | Mar 10, 1983 |

DIVALPROEX SODIUM

| | | | | | |
|-----------|--|-------------------------|-------------------------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS LABS FL INC | <u>EQ 500MG VALPROIC ACID</u> | <u>A079080 001</u> | Feb 25, 2011 |
| <u>AB</u> | | ANCHEN PHARMS | <u>EQ 500MG VALPROIC ACID</u> | <u>A078411 001</u> | Nov 03, 2008 |
| <u>AB</u> | | APOTEX INC | <u>EQ 125MG VALPROIC ACID</u> | <u>A077615 003</u> | Jul 29, 2008 |
| <u>AB</u> | | | <u>EQ 250MG VALPROIC ACID</u> | <u>A077615 002</u> | Jul 29, 2008 |
| <u>AB</u> | | | <u>EQ 500MG VALPROIC ACID</u> | <u>A077615 001</u> | Jul 29, 2008 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>EQ 125MG VALPROIC ACID</u> | <u>A090554 001</u> | Apr 21, 2011 |
| <u>AB</u> | | | <u>EQ 250MG VALPROIC ACID</u> | <u>A090554 002</u> | Apr 21, 2011 |
| <u>AB</u> | | | <u>EQ 500MG VALPROIC ACID</u> | <u>A090554 003</u> | Apr 21, 2011 |
| <u>AB</u> | | DR REDDYS LABS LTD | <u>EQ 125MG VALPROIC ACID</u> | <u>A078755 001</u> | Jul 29, 2008 |
| <u>AB</u> | | | <u>EQ 250MG VALPROIC ACID</u> | <u>A078755 002</u> | Jul 29, 2008 |
| <u>AB</u> | | | <u>EQ 500MG VALPROIC ACID</u> | <u>A078755 003</u> | Jul 29, 2008 |
| <u>AB</u> | | LUPIN | <u>EQ 125MG VALPROIC ACID</u> | <u>A078790 001</u> | Jul 29, 2008 |
| <u>AB</u> | | | <u>EQ 250MG VALPROIC ACID</u> | <u>A078790 002</u> | Jul 29, 2008 |
| <u>AB</u> | | | <u>EQ 500MG VALPROIC ACID</u> | <u>A078790 003</u> | Jul 29, 2008 |
| <u>AB</u> | | MYLAN | <u>EQ 125MG VALPROIC ACID</u> | <u>A090062 001</u> | Mar 17, 2009 |
| <u>AB</u> | | | <u>EQ 250MG VALPROIC ACID</u> | <u>A090062 002</u> | Mar 17, 2009 |
| <u>AB</u> | | | <u>EQ 500MG VALPROIC ACID</u> | <u>A090062 003</u> | Mar 17, 2009 |
| <u>AB</u> | | ORCHID HLTHCARE | <u>EQ 125MG VALPROIC ACID</u> | <u>A078853 001</u> | Nov 25, 2008 |
| <u>AB</u> | | | <u>EQ 250MG VALPROIC ACID</u> | <u>A078853 002</u> | Nov 25, 2008 |

PRESCRIPTION DRUG PRODUCT LIST

DIVALPROEX SODIUM

TABLET, DELAYED RELEASE;ORAL

DIVALPROEX SODIUM

| | | | | |
|-----------|-------------------------|-------------------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A078853 003</u> | Nov 25, 2008 |
| <u>AB</u> | PRINSTON INC | <u>EQ 125MG VALPROIC ACID</u> | <u>A090210 001</u> | Nov 30, 2009 |
| <u>AB</u> | | <u>EQ 250MG VALPROIC ACID</u> | <u>A090210 002</u> | Nov 30, 2009 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A090210 003</u> | Nov 30, 2009 |
| <u>AB</u> | SANDOZ | <u>EQ 125MG VALPROIC ACID</u> | <u>A078290 003</u> | Jul 29, 2008 |
| <u>AB</u> | | <u>EQ 250MG VALPROIC ACID</u> | <u>A078290 002</u> | Jul 29, 2008 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A078290 001</u> | Jul 29, 2008 |
| <u>AB</u> | SUN PHARM INDS | <u>EQ 125MG VALPROIC ACID</u> | <u>A078597 001</u> | Jul 29, 2008 |
| <u>AB</u> | | <u>EQ 250MG VALPROIC ACID</u> | <u>A078597 002</u> | Jul 29, 2008 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A078597 003</u> | Jul 29, 2008 |
| <u>AB</u> | TEVA | <u>EQ 125MG VALPROIC ACID</u> | <u>A076941 001</u> | Jul 29, 2008 |
| <u>AB</u> | | <u>EQ 250MG VALPROIC ACID</u> | <u>A076941 002</u> | Jul 29, 2008 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A076941 003</u> | Jul 29, 2008 |
| <u>AB</u> | UNICHEM LABS LTD | <u>EQ 125MG VALPROIC ACID</u> | <u>A079163 001</u> | Apr 05, 2011 |
| <u>AB</u> | | <u>EQ 250MG VALPROIC ACID</u> | <u>A079163 002</u> | Apr 05, 2011 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A079163 003</u> | Apr 05, 2011 |
| <u>AB</u> | UPSHER SMITH LABS | <u>EQ 125MG VALPROIC ACID</u> | <u>A078182 001</u> | Jul 29, 2008 |
| <u>AB</u> | | <u>EQ 250MG VALPROIC ACID</u> | <u>A078182 002</u> | Jul 29, 2008 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A078182 003</u> | Jul 29, 2008 |
| <u>AB</u> | WOCKHARDT | <u>EQ 125MG VALPROIC ACID</u> | <u>A077296 001</u> | Jul 31, 2008 |
| <u>AB</u> | | <u>EQ 250MG VALPROIC ACID</u> | <u>A077296 002</u> | Jul 31, 2008 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A077296 003</u> | Jul 31, 2008 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 125MG VALPROIC ACID</u> | <u>A077100 001</u> | Mar 05, 2009 |
| <u>AB</u> | | <u>EQ 250MG VALPROIC ACID</u> | <u>A077100 002</u> | Mar 05, 2009 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A077100 003</u> | Mar 05, 2009 |

TABLET, EXTENDED RELEASE;ORAL

DEPAKOTE ER

| | | | | | |
|-----------|---|--------|-------------------------------|--------------------|--------------|
| <u>AB</u> | + | ABEVIE | <u>EQ 250MG VALPROIC ACID</u> | <u>N021168 002</u> | May 31, 2002 |
| <u>AB</u> | + | ! | <u>EQ 500MG VALPROIC ACID</u> | <u>N021168 001</u> | Aug 04, 2000 |

DIVALPROEX SODIUM

| | | | | |
|-----------|-------------------------|-------------------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 250MG VALPROIC ACID</u> | <u>A203730 001</u> | May 29, 2015 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A203730 002</u> | May 29, 2015 |
| <u>AB</u> | ANCHEN PHARMS | <u>EQ 250MG VALPROIC ACID</u> | <u>A078445 001</u> | Feb 26, 2009 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A078445 002</u> | Aug 04, 2009 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 250MG VALPROIC ACID</u> | <u>A202419 001</u> | Jun 02, 2014 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A202419 002</u> | Jun 02, 2014 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 250MG VALPROIC ACID</u> | <u>A090161 001</u> | Mar 15, 2012 |
| <u>AB</u> | IMPAX LABS | <u>EQ 250MG VALPROIC ACID</u> | <u>A078791 001</u> | May 06, 2009 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A078791 002</u> | Aug 04, 2009 |
| <u>AB</u> | MYLAN | <u>EQ 250MG VALPROIC ACID</u> | <u>A077567 001</u> | Jan 29, 2009 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A077567 002</u> | Jan 29, 2009 |
| <u>AB</u> | REDDYS | <u>EQ 500MG VALPROIC ACID</u> | <u>A090070 001</u> | Mar 12, 2012 |
| <u>AB</u> | WOCKHARDT | <u>EQ 250MG VALPROIC ACID</u> | <u>A078705 002</u> | Feb 10, 2009 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A078705 001</u> | Aug 04, 2009 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 250MG VALPROIC ACID</u> | <u>A078239 001</u> | Feb 27, 2009 |
| <u>AB</u> | | <u>EQ 500MG VALPROIC ACID</u> | <u>A078239 002</u> | Aug 04, 2009 |

DOBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOBUTAMINE HYDROCHLORIDE

| | | | | | |
|-----------|---|-------------------------|--------------------------|--------------------|--------------|
| <u>AP</u> | | HOSPIRA | <u>EQ 12.5MG BASE/ML</u> | <u>A074086 001</u> | Nov 29, 1993 |
| <u>AP</u> | ! | | <u>EQ 12.5MG BASE/ML</u> | <u>A074292 001</u> | Feb 16, 1995 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>EQ 12.5MG BASE/ML</u> | <u>A074277 001</u> | Oct 31, 1994 |

DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | |
|-----------|---|---------|----------------------------|--------------------|--------------|
| <u>AP</u> | + | ! | <u>EQ 50MG BASE/100ML</u> | <u>N020255 001</u> | Oct 19, 1993 |
| <u>AP</u> | + | ! | <u>EQ 100MG BASE/100ML</u> | <u>N020255 003</u> | Oct 19, 1993 |
| <u>AP</u> | + | ! | <u>EQ 200MG BASE/100ML</u> | <u>N020255 004</u> | Oct 19, 1993 |
| <u>AP</u> | + | ! | <u>EQ 400MG BASE/100ML</u> | <u>N020255 005</u> | Oct 19, 1993 |
| <u>AP</u> | + | HOSPIRA | <u>EQ 50MG BASE/100ML</u> | <u>N020201 003</u> | Oct 19, 1993 |
| <u>AP</u> | + | ! | <u>EQ 100MG BASE/100ML</u> | <u>N020201 002</u> | Oct 19, 1993 |
| <u>AP</u> | + | ! | <u>EQ 200MG BASE/100ML</u> | <u>N020201 001</u> | Oct 19, 1993 |
| <u>AP</u> | + | ! | <u>EQ 400MG BASE/100ML</u> | <u>N020201 006</u> | Jul 07, 1994 |

PRESCRIPTION DRUG PRODUCT LIST

DOCETAXEL

INJECTABLE; INJECTION

DOCETAXEL

| | | | | | |
|-----------|---|---------------------|-----------------------------|--------------------|--------------|
| <u>AP</u> | + | ACCORD HLTHCARE | <u>20MG/ML (20MG/ML)</u> | <u>N201195 003</u> | Apr 20, 2012 |
| <u>AP</u> | + | | <u>80MG/4ML (20MG/ML)</u> | <u>N201195 004</u> | Apr 20, 2012 |
| <u>AP</u> | + | | <u>160MG/8ML (20MG/ML)</u> | <u>N201195 005</u> | Apr 20, 2012 |
| <u>AP</u> | | ACTAVIS LLC | <u>20MG/ML (20MG/ML)</u> | <u>N203551 001</u> | Apr 12, 2013 |
| <u>AP</u> | | | <u>80MG/4ML (20MG/ML)</u> | <u>N203551 002</u> | Apr 12, 2013 |
| <u>AP</u> | | AMNEAL PHARMS CO | <u>20MG/ML (20MG/ML)</u> | <u>A209640 001</u> | Jan 19, 2018 |
| <u>AP</u> | | | <u>80MG/4ML (20MG/ML)</u> | <u>A209640 002</u> | Jan 19, 2018 |
| <u>AP</u> | | | <u>160MG/8ML (20MG/ML)</u> | <u>A209640 003</u> | Jan 19, 2018 |
| <u>AP</u> | | CIPLA | <u>20MG/2ML (10MG/ML)</u> | <u>A209634 001</u> | Aug 24, 2018 |
| <u>AP</u> | | | <u>80MG/8ML (10MG/ML)</u> | <u>A209634 002</u> | Aug 24, 2018 |
| <u>AP</u> | | | <u>160MG/16ML (10MG/ML)</u> | <u>A209634 003</u> | Aug 24, 2018 |
| <u>AP</u> | | DFB ONCOLOGY LTD | <u>20MG/ML (20MG/ML)</u> | <u>A206177 001</u> | Jan 20, 2017 |
| <u>AP</u> | | | <u>80MG/4ML (20MG/ML)</u> | <u>A206177 002</u> | Jan 20, 2017 |
| <u>AP</u> | | DR REDDYS LABS LTD | <u>20MG/ML (20MG/ML)</u> | <u>A204193 001</u> | Nov 05, 2014 |
| <u>AP</u> | | | <u>80MG/4ML (20MG/ML)</u> | <u>A204193 002</u> | Nov 05, 2014 |
| <u>AP</u> | | EAGLE PHARMS | <u>20MG/ML (20MG/ML)</u> | <u>N205934 001</u> | Dec 22, 2015 |
| <u>AP</u> | | | <u>80MG/4ML (20MG/ML)</u> | <u>N205934 002</u> | Dec 22, 2015 |
| <u>AP</u> | | | <u>160MG/8ML (20MG/ML)</u> | <u>N205934 003</u> | Dec 22, 2015 |
| <u>AP</u> | + | HOSPIRA INC | <u>20MG/2ML (10MG/ML)</u> | <u>N022234 001</u> | Mar 08, 2011 |
| <u>AP</u> | + | | <u>80MG/8ML (10MG/ML)</u> | <u>N022234 002</u> | Mar 08, 2011 |
| <u>AP</u> | + | | <u>160MG/16ML (10MG/ML)</u> | <u>N022234 003</u> | Mar 08, 2011 |
| <u>AP</u> | | INGENUS PHARMS LLC | <u>20MG/2ML (10MG/ML)</u> | <u>A207563 001</u> | Aug 31, 2017 |
| <u>AP</u> | | | <u>80MG/8ML (10MG/ML)</u> | <u>A207563 002</u> | Aug 31, 2017 |
| <u>AP</u> | | | <u>160MG/16ML (10MG/ML)</u> | <u>A207563 003</u> | Aug 31, 2017 |
| <u>AP</u> | | JIANGSU HENGRUI MED | <u>20MG/ML (20MG/ML)</u> | <u>A207252 001</u> | Aug 09, 2017 |
| <u>AP</u> | | | <u>80MG/4ML (20MG/ML)</u> | <u>A207252 002</u> | Aug 09, 2017 |
| <u>AP</u> | | | <u>160MG/8ML (20MG/ML)</u> | <u>A207252 003</u> | Aug 09, 2017 |
| <u>AP</u> | | MYLAN LABS LTD | <u>20MG/2ML (10MG/ML)</u> | <u>A210072 001</u> | Jul 02, 2018 |
| <u>AP</u> | | | <u>80MG/8ML (10MG/ML)</u> | <u>A210848 001</u> | Jul 06, 2018 |
| <u>AP</u> | | SANDOZ | <u>20MG/2ML (10MG/ML)</u> | <u>N201525 001</u> | Jun 29, 2011 |
| <u>AP</u> | | | <u>80MG/8ML (10MG/ML)</u> | <u>N201525 002</u> | Jun 29, 2011 |
| <u>AP</u> | | | <u>160MG/16ML (10MG/ML)</u> | <u>N201525 003</u> | Jun 29, 2011 |
| <u>AP</u> | | TEVA PHARMS USA | <u>20MG/ML (20MG/ML)</u> | <u>A203877 001</u> | Sep 16, 2015 |
| <u>AP</u> | | | <u>80MG/4ML (20MG/ML)</u> | <u>A203877 002</u> | Sep 16, 2015 |

TAXOTERE

| | | | | | |
|-----------|---|-------------------|----------------------------|--------------------|--------------|
| <u>AP</u> | + | SANOFI AVENTIS US | <u>20MG/ML (20MG/ML)</u> | <u>N020449 003</u> | Aug 03, 2010 |
| <u>AP</u> | + | | <u>80MG/4ML (20MG/ML)</u> | <u>N020449 004</u> | Aug 02, 2010 |
| <u>AP</u> | + | | <u>160MG/8ML (20MG/ML)</u> | <u>N020449 005</u> | Apr 13, 2012 |

DOCETAXEL

| | | | | | |
|--|---|---------------------|----------------------|-------------|--------------|
| | | ACTAVIS LLC | 140MG/7ML (20MG/ML) | N203551 003 | Apr 12, 2013 |
| | | DFB ONCOLOGY LTD | 200MG/10ML (20MG/ML) | A206177 003 | Jan 20, 2017 |
| | + | HOSPIRA INC | 20MG/ML (20MG/ML) | N022234 004 | Jun 23, 2016 |
| | + | | 80MG/4ML (20MG/ML) | N022234 005 | Jun 23, 2016 |
| | + | | 160MG/8ML (20MG/ML) | N022234 007 | Jan 24, 2017 |
| | ! | JIANGSU HENGRUI MED | 40MG/ML | A203170 001 | Feb 15, 2017 |

DOCETAXEL

INJECTABLE; INJECTION

DOCETAXEL

| | | | | | |
|-----------|--|----------------|-----------------------------|--------------------|--------------|
| <u>AP</u> | | MYLAN LABS LTD | <u>160MG/16ML (10MG/ML)</u> | <u>A208859 001</u> | Apr 30, 2018 |
|-----------|--|----------------|-----------------------------|--------------------|--------------|

DOFETILIDE

CAPSULE; ORAL

DOFETILIDE

| | | | | | |
|-----------|--|---------------------|----------------|--------------------|--------------|
| <u>AB</u> | | BIONPHARMA INC | <u>0.125MG</u> | <u>A208625 001</u> | Apr 10, 2018 |
| <u>AB</u> | | | <u>0.25MG</u> | <u>A208625 002</u> | Apr 10, 2018 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A208625 003</u> | Apr 10, 2018 |
| <u>AB</u> | | MAYNE PHARMA INC | <u>0.125MG</u> | <u>A207058 001</u> | Jun 06, 2016 |
| <u>AB</u> | | | <u>0.25MG</u> | <u>A207058 002</u> | Jun 06, 2016 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A207058 003</u> | Jun 06, 2016 |
| <u>AB</u> | | PAR PHARM INC | <u>0.125MG</u> | <u>A208519 001</u> | Oct 09, 2018 |
| <u>AB</u> | | | <u>0.25MG</u> | <u>A208519 002</u> | Oct 09, 2018 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A208519 003</u> | Oct 09, 2018 |
| <u>AB</u> | | SIGMAPHARM LABS LLC | <u>0.125MG</u> | <u>A207746 001</u> | Mar 26, 2018 |
| <u>AB</u> | | | <u>0.25MG</u> | <u>A207746 002</u> | Mar 26, 2018 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A207746 003</u> | Mar 26, 2018 |
| <u>AB</u> | | SUN PHARMA GLOBAL | <u>0.125MG</u> | <u>A210466 001</u> | Oct 09, 2018 |
| <u>AB</u> | | | <u>0.25MG</u> | <u>A210466 002</u> | Oct 09, 2018 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A210466 003</u> | Oct 09, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

DOFETILIDE

CAPSULE;ORAL

TIKOSYN

| | | | | | | |
|-----------|---|--------|----------------|----------------|------------|--------------|
| <u>AB</u> | + | PFIZER | <u>0.125MG</u> | <u>N020931</u> | <u>001</u> | Oct 01, 1999 |
| <u>AB</u> | + | | <u>0.25MG</u> | <u>N020931</u> | <u>002</u> | Oct 01, 1999 |
| <u>AB</u> | + | ! | <u>0.5MG</u> | <u>N020931</u> | <u>003</u> | Oct 01, 1999 |

DOLUTEGRAVIR SODIUM

TABLET;ORAL

TIVICAY

| | | | | | |
|---|---------------|--------------|---------|-----|--------------|
| + | VIIV HLTHCARE | EQ 10MG BASE | N204790 | 002 | Jun 09, 2016 |
| + | | EQ 25MG BASE | N204790 | 003 | Jun 09, 2016 |
| + | ! | EQ 50MG BASE | N204790 | 001 | Aug 12, 2013 |

DOLUTEGRAVIR SODIUM; RILPIVIRINE HYDROCHLORIDE

TABLET;ORAL

JULUCA

| | | | | | | |
|---|---|---------------|---------------------------|---------|-----|--------------|
| + | ! | VIIV HLTHCARE | EQ 50MG BASE;EQ 25MG BASE | N210192 | 001 | Nov 21, 2017 |
|---|---|---------------|---------------------------|---------|-----|--------------|

DONEPEZIL HYDROCHLORIDE

TABLET;ORAL

ARICEPT

| | | | | | | |
|-----------|---|-----------|-------------|----------------|------------|--------------|
| <u>AB</u> | + | EISAI INC | <u>5MG</u> | <u>N020690</u> | <u>002</u> | Nov 25, 1996 |
| <u>AB</u> | + | ! | <u>10MG</u> | <u>N020690</u> | <u>001</u> | Nov 25, 1996 |
| <u>AB</u> | + | ! | <u>23MG</u> | <u>N022568</u> | <u>001</u> | Jul 23, 2010 |

DONEPEZIL HYDROCHLORIDE

| | | | | | | |
|-----------|--|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | ACI HEALTHCARE LTD | <u>5MG</u> | <u>A078662</u> | <u>001</u> | May 31, 2011 |
| <u>AB</u> | | | <u>10MG</u> | <u>A078662</u> | <u>002</u> | May 31, 2011 |
| <u>AB</u> | | ACTAVIS ELIZABETH | <u>23MG</u> | <u>A202415</u> | <u>001</u> | Dec 17, 2015 |
| <u>AB</u> | | ALEMBIC PHARMS LTD | <u>5MG</u> | <u>A201724</u> | <u>001</u> | Feb 25, 2013 |
| <u>AB</u> | | | <u>10MG</u> | <u>A201724</u> | <u>002</u> | Feb 25, 2013 |
| <u>AB</u> | | AUROBINDO | <u>5MG</u> | <u>A090056</u> | <u>001</u> | May 31, 2011 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090056</u> | <u>002</u> | May 31, 2011 |
| <u>AB</u> | | CADILA PHARMS LTD | <u>5MG</u> | <u>A204609</u> | <u>001</u> | Sep 19, 2017 |
| <u>AB</u> | | | <u>10MG</u> | <u>A204609</u> | <u>002</u> | Sep 19, 2017 |
| <u>AB</u> | | CIPLA LTD | <u>5MG</u> | <u>A077518</u> | <u>001</u> | May 31, 2011 |
| <u>AB</u> | | | <u>10MG</u> | <u>A077518</u> | <u>002</u> | May 31, 2011 |
| <u>AB</u> | | CSPC OUYI PHARM CO | <u>5MG</u> | <u>A202114</u> | <u>001</u> | Jul 05, 2013 |
| <u>AB</u> | | | <u>10MG</u> | <u>A202114</u> | <u>002</u> | Jul 05, 2013 |
| <u>AB</u> | | DEXCEL PHARMA | <u>23MG</u> | <u>A203713</u> | <u>001</u> | Feb 19, 2016 |
| <u>AB</u> | | DR REDDYS LABS LTD | <u>5MG</u> | <u>A201001</u> | <u>001</u> | May 31, 2011 |
| <u>AB</u> | | | <u>10MG</u> | <u>A201001</u> | <u>002</u> | May 31, 2011 |
| <u>AB</u> | | | <u>23MG</u> | <u>A202723</u> | <u>001</u> | Jul 24, 2013 |
| <u>AB</u> | | HETERO LABS LTD V | <u>5MG</u> | <u>A203034</u> | <u>001</u> | Jan 30, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A203034</u> | <u>002</u> | Jan 30, 2015 |
| <u>AB</u> | | HISUN PHARM HANGZHOU | <u>23MG</u> | <u>A202410</u> | <u>001</u> | Mar 24, 2017 |
| <u>AB</u> | | INDICUS PHARMA | <u>5MG</u> | <u>A201634</u> | <u>001</u> | Jun 13, 2012 |
| <u>AB</u> | | | <u>10MG</u> | <u>A201634</u> | <u>002</u> | Jun 13, 2012 |
| <u>AB</u> | | | <u>23MG</u> | <u>A203419</u> | <u>001</u> | Apr 12, 2016 |
| <u>AB</u> | | JUBILANT GENERICS | <u>5MG</u> | <u>A090768</u> | <u>001</u> | May 31, 2011 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090768</u> | <u>002</u> | May 31, 2011 |
| <u>AB</u> | | LUPIN LTD | <u>23MG</u> | <u>A202782</u> | <u>001</u> | Oct 30, 2015 |
| <u>AB</u> | | MACLEODS PHARMS LTD | <u>5MG</u> | <u>A201146</u> | <u>001</u> | Aug 17, 2012 |
| <u>AB</u> | | | <u>10MG</u> | <u>A201146</u> | <u>002</u> | Aug 17, 2012 |
| <u>AB</u> | | | <u>23MG</u> | <u>A202631</u> | <u>001</u> | Jan 22, 2014 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>5MG</u> | <u>A090521</u> | <u>001</u> | May 31, 2011 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090521</u> | <u>002</u> | May 31, 2011 |
| <u>AB</u> | | | <u>23MG</u> | <u>A202656</u> | <u>001</u> | Oct 22, 2015 |
| <u>AB</u> | | OSMOTICA PHARM US | <u>23MG</u> | <u>A203114</u> | <u>001</u> | Jan 26, 2016 |
| <u>AB</u> | | PAR PHARM | <u>23MG</u> | <u>A202542</u> | <u>001</u> | Jul 24, 2013 |
| <u>AB</u> | | PLIVA HRVATSKA DOO | <u>5MG</u> | <u>A090425</u> | <u>001</u> | May 31, 2011 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090425</u> | <u>002</u> | May 31, 2011 |
| <u>AB</u> | | PRINSTON INC | <u>5MG</u> | <u>A200292</u> | <u>001</u> | May 31, 2011 |
| <u>AB</u> | | | <u>10MG</u> | <u>A200292</u> | <u>002</u> | May 31, 2011 |
| <u>AB</u> | | SANDOZ | <u>5MG</u> | <u>A090290</u> | <u>001</u> | May 31, 2011 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090290</u> | <u>002</u> | May 31, 2011 |
| <u>AB</u> | | SCIEGEN PHARMS INC | <u>5MG</u> | <u>A203907</u> | <u>001</u> | Oct 29, 2014 |
| <u>AB</u> | | | <u>10MG</u> | <u>A203907</u> | <u>002</u> | Oct 29, 2014 |
| <u>AB</u> | | STRIDES VIVIMED | <u>5MG</u> | <u>A090551</u> | <u>001</u> | May 31, 2011 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090551</u> | <u>002</u> | May 31, 2011 |
| <u>AB</u> | | SUN PHARM INDS | <u>5MG</u> | <u>A090493</u> | <u>001</u> | May 31, 2011 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090493</u> | <u>002</u> | May 31, 2011 |
| <u>AB</u> | | SUN PHARM INDS LTD | <u>23MG</u> | <u>A204293</u> | <u>001</u> | Jun 05, 2015 |

PRESCRIPTION DRUG PRODUCT LIST

DONEPEZIL HYDROCHLORIDE

TABLET; ORAL

DONEPEZIL HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|-------------|--------------------|--------------|
| <u>AB</u> | TEVA | <u>5MG</u> | <u>A077344 001</u> | May 31, 2011 |
| <u>AB</u> | | <u>10MG</u> | <u>A077344 002</u> | May 31, 2011 |
| <u>AB</u> | TORRENT PHARMS | <u>5MG</u> | <u>A090686 001</u> | May 31, 2011 |
| <u>AB</u> | | <u>10MG</u> | <u>A090686 002</u> | May 31, 2011 |
| <u>AB</u> | TWI PHARMS | <u>23MG</u> | <u>A203104 001</u> | Oct 29, 2014 |
| <u>AB</u> | UNICHEM LABS LTD | <u>5MG</u> | <u>A203656 001</u> | Jun 23, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A203656 002</u> | Jun 23, 2016 |
| <u>AB</u> | WOCKHARDT | <u>5MG</u> | <u>A091267 001</u> | May 31, 2011 |
| <u>AB</u> | | <u>10MG</u> | <u>A091267 002</u> | May 31, 2011 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>5MG</u> | <u>A090100 001</u> | Oct 24, 2012 |
| <u>AB</u> | | <u>10MG</u> | <u>A090100 002</u> | Oct 24, 2012 |
| <u>AB</u> | | <u>23MG</u> | <u>A203162 001</u> | Aug 31, 2017 |

TABLET, ORALLY DISINTEGRATING; ORAL

DONEPEZIL HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|-------------|--------------------|--------------|
| <u>AB</u> | HISUN PHARM HANGZHOU | <u>5MG</u> | <u>A205269 001</u> | Jul 27, 2018 |
| <u>AB</u> | | <u>10MG</u> | <u>A205269 002</u> | Jul 27, 2018 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>5MG</u> | <u>A201787 001</u> | Dec 14, 2012 |
| <u>AB</u> | | <u>10MG</u> | <u>A201787 002</u> | Dec 14, 2012 |
| <u>AB</u> | SANDOZ | <u>5MG</u> | <u>A091198 001</u> | May 10, 2011 |
| <u>AB</u> | ! | <u>10MG</u> | <u>A091198 002</u> | May 10, 2011 |
| <u>AB</u> | UNICHEM LABS LTD | <u>5MG</u> | <u>A204831 001</u> | Nov 10, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A204831 002</u> | Nov 10, 2016 |

DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

NAMZARIC

| | | | | |
|---|--------------------|------------|-------------|--------------|
| + | ALLERGAN SALES LLC | 10MG; 7MG | N206439 003 | Jul 18, 2016 |
| + | | 10MG; 14MG | N206439 001 | Dec 23, 2014 |
| + | | 10MG; 21MG | N206439 004 | Jul 18, 2016 |
| + | ! | 10MG; 28MG | N206439 002 | Dec 23, 2014 |

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOPAMINE HYDROCHLORIDE

| | | | | |
|-----------|-------------------|--------------------|--------------------|--------------|
| <u>AP</u> | HIKMA INTL PHARMS | <u>40MG/ML</u> | <u>A207707 001</u> | Apr 11, 2018 |
| <u>AP</u> | | <u>80MG/ML</u> | <u>A207707 002</u> | Apr 11, 2018 |
| <u>AP</u> | +! HOSPIRA | <u>40MG/ML</u> | <u>N018132 001</u> | |
| <u>AP</u> | +! | <u>80MG/100ML</u> | <u>N018132 002</u> | Feb 04, 1982 |
| <u>AP</u> | +! | <u>80MG/ML</u> | <u>N018132 004</u> | Jul 09, 1982 |
| <u>AP</u> | +! | <u>160MG/100ML</u> | <u>N018132 003</u> | Feb 04, 1982 |
| <u>AP</u> | ! LUITPOLD | <u>40MG/ML</u> | <u>A070799 001</u> | Feb 11, 1987 |
| <u>AP</u> | ! | <u>80MG/ML</u> | <u>A070820 001</u> | Feb 11, 1987 |

DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%

| | | | | |
|-----------|------------|--------------------|--------------------|--------------|
| <u>AP</u> | +! B BRAUN | <u>80MG/100ML</u> | <u>N019099 002</u> | Oct 15, 1986 |
| <u>AP</u> | +! | <u>320MG/100ML</u> | <u>N019099 004</u> | Oct 15, 1986 |

DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|-----------|------------|--------------------|--------------------|--------------|
| <u>AP</u> | +! B BRAUN | <u>160MG/100ML</u> | <u>N019099 003</u> | Oct 15, 1986 |
|-----------|------------|--------------------|--------------------|--------------|

DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|-----------|--------------------|--------------------|--------------------|--------------|
| <u>AP</u> | +! BAXTER HLTHCARE | <u>80MG/100ML</u> | <u>N019615 001</u> | Mar 27, 1987 |
| <u>AP</u> | +! | <u>160MG/100ML</u> | <u>N019615 002</u> | Mar 27, 1987 |
| <u>AP</u> | +! | <u>320MG/100ML</u> | <u>N019615 003</u> | Mar 27, 1987 |
| <u>AP</u> | +! HOSPIRA | <u>80MG/100ML</u> | <u>N018826 001</u> | Sep 30, 1983 |
| <u>AP</u> | +! | <u>160MG/100ML</u> | <u>N018826 002</u> | Sep 30, 1983 |
| <u>AP</u> | +! | <u>320MG/100ML</u> | <u>N018826 003</u> | Sep 30, 1983 |

DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|---|---------|------------|-------------|--------------|
| + | B BRAUN | 40MG/100ML | N019099 001 | Oct 15, 1986 |
|---|---------|------------|-------------|--------------|

DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|---|-----------------|-------------|-------------|--------------|
| + | BAXTER HLTHCARE | 640MG/100ML | N019615 004 | Mar 27, 1987 |
|---|-----------------|-------------|-------------|--------------|

DORAVIRINE

TABLET; ORAL

PIFELTRO

| | | | | |
|---|--------------|-------|-------------|--------------|
| + | MSD MERCK CO | 100MG | N210806 001 | Aug 30, 2018 |
|---|--------------|-------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

DORAVIRINE; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

DELSTRIGO

+! MSD MERCK CO 100MG; 300MG; 300MG N210807 001 Aug 30, 2018

DORZOLAMIDE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE

| | | | | |
|-----------|-----------------|-------------------|--------------------|--------------|
| <u>AT</u> | BAUSCH AND LOMB | <u>EQ 2% BASE</u> | <u>A090143 001</u> | Jun 25, 2009 |
| <u>AT</u> | HI TECH PHARMA | <u>EQ 2% BASE</u> | <u>A077846 001</u> | Oct 28, 2008 |
| <u>AT</u> | LUITPOLD | <u>EQ 2% BASE</u> | <u>A079186 001</u> | Nov 18, 2009 |
| <u>AT</u> | SANDOZ INC | <u>EQ 2% BASE</u> | <u>A078748 001</u> | Nov 06, 2008 |
| <u>AT</u> | | <u>EQ 2% BASE</u> | <u>A078981 001</u> | Apr 13, 2009 |

TRUSOPT

| | | | | |
|-----------|----------|-------------------|--------------------|--------------|
| <u>AT</u> | +! MERCK | <u>EQ 2% BASE</u> | <u>N020408 001</u> | Dec 09, 1994 |
|-----------|----------|-------------------|--------------------|--------------|

DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

COSOPT

| | | | | |
|-----------|-------------------|---------------------------------|--------------------|--------------|
| <u>AT</u> | +! OAK PHARMS INC | <u>EQ 2% BASE; EQ 0.5% BASE</u> | <u>N020869 001</u> | Apr 07, 1998 |
|-----------|-------------------|---------------------------------|--------------------|--------------|

COSOPT PF

| | | | | |
|-----------|-------------------|---------------------------------|--------------------|--------------|
| <u>AT</u> | +! OAK PHARMS INC | <u>EQ 2% BASE; EQ 0.5% BASE</u> | <u>N202667 001</u> | Feb 01, 2012 |
|-----------|-------------------|---------------------------------|--------------------|--------------|

DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE

| | | | | |
|-----------|----------------------|---------------------------------|--------------------|--------------|
| <u>AT</u> | AKORN INC | <u>EQ 2% BASE; EQ 0.5% BASE</u> | <u>A203058 001</u> | Sep 22, 2014 |
| <u>AT</u> | AUROBINDO PHARMA LTD | <u>EQ 2% BASE; EQ 0.5% BASE</u> | <u>A207630 001</u> | Jul 24, 2018 |
| <u>AT</u> | BAUSCH AND LOMB | <u>EQ 2% BASE; EQ 0.5% BASE</u> | <u>A090037 001</u> | Jul 14, 2009 |
| <u>AT</u> | HI TECH PHARMA | <u>EQ 2% BASE; EQ 0.5% BASE</u> | <u>A077847 001</u> | Oct 28, 2008 |
| <u>AT</u> | SANDOZ | <u>EQ 2% BASE; EQ 0.5% BASE</u> | <u>A078749 001</u> | Nov 06, 2008 |
| <u>AT</u> | SANDOZ INC | <u>EQ 2% BASE; EQ 0.5% BASE</u> | <u>A090604 001</u> | Nov 18, 2009 |
| <u>AT</u> | TEVA PHARMS | <u>EQ 2% BASE; EQ 0.5% BASE</u> | <u>A078704 001</u> | Sep 28, 2009 |

DOXAPRAM HYDROCHLORIDE

INJECTABLE; INJECTION

DOPRAM

| | | | | |
|-----------|-------------------------|----------------|--------------------|--|
| <u>AP</u> | +! WEST-WARD PHARMS INT | <u>20MG/ML</u> | <u>N014879 001</u> | |
|-----------|-------------------------|----------------|--------------------|--|

DOXAPRAM HYDROCHLORIDE

| | | | | |
|-----------|-------------|----------------|--------------------|--------------|
| <u>AP</u> | ATHENEX INC | <u>20MG/ML</u> | <u>A076266 001</u> | Jan 10, 2003 |
|-----------|-------------|----------------|--------------------|--------------|

DOXAZOSIN MESYLATE

TABLET; ORAL

CARDURA

| | | | | |
|-----------|-----------|--------------------|--------------------|--------------|
| <u>AB</u> | +! PFIZER | <u>EQ 1MG BASE</u> | <u>N019668 001</u> | Nov 02, 1990 |
| <u>AB</u> | + | <u>EQ 2MG BASE</u> | <u>N019668 002</u> | Nov 02, 1990 |
| <u>AB</u> | + | <u>EQ 4MG BASE</u> | <u>N019668 003</u> | Nov 02, 1990 |
| <u>AB</u> | + | <u>EQ 8MG BASE</u> | <u>N019668 004</u> | Nov 02, 1990 |

DOXAZOSIN MESYLATE

| | | | | |
|-----------|-----------------|--------------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>EQ 1MG BASE</u> | <u>A202824 001</u> | Jun 11, 2014 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A202824 002</u> | Jun 11, 2014 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A202824 003</u> | Jun 11, 2014 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A202824 004</u> | Jun 11, 2014 |
| <u>AB</u> | ANI PHARMS INC | <u>EQ 1MG BASE</u> | <u>A075432 001</u> | Oct 18, 2000 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A075432 002</u> | Oct 18, 2000 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A075432 003</u> | Oct 18, 2000 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A075432 004</u> | Oct 18, 2000 |
| <u>AB</u> | APOTEX | <u>EQ 1MG BASE</u> | <u>A075580 001</u> | Oct 18, 2000 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A075580 002</u> | Oct 18, 2000 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A075580 003</u> | Oct 18, 2000 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A075580 004</u> | Oct 18, 2000 |
| <u>AB</u> | DAVA PHARMS INC | <u>EQ 1MG BASE</u> | <u>A076161 001</u> | Jun 10, 2004 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A076161 002</u> | Jun 10, 2004 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A076161 003</u> | Jun 10, 2004 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A076161 004</u> | Jun 10, 2004 |
| <u>AB</u> | HERITAGE PHARMA | <u>EQ 1MG BASE</u> | <u>A205210 001</u> | Feb 13, 2018 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A205210 002</u> | Feb 13, 2018 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A205210 003</u> | Feb 13, 2018 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A205210 004</u> | Feb 13, 2018 |
| <u>AB</u> | MYLAN | <u>EQ 1MG BASE</u> | <u>A075509 001</u> | Oct 19, 2000 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A075509 002</u> | Oct 19, 2000 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A075509 003</u> | Oct 19, 2000 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A075509 004</u> | Oct 19, 2000 |
| <u>AB</u> | PLIVA | <u>EQ 1MG BASE</u> | <u>A075750 001</u> | Jun 08, 2001 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A075750 002</u> | Jun 08, 2001 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A075750 003</u> | Jun 08, 2001 |

PRESCRIPTION DRUG PRODUCT LIST

DOXAZOSIN MESYLATE

TABLET; ORAL

DOXAZOSIN MESYLATE

| | | | | |
|-----------|-------------------------|--------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A075750 004</u> | Jun 08, 2001 |
| <u>AB</u> | TEVA | <u>EQ 1MG BASE</u> | <u>A075536 001</u> | Oct 18, 2000 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A075536 002</u> | Oct 18, 2000 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A075536 003</u> | Oct 18, 2000 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A075536 004</u> | Oct 18, 2000 |
| <u>AB</u> | UPSHER SMITH LABS | <u>EQ 1MG BASE</u> | <u>A209013 001</u> | Apr 17, 2018 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A209013 002</u> | Apr 17, 2018 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A209013 003</u> | Apr 17, 2018 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A209013 004</u> | Apr 17, 2018 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 1MG BASE</u> | <u>A208719 001</u> | Jul 07, 2017 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A208719 002</u> | Jul 07, 2017 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A208719 003</u> | Jul 07, 2017 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A208719 004</u> | Jul 07, 2017 |

TABLET, EXTENDED RELEASE; ORAL

CARDURA XL

| | | | | |
|---|--------|-------------|-------------|--------------|
| + | PFIZER | EQ 4MG BASE | N021269 001 | Feb 22, 2005 |
| + | ! | EQ 8MG BASE | N021269 002 | Feb 22, 2005 |

DOXEPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPIN HYDROCHLORIDE

| | | | | |
|-----------|------------------|----------------------|--------------------|--------------|
| <u>AB</u> | AMNEAL PHARMS CO | <u>EQ 10MG BASE</u> | <u>A207482 001</u> | Jun 28, 2017 |
| <u>AB</u> | | <u>EQ 25MG BASE</u> | <u>A207482 002</u> | Jun 28, 2017 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A207482 003</u> | Jun 28, 2017 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A207482 004</u> | Jun 28, 2017 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A207482 005</u> | Jun 28, 2017 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 10MG BASE</u> | <u>A070791 002</u> | May 13, 1986 |
| <u>AB</u> | ! | <u>EQ 25MG BASE</u> | <u>A070791 003</u> | May 13, 1986 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A070791 001</u> | May 13, 1986 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A070791 004</u> | May 13, 1986 |
| <u>AB</u> | ! | <u>EQ 100MG BASE</u> | <u>A070791 005</u> | May 13, 1986 |
| <u>AB</u> | PAR PHARM | <u>EQ 50MG BASE</u> | <u>A071595 001</u> | Nov 09, 1987 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A071422 001</u> | Nov 09, 1987 |
| <u>AB</u> | ! | EQ 150MG BASE | A071669 001 | Nov 09, 1987 |

CONCENTRATE; ORAL

DOXEPIN HYDROCHLORIDE

| | | | | |
|-----------|------------------|------------------------|--------------------|--------------|
| <u>AA</u> | LANNETT CO INC | <u>EQ 10MG BASE/ML</u> | <u>A074721 001</u> | Dec 29, 1998 |
| <u>AA</u> | ! | <u>EQ 10MG BASE/ML</u> | <u>A071609 001</u> | Nov 09, 1987 |
| <u>AA</u> | WOCKHARDT BIO AG | <u>EQ 10MG BASE/ML</u> | <u>A071918 001</u> | Jul 20, 1988 |

CREAM; TOPICAL

ZONALON

| | | | | | |
|---|---|------------------|----|-------------|--------------|
| + | ! | MYLAN PHARMS INC | 5% | N020126 001 | Apr 01, 1994 |
|---|---|------------------|----|-------------|--------------|

TABLET; ORAL

SILENOR

| | | | | | |
|---|---|--------------------|-------------|-------------|--------------|
| + | | PERNIX THERAPS LLC | EQ 3MG BASE | N022036 001 | Mar 17, 2010 |
| + | ! | | EQ 6MG BASE | N022036 002 | Mar 17, 2010 |

DOXERCALCIFEROL

CAPSULE; ORAL

DOXERCALCIFEROL

| | | | | |
|-----------|-------------------------|---------------|--------------------|--------------|
| <u>AB</u> | RISING PHARMS | <u>0.5MCG</u> | <u>A201518 001</u> | Sep 09, 2016 |
| <u>AB</u> | | <u>1MCG</u> | <u>A201518 002</u> | Sep 09, 2016 |
| <u>AB</u> | | <u>2.5MCG</u> | <u>A201518 003</u> | Sep 09, 2016 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>0.5MCG</u> | <u>A091433 001</u> | Sep 23, 2011 |
| <u>AB</u> | | <u>1MCG</u> | <u>A091433 002</u> | Jan 14, 2014 |
| <u>AB</u> | | <u>2.5MCG</u> | <u>A091433 003</u> | Jan 14, 2014 |

HECTOROL

| | | | | | |
|-----------|---|--------|---------------|--------------------|--------------|
| <u>AB</u> | + | SANOFI | <u>0.5MCG</u> | <u>N020862 002</u> | Apr 23, 2004 |
| <u>AB</u> | + | | <u>1MCG</u> | <u>N020862 003</u> | Jul 13, 2009 |
| <u>AB</u> | + | ! | <u>2.5MCG</u> | <u>N020862 001</u> | Jun 09, 1999 |

INJECTABLE; INJECTION

DOXERCALCIFEROL

| | | | | | |
|-----------|------------------|---------------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | AKORN INC | <u>2MCG/ML (2MCG/ML)</u> | <u>A203929 002</u> | Mar 28, 2016 | |
| <u>AP</u> | | <u>4MCG/2ML (2MCG/ML)</u> | <u>A203929 001</u> | May 07, 2015 | |
| <u>AP</u> | AMNEAL PHARMS CO | <u>2MCG/ML (2MCG/ML)</u> | <u>A208974 001</u> | May 24, 2017 | |
| <u>AP</u> | | <u>4MCG/2ML (2MCG/ML)</u> | <u>A208974 002</u> | May 24, 2017 | |
| <u>AP</u> | | <u>4MCG/2ML (2MCG/ML)</u> | <u>A208975 001</u> | May 24, 2017 | |
| <u>AP</u> | HIKMA PHARMS | <u>4MCG/2ML (2MCG/ML)</u> | <u>A091101 001</u> | Aug 30, 2013 | |
| <u>AP</u> | + | HOSPIRA INC | <u>4MCG/2ML (2MCG/ML)</u> | <u>N208614 001</u> | Jul 24, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

DOXERCALCIFEROL

INJECTABLE; INJECTION

DOXERCALCIFEROL

| | | | | |
|-----------------|----------------|---------------------------|--------------------|--------------|
| <u>AP</u> | LUPIN LTD | <u>4MCG/2ML (2MCG/ML)</u> | <u>A210801 001</u> | Nov 01, 2018 |
| <u>AP</u> | SANDOZ INC | <u>4MCG/2ML (2MCG/ML)</u> | <u>A091333 001</u> | May 05, 2014 |
| <u>AP</u> | | <u>4MCG/2ML (2MCG/ML)</u> | <u>A200926 001</u> | Feb 04, 2014 |
| <u>HECTOROL</u> | | | | |
| <u>AP</u> | + SANOFI | <u>2MCG/ML (2MCG/ML)</u> | <u>N021027 002</u> | Apr 06, 2000 |
| <u>AP</u> | +! | <u>4MCG/2ML (2MCG/ML)</u> | <u>N021027 001</u> | Apr 06, 2000 |
| DOXERCALCIFEROL | | | | |
| | +! HOSPIRA INC | 10MCG/5ML (2MCG/ML) | N208614 002 | Jul 24, 2018 |

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

DOXORUBICIN HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|--------------------|--------------------|--------------|
| <u>AP</u> | ACTAVIS INC | <u>2MG/ML</u> | <u>A203622 001</u> | Jun 27, 2014 |
| <u>AP</u> | | <u>200MG/100ML</u> | <u>A203622 002</u> | Jun 27, 2014 |
| <u>AP</u> | AMNEAL PHARMS CO | <u>20MG/VIAL</u> | <u>A208888 001</u> | Feb 17, 2017 |
| <u>AP</u> | | <u>50MG/VIAL</u> | <u>A208888 002</u> | Feb 17, 2017 |
| <u>AP</u> | FRESENIUS KABI USA | <u>2MG/ML</u> | <u>A063277 001</u> | Oct 26, 1995 |
| <u>AP</u> | GLAND PHARMA LTD | <u>2MG/ML</u> | <u>A209825 001</u> | Aug 11, 2017 |
| <u>AP</u> | MYLAN LABS LTD | <u>2MG/ML</u> | <u>A200901 001</u> | Feb 14, 2012 |
| <u>AP</u> | | <u>50MG/VIAL</u> | <u>A200170 002</u> | Oct 28, 2011 |
| <u>AP</u> | PHARMACHEMIE BV | <u>2MG/ML</u> | <u>A063336 001</u> | Feb 28, 1995 |
| <u>AP</u> | | <u>10MG/VIAL</u> | <u>A063097 001</u> | May 21, 1990 |
| <u>AP</u> | | <u>20MG/VIAL</u> | <u>A063097 002</u> | May 21, 1990 |
| <u>AP</u> | | <u>50MG/VIAL</u> | <u>A063097 003</u> | May 21, 1990 |
| <u>AP</u> | | <u>200MG/100ML</u> | <u>A063336 004</u> | Feb 28, 1995 |
| <u>AP</u> | +! PHARMACIA AND UPJOHN | <u>2MG/ML</u> | <u>N050629 001</u> | Dec 23, 1987 |
| <u>AP</u> | +! | <u>200MG/100ML</u> | <u>N050629 002</u> | May 03, 1988 |
| <u>AP</u> | SAGENT PHARMS | <u>2MG/ML</u> | <u>A091495 001</u> | Mar 18, 2013 |
| <u>AP</u> | SUN PHARM INDS | <u>2MG/ML</u> | <u>A091418 001</u> | Feb 15, 2012 |
| <u>AP</u> | TEVA PHARMS USA | <u>2MG/ML</u> | <u>A064140 001</u> | Jul 28, 1995 |
| <u>AP</u> | | <u>200MG/100ML</u> | <u>A064140 002</u> | Jul 28, 1995 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>2MG/ML</u> | <u>A062975 001</u> | Mar 17, 1989 |
| <u>AP</u> | ! | <u>10MG/VIAL</u> | <u>A062921 001</u> | Mar 17, 1989 |
| <u>AP</u> | ! | <u>20MG/VIAL</u> | <u>A062921 002</u> | Mar 17, 1989 |
| <u>AP</u> | ! | <u>50MG/VIAL</u> | <u>A062921 003</u> | Mar 17, 1989 |
| <u>AP</u> | | <u>200MG/100ML</u> | <u>A064097 001</u> | Sep 13, 1994 |
| | + PHARMACIA AND UPJOHN | 150MG/75ML | N050629 003 | Mar 28, 2011 |

INJECTABLE, LIPOSOMAL; INJECTION

DOXIL (LIPOSOMAL)

| | | | | |
|-----------|-----------------------|---------------------------|--------------------|--------------|
| <u>AB</u> | + JANSSEN RES AND DEV | <u>20MG/10ML (2MG/ML)</u> | <u>N050718 001</u> | Nov 17, 1995 |
| <u>AB</u> | + | <u>50MG/25ML (2MG/ML)</u> | <u>N050718 002</u> | Jun 13, 2000 |

DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL)

| | | | | |
|-----------|--------------------|---------------------------|--------------------|--------------|
| <u>AB</u> | DR REDDYS LABS LTD | <u>20MG/10ML (2MG/ML)</u> | <u>A208657 001</u> | May 15, 2017 |
| <u>AB</u> | | <u>50MG/25ML (2MG/ML)</u> | <u>A208657 002</u> | May 15, 2017 |
| <u>AB</u> | ! | <u>20MG/10ML (2MG/ML)</u> | <u>A203263 001</u> | Feb 04, 2013 |
| <u>AB</u> | ! | <u>50MG/25ML (2MG/ML)</u> | <u>A203263 002</u> | Feb 04, 2013 |

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

| | | | | |
|-----------|--------------------|----------------------|--------------------|--------------|
| <u>AB</u> | ALEMBIC PHARMS LTD | <u>EQ 75MG BASE</u> | <u>A209165 001</u> | Jul 28, 2017 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A209165 002</u> | Jul 28, 2017 |
| <u>AB</u> | G AND W LABS INC | <u>EQ 50MG BASE</u> | <u>A204446 001</u> | May 28, 2015 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A204446 002</u> | May 28, 2015 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A204446 003</u> | May 28, 2015 |
| <u>AB</u> | IMPAX LABS INC | <u>EQ 150MG BASE</u> | <u>A200065 001</u> | Feb 17, 2011 |
| <u>AB</u> | LUPIN LTD | <u>EQ 50MG BASE</u> | <u>A204234 001</u> | Mar 05, 2014 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A204234 002</u> | Mar 05, 2014 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A204234 003</u> | Mar 05, 2014 |
| <u>AB</u> | MAYNE PHARMA INC | <u>EQ 50MG BASE</u> | <u>A209396 001</u> | Sep 29, 2017 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A209396 002</u> | Sep 29, 2017 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A209396 003</u> | Sep 29, 2017 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 150MG BASE</u> | <u>A202778 001</u> | Jun 08, 2012 |
| <u>AB</u> | PAR PHARM | <u>EQ 50MG BASE</u> | <u>A065055 001</u> | Dec 01, 2000 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A065055 002</u> | Dec 01, 2000 |
| <u>AB</u> | ! | <u>EQ 150MG BASE</u> | <u>A065055 003</u> | Jul 15, 2005 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>EQ 50MG BASE</u> | <u>A065053 001</u> | Nov 22, 2000 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A065053 003</u> | Sep 10, 2003 |

PRESCRIPTION DRUG PRODUCT LIST

DOXYCYCLINE

CAPSULE;ORAL

DOXYCYCLINE

| | | | | |
|-----------|-------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A065053 002</u> | Nov 22, 2000 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 50MG BASE</u> | <u>A205115 001</u> | Feb 18, 2016 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A205115 002</u> | Feb 18, 2016 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A205115 003</u> | Feb 18, 2016 |

MONODOX

| | | | | | |
|-----------|---------------------|------------------|----------------------|--------------------|--------------|
| <u>AB</u> | + | AQUA PHARMS | <u>EQ 50MG BASE</u> | <u>N050641 002</u> | Feb 10, 1992 |
| <u>AB</u> | + | | <u>EQ 75MG BASE</u> | <u>N050641 003</u> | Oct 18, 2006 |
| <u>AB</u> | + | | <u>EQ 100MG BASE</u> | <u>N050641 001</u> | Dec 29, 1989 |
| | ORACEA | | | | |
| | + | GALDERMA LABS LP | 40MG | N050805 001 | May 26, 2006 |
| | FOR SUSPENSION;ORAL | | | | |

DOXYCYCLINE

| | | | | | |
|-----------|--|--------------------|-------------------------|--------------------|--------------|
| <u>AB</u> | | CHARTWELL LIFE SCI | <u>EQ 25MG BASE/5ML</u> | <u>A065454 001</u> | Jul 16, 2008 |
| <u>AB</u> | | LUPIN LTD | <u>EQ 25MG BASE/5ML</u> | <u>A201678 001</u> | Mar 18, 2013 |

VIBRAMYCIN

| | | | | | |
|-----------|-------------|--------|-------------------------|--------------------|--|
| <u>AB</u> | + | PFIZER | <u>EQ 25MG BASE/5ML</u> | <u>N050006 001</u> | |
| | TABLET;ORAL | | | | |

DOXYCYCLINE

| | | | | | |
|-----------|---|-------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | HERITAGE PHARMS INC | <u>EQ 50MG BASE</u> | <u>A091605 001</u> | Dec 20, 2011 |
| <u>AB</u> | | | <u>EQ 75MG BASE</u> | <u>A091605 002</u> | Dec 20, 2011 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A091605 003</u> | Dec 20, 2011 |
| <u>AB</u> | | | <u>EQ 150MG BASE</u> | <u>A091605 004</u> | Dec 20, 2011 |
| <u>AB</u> | | LANNETT CO INC | <u>EQ 50MG BASE</u> | <u>A065285 001</u> | Dec 08, 2005 |
| <u>AB</u> | | | <u>EQ 75MG BASE</u> | <u>A065285 003</u> | Jul 30, 2008 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A065285 002</u> | Dec 08, 2005 |
| <u>AB</u> | | | <u>EQ 150MG BASE</u> | <u>A065285 004</u> | Jul 30, 2008 |
| <u>AB</u> | | MYLAN | <u>EQ 50MG BASE</u> | <u>A065377 001</u> | Nov 07, 2006 |
| <u>AB</u> | | | <u>EQ 75MG BASE</u> | <u>A065377 002</u> | Nov 07, 2006 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A065377 003</u> | Nov 07, 2006 |
| <u>AB</u> | | | <u>EQ 150MG BASE</u> | <u>A065427 001</u> | Jun 07, 2007 |
| <u>AB</u> | | PAR PHARM | <u>EQ 50MG BASE</u> | <u>A065070 001</u> | Dec 15, 2000 |
| <u>AB</u> | | | <u>EQ 75MG BASE</u> | <u>A065070 003</u> | Dec 30, 2002 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A065070 002</u> | Dec 15, 2000 |
| <u>AB</u> | ! | | <u>EQ 150MG BASE</u> | <u>A065070 004</u> | Jul 14, 2005 |
| <u>AB</u> | | SUN PHARM INDS LTD | <u>EQ 50MG BASE</u> | <u>A065356 001</u> | May 31, 2006 |
| <u>AB</u> | | | <u>EQ 75MG BASE</u> | <u>A065356 002</u> | May 31, 2006 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A065356 003</u> | May 31, 2006 |
| <u>AB</u> | | | <u>EQ 150MG BASE</u> | <u>A065356 004</u> | Jul 29, 2010 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>EQ 50MG BASE</u> | <u>A209582 001</u> | Sep 28, 2017 |
| <u>AB</u> | | | <u>EQ 75MG BASE</u> | <u>A209582 002</u> | Sep 28, 2017 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A209582 003</u> | Sep 28, 2017 |
| <u>AB</u> | | | <u>EQ 150MG BASE</u> | <u>A209582 004</u> | Sep 28, 2017 |

DOXYCYCLINE CALCIUM

SUSPENSION;ORAL

VIBRAMYCIN

| | | | | | |
|--|---|--------|------------------|-------------|--|
| | + | PFIZER | EQ 50MG BASE/5ML | N050480 001 | |
|--|---|--------|------------------|-------------|--|

DOXYCYCLINE HYCLATE

CAPSULE;ORAL

DOXYCYCLINE HYCLATE

| | | | | | |
|-----------|--|-------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS LABS FL INC | <u>EQ 50MG BASE</u> | <u>A062031 002</u> | Oct 13, 1982 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A062031 001</u> | |
| <u>AB</u> | | AJANTA PHARMA LTD | <u>EQ 50MG BASE</u> | <u>A211012 001</u> | Sep 24, 2018 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A211012 002</u> | Sep 24, 2018 |
| <u>AB</u> | | ALEMBIC PHARMS LTD | <u>EQ 50MG BASE</u> | <u>A210527 001</u> | Jun 13, 2018 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A210527 002</u> | Jun 13, 2018 |
| <u>AB</u> | | AMNEAL PHARMS | <u>EQ 100MG BASE</u> | <u>A207289 001</u> | Jun 27, 2016 |
| <u>AB</u> | | CHARTWELL LIFE SCI | <u>EQ 50MG BASE</u> | <u>A062500 001</u> | Sep 11, 1984 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A062500 002</u> | Sep 11, 1984 |
| <u>AB</u> | | HIKMA INTL PHARMS | <u>EQ 50MG BASE</u> | <u>A062396 002</u> | Nov 07, 1984 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A062396 001</u> | May 07, 1984 |
| <u>AB</u> | | MYLAN | <u>EQ 50MG BASE</u> | <u>A062337 001</u> | Mar 29, 1982 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A062337 002</u> | Mar 29, 1982 |
| <u>AB</u> | | SUN PHARM INDUSTRIES | <u>EQ 50MG BASE</u> | <u>A062676 002</u> | Jul 10, 1986 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A062676 001</u> | Jul 10, 1986 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>EQ 50MG BASE</u> | <u>A207774 001</u> | May 31, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

DOXYCYCLINE HYCLATE**AB** EQ 100MG BASE **A207774 002** May 31, 2018VIBRAMYCIN**AB** +! PFIZER EQ 100MG BASE **N050007 002**

INJECTABLE; INJECTION

DOXY 100**AP** ! FRESENIUS KABI USA EQ 100MG BASE/VIAL **A062475 001** Dec 09, 1983DOXY 200**AP** ! FRESENIUS KABI USA EQ 200MG BASE/VIAL **A062475 002** Dec 09, 1983DOXYCYCLINE**AP** MYLAN LABS LTD EQ 100MG BASE/VIAL **A091406 001** Aug 21, 2012**AP** ! WEST-WARD PHARMS EQ 100MG BASE/VIAL **A062569 001** Mar 09, 1988

INT

AP ZYDUS PHARMS USA EQ 100MG BASE/VIAL **A207757 001** Sep 28, 2017

INC

AP EQ 200MG BASE/VIAL **A207757 002** Sep 28, 2017

SYSTEM, EXTENDED RELEASE; PERIODONTAL

ATRIDOX

+! TOLMAR

50MG

N050751 001 Sep 03, 1998

TABLET; ORAL

ACTICLATE**AB** + AQUA PHARMS LLC EQ 75MG BASE **N205931 001** Jul 25, 2014**AB** +! EQ 150MG BASE **N205931 002** Jul 25, 2014DOXYCYCLINE HYCLATE**AB** ACTAVIS LABS FL INC EQ 100MG BASE **A062421 001** Feb 02, 1983**AB** AMNEAL PHARMS CO EQ 75MG BASE **A209372 001** Oct 06, 2017**AB** EQ 150MG BASE **A209372 002** Oct 06, 2017**AB** CARIBE HOLDINGS EQ 100MG BASE **A062269 002** Nov 08, 1982**AB** CHARTWELL LIFE SCI EQ 100MG BASE **A062505 001** Sep 11, 1984**AB** EMCURE PHARMS LTD EQ 100MG BASE **A209969 001** Nov 09, 2018**AB** ! HIKMA INTL PHARMS EQ 100MG BASE **A065095 001** Jul 02, 2003**AB** IVAX SUB TEVA EQ 20MG BASE **A065163 001** May 13, 2005

PHARMS

AB ! LANNETT CO INC EQ 20MG BASE **A065277 001** Nov 10, 2005**AB** LARKEN LABS EQ 20MG BASE **A065287 001** Feb 28, 2006**AB** LUPIN LTD EQ 75MG BASE **A208818 001** Sep 27, 2017**AB** EQ 150MG BASE **A208818 002** Sep 27, 2017**AB** MAYNE PHARMA INC EQ 75MG BASE **A208765 001** Jun 14, 2017**AB** EQ 150MG BASE **A208765 002** Jun 14, 2017**AB** MYLAN EQ 100MG BASE **A062432 001** Feb 15, 1983**AB** NOVEL LABS INC EQ 100MG BASE **A207558 001** Sep 06, 2017**AB** SUN PHARM EQ 20MG BASE **A065134 001** May 13, 2005

INDUSTRIES

AB EQ 100MG BASE **A062677 001** Jul 10, 1986**AB** ZYDUS PHARMS USA EQ 100MG BASE **A207773 001** Oct 30, 2017

INC

CARIBE HOLDINGS

EQ 50MG BASE

A062269 003 Oct 05, 1983

TABLET, DELAYED RELEASE; ORAL

DORYX**AB** + MAYNE PHARMA EQ 50MG BASE **N050795 006** Dec 19, 2014**AB** + EQ 75MG BASE **N050795 001** May 06, 2005**AB** + EQ 100MG BASE **N050795 002** May 06, 2005**AB** + EQ 150MG BASE **N050795 003** Jun 20, 2008**AB** +! EQ 200MG BASE **N050795 005** Apr 11, 2013DOXYCYCLINE HYCLATE**AB** ACTAVIS ELIZABETH EQ 50MG BASE **A090134 003** May 22, 2018**AB** EQ 75MG BASE **A090134 001** Dec 14, 2011**AB** EQ 100MG BASE **A090134 002** Dec 14, 2011**AB** EQ 200MG BASE **A090134 004** May 22, 2018**AB** HERITAGE PHARMS INC EQ 75MG BASE **A200856 001** Apr 30, 2013**AB** EQ 100MG BASE **A200856 002** Apr 30, 2013**AB** EQ 150MG BASE **A200856 003** Apr 30, 2013**AB** EQ 200MG BASE **A200856 004** Nov 13, 2018**AB** MYLAN EQ 50MG BASE **A090431 003** May 23, 2016**AB** EQ 75MG BASE **A090431 001** Dec 28, 2010**AB** EQ 100MG BASE **A090431 002** Dec 28, 2010**AB** EQ 200MG BASE **A090431 005** May 19, 2016**AB** MYLAN PHARMS INC EQ 150MG BASE **A091052 001** Feb 08, 2012**AB** PRINSTON INC EQ 150MG BASE **A207494 001** Nov 15, 2016**AB** EQ 200MG BASE **A207494 002** Nov 15, 2016**AB** ZYDUS PHARMS USA EQ 75MG BASE **A206772 001** Dec 21, 2018

PRESCRIPTION DRUG PRODUCT LIST

DOXYCYCLINE HYCLATE

TABLET, DELAYED RELEASE;ORAL

DOXYCYCLINE HYCLATE

INC

| | | | | |
|-----------|-----------------|-----------------------------|---------------------------|--------------|
| AB | | <u>EQ 100MG BASE</u> | <u>A206772 002</u> | Dec 21, 2018 |
| AB | | <u>EQ 150MG BASE</u> | <u>A206772 003</u> | Dec 21, 2018 |
| | DORYX MPC | | | |
| | +! MAYNE PHARMA | EQ 120MG BASE | N050795 008 | May 20, 2016 |

DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE

TABLET, DELAYED RELEASE;ORAL

DICLEGIS

| | | | | |
|-----------|--------------|-------------------------|---------------------------|--------------|
| AB | +! DUCHESNAY | <u>10MG;10MG</u> | <u>N021876 001</u> | Apr 08, 2013 |
|-----------|--------------|-------------------------|---------------------------|--------------|

DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE

| | | | | |
|-----------|---------------------|-------------------------|---------------------------|--------------|
| AB | ACTAVIS LABS FL INC | <u>10MG;10MG</u> | <u>A205811 001</u> | Aug 19, 2016 |
| AB | PAR PHARM INC | <u>10MG;10MG</u> | <u>A208518 001</u> | Dec 06, 2017 |

TABLET, EXTENDED RELEASE;ORAL

BONJESTA

| | | | | |
|--|--------------|-----------|-------------|--------------|
| | +! DUCHESNAY | 20MG;20MG | N209661 001 | Nov 07, 2016 |
|--|--------------|-----------|-------------|--------------|

DRONABINOL

CAPSULE;ORAL

DRONABINOL

| | | | | |
|-----------|----------------|---------------------|---------------------------|--------------|
| AB | AKORN INC | <u>2.5MG</u> | <u>A079217 001</u> | Jun 20, 2014 |
| AB | | <u>5MG</u> | <u>A079217 002</u> | Jun 20, 2014 |
| AB | | <u>10MG</u> | <u>A079217 003</u> | Jun 20, 2014 |
| AB | LANNETT CO INC | <u>2.5MG</u> | <u>A201463 001</u> | May 18, 2018 |
| AB | | <u>5MG</u> | <u>A201463 002</u> | May 18, 2018 |
| AB | | <u>10MG</u> | <u>A201463 003</u> | May 18, 2018 |
| AB | SVC PHARMA | <u>2.5MG</u> | <u>A078292 001</u> | Jun 27, 2008 |
| AB | | <u>5MG</u> | <u>A078292 002</u> | Jun 27, 2008 |
| AB | | <u>10MG</u> | <u>A078292 003</u> | Jun 27, 2008 |

MARINOL

| | | | | |
|-----------|-----------|---------------------|---------------------------|--------------|
| AB | + ABBEVIE | <u>2.5MG</u> | <u>N018651 001</u> | May 31, 1985 |
| AB | +! | <u>5MG</u> | <u>N018651 002</u> | May 31, 1985 |
| AB | + | <u>10MG</u> | <u>N018651 003</u> | May 31, 1985 |

SOLUTION;ORAL

SYNDROS

| | | | | |
|--|---------------------|--------|-------------|--------------|
| | +! INSYS DEV CO INC | 5MG/ML | N205525 001 | Mar 23, 2017 |
|--|---------------------|--------|-------------|--------------|

DRONEDARONE HYDROCHLORIDE

TABLET;ORAL

MULTAQ

| | | | | |
|--|----------------------|---------------|-------------|--------------|
| | +! SANOFI AVENTIS US | EQ 400MG BASE | N022425 001 | Jul 01, 2009 |
|--|----------------------|---------------|-------------|--------------|

DROPERIDOL

INJECTABLE; INJECTION

DROPERIDOL

| | | | | |
|-----------|--------------------|------------------------|---------------------------|--------------|
| AP | EUROHLTH INTL SARL | <u>2.5MG/ML</u> | <u>A208197 001</u> | Dec 14, 2017 |
| AP | HOSPIRA | <u>2.5MG/ML</u> | <u>A071981 001</u> | Feb 29, 1988 |
| AP | LUITPOLD | <u>2.5MG/ML</u> | <u>A072123 001</u> | Oct 24, 1988 |

INAPSINE

| | | | | |
|-----------|--------------|------------------------|---------------------------|--|
| AP | +! AKORN INC | <u>2.5MG/ML</u> | <u>N016796 001</u> | |
|-----------|--------------|------------------------|---------------------------|--|

DROSPIRENONE; ESTRADIOL

TABLET;ORAL

ANGELIQ

| | | | | |
|--|------------------|--------------|-------------|--------------|
| | + BAYER HLTHCARE | 0.25MG;0.5MG | N021355 001 | Feb 29, 2012 |
| | +! | 0.5MG;1MG | N021355 002 | Sep 28, 2005 |

DROSPIRENONE; ETHINYL ESTRADIOL

TABLET;ORAL

DROSPIRENONE AND ETHINYL ESTRADIOL

| | | | | |
|-----------|---------------------|--------------------------|---------------------------|--------------|
| AB | BARR | <u>3MG;0.02MG</u> | <u>A078515 001</u> | Mar 30, 2009 |
| AB | GLENMARK PHARMS LTD | <u>3MG;0.02MG</u> | <u>A204296 001</u> | Aug 17, 2015 |
| AB | JUBILANT CADISTA | <u>3MG;0.02MG</u> | <u>A209423 001</u> | Dec 22, 2017 |
| AB | MYLAN LABS LTD | <u>3MG;0.02MG</u> | <u>A202594 001</u> | Oct 22, 2015 |
| AB | PII | <u>3MG;0.02MG</u> | <u>A203291 001</u> | Jul 18, 2017 |
| AB | WATSON LABS | <u>3MG;0.02MG</u> | <u>A078833 001</u> | Nov 28, 2011 |

LO-ZUMANDIMINE

| | | | | |
|-----------|----------------------|--------------------------|---------------------------|--------------|
| AB | AUROBINDO PHARMA LTD | <u>3MG;0.02MG</u> | <u>A209632 001</u> | Feb 27, 2018 |
|-----------|----------------------|--------------------------|---------------------------|--------------|

LORYNA

| | | | | |
|-----------|-----------------|--------------------------|---------------------------|--------------|
| AB | LABS LEON FARMA | <u>3MG;0.02MG</u> | <u>A079221 001</u> | Mar 28, 2011 |
|-----------|-----------------|--------------------------|---------------------------|--------------|

PRESCRIPTION DRUG PRODUCT LISTDROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL

MELAMISA

| | | | | |
|--------------|--------------------------|-------------------|--------------------|--------------|
| AB | NOVAST LABS | 3MG;0.02MG | A202016 001 | Jan 26, 2016 |
| NIKKI | | | | |
| AB | LUPIN LTD | 3MG;0.02MG | A201661 001 | May 27, 2014 |
| YAZ | | | | |
| AB | +! BAYER HLTHCARE | 3MG;0.02MG | N021676 001 | Mar 16, 2006 |

TABLET; ORAL-28

DROSPIRENONE AND ETHINYL ESTRADIOL

| | | | | |
|--------------------|-----------------------------|-------------------|--------------------|--------------|
| AB | ACCORD HLTHCARE | 3MG;0.03MG | A207245 001 | Nov 22, 2016 |
| AB | APOTEX INC | 3MG;0.03MG | A205876 001 | Sep 21, 2016 |
| AB | BARR | 3MG;0.03MG | A077527 001 | May 09, 2008 |
| AB | GLENMARK PHARMS LTD | 3MG;0.03MG | A204848 001 | Mar 25, 2016 |
| AB | JUBILANT CADISTA | 3MG;0.03MG | A210017 001 | Sep 10, 2018 |
| AB | LUPIN LTD | 3MG;0.03MG | A201663 001 | Dec 18, 2012 |
| AB | MAYNE PHARMA | 3MG;0.03MG | A090081 001 | Sep 07, 2010 |
| AB | MYLAN LABS LTD | 3MG;0.03MG | A202131 001 | May 04, 2015 |
| SYEDA | | | | |
| AB | LABS LEON FARMA | 3MG;0.03MG | A090114 001 | Mar 28, 2011 |
| YAELA | | | | |
| AB | NOVAST LABS | 3MG;0.03MG | A202015 001 | Nov 19, 2014 |
| YASMIN | | | | |
| AB | +! BAYER HLTHCARE | 3MG;0.03MG | N021098 001 | May 11, 2001 |
| ZUMANDIMINE | | | | |
| AB | AUROBINDO PHARMA LTD | 3MG;0.03MG | A209407 001 | Mar 26, 2018 |

DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM

TABLET; ORAL

BEYAZ

| | | | | |
|-----------|-----------------------|-------------------------------------------|--------------------|--------------|
| AB | BAYER HLTHCARE | 3MG,N/A;0.02MG,N/A;0.451MG,0.451MG | N022532 001 | Sep 24, 2010 |
|-----------|-----------------------|-------------------------------------------|--------------------|--------------|

DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM

| | | | | |
|----------------|--------------------------|-------------------------------------------|--------------------|--------------|
| AB | LUPIN LTD | 3MG,N/A;0.02MG,N/A;0.451MG,0.451MG | A205947 001 | Jun 13, 2018 |
| AB | WATSON LABS INC | 3MG,N/A;0.02MG,N/A;0.451MG,0.451MG | A203593 001 | Oct 11, 2016 |
| AB | | 3MG,N/A;0.03MG,N/A;0.451MG,0.451MG | A203594 001 | Oct 11, 2016 |
| SAFYRAL | | | | |
| AB | +! BAYER HLTHCARE | 3MG,N/A;0.03MG,N/A;0.451MG,0.451MG | N022574 001 | Dec 16, 2010 |
| TYDEMY | | | | |
| AB | LUPIN LTD | 3MG,N/A;0.03MG,N/A;0.451MG,0.451MG | A205948 001 | Dec 12, 2017 |

DROXIDOPA

CAPSULE; ORAL

NORTHERA

| | | | | |
|-----------|------------------------|--------------|--------------------|--------------|
| + | LUNDBECK NA LTD | 100MG | N203202 001 | Feb 18, 2014 |
| + | | 200MG | N203202 002 | Feb 18, 2014 |
| +! | | 300MG | N203202 003 | Feb 18, 2014 |

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS; ORAL

CYMBALTA

| | | | | |
|-----------|----------------|---------------------|--------------------|--------------|
| AB | + LILLY | EQ 20MG BASE | N021427 001 | Aug 03, 2004 |
| AB | + | EQ 30MG BASE | N021427 002 | Aug 03, 2004 |
| AB | +! | EQ 60MG BASE | N021427 004 | Aug 03, 2004 |

DULOXETINE HYDROCHLORIDE

| | | | | |
|-----------|---------------------------|---------------------|--------------------|--------------|
| AB | ACTAVIS ELIZABETH | EQ 20MG BASE | A090776 001 | Dec 17, 2013 |
| AB | | EQ 30MG BASE | A090776 002 | Dec 17, 2013 |
| AB | | EQ 60MG BASE | A090776 003 | Dec 17, 2013 |
| AB | AJANTA PHARMA LTD | EQ 20MG BASE | A208706 001 | Jan 06, 2017 |
| AB | | EQ 30MG BASE | A208706 002 | Jan 06, 2017 |
| AB | | EQ 60MG BASE | A208706 003 | Jan 06, 2017 |
| AB | ALEMBIC PHARMS LTD | EQ 20MG BASE | A202949 001 | Jun 09, 2014 |
| AB | | EQ 30MG BASE | A202949 002 | Jun 09, 2014 |
| AB | | EQ 60MG BASE | A202949 003 | Jun 09, 2014 |
| AB | ALKEM LABS LTD | EQ 20MG BASE | A203197 001 | Aug 26, 2015 |
| AB | | EQ 30MG BASE | A203197 002 | Aug 26, 2015 |
| AB | | EQ 60MG BASE | A203197 003 | Aug 26, 2015 |
| AB | ANCHEN PHARMS | EQ 20MG BASE | A090780 001 | Oct 28, 2015 |
| AB | | EQ 30MG BASE | A090780 002 | Oct 28, 2015 |
| AB | | EQ 60MG BASE | A090780 003 | Oct 28, 2015 |
| AB | APOTEX INC | EQ 20MG BASE | A202045 001 | Jun 11, 2014 |
| AB | | EQ 30MG BASE | A202045 002 | Jun 11, 2014 |
| AB | | EQ 60MG BASE | A202045 003 | Jun 11, 2014 |
| AB | AUROBINDO PHARMA | EQ 20MG BASE | A090778 001 | Dec 11, 2013 |

PRESCRIPTION DRUG PRODUCT LIST

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS;ORAL

DULOXETINE HYDROCHLORIDE

LTD

| | | | | |
|-----------|-------------------------|---------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A090778 002</u> | Dec 11, 2013 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A090778 003</u> | Dec 11, 2013 |
| <u>AB</u> | BRECKENRIDGE PHARM | <u>EQ 20MG BASE</u> | <u>A203088 001</u> | Jun 11, 2014 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A203088 002</u> | Jun 11, 2014 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A203088 004</u> | May 18, 2018 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A203088 003</u> | Jun 11, 2014 |
| <u>AB</u> | CSPC OUYI PHARM CO | <u>EQ 20MG BASE</u> | <u>A211310 001</u> | Oct 16, 2018 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A211310 002</u> | Oct 16, 2018 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A211310 003</u> | Oct 16, 2018 |
| <u>AB</u> | HETERO LABS LTD III | <u>EQ 20MG BASE</u> | <u>A204343 001</u> | Aug 03, 2016 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A204343 002</u> | Aug 03, 2016 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A204343 003</u> | Aug 03, 2016 |
| <u>AB</u> | INVENTIA HLTHCARE | <u>EQ 20MG BASE</u> | <u>A202336 001</u> | Oct 28, 2015 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A202336 002</u> | Oct 28, 2015 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A202336 003</u> | Oct 28, 2015 |
| <u>AB</u> | LUPIN LTD | <u>EQ 20MG BASE</u> | <u>A090694 001</u> | Dec 11, 2013 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A090694 002</u> | Dec 11, 2013 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A090694 003</u> | Dec 11, 2013 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A090694 004</u> | Dec 11, 2013 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>EQ 20MG BASE</u> | <u>A204815 001</u> | Mar 23, 2017 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A204815 002</u> | Mar 23, 2017 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A204815 003</u> | Mar 23, 2017 |
| <u>AB</u> | MARKSANS PHARMA | <u>EQ 20MG BASE</u> | <u>A090723 001</u> | Dec 11, 2013 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A090723 002</u> | Dec 11, 2013 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A090723 003</u> | Dec 11, 2013 |
| <u>AB</u> | PRINSTON INC | <u>EQ 20MG BASE</u> | <u>A206653 001</u> | May 18, 2017 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A206653 002</u> | May 18, 2017 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A206653 003</u> | May 18, 2017 |
| <u>AB</u> | SUN PHARMA GLOBAL | <u>EQ 20MG BASE</u> | <u>A090745 001</u> | Dec 11, 2013 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A090745 002</u> | Dec 11, 2013 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A090745 003</u> | Dec 11, 2013 |
| <u>AB</u> | TEVA PHARMS USA | <u>EQ 20MG BASE</u> | <u>A090783 001</u> | Dec 11, 2013 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A090783 002</u> | Dec 11, 2013 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A090783 003</u> | Dec 11, 2013 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>EQ 20MG BASE</u> | <u>A090774 001</u> | Dec 11, 2013 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A090774 002</u> | Dec 11, 2013 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A090774 003</u> | Dec 11, 2013 |
| <u>AB</u> | ZYDUS HLTHCARE | <u>EQ 20MG BASE</u> | <u>A090739 001</u> | Jan 08, 2014 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A090739 002</u> | Jan 08, 2014 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A090739 003</u> | Jan 08, 2014 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 20MG BASE</u> | <u>A090728 001</u> | Jan 08, 2014 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A090728 002</u> | Jan 08, 2014 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A090728 003</u> | Jan 08, 2014 |

DUTASTERIDE

CAPSULE;ORAL

AVODART

| | | | | | |
|-----------|----|-----------------|--------------|--------------------|--------------|
| <u>AB</u> | +! | GLAXOSMITHKLINE | <u>0.5MG</u> | <u>N021319 001</u> | Nov 20, 2001 |
|-----------|----|-----------------|--------------|--------------------|--------------|

DUTASTERIDE

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS LABS FL INC | <u>0.5MG</u> | <u>A202808 001</u> | Nov 20, 2015 |
| <u>AB</u> | AMNEAL PHARMS | <u>0.5MG</u> | <u>A203118 001</u> | Nov 20, 2015 |
| <u>AB</u> | APOTEX INC | <u>0.5MG</u> | <u>A204292 001</u> | Nov 24, 2015 |
| <u>AB</u> | ASCENT PHARMS INC | <u>0.5MG</u> | <u>A206574 001</u> | Oct 21, 2016 |
| <u>AB</u> | AUROLIFE PHARMA LLC | <u>0.5MG</u> | <u>A202660 001</u> | Nov 20, 2015 |
| <u>AB</u> | BARR | <u>0.5MG</u> | <u>A090095 001</u> | Dec 21, 2010 |
| <u>AB</u> | BIONPHARMA INC | <u>0.5MG</u> | <u>A200899 001</u> | Nov 20, 2015 |
| <u>AB</u> | BRECKENRIDGE PHARM | <u>0.5MG</u> | <u>A204705 001</u> | Nov 20, 2015 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A208227 001</u> | Jun 22, 2018 |
| <u>AB</u> | HERITAGE PHARMS INC | <u>0.5MG</u> | <u>A207935 001</u> | Oct 13, 2017 |
| <u>AB</u> | HUMANWELL PURACAP | <u>0.5MG</u> | <u>A209909 001</u> | Nov 21, 2017 |
| <u>AB</u> | INTERGEL PHARMS INC | <u>0.5MG</u> | <u>A206373 001</u> | Mar 17, 2016 |
| <u>AB</u> | MARKSANS PHARMA | <u>0.5MG</u> | <u>A204376 001</u> | Apr 07, 2017 |
| <u>AB</u> | RISING PHARMS | <u>0.5MG</u> | <u>A202530 001</u> | Nov 20, 2015 |
| <u>AB</u> | STRIDES PHARMA | <u>0.5MG</u> | <u>A204262 001</u> | Nov 20, 2015 |
| <u>AB</u> | VINTAGE PHARMS LLC | <u>0.5MG</u> | <u>A202421 001</u> | Nov 20, 2015 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>0.5MG</u> | <u>A202204 001</u> | Nov 23, 2015 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>0.5MG</u> | <u>A204373 001</u> | Oct 04, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

DUTASTERIDE

CAPSULE; ORAL

DUTASTERIDE

INC

DUTASTERIDE; TAMSULOSIN HYDROCHLORIDE

CAPSULE; ORAL

DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|--------------------|--------------------|--------------|
| AB | ACTAVIS LABS FL INC | 0.5MG;0.4MG | A202975 001 | Nov 20, 2015 |
| AB | ANCHEN PHARMS | 0.5MG;0.4MG | A202509 001 | Feb 26, 2014 |
| AB | ZYDUS PHARMS USA INC | 0.5MG;0.4MG | A207769 001 | May 24, 2018 |

JALYN

| | | | | | |
|-----------|----------|-----------------|--------------------|--------------------|--------------|
| AB | + | GLAXOSMITHKLINE | 0.5MG;0.4MG | N022460 001 | Jun 14, 2010 |
|-----------|----------|-----------------|--------------------|--------------------|--------------|

DUVELISIB

CAPSULE; ORAL

COPIKTRA

| | | | | | |
|--|----------|--------------|------|-------------|--------------|
| | + | VERASTEM INC | 15MG | N211155 001 | Sep 24, 2018 |
| | + | ! | 25MG | N211155 002 | Sep 24, 2018 |

DYCLONINE HYDROCHLORIDE

SOLUTION; TOPICAL

DYCLOPRO

| | | | | | |
|--|----------|-------------|------|-------------|--------------|
| | ! | NOVOCOL INC | 0.5% | A200480 001 | Nov 20, 2018 |
| | ! | | 1% | A200480 002 | Nov 20, 2018 |

ECHOTHIOPHATE IODIDE

FOR SOLUTION; OPHTHALMIC

PHOSPHOLINE IODIDE

| | | | | | |
|--|----------|--------------|--------|-------------|--|
| | + | WYETH PHARMS | 0.125% | N011963 001 | |
|--|----------|--------------|--------|-------------|--|

ECONAZOLE NITRATE

AEROSOL, FOAM; TOPICAL

ECOZA

| | | | | | |
|--|----------|----------|----|-------------|--------------|
| | + | GLENMARK | 1% | N205175 001 | Oct 24, 2013 |
|--|----------|----------|----|-------------|--------------|

CREAM; TOPICAL

ECONAZOLE NITRATE

| | | | | | |
|-----------|----------|----------------------|-----------|--------------------|--------------|
| AB | | CASI PHARMS INC | 1% | A076075 001 | Nov 26, 2002 |
| AB | | MYLAN PHARMS INC | 1% | A210364 001 | Apr 18, 2018 |
| AB | ! | PERRIGO NEW YORK | 1% | A076479 001 | Jun 23, 2004 |
| AB | | TARO | 1% | A076005 001 | Nov 26, 2002 |
| AB | | TELLIGENT PHARMA INC | 1% | A076574 001 | Dec 17, 2004 |

SPECTAZOLE

| | | | | | |
|-----------|----------|---------------|-----------|--------------------|--------------|
| AB | + | ALVOGEN MALTA | 1% | N018751 001 | Dec 23, 1982 |
|-----------|----------|---------------|-----------|--------------------|--------------|

EDARAVONE

SOLUTION; INTRAVENOUS

RADICAVA

| | | | | | |
|--|----------|-------------------|-----------------------|-------------|--------------|
| | + | MITSUBISHI TANABE | 30MG/100ML (0.3MG/ML) | N209176 001 | May 05, 2017 |
| | + | | 60MG/100ML (0.6MG/ML) | N209176 002 | Nov 15, 2018 |

EDETATE CALCIUM DISODIUM

INJECTABLE; INJECTION

CALCIUM DISODIUM VERSENATE

| | | | | | |
|--|----------|---------|----------|-------------|--|
| | + | MEDICIS | 200MG/ML | N008922 001 | |
|--|----------|---------|----------|-------------|--|

EDOXYBAN TOSYLATE

TABLET; ORAL

SAVAYSA

| | | | | | |
|--|----------|--------------------|--------------|-------------|--------------|
| | + | DAIICHI SANKYO INC | EQ 15MG BASE | N206316 001 | Jan 08, 2015 |
| | + | | EQ 30MG BASE | N206316 002 | Jan 08, 2015 |
| | + | | EQ 60MG BASE | N206316 003 | Jan 08, 2015 |

EFAVIRENZ

CAPSULE; ORAL

EFAVIRENZ

| | | | | | |
|-----------|--|-------------------------|-------------|--------------------|--------------|
| AB | | AUROBINDO PHARMA LTD | 50MG | A078064 001 | Dec 15, 2017 |
|-----------|--|-------------------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 200MG | A078064 003 | Dec 15, 2017 |
|-----------|--|--|--------------|--------------------|--------------|

SUSTIVA

| | | | | | |
|-----------|----------|-------------------------|-------------|--------------------|--------------|
| AB | + | BRISTOL MYERS SQUIBB | 50MG | N020972 001 | Sep 17, 1998 |
|-----------|----------|-------------------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|----------|---|--------------|--------------------|--------------|
| AB | + | ! | 200MG | N020972 003 | Sep 17, 1998 |
|-----------|----------|---|--------------|--------------------|--------------|

EFAVIRENZ

| | | | | | |
|--|--|-------------------------|-------|-------------|--------------|
| | | AUROBINDO PHARMA LTD | 100MG | A078064 002 | Dec 15, 2017 |
|--|--|-------------------------|-------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

EFAVIRENZ

TABLET;ORAL

EFAVIRENZ

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| AB | AUROBINDO PHARMA LTD | 600MG | A077673 001 | Sep 21, 2018 |
| AB | | 600MG | A205322 001 | Aug 30, 2018 |
| AB | CIPLA | 600MG | A204766 001 | Jun 15, 2018 |
| AB | HETERO LABS LTD III | 600MG | A078886 001 | Apr 27, 2018 |
| AB | MYLAN PHARMS INC | 600MG | A091471 001 | Feb 17, 2016 |
| AB | STRIDES PHARMA | 600MG | A204869 001 | Mar 12, 2018 |

SUSTIVA

| | | | | |
|-----------|---------------------------------|--------------|--------------------|--------------|
| AB | + ! BRISTOL MYERS SQUIBB | 600MG | N021360 002 | Feb 01, 2002 |
|-----------|---------------------------------|--------------|--------------------|--------------|

EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

ATRIPLA

| | | | | |
|-----------|----------------------------|--------------------------|--------------------|--------------|
| AB | + ! GILEAD SCIENCES | 600MG;200MG;300MG | N021937 001 | Jul 12, 2006 |
|-----------|----------------------------|--------------------------|--------------------|--------------|

EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE

| | | | | |
|-----------|----------------------|--------------------------|--------------------|--------------|
| AB | AUROBINDO PHARMA LTD | 600MG;200MG;300MG | A203041 001 | Sep 04, 2018 |
| AB | TEVA PHARMS USA | 600MG;200MG;300MG | A091215 001 | Nov 09, 2018 |

EFAVIRENZ; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

SYMFI

| | | | | |
|------------|----------------|-------------------|-------------|--------------|
| + ! | MYLAN LABS LTD | 600MG;300MG;300MG | N022142 001 | Mar 22, 2018 |
|------------|----------------|-------------------|-------------|--------------|

SYMFI LO

| | | | | |
|------------|------------------|-------------------|-------------|--------------|
| + ! | MYLAN PHARMS INC | 400MG;300MG;300MG | N208255 001 | Feb 05, 2018 |
|------------|------------------|-------------------|-------------|--------------|

EFINACONAZOLE

SOLUTION;TOPICAL

JUBLIA

| | | | | |
|------------|---------------------|-----|-------------|--------------|
| + ! | VALEANT PHARMS INTL | 10% | N203567 001 | Jun 06, 2014 |
|------------|---------------------|-----|-------------|--------------|

EFLORNITHINE HYDROCHLORIDE

CREAM;TOPICAL

VANIQA

| | | | | |
|------------|------------|-------|-------------|--------------|
| + ! | SKINMEDICA | 13.9% | N021145 001 | Jul 27, 2000 |
|------------|------------|-------|-------------|--------------|

ELAGOLIX SODIUM

TABLET;ORAL

ORILISSA

| | | | | |
|----------|------------|---------------|-------------|--------------|
| + | ABBVIE INC | EQ 150MG BASE | N210450 001 | Jul 23, 2018 |
|----------|------------|---------------|-------------|--------------|

| | | | | |
|------------|--|---------------|-------------|--------------|
| + ! | | EQ 200MG BASE | N210450 002 | Jul 23, 2018 |
|------------|--|---------------|-------------|--------------|

ELBASVIR; GRAZOPREVIR

TABLET;ORAL

ZEPATIER

| | | | | |
|------------|-------------------|------------|-------------|--------------|
| + ! | MERCK SHARP DOHME | 50MG;100MG | N208261 001 | Jan 28, 2016 |
|------------|-------------------|------------|-------------|--------------|

ELETRIPTAN HYDROBROMIDE

TABLET;ORAL

ELETRIPTAN HYDROBROMIDE

| | | | | | |
|-----------|----------------------|---------------------|---------------------|--------------------|--------------|
| AB | AJANTA PHARMA LTD | EQ 20MG BASE | A205186 001 | Aug 29, 2017 | |
| AB | | EQ 40MG BASE | A205186 002 | Aug 29, 2017 | |
| AB | AMNEAL PHARMS CO | EQ 20MG BASE | A206787 001 | May 25, 2018 | |
| AB | | EQ 40MG BASE | A206787 002 | May 25, 2018 | |
| AB | AUROBINDO PHARMA LTD | EQ 20MG BASE | A210708 001 | Jan 15, 2019 | |
| AB | | EQ 40MG BASE | A210708 002 | Jan 15, 2019 | |
| AB | MYLAN PHARMS INC | EQ 20MG BASE | A205152 001 | Aug 11, 2017 | |
| AB | | EQ 40MG BASE | A205152 002 | Aug 11, 2017 | |
| AB | TEVA PHARMS USA | EQ 20MG BASE | A202040 001 | Jun 27, 2017 | |
| AB | | EQ 40MG BASE | A202040 002 | Jun 27, 2017 | |
| AB | ZYDUS PHARMS USA INC | EQ 20MG BASE | A206409 001 | Jun 16, 2017 | |
| AB | | EQ 40MG BASE | A206409 002 | Jun 16, 2017 | |
| | RELPAK | | | | |
| AB | + | PFIZER IRELAND | EQ 20MG BASE | N021016 001 | Dec 26, 2002 |
| AB | + ! | | EQ 40MG BASE | N021016 002 | Dec 26, 2002 |

PRESCRIPTION DRUG PRODUCT LIST

ELIGLUSTAT TARTRATE

CAPSULE; ORAL

CERDELGA

+! GENZYME CORP EQ 84MG BASE N205494 001 Aug 19, 2014

ELTROMBOPAG OLAMINE

FOR SUSPENSION; ORAL

PROMACTA KIT

+ NOVARTIS PHARMS EQ 12.5MG ACID/PACKET N207027 002 Sep 27, 2018

CORP

+! EQ 25MG ACID/PACKET N207027 001 Aug 24, 2015

TABLET; ORAL

PROMACTA

+ NOVARTIS PHARMS EQ 12.5MG ACID N022291 004 Oct 20, 2011

CORP

+ EQ 25MG ACID N022291 001 Nov 20, 2008

+ EQ 50MG ACID N022291 002 Nov 20, 2008

+! EQ 75MG ACID N022291 003 Sep 08, 2009

+! EQ 100MG ACID N022291 005 Nov 16, 2012

ELUXADOLINE

TABLET; ORAL

VIBERZI

+ ALLERGAN HOLDINGS 75MG N206940 001 May 27, 2015

+! 100MG N206940 002 May 27, 2015

EMEDASTINE DIFUMARATE

SOLUTION/DROPS; OPHTHALMIC

EMADINE

+! NOVARTIS PHARMS 0.05% N020706 001 Dec 29, 1997

CORP

EMPAGLIFLOZIN

TABLET; ORAL

JARDIANCE

+ BOEHRINGER 10MG N204629 001 Aug 01, 2014

INGELHEIM

+! 25MG N204629 002 Aug 01, 2014

EMPAGLIFLOZIN; LINAGLIPTIN

TABLET; ORAL

GLYXAMBI

+ BOEHRINGER 10MG; 5MG N206073 001 Jan 30, 2015

INGELHEIM

+! 25MG; 5MG N206073 002 Jan 30, 2015

EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

SYNJARDY

+ BOEHRINGER 5MG; 500MG N206111 001 Aug 26, 2015

INGELHEIM

+ 5MG; 1GM N206111 002 Aug 26, 2015

+ 12.5MG; 500MG N206111 003 Aug 26, 2015

+! 12.5MG; 1GM N206111 004 Aug 26, 2015

TABLET, EXTENDED RELEASE; ORAL

SYNJARDY XR

+ BOEHRINGER 5MG; 1GM N208658 001 Dec 09, 2016

INGELHEIM

+ 10MG; 1GM N208658 002 Dec 09, 2016

+ 12.5MG; 1GM N208658 003 Dec 09, 2016

+! 25MG; 1GM N208658 004 Dec 09, 2016

EMTRICITABINE

CAPSULE; ORAL

EMTRICITABINE**AB** CIPLA **200MG** **A091168 001** Jul 02, 2018**EMTRIVA****AB** +! GILEAD **200MG** **N021500 001** Jul 02, 2003

SOLUTION; ORAL

EMTRIVA

+! GILEAD 10MG/ML N021896 001 Sep 28, 2005

PRESCRIPTION DRUG PRODUCT LIST

EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

ODEFSEY

+! GILEAD SCIENCES INC 200MG;EQ 25MG BASE;EQ 25MG BASE N208351 001 Mar 01, 2016

EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

COMPLERA

+! GILEAD SCIENCES INC 200MG;EQ 25MG BASE;300MG N202123 001 Aug 10, 2011

EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

DESCOVY

+! GILEAD SCIENCES INC 200MG;EQ 25MG BASE N208215 001 Apr 04, 2016

EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE

| | | | | |
|----------------|----------------------|--------------------|--------------------|--------------|
| <u>AB</u> | AMNEAL PHARMS CO | <u>100MG;150MG</u> | <u>A209721 001</u> | Aug 22, 2018 |
| <u>AB</u> | | <u>133MG;200MG</u> | <u>A209721 002</u> | Aug 22, 2018 |
| <u>AB</u> | | <u>167MG;250MG</u> | <u>A209721 003</u> | Aug 22, 2018 |
| <u>AB</u> | | <u>200MG;300MG</u> | <u>A209721 004</u> | Aug 22, 2018 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>200MG;300MG</u> | <u>A090513 001</u> | Jan 26, 2018 |
| <u>AB</u> | MYLAN PHARMS INC | <u>200MG;300MG</u> | <u>A206436 001</u> | Apr 09, 2018 |
| <u>AB</u> | TEVA PHARMS USA | <u>200MG;300MG</u> | <u>A090894 001</u> | Jun 08, 2017 |
| <u>TRUVADA</u> | | | | |
| <u>AB</u> | + GILEAD | <u>100MG;150MG</u> | <u>N021752 002</u> | Mar 10, 2016 |
| <u>AB</u> | + | <u>133MG;200MG</u> | <u>N021752 003</u> | Mar 10, 2016 |
| <u>AB</u> | + | <u>167MG;250MG</u> | <u>N021752 004</u> | Mar 10, 2016 |
| <u>AB</u> | +! | <u>200MG;300MG</u> | <u>N021752 001</u> | Aug 02, 2004 |

ENALAPRIL MALEATE

FOR SOLUTION;ORAL

EPANED KIT

+! SILVERGATE PHARMS 1MG/ML N204308 001 Aug 13, 2013

SOLUTION;ORAL

EPANED

+! SILVERGATE PHARMS 1MG/ML N208686 001 Sep 20, 2016

TABLET;ORAL

ENALAPRIL MALEATE

| | | | | |
|----------------|------------------------|--------------|--------------------|--------------|
| <u>AB</u> | APOTEX | <u>2.5MG</u> | <u>A075178 002</u> | Mar 23, 2001 |
| <u>AB</u> | | <u>5MG</u> | <u>A075178 001</u> | Mar 23, 2001 |
| <u>AB</u> | | <u>10MG</u> | <u>A075178 003</u> | Mar 23, 2001 |
| <u>AB</u> | | <u>20MG</u> | <u>A075178 004</u> | Mar 23, 2001 |
| <u>AB</u> | MYLAN | <u>5MG</u> | <u>A075480 002</u> | Aug 22, 2000 |
| <u>AB</u> | | <u>10MG</u> | <u>A075480 003</u> | Aug 22, 2000 |
| <u>AB</u> | SANDOZ INC | <u>2.5MG</u> | <u>A075496 001</u> | Aug 22, 2000 |
| <u>AB</u> | | <u>5MG</u> | <u>A075496 002</u> | Aug 22, 2000 |
| <u>AB</u> | | <u>10MG</u> | <u>A075459 001</u> | Aug 22, 2000 |
| <u>AB</u> | | <u>20MG</u> | <u>A075459 002</u> | Aug 22, 2000 |
| <u>AB</u> | TARO | <u>2.5MG</u> | <u>A075657 001</u> | Jan 23, 2001 |
| <u>AB</u> | | <u>5MG</u> | <u>A075657 002</u> | Jan 23, 2001 |
| <u>AB</u> | | <u>10MG</u> | <u>A075657 003</u> | Jan 23, 2001 |
| <u>AB</u> | | <u>20MG</u> | <u>A075657 004</u> | Jan 23, 2001 |
| <u>AB</u> | TEVA | <u>2.5MG</u> | <u>A075479 001</u> | Aug 22, 2000 |
| <u>AB</u> | | <u>5MG</u> | <u>A075479 002</u> | Aug 22, 2000 |
| <u>AB</u> | | <u>10MG</u> | <u>A075479 003</u> | Aug 22, 2000 |
| <u>AB</u> | | <u>20MG</u> | <u>A075479 004</u> | Aug 22, 2000 |
| <u>AB</u> | WOCKHARDT LTD | <u>2.5MG</u> | <u>A075483 001</u> | Aug 22, 2000 |
| <u>AB</u> | | <u>5MG</u> | <u>A075483 002</u> | Aug 22, 2000 |
| <u>AB</u> | | <u>10MG</u> | <u>A075483 003</u> | Aug 22, 2000 |
| <u>AB</u> | | <u>20MG</u> | <u>A075483 004</u> | Aug 22, 2000 |
| <u>VASOTEC</u> | | | | |
| <u>AB</u> | + VALEANT PHARMS NORTH | <u>2.5MG</u> | <u>N018998 005</u> | Jul 26, 1988 |
| <u>AB</u> | + | <u>5MG</u> | <u>N018998 001</u> | Dec 24, 1985 |
| <u>AB</u> | + | <u>10MG</u> | <u>N018998 002</u> | Dec 24, 1985 |
| <u>AB</u> | +! | <u>20MG</u> | <u>N018998 003</u> | Dec 24, 1985 |

PRESCRIPTION DRUG PRODUCT LIST

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE

| | | | | |
|------------------|--------------------|-------------------|--------------------|--------------|
| <u>AB</u> | APOTEX INC | <u>5MG;12.5MG</u> | <u>A076486 001</u> | Oct 27, 2004 |
| <u>AB</u> | | <u>10MG;25MG</u> | <u>A076486 002</u> | Oct 27, 2004 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>5MG;12.5MG</u> | <u>A075909 001</u> | Oct 15, 2001 |
| <u>AB</u> | | <u>10MG;25MG</u> | <u>A075909 002</u> | Oct 15, 2001 |
| <u>AB</u> | G AND W LABS INC | <u>5MG;12.5MG</u> | <u>A075727 001</u> | Sep 18, 2001 |
| <u>AB</u> | | <u>10MG;25MG</u> | <u>A075727 002</u> | Sep 18, 2001 |
| <u>AB</u> | MYLAN | <u>5MG;12.5MG</u> | <u>A075624 001</u> | Sep 18, 2001 |
| <u>AB</u> | | <u>10MG;25MG</u> | <u>A075624 002</u> | Sep 18, 2001 |
| <u>AB</u> | TARO PHARM INDS | <u>5MG;12.5MG</u> | <u>A075788 001</u> | Sep 18, 2001 |
| <u>AB</u> | | <u>10MG;25MG</u> | <u>A075788 002</u> | Sep 18, 2001 |
| <u>VASERETIC</u> | | | | |
| <u>AB</u> | + VALEANT INTL | <u>5MG;12.5MG</u> | <u>N019221 003</u> | Jul 12, 1995 |
| <u>AB</u> | +! | <u>10MG;25MG</u> | <u>N019221 001</u> | Oct 31, 1986 |

ENALAPRILAT

INJECTABLE; INJECTION

ENALAPRILAT

| | | | | |
|-----------|--------------------|------------------|--------------------|--------------|
| <u>AP</u> | ! ATHENEX INC | <u>1.25MG/ML</u> | <u>A075634 001</u> | Aug 22, 2000 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>1.25MG/ML</u> | <u>A078687 001</u> | Dec 23, 2008 |
| <u>AP</u> | ! HOSPIRA | <u>1.25MG/ML</u> | <u>A075458 001</u> | Aug 22, 2000 |
| <u>AP</u> | TEVA PHARMS USA | <u>1.25MG/ML</u> | <u>A075578 001</u> | Aug 22, 2000 |

ENASIDENIB MESYLATE

TABLET; ORAL

IDHIFA

| | | | | |
|---|--------------|---------------|-------------|--------------|
| + | CELGENE CORP | EQ 50MG BASE | N209606 001 | Aug 01, 2017 |
| + | ! | EQ 100MG BASE | N209606 002 | Aug 01, 2017 |

ENCORAFENIB

CAPSULE; ORAL

BRAFTOVI

| | | | | |
|---|---------------------|------|-------------|--------------|
| + | ARRAY BIOPHARMA INC | 50MG | N210496 001 | Jun 27, 2018 |
| + | ! | 75MG | N210496 002 | Jun 27, 2018 |

ENFUVRTIDE

INJECTABLE; SUBCUTANEOUS

FUZEON

| | | | | |
|---|---------|-----------|-------------|--------------|
| + | ! ROCHE | 90MG/VIAL | N021481 001 | Mar 13, 2003 |
|---|---------|-----------|-------------|--------------|

ENOXAPARIN SODIUM

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

ENOXAPARIN SODIUM

| | | | | |
|-----------|------------|-----------------------------|--------------------|--------------|
| <u>AB</u> | SANDOZ INC | <u>300MG/3ML (100MG/ML)</u> | <u>A078660 001</u> | Nov 28, 2011 |
|-----------|------------|-----------------------------|--------------------|--------------|

LOVENOX

| | | | | |
|-----------|---------------------|-----------------------------|--------------------|--------------|
| <u>AB</u> | + SANOFI AVENTIS US | <u>300MG/3ML (100MG/ML)</u> | <u>N020164 009</u> | Jan 23, 2003 |
|-----------|---------------------|-----------------------------|--------------------|--------------|

INJECTABLE; SUBCUTANEOUS

ENOXAPARIN SODIUM (PRESERVATIVE FREE)

| | | | | |
|-----------|-----------------|-------------------------------|--------------------|--------------|
| <u>AP</u> | AMPHASTAR PHARM | <u>30MG/0.3ML (100MG/ML)</u> | <u>A076684 001</u> | Sep 19, 2011 |
| <u>AP</u> | | <u>40MG/0.4ML (100MG/ML)</u> | <u>A076684 002</u> | Sep 19, 2011 |
| <u>AP</u> | | <u>60MG/0.6ML (100MG/ML)</u> | <u>A076684 003</u> | Sep 19, 2011 |
| <u>AP</u> | | <u>80MG/0.8ML (100MG/ML)</u> | <u>A076684 004</u> | Sep 19, 2011 |
| <u>AP</u> | | <u>100MG/ML (100MG/ML)</u> | <u>A076684 005</u> | Sep 19, 2011 |
| <u>AP</u> | | <u>120MG/0.8ML (150MG/ML)</u> | <u>A076684 006</u> | Sep 19, 2011 |
| <u>AP</u> | | <u>150MG/ML (150MG/ML)</u> | <u>A076684 007</u> | Sep 19, 2011 |
| <u>AP</u> | APOTEX INC | <u>30MG/0.3ML (100MG/ML)</u> | <u>A078990 001</u> | Sep 28, 2018 |
| <u>AP</u> | | <u>40MG/0.4ML (100MG/ML)</u> | <u>A078990 002</u> | Sep 28, 2018 |
| <u>AP</u> | | <u>60MG/0.6ML (100MG/ML)</u> | <u>A078990 003</u> | Sep 28, 2018 |
| <u>AP</u> | | <u>80MG/0.8ML (100MG/ML)</u> | <u>A078990 004</u> | Sep 28, 2018 |
| <u>AP</u> | | <u>100MG/ML (100MG/ML)</u> | <u>A078990 005</u> | Sep 28, 2018 |
| <u>AP</u> | | <u>120MG/0.8ML (150MG/ML)</u> | <u>A078990 006</u> | Sep 28, 2018 |
| <u>AP</u> | | <u>150MG/ML (150MG/ML)</u> | <u>A078990 007</u> | Sep 28, 2018 |
| <u>AP</u> | SANDOZ | <u>30MG/0.3ML (100MG/ML)</u> | <u>A077857 002</u> | Jul 23, 2010 |
| <u>AP</u> | | <u>40MG/0.4ML (100MG/ML)</u> | <u>A077857 003</u> | Jul 23, 2010 |
| <u>AP</u> | | <u>60MG/0.6ML (100MG/ML)</u> | <u>A077857 004</u> | Jul 23, 2010 |
| <u>AP</u> | | <u>80MG/0.8ML (100MG/ML)</u> | <u>A077857 005</u> | Jul 23, 2010 |
| <u>AP</u> | | <u>100MG/ML (100MG/ML)</u> | <u>A077857 001</u> | Jul 23, 2010 |
| <u>AP</u> | | <u>120MG/0.8ML (150MG/ML)</u> | <u>A077857 006</u> | Jul 23, 2010 |
| <u>AP</u> | | <u>150MG/ML (150MG/ML)</u> | <u>A077857 007</u> | Jul 23, 2010 |
| <u>AP</u> | TEVA | <u>30MG/0.3ML (100MG/ML)</u> | <u>A076726 001</u> | Jun 23, 2014 |
| <u>AP</u> | | <u>40MG/0.4ML (100MG/ML)</u> | <u>A076726 002</u> | Jun 23, 2014 |
| <u>AP</u> | | <u>60MG/0.6ML (100MG/ML)</u> | <u>A076726 003</u> | Jun 23, 2014 |
| <u>AP</u> | | <u>80MG/0.8ML (100MG/ML)</u> | <u>A076726 004</u> | Jun 23, 2014 |

PRESCRIPTION DRUG PRODUCT LIST

ENOXAPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

ENOXAPARIN SODIUM (PRESERVATIVE FREE)

| | | | | |
|------------------------------------|---|-------------------------------|-------------------------------|---------------------------------|
| <u>AP</u> | | <u>100MG/ML (100MG/ML)</u> | <u>A076726 005</u> | Jun 23, 2014 |
| <u>AP</u> | | <u>120MG/0.8ML (150MG/ML)</u> | <u>A076726 006</u> | Jun 23, 2014 |
| <u>AP</u> | | <u>150MG/ML (150MG/ML)</u> | <u>A076726 007</u> | Jun 23, 2014 |
| <u>LOVENOX (PRESERVATIVE FREE)</u> | | | | |
| <u>AP</u> | + | SANOFI AVENTIS US | <u>30MG/0.3ML (100MG/ML)</u> | <u>N020164 001</u> Mar 29, 1993 |
| <u>AP</u> | + | | <u>40MG/0.4ML (100MG/ML)</u> | <u>N020164 002</u> Jan 30, 1998 |
| <u>AP</u> | + | | <u>60MG/0.6ML (100MG/ML)</u> | <u>N020164 003</u> Mar 27, 1998 |
| <u>AP</u> | + | | <u>80MG/0.8ML (100MG/ML)</u> | <u>N020164 004</u> Mar 27, 1998 |
| <u>AP</u> | + | | <u>100MG/ML (100MG/ML)</u> | <u>N020164 005</u> Mar 27, 1998 |
| <u>AP</u> | + | | <u>120MG/0.8ML (150MG/ML)</u> | <u>N020164 007</u> Jun 02, 2000 |
| <u>AP</u> | + | | <u>150MG/ML (150MG/ML)</u> | <u>N020164 008</u> Jun 02, 2000 |

ENTACAPONE

TABLET; ORAL

COMTAN

| | | | | |
|-----------|---|--------------|--------------|---------------------------------|
| <u>AB</u> | + | ORION PHARMA | <u>200MG</u> | <u>N020796 001</u> Oct 19, 1999 |
|-----------|---|--------------|--------------|---------------------------------|

ENTACAPONE

| | | | | |
|-----------|--|----------------------|--------------|---------------------------------|
| <u>AB</u> | | AJANTA PHARMA LTD | <u>200MG</u> | <u>A205792 001</u> Aug 31, 2017 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>200MG</u> | <u>A203437 001</u> Jun 19, 2015 |
| <u>AB</u> | | MACLEODS PHARMS LTD | <u>200MG</u> | <u>A207210 001</u> Jun 05, 2017 |
| <u>AB</u> | | SUN PHARMA GLOBAL | <u>200MG</u> | <u>A090690 001</u> Jul 16, 2012 |
| <u>AB</u> | | SUNSHINE LAKE | <u>200MG</u> | <u>A206669 001</u> Oct 03, 2018 |
| <u>AB</u> | | WOCKHARDT LTD | <u>200MG</u> | <u>A078941 001</u> Aug 16, 2012 |

ENTECAVIR

SOLUTION; ORAL

BARACLUE

| | | | | |
|--|---|----------------------|-----------|--------------------------|
| | + | BRISTOL MYERS SQUIBB | 0.05MG/ML | N021798 001 Mar 29, 2005 |
|--|---|----------------------|-----------|--------------------------|

TABLET; ORAL

BARACLUE

| | | | | |
|-----------|---|----------------------|--------------|---------------------------------|
| <u>AB</u> | + | BRISTOL MYERS SQUIBB | <u>0.5MG</u> | <u>N021797 001</u> Mar 29, 2005 |
| <u>AB</u> | + | | <u>1MG</u> | <u>N021797 002</u> Mar 29, 2005 |

ENTECAVIR

| | | | | |
|-----------|--|----------------------|--------------|---------------------------------|
| <u>AB</u> | | ACCORD HLTHCARE | <u>0.5MG</u> | <u>A205824 001</u> Aug 25, 2017 |
| <u>AB</u> | | | <u>1MG</u> | <u>A205824 002</u> Aug 25, 2017 |
| <u>AB</u> | | AMNEAL PHARMS | <u>0.5MG</u> | <u>A206652 001</u> Nov 12, 2015 |
| <u>AB</u> | | | <u>1MG</u> | <u>A206652 002</u> Nov 12, 2015 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>0.5MG</u> | <u>A206217 001</u> Aug 26, 2015 |
| <u>AB</u> | | | <u>1MG</u> | <u>A206217 002</u> Aug 26, 2015 |
| <u>AB</u> | | BRECKENRIDGE PHARM | <u>0.5MG</u> | <u>A208721 001</u> Mar 15, 2018 |
| <u>AB</u> | | | <u>1MG</u> | <u>A208721 002</u> Mar 15, 2018 |
| <u>AB</u> | | CASI PHARMS INC | <u>0.5MG</u> | <u>A206672 001</u> May 11, 2017 |
| <u>AB</u> | | | <u>1MG</u> | <u>A206672 002</u> May 11, 2017 |
| <u>AB</u> | | CIPLA | <u>0.5MG</u> | <u>A206872 001</u> Dec 06, 2016 |
| <u>AB</u> | | | <u>1MG</u> | <u>A206872 002</u> Dec 06, 2016 |
| <u>AB</u> | | HETERO LABS LTD V | <u>0.5MG</u> | <u>A205740 001</u> Aug 21, 2015 |
| <u>AB</u> | | | <u>1MG</u> | <u>A205740 002</u> Aug 21, 2015 |
| <u>AB</u> | | PAR PHARM INC | <u>0.5MG</u> | <u>A206294 001</u> Nov 23, 2016 |
| <u>AB</u> | | | <u>1MG</u> | <u>A206294 002</u> Nov 23, 2016 |
| <u>AB</u> | | PRINSTON INC | <u>0.5MG</u> | <u>A208782 001</u> Oct 10, 2017 |
| <u>AB</u> | | | <u>1MG</u> | <u>A208782 002</u> Oct 10, 2017 |
| <u>AB</u> | | TEVA PHARMS USA | <u>0.5MG</u> | <u>A202122 001</u> Aug 26, 2014 |
| <u>AB</u> | | | <u>1MG</u> | <u>A202122 002</u> Aug 26, 2014 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>0.5MG</u> | <u>A206745 001</u> Jun 23, 2017 |
| <u>AB</u> | | | <u>1MG</u> | <u>A206745 002</u> Jun 23, 2017 |

ENZALUTAMIDE

CAPSULE; ORAL

XTANDI

| | | | | |
|--|---|----------|------|--------------------------|
| | + | ASTELLAS | 40MG | N203415 001 Aug 31, 2012 |
|--|---|----------|------|--------------------------|

PRESCRIPTION DRUG PRODUCT LIST

EPHEDRINE SULFATE

SOLUTION; INTRAVENOUS

AKOVAZ

| | | | | | |
|-----------|-----------|--------------------|--------------------------|--------------------|--------------|
| AP | +! | FLAMEL IRELAND LTD | 50MG/ML (50MG/ML) | N208289 001 | Apr 29, 2016 |
|-----------|-----------|--------------------|--------------------------|--------------------|--------------|

CORPHEDRA

| | | | | | |
|-----------|--|----------------------|--------------------------|--------------------|--------------|
| AP | | PAR STERILE PRODUCTS | 50MG/ML (50MG/ML) | N208943 001 | Jan 27, 2017 |
|-----------|--|----------------------|--------------------------|--------------------|--------------|

EPHEDRINE SULFATE

| | | | | | |
|-----------|--|-----------|--------------------------|--------------------|--------------|
| AP | | AKORN INC | 50MG/ML (50MG/ML) | N208609 001 | Mar 01, 2017 |
|-----------|--|-----------|--------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------|--------------------------|--------------------|--------------|
| AP | | SANDOZ INC | 50MG/ML (50MG/ML) | A209784 001 | Aug 23, 2017 |
|-----------|--|------------|--------------------------|--------------------|--------------|

EPINASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

ELESTAT

| | | | | | |
|-----------|-----------|----------|--------------|--------------------|--------------|
| AT | +! | ALLERGAN | 0.05% | N021565 001 | Oct 16, 2003 |
|-----------|-----------|----------|--------------|--------------------|--------------|

EPINASTINE HYDROCHLORIDE

| | | | | | |
|-----------|--|-------|--------------|--------------------|--------------|
| AT | | AKORN | 0.05% | A204055 001 | May 05, 2017 |
|-----------|--|-------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------|--------------|--------------------|--------------|
| AT | | APOTEX | 0.05% | A090919 001 | Oct 31, 2011 |
|-----------|--|--------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------------|--------------|--------------------|--------------|
| AT | | BRECKENRIDGE PHARM | 0.05% | A090870 001 | Mar 14, 2011 |
|-----------|--|--------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|-----------------|--------------|--------------------|--------------|
| AT | | CASI PHARMS INC | 0.05% | A203384 001 | Dec 07, 2016 |
|-----------|--|-----------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------------|--------------|--------------------|--------------|
| AT | | SOMERSET THERAPS LLC | 0.05% | A090951 001 | Oct 31, 2011 |
|-----------|--|----------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------|--------------|--------------------|--------------|
| AT | | SUN PHARM INDS | 0.05% | A091626 001 | Oct 31, 2011 |
|-----------|--|----------------|--------------|--------------------|--------------|

EPINEPHRINE

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

EPINEPHRINE (AUTOINJECTOR)

| | | | | | |
|-----------|--|-----------------|------------------------|--------------------|--------------|
| AB | | TEVA PHARMS USA | 0.15MG/DELIVERY | A090589 002 | Aug 16, 2018 |
|-----------|--|-----------------|------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-----------------------|--------------------|--------------|
| AB | | | 0.3MG/DELIVERY | A090589 001 | Aug 16, 2018 |
|-----------|--|--|-----------------------|--------------------|--------------|

EPIPEN

| | | | | | |
|-----------|-----------|---------------------|-----------------------|--------------------|--------------|
| AB | +! | MYLAN SPECIALITY LP | 0.3MG/DELIVERY | N019430 001 | Dec 22, 1987 |
|-----------|-----------|---------------------|-----------------------|--------------------|--------------|

EPIPEN JR.

| | | | | | |
|-----------|-----------|---------------------|------------------------|--------------------|--------------|
| AB | +! | MYLAN SPECIALITY LP | 0.15MG/DELIVERY | N019430 002 | Dec 22, 1987 |
|-----------|-----------|---------------------|------------------------|--------------------|--------------|

ADRENACLICK

| | | | | | |
|-----------|-----------|-------|--------------------|-------------|--------------|
| BX | +! | IMPAX | EQ 0.15MG/DELIVERY | N020800 003 | Nov 25, 2009 |
|-----------|-----------|-------|--------------------|-------------|--------------|

| | | | | | |
|-----------|-----------|--|-------------------|-------------|--------------|
| BX | +! | | EQ 0.3MG/DELIVERY | N020800 004 | Nov 25, 2009 |
|-----------|-----------|--|-------------------|-------------|--------------|

SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS

ADRENALIN

| | | | | | |
|-----------|-----------|----------------------|----------------------------------------|--------------------|--------------|
| AP | +! | PAR STERILE PRODUCTS | EQ 1MG BASE/ML (EQ 1MG BASE/ML) | N204200 001 | Dec 07, 2012 |
|-----------|-----------|----------------------|----------------------------------------|--------------------|--------------|

EPINEPHRINE

| | | | | | |
|-----------|--|----------|----------------------------------------|--------------------|--------------|
| AP | | LUITPOLD | EQ 1MG BASE/ML (EQ 1MG BASE/ML) | A207568 001 | Jul 06, 2018 |
|-----------|--|----------|----------------------------------------|--------------------|--------------|

AUVI-Q

| | | | | | |
|-----------|-----------|-----------|--------------------|-------------|--------------|
| BX | +! | KALEO INC | EQ 0.15MG/DELIVERY | N201739 002 | Aug 10, 2012 |
|-----------|-----------|-----------|--------------------|-------------|--------------|

| | | | | | |
|-----------|----------|--|-------------------|-------------|--------------|
| BX | + | | EQ 0.3MG/DELIVERY | N201739 001 | Aug 10, 2012 |
|-----------|----------|--|-------------------|-------------|--------------|

ADRENALIN

| | | | | | |
|--|-----------|----------------------|------------------------------------|-------------|--------------|
| | +! | PAR STERILE PRODUCTS | EQ 30MG BASE/30ML (EQ 1MG BASE/ML) | N204640 001 | Dec 18, 2013 |
|--|-----------|----------------------|------------------------------------|-------------|--------------|

AUVI-Q

| | | | | | |
|--|----------|-----------|-------------------|-------------|--------------|
| | + | KALEO INC | EQ 0.1MG/DELIVERY | N201739 003 | Nov 17, 2017 |
|--|----------|-----------|-------------------|-------------|--------------|

SYMJEPI

| | | | | | |
|--|-----------|--------------------|-----------------------------|-------------|--------------|
| | +! | ADAMIS PHARMS CORP | 0.15MG/0.3ML (0.15MG/0.3ML) | N207534 002 | Sep 27, 2018 |
|--|-----------|--------------------|-----------------------------|-------------|--------------|

| | | | | | |
|--|-----------|--|---------------------------|-------------|--------------|
| | +! | | 0.3MG/0.3ML (0.3MG/0.3ML) | N207534 001 | Jun 15, 2017 |
|--|-----------|--|---------------------------|-------------|--------------|

SOLUTION; IV (INFUSION), INTRAOCULAR, INTRAMUSCULAR, SUBCUTANEOUS

EPINEPHRINE

| | | | | | |
|--|-----------|--------------------|---------------------------------|-------------|--------------|
| | +! | BELCHER PHARMS LLC | EQ 1MG BASE/ML (EQ 1MG BASE/ML) | N205029 001 | Jul 29, 2014 |
|--|-----------|--------------------|---------------------------------|-------------|--------------|

EPINEPHRINE BITARTRATE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIGNOSPAN FORTE

| | | | | | |
|--|----------|---------|-----------------------|-------------|--------------|
| | ! | DEPROCO | EQ 0.02MG BASE/ML; 2% | A088389 001 | Jan 22, 1985 |
|--|----------|---------|-----------------------|-------------|--------------|

LIGNOSPAN STANDARD

| | | | | | |
|--|----------|---------|-----------------------|-------------|--------------|
| | ! | DEPROCO | EQ 0.01MG BASE/ML; 2% | A088390 001 | Jan 22, 1985 |
|--|----------|---------|-----------------------|-------------|--------------|

EPINEPHRINE BITARTRATE; PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST FORTE DENTAL

| | | | | | |
|-----------|-----------|----------------|-----------------------|--------------------|--|
| AP | +! | DENTSPLY PHARM | 0.005MG/ML; 4% | N021383 001 | |
|-----------|-----------|----------------|-----------------------|--------------------|--|

PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE

| | | | | | |
|-----------|--|---------------|-----------------------|--------------------|--------------|
| AP | | SEPTODONT INC | 0.005MG/ML; 4% | A078959 001 | Aug 30, 2011 |
|-----------|--|---------------|-----------------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

| | | | | |
|---------------------------------|---------|------------------------|------------------------|---------------------------------|
| <u>AP</u> | HOSPIRA | <u>0.005MG/ML;0.5%</u> | <u>A089635 001</u> | Jun 21, 1988 |
| <u>AP</u> | | <u>0.005MG/ML;1.5%</u> | <u>A088571 001</u> | Sep 13, 1985 |
| <u>AP</u> | | <u>0.005MG/ML;1.5%</u> | <u>A089645 001</u> | Jun 21, 1988 |
| <u>AP</u> | | <u>0.005MG/ML;2%</u> | <u>A089651 001</u> | Jun 21, 1988 |
| <u>AP</u> | | <u>0.01MG/ML;1%</u> | <u>A089644 001</u> | Jun 21, 1988 |
| <u>AP</u> | ! | <u>0.01MG/ML;2%</u> | <u>A089646 001</u> | Jun 21, 1988 |
| <u>XYLOCAINE W/ EPINEPHRINE</u> | | | | |
| <u>AP</u> | +! | FRESENIUS KABI USA | <u>0.005MG/ML;0.5%</u> | <u>N006488 012</u> |
| <u>AP</u> | +! | | <u>0.005MG/ML;1.5%</u> | <u>N006488 017</u> |
| <u>AP</u> | +! | | <u>0.005MG/ML;2%</u> | <u>N006488 019</u> Nov 13, 1986 |
| <u>AP</u> | +! | | <u>0.01MG/ML;1%</u> | <u>N006488 004</u> |
| <u>AP</u> | +! | | <u>0.02MG/ML;2%</u> | <u>N006488 005</u> |
| | +! | | 0.005MG/ML;1% | N006488 018 Nov 13, 1986 |

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ELLEENCE

| | | | | |
|-----------|----|------------|-----------------------------|---------------------------------|
| <u>AP</u> | +! | PFIZER INC | <u>200MG/100ML (2MG/ML)</u> | <u>N050778 001</u> Sep 15, 1999 |
| <u>AP</u> | + | | <u>50MG/25ML (2MG/ML)</u> | <u>N050778 002</u> Sep 15, 1999 |

EPIRUBICIN HYDROCHLORIDE

| | | | | |
|-----------|--|-------------------------|-----------------------------|---------------------------------|
| <u>AP</u> | | ACTAVIS TOTOWA | <u>10MG/5ML (2MG/ML)</u> | <u>A065445 001</u> Sep 18, 2008 |
| <u>AP</u> | | | <u>50MG/25ML (2MG/ML)</u> | <u>A065445 002</u> Sep 18, 2008 |
| <u>AP</u> | | | <u>200MG/100ML (2MG/ML)</u> | <u>A065445 003</u> Sep 18, 2008 |
| <u>AP</u> | | AKORN INC | <u>50MG/25ML (2MG/ML)</u> | <u>A090163 001</u> Jun 24, 2009 |
| <u>AP</u> | | CIPLA LTD | <u>50MG/25ML (2MG/ML)</u> | <u>A065361 001</u> Oct 22, 2007 |
| <u>AP</u> | | | <u>200MG/100ML (2MG/ML)</u> | <u>A065361 002</u> Oct 22, 2007 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>10MG/5ML (2MG/ML)</u> | <u>A065408 001</u> Oct 15, 2007 |
| <u>AP</u> | | | <u>50MG/25ML (2MG/ML)</u> | <u>A065408 002</u> Oct 15, 2007 |
| <u>AP</u> | | | <u>150MG/75ML (2MG/ML)</u> | <u>A065408 003</u> Oct 15, 2007 |
| <u>AP</u> | | | <u>200MG/100ML (2MG/ML)</u> | <u>A065408 004</u> Oct 15, 2007 |
| <u>AP</u> | | | <u>200MG/100ML (2MG/ML)</u> | <u>A065411 001</u> Aug 20, 2007 |
| <u>AP</u> | | | <u>50MG/25ML (2MG/ML)</u> | <u>A065411 002</u> Aug 20, 2007 |
| <u>AP</u> | | HISUN PHARM HANGZHOU | <u>50MG/25ML (2MG/ML)</u> | <u>A090075 001</u> Mar 25, 2010 |
| <u>AP</u> | | | <u>200MG/100ML (2MG/ML)</u> | <u>A090075 002</u> Mar 25, 2010 |
| <u>AP</u> | | HOSPIRA | <u>10MG/5ML (2MG/ML)</u> | <u>A065343 001</u> Apr 19, 2007 |
| <u>AP</u> | | | <u>150MG/75ML (2MG/ML)</u> | <u>A065343 003</u> Apr 19, 2007 |
| <u>AP</u> | | | <u>200MG/100ML (2MG/ML)</u> | <u>A065343 004</u> Apr 19, 2007 |
| <u>AP</u> | | IMPAX LABS INC | <u>50MG/25ML (2MG/ML)</u> | <u>A065331 001</u> Aug 09, 2007 |
| <u>AP</u> | | | <u>200MG/100ML (2MG/ML)</u> | <u>A065331 002</u> Aug 09, 2007 |
| <u>AP</u> | | MYLAN LABS LTD | <u>50MG/25ML (2MG/ML)</u> | <u>A091599 001</u> Mar 12, 2012 |
| <u>AP</u> | | | <u>200MG/100ML (2MG/ML)</u> | <u>A091599 002</u> Mar 12, 2012 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>50MG/25ML (2MG/ML)</u> | <u>A065289 001</u> Jun 27, 2007 |
| <u>AP</u> | | | <u>200MG/100ML (2MG/ML)</u> | <u>A065289 002</u> Jun 27, 2007 |

EPLERENONE

TABLET; ORAL

EPLERENONE

| | | | | |
|----------------|----|--------------------|-------------|---------------------------------|
| <u>AB</u> | | ACCORD HLTHCARE | <u>25MG</u> | <u>A206922 001</u> Jul 13, 2017 |
| <u>AB</u> | | | <u>50MG</u> | <u>A206922 002</u> Jul 13, 2017 |
| <u>AB</u> | | APOTEX | <u>25MG</u> | <u>A078482 001</u> Jul 30, 2008 |
| <u>AB</u> | | | <u>50MG</u> | <u>A078482 002</u> Jul 30, 2008 |
| <u>AB</u> | | BRECKENRIDGE PHARM | <u>25MG</u> | <u>A208283 001</u> Sep 14, 2018 |
| <u>AB</u> | | | <u>50MG</u> | <u>A208283 002</u> Sep 14, 2018 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>25MG</u> | <u>A203896 001</u> Feb 02, 2017 |
| <u>AB</u> | | | <u>50MG</u> | <u>A203896 002</u> Feb 02, 2017 |
| <u>AB</u> | | SANDOZ | <u>25MG</u> | <u>A078510 001</u> Aug 01, 2008 |
| <u>AB</u> | | | <u>50MG</u> | <u>A078510 002</u> Aug 01, 2008 |
| <u>INSPIRA</u> | | | | |
| <u>AB</u> | + | GD SEARLE LLC | <u>25MG</u> | <u>N021437 001</u> Sep 27, 2002 |
| <u>AB</u> | +! | | <u>50MG</u> | <u>N021437 002</u> Sep 27, 2002 |

EPOPROSTENOL SODIUM

INJECTABLE; INJECTION

EPOPROSTENOL SODIUM

| | | | | |
|---------------|----|---------------------|---------------------------|---------------------------------|
| <u>AP</u> | | TEVA PHARMS USA | <u>EQ 0.5MG BASE/VIAL</u> | <u>A078396 001</u> Apr 23, 2008 |
| <u>AP</u> | | | <u>EQ 1.5MG BASE/VIAL</u> | <u>A078396 002</u> Apr 23, 2008 |
| <u>FLOLAN</u> | | | | |
| <u>AP</u> | +! | GLAXOSMITHKLINE LLC | <u>EQ 0.5MG BASE/VIAL</u> | <u>N020444 001</u> Sep 20, 1995 |
| <u>AP</u> | +! | | <u>EQ 1.5MG BASE/VIAL</u> | <u>N020444 002</u> Sep 20, 1995 |

PRESCRIPTION DRUG PRODUCT LIST

EPOPROSTENOL SODIUM

INJECTABLE; INJECTION

VELETRI

| | | | | | |
|---|---------------------|--------------------|---------|-----|--------------|
| + | ACTELION PHARMS LTD | EQ 0.5MG BASE/VIAL | N022260 | 002 | Jun 28, 2012 |
| + | ! | EQ 1.5MG BASE/VIAL | N022260 | 001 | Jun 27, 2008 |

EPROSARTAN MESYLATE

TABLET; ORAL

EPROSARTAN MESYLATE

| | | | | | |
|-----------|------------------|----------------------|----------------|------------|--------------|
| AB | MYLAN PHARMS INC | EQ 400MG BASE | A202012 | 001 | Nov 16, 2011 |
| AB | ! | EQ 600MG BASE | A202012 | 002 | Nov 16, 2011 |

EPTIFIBATIDE

INJECTABLE; INJECTION

EPTIFIBATIDE

| | | | | | |
|-------------------|----------------------|-------------------|-------------------|----------------|-------------------------|
| AP | ACCORD HLTHCARE | 2MG/ML | A205557 | 001 | Nov 06, 2017 |
| AP | | 75MG/100ML | A205557 | 002 | Nov 06, 2017 |
| AP | AKORN | 2MG/ML | A204589 | 001 | Apr 18, 2017 |
| AP | | 75MG/100ML | A204589 | 002 | Apr 18, 2017 |
| AP | AMNEAL PHARMS | 2MG/ML | A205581 | 001 | Dec 08, 2016 |
| AP | | 75MG/100ML | A205581 | 002 | Dec 08, 2016 |
| AP | AUROBINDO PHARMA LTD | 2MG/ML | A206127 | 001 | Dec 08, 2015 |
| AP | | 75MG/100ML | A206127 | 002 | Dec 08, 2015 |
| AP | CELERITY PHARMS | 2MG/ML | A208554 | 001 | Nov 23, 2018 |
| AP | | 75MG/100ML | A208554 | 002 | Nov 23, 2018 |
| AP | MYLAN LABS LTD | 2MG/ML | A203258 | 001 | Jul 20, 2018 |
| AP | | 75MG/100ML | A203258 | 002 | Jul 20, 2018 |
| AP | SAGENT PHARMS | 2MG/ML | A204693 | 001 | Mar 07, 2018 |
| AP | | 75MG/100ML | A204693 | 002 | Mar 07, 2018 |
| AP | TEVA PHARMS USA | 2MG/ML | A090854 | 001 | Jun 12, 2015 |
| INTEGRILIN | | | | | |
| AP | + | SCHERING | 2MG/ML | N020718 | 001 May 18, 1998 |
| AP | + | ! | 75MG/100ML | N020718 | 002 May 18, 1998 |

ERAVACYCLINE DIHYDROCHLORIDE

POWDER; INTRAVENOUS

XERAVA

| | | | | | |
|---|-------------------|-------------------|---------|-----|--------------|
| + | TETRAPHASE PHARMS | EQ 50MG BASE/VIAL | N211109 | 001 | Aug 27, 2018 |
|---|-------------------|-------------------|---------|-----|--------------|

ERGOCALCIFEROL

CAPSULE; ORAL

DRISDOL

| | | | | | |
|-----------|---|-------------------|------------------|----------------|------------|
| AA | + | US PHARM HOLDINGS | 50,000 IU | N003444 | 001 |
|-----------|---|-------------------|------------------|----------------|------------|

ERGOCALCIFEROL

| | | | | | |
|-----------|---------------------|------------------|----------------|------------|--------------|
| AA | ORIT LABS LLC | 50,000 IU | A040833 | 001 | May 20, 2009 |
| AA | PURACAP PHARM LLC | 50000IU | A204276 | 001 | Dec 07, 2018 |
| AA | SIGMAPHARM LABS LLC | 50,000 IU | A091004 | 001 | Jul 14, 2010 |
| AA | STRIDES PHARMA | 50,000 IU | A090455 | 001 | Aug 03, 2010 |
| AA | SUN PHARM INDS INC | 50,000 IU | A040865 | 001 | Dec 29, 2009 |

VITAMIN D

| | | | | | |
|-----------|----------------|------------------|----------------|------------|--|
| AA | BIONPHARMA INC | 50,000 IU | A080704 | 001 | |
|-----------|----------------|------------------|----------------|------------|--|

ERGOLOID MESYLATES

TABLET; ORAL

ERGOLOID MESYLATES

| | | | | | |
|---|----------------------|-----|---------|-----|--------------|
| ! | SUN PHARM INDUSTRIES | 1MG | A081113 | 001 | Oct 31, 1991 |
|---|----------------------|-----|---------|-----|--------------|

ERGOTAMINE TARTRATE

TABLET; SUBLINGUAL

ERGOMAR

| | | | | | |
|---|---------------------|-----|---------|-----|--------------|
| ! | TERSERA THERAPS LLC | 2MG | A087693 | 001 | Feb 24, 1983 |
|---|---------------------|-----|---------|-----|--------------|

ERIBULIN MESYLATE

SOLUTION; INTRAVENOUS

HALAVEN

| | | | | | |
|---|-----------|--------------------|---------|-----|--------------|
| + | EISAI INC | 1MG/2ML (0.5MG/ML) | N201532 | 001 | Nov 15, 2010 |
|---|-----------|--------------------|---------|-----|--------------|

ERLOTINIB HYDROCHLORIDE

TABLET; ORAL

TARCEVA

| | | | | | |
|---|------------|---------------|---------|-----|--------------|
| + | OSI PHARMS | EQ 25MG BASE | N021743 | 001 | Nov 18, 2004 |
| + | | EQ 100MG BASE | N021743 | 002 | Nov 18, 2004 |
| + | ! | EQ 150MG BASE | N021743 | 003 | Nov 18, 2004 |

PRESCRIPTION DRUG PRODUCT LIST

ERTAPENEM SODIUM

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

ERTAPENEM SODIUM

| | | | | |
|-----------|----------------------|--------------------------------|---------------------------|--------------|
| AP | ACS DOBFAR SPA | <u>EQ 1GM BASE/VIAL</u> | <u>A208790 001</u> | Apr 16, 2018 |
| AP | AUROBINDO PHARMA LTD | <u>EQ 1GM BASE/VIAL</u> | <u>A209133 001</u> | Jun 25, 2018 |

INVANZ

| | | | | | |
|-----------|------------|-------------------|--------------------------------|---------------------------|--------------|
| AP | + ! | MERCK SHARP DOHME | <u>EQ 1GM BASE/VIAL</u> | <u>N021337 001</u> | Nov 21, 2001 |
|-----------|------------|-------------------|--------------------------------|---------------------------|--------------|

ERTUGLIFLOZIN

TABLET; ORAL

STEGLATRO

| | | | | |
|------------|-------------------|------|-------------|--------------|
| + | MERCK SHARP DOHME | 5MG | N209803 001 | Dec 19, 2017 |
| + ! | | 15MG | N209803 002 | Dec 19, 2017 |

ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

SEGLUROMET

| | | | | |
|------------|-------------------|--------------|-------------|--------------|
| + | MERCK SHARP DOHME | 2.5MG; 500MG | N209806 001 | Dec 19, 2017 |
| + | | 2.5MG; 1GM | N209806 002 | Dec 19, 2017 |
| + | | 7.5MG; 500MG | N209806 003 | Dec 19, 2017 |
| + ! | | 7.5MG; 1GM | N209806 004 | Dec 19, 2017 |

ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE

TABLET; ORAL

STEGLUJAN

| | | | | |
|------------|-------------------|---------------------|-------------|--------------|
| + | MERCK SHARP DOHME | 5MG; EQ 100MG BASE | N209805 001 | Dec 19, 2017 |
| + ! | | 15MG; EQ 100MG BASE | N209805 002 | Dec 19, 2017 |

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS; ORAL

ERYC

| | | | | |
|-----------|------------|--------------|---------------------|---------------------------|
| AB | + ! | MAYNE PHARMA | <u>250MG</u> | <u>N050536 001</u> |
|-----------|------------|--------------|---------------------|---------------------------|

ERYTHROMYCIN

| | | | | | |
|-----------|--|------------------|---------------------|---------------------------|--------------|
| AB | | ARBOR PHARMS LLC | <u>250MG</u> | <u>A062746 001</u> | Dec 22, 1986 |
|-----------|--|------------------|---------------------|---------------------------|--------------|

GEL; TOPICAL

ERYGEL

| | | | | | |
|-----------|------------|------------------|------------------|---------------------------|--------------|
| AT | + ! | MYLAN PHARMS INC | <u>2%</u> | <u>N050617 001</u> | Oct 21, 1987 |
|-----------|------------|------------------|------------------|---------------------------|--------------|

ERYTHROMYCIN

| | | | | | |
|-----------|--|----------------|------------------|---------------------------|--------------|
| AT | | FOUGERA PHARMS | <u>2%</u> | <u>A064184 001</u> | Sep 30, 1997 |
|-----------|--|----------------|------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|------------|------------------|---------------------------|--------------|
| AT | | PERRIGO CO | <u>2%</u> | <u>A063211 001</u> | Jan 29, 1993 |
|-----------|--|------------|------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|---------------------|------------------|---------------------------|--------------|
| AT | | TELIGENT PHARMA INC | <u>2%</u> | <u>A208154 001</u> | Jul 19, 2017 |
|-----------|--|---------------------|------------------|---------------------------|--------------|

OINTMENT; OPHTHALMIC

ERYTHROMYCIN

| | | | | | |
|-----------|--|-------|--------------------|---------------------------|--------------|
| AT | | AKORN | <u>0.5%</u> | <u>A064030 001</u> | Jul 18, 1996 |
|-----------|--|-------|--------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|-----------------|--------------------|---------------------------|--------------|
| AT | | BAUSCH AND LOMB | <u>0.5%</u> | <u>A064067 001</u> | Jul 29, 1994 |
|-----------|--|-----------------|--------------------|---------------------------|--------------|

| | | | | | |
|-----------|----------|-------------------------|--------------------|---------------------------|--------------|
| AT | ! | PERRIGO CO TENNESSEE | <u>0.5%</u> | <u>A062447 001</u> | Sep 26, 1983 |
|-----------|----------|-------------------------|--------------------|---------------------------|--------------|

SOLUTION; TOPICAL

ERYTHROMYCIN

| | | | | | |
|-----------|----------|------------------|------------------|---------------------------|--------------|
| AT | ! | PERRIGO NEW YORK | <u>2%</u> | <u>A063038 001</u> | Jan 11, 1991 |
|-----------|----------|------------------|------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|---------------------|------------------|---------------------------|--------------|
| AT | | TELIGENT PHARMA INC | <u>2%</u> | <u>A208100 001</u> | Nov 20, 2017 |
|-----------|--|---------------------|------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|------------------|------------------|---------------------------|--------------|
| AT | | WOCKHARDT BIO AG | <u>2%</u> | <u>A062825 001</u> | Oct 23, 1987 |
|-----------|--|------------------|------------------|---------------------------|--------------|

SWAB; TOPICAL

ERYTHROMYCIN

| | | | | | |
|-----------|--|-------|------------------|---------------------------|--------------|
| AT | | AKORN | <u>2%</u> | <u>A090215 001</u> | May 12, 2010 |
|-----------|--|-------|------------------|---------------------------|--------------|

| | | | | | |
|-----------|----------|------------|------------------|---------------------------|--------------|
| AT | ! | PERRIGO CO | <u>2%</u> | <u>A064126 001</u> | Jul 03, 1996 |
|-----------|----------|------------|------------------|---------------------------|--------------|

TABLET; ORAL

ERYTHROMYCIN

| | | | | | |
|-----------|--|------------------|---------------------|---------------------------|--------------|
| AB | | AMNEAL PHARMS CO | <u>250MG</u> | <u>A209720 001</u> | Mar 09, 2018 |
|-----------|--|------------------|---------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|--|---------------------|---------------------------|--------------|
| AB | | | <u>500MG</u> | <u>A209720 002</u> | Mar 09, 2018 |
|-----------|--|--|---------------------|---------------------------|--------------|

| | | | | |
|-----------|--|------------------|---------------------|---------------------------|
| AB | | ARBOR PHARMS LLC | <u>250MG</u> | <u>A061621 001</u> |
|-----------|--|------------------|---------------------|---------------------------|

| | | | | |
|-----------|----------|--|---------------------|---------------------------|
| AB | ! | | <u>500MG</u> | <u>A061621 002</u> |
|-----------|----------|--|---------------------|---------------------------|

TABLET, DELAYED RELEASE; ORAL

ERY-TAB

| | | | | |
|--|--|------------------|-------|-------------|
| | | ARBOR PHARMS LLC | 250MG | A062298 001 |
|--|--|------------------|-------|-------------|

| | | | | | |
|--|--|--|-------|-------------|--------------|
| | | | 333MG | A062298 003 | Mar 29, 1982 |
|--|--|--|-------|-------------|--------------|

| | | | | |
|----------|--|--|-------|-------------|
| ! | | | 500MG | A062298 002 |
|----------|--|--|-------|-------------|

PRESCRIPTION DRUG PRODUCT LISTERYTHROMYCIN ETHYLSUCCINATE

GRANULE; ORAL

ERYPED**AB +!** ARBOR PHARMS LLC **EQ 400MG BASE/5ML** **N050207 002**ERYTHROMYCIN ETHYLSUCCINATE**AB** ANI PHARMS INC **EQ 400MG BASE/5ML** **A062055 002** Nov 02, 2018E.E.S.**AB1 +** ARBOR PHARMS LLC **EQ 200MG BASE/5ML** **N050207 001**ERYTHROMYCIN ETHYLSUCCINATE**AB1** ANI PHARMS INC **EQ 200MG BASE/5ML** **A062055 001**ERYPED**AB2 +** ARBOR PHARMS LLC **EQ 200MG BASE/5ML** **N050207 003** Mar 30, 1987ERYTHROMYCIN ETHYLSUCCINATE**AB2** ANI PHARMS INC **EQ 200MG BASE/5ML** **A062055 003** Nov 02, 2018

TABLET; ORAL

E.E.S. 400

BX ! ARBOR PHARMS LLC **EQ 400MG BASE** **A061905 002** Aug 12, 1982

ERYTHROMYCIN ETHYLSUCCINATE

BX ! ARBOR PHARMS LLC **EQ 400MG BASE** **A061904 001**ERYTHROMYCIN LACTOBIONATE

INJECTABLE; INJECTION

ERYTHROCIN**AP** HOSPIRA **EQ 500MG BASE/VIAL** **A062638 001** Oct 31, 1986**AP +!** **EQ 500MG BASE/VIAL** **N050609 001** Sep 24, 1986ERYTHROMYCIN STEARATE

TABLET; ORAL

ERYTHROCIN STEARATE

! ARBOR PHARMS LLC **EQ 250MG BASE** **A060359 001**ESCITALOPRAM OXALATE

SOLUTION; ORAL

ESCITALOPRAM OXALATE**AA** AMNEAL PHARMS **EQ 5MG BASE/5ML** **A202227 001** Mar 14, 2012**AA** ANTRIM PHARMS LLC **EQ 5MG BASE/5ML** **A203967 001** May 26, 2015**AA** AUROBINDO PHARMA **EQ 5MG BASE/5ML** **A079062 001** Apr 02, 2012

LTD

AA HETERO LABS LTD III **EQ 5MG BASE/5ML** **A202221 001** Jun 12, 2012**AA** LANNETT CO INC **EQ 5MG BASE/5ML** **A090477 001** Jun 12, 2013**AA** MACLEODS PHARMS LTD **EQ 5MG BASE/5ML** **A202754 001** Mar 31, 2016**AA** TARO **EQ 5MG BASE/5ML** **A079121 001** May 03, 2012LEXAPRO**AA +!** ALLERGAN SALES LLC **EQ 5MG BASE/5ML** **N021365 001** Nov 27, 2002

TABLET; ORAL

ESCITALOPRAM OXALATE**AB** ACCORD HLTHCARE **EQ 5MG BASE** **A202389 001** Sep 11, 2012**AB** **EQ 10MG BASE** **A202389 002** Sep 11, 2012**AB** **EQ 20MG BASE** **A202389 003** Sep 11, 2012**AB** AMNEAL PHARMS **EQ 5MG BASE** **A205619 001** May 17, 2017**AB** **EQ 10MG BASE** **A205619 002** May 17, 2017**AB** **EQ 20MG BASE** **A205619 003** May 17, 2017**AB** APOTEX INC **EQ 5MG BASE** **A078777 001** Sep 11, 2012**AB** **EQ 10MG BASE** **A078777 002** Sep 11, 2012**AB** **EQ 20MG BASE** **A078777 003** Sep 11, 2012**AB** AUROBINDO PHARMA **EQ 5MG BASE** **A090432 001** Sep 11, 2012

LTD

AB **EQ 10MG BASE** **A090432 002** Sep 11, 2012**AB** **EQ 20MG BASE** **A090432 003** Sep 11, 2012**AB** HIKMA PHARMS **EQ 5MG BASE** **A078766 001** Sep 11, 2012**AB** **EQ 10MG BASE** **A078766 002** Sep 11, 2012**AB** **EQ 20MG BASE** **A078766 003** Sep 11, 2012**AB** INVAGEN PHARMS **EQ 5MG BASE** **A078604 001** Sep 11, 2012**AB** **EQ 10MG BASE** **A078604 002** Sep 11, 2012**AB** **EQ 20MG BASE** **A078604 003** Sep 11, 2012**AB** JUBILANT GENERICS **EQ 5MG BASE** **A202280 001** Sep 12, 2012**AB** **EQ 10MG BASE** **A202280 002** Sep 12, 2012**AB** **EQ 20MG BASE** **A202280 003** Sep 12, 2012**AB** LUPIN LTD **EQ 5MG BASE** **A078169 001** Sep 11, 2012**AB** **EQ 10MG BASE** **A078169 002** Sep 11, 2012**AB** **EQ 20MG BASE** **A078169 003** Sep 11, 2012**AB** MACLEODS PHARMS LTD **EQ 5MG BASE** **A202210 001** Sep 11, 2012**AB** **EQ 10MG BASE** **A202210 002** Sep 11, 2012**AB** **EQ 20MG BASE** **A202210 003** Sep 11, 2012

PRESCRIPTION DRUG PRODUCT LIST

ESCITALOPRAM OXALATE

TABLET; ORAL

ESCITALOPRAM OXALATE

| | | | | |
|----------------|-------------------------|---------------------|--------------------|--------------|
| <u>AB</u> | PHARM ASSOC | <u>EQ 5MG BASE</u> | <u>A077512 001</u> | Sep 12, 2012 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A077512 002</u> | Sep 12, 2012 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077512 003</u> | Sep 12, 2012 |
| <u>AB</u> | PRINSTON INC | <u>EQ 5MG BASE</u> | <u>A078032 001</u> | Aug 28, 2015 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A078032 002</u> | Aug 28, 2015 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A078032 003</u> | Aug 28, 2015 |
| <u>AB</u> | TEVA PHARMS USA | <u>EQ 5MG BASE</u> | <u>A076765 001</u> | Mar 14, 2012 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A076765 002</u> | Mar 14, 2012 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A076765 003</u> | Mar 14, 2012 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>EQ 5MG BASE</u> | <u>A090939 001</u> | Sep 11, 2012 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A090939 002</u> | Sep 11, 2012 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A090939 003</u> | Sep 11, 2012 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 5MG BASE</u> | <u>A077734 001</u> | Sep 11, 2012 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A077734 002</u> | Sep 11, 2012 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077734 003</u> | Sep 11, 2012 |
| <u>LEXAPRO</u> | | | | |
| <u>AB</u> | + ALLERGAN SALES LLC | <u>EQ 5MG BASE</u> | <u>N021323 001</u> | Aug 14, 2002 |
| <u>AB</u> | + | <u>EQ 10MG BASE</u> | <u>N021323 002</u> | Aug 14, 2002 |
| <u>AB</u> | +! | <u>EQ 20MG BASE</u> | <u>N021323 003</u> | Aug 14, 2002 |

ESLICARBAZEPINE ACETATE

TABLET; ORAL

APTIOM

| | | | | |
|----|---------------------|-------|-------------|--------------|
| + | SUNOVION PHARMS INC | 200MG | N022416 001 | Nov 08, 2013 |
| + | | 400MG | N022416 002 | Nov 08, 2013 |
| + | | 600MG | N022416 003 | Nov 08, 2013 |
| +! | | 800MG | N022416 004 | Nov 08, 2013 |

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

BREVIBLOC

| | | | | |
|------------------------------------------------------------|-------------------------|-----------------------|--------------------|--------------|
| <u>AP</u> | +! BAXTER HLTHCARE | <u>10MG/ML</u> | <u>N019386 006</u> | Feb 25, 2003 |
| <u>ESMOLOL HYDROCHLORIDE</u> | | | | |
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>10MG/ML</u> | <u>A205520 001</u> | Jul 23, 2015 |
| <u>AP</u> | FRESENIUS KABI USA | <u>10MG/ML</u> | <u>A076573 001</u> | May 02, 2005 |
| <u>AP</u> | LUITPOLD | <u>10MG/ML</u> | <u>A201126 001</u> | Feb 20, 2015 |
| <u>AP</u> | MYLAN INSTITUTIONAL | <u>10MG/ML</u> | <u>A076474 001</u> | May 02, 2005 |
| <u>AP</u> | MYLAN LABS LTD | <u>10MG/ML</u> | <u>A206608 001</u> | Jun 08, 2018 |
| <u>AP</u> | | <u>20MG/ML</u> | <u>A206608 002</u> | Jun 08, 2018 |
| <u>AP</u> | SAGENT PHARMS | <u>10MG/ML</u> | <u>A207107 001</u> | Jun 08, 2018 |
| <u>AP</u> | | <u>20MG/ML</u> | <u>A207107 002</u> | Jun 08, 2018 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>10MG/ML</u> | <u>A076323 001</u> | Aug 10, 2004 |
| BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER | | | | |
| + | +! BAXTER HLTHCARE | 2GM/100ML | N019386 005 | Jan 27, 2003 |
| BREVIBLOC IN PLASTIC CONTAINER | | | | |
| + | +! BAXTER HLTHCARE | 1GM/100ML | N019386 004 | Feb 16, 2001 |
| SOLUTION; INTRAVENOUS | | | | |
| ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER | | | | |
| + | +! HQ SPCLT PHARMA | 2GM/100ML (20MG/ML) | N205703 002 | Apr 07, 2016 |
| ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER | | | | |
| + | +! HQ SPCLT PHARMA | 2.5GM/250ML (10MG/ML) | N205703 001 | Apr 07, 2016 |

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED REL PELLETS; ORAL

ESOMEPRAZOLE MAGNESIUM

| | | | | |
|-----------|-------------------------|---------------------|--------------------|--------------|
| <u>AB</u> | ALKEM LABS LTD | <u>EQ 20MG BASE</u> | <u>A208333 001</u> | Oct 20, 2017 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A208333 002</u> | Oct 20, 2017 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 20MG BASE</u> | <u>A205606 001</u> | Apr 21, 2016 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A205606 002</u> | Apr 21, 2016 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 20MG BASE</u> | <u>A078279 001</u> | Sep 25, 2015 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A078279 002</u> | Sep 25, 2015 |
| <u>AB</u> | HEC PHARM | <u>EQ 20MG BASE</u> | <u>A207265 002</u> | May 18, 2018 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A207265 001</u> | May 18, 2018 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>EQ 20MG BASE</u> | <u>A078003 001</u> | Jan 26, 2015 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A078003 002</u> | Jan 26, 2015 |
| <u>AB</u> | LANNETT CO INC | <u>EQ 20MG BASE</u> | <u>A205563 001</u> | Sep 01, 2017 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A205563 002</u> | Sep 01, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED REL PELLETS;ORAL

ESOMEPRAZOLE MAGNESIUM

| | | | | |
|-----------|--------------------|---------------------|--------------------|--------------|
| AB | MYLAN PHARMS INC | EQ 20MG BASE | A078936 001 | Aug 02, 2015 |
| AB | | EQ 40MG BASE | A078936 002 | Aug 03, 2015 |
| AB | SUN PHARM INDS LTD | EQ 20MG BASE | A209735 001 | Apr 30, 2018 |
| AB | | EQ 40MG BASE | A209735 002 | Apr 30, 2018 |
| AB | TORRENT PHARMS LTD | EQ 20MG BASE | A203636 001 | Oct 19, 2015 |
| AB | | EQ 40MG BASE | A203636 002 | Oct 19, 2015 |

NEXIUM

| | | | | | |
|-----------|---|--------------------|---------------------|--------------------|--------------|
| AB | + | ASTRAZENECA PHARMS | EQ 20MG BASE | N021153 001 | Feb 20, 2001 |
| AB | + | ! | EQ 40MG BASE | N021153 002 | Feb 20, 2001 |

ESOMEPRAZOLE MAGNESIUM

| | | | | |
|----|---------------------|--------------|-------------|--------------|
| BX | HETERO LABS LTD III | EQ 20MG BASE | A202784 001 | Sep 21, 2015 |
| BX | | EQ 40MG BASE | A202784 002 | Sep 21, 2015 |

FOR SUSPENSION, DELAYED RELEASE;ORAL

NEXIUM

| | | | | |
|---|--------------------|----------------------|-------------|--------------|
| + | ASTRAZENECA PHARMS | EQ 2.5MG BASE/PACKET | N021957 003 | Dec 15, 2011 |
| + | | EQ 5MG BASE/PACKET | N021957 004 | Dec 15, 2011 |
| + | | EQ 10MG BASE/PACKET | N022101 001 | Feb 27, 2008 |
| + | | EQ 20MG BASE/PACKET | N021957 001 | Oct 20, 2006 |
| + | ! | EQ 40MG BASE/PACKET | N021957 002 | Oct 20, 2006 |

ESOMEPRAZOLE MAGNESIUM; NAPROXEN

TABLET, DELAYED RELEASE;ORAL

VIMOVO

| | | | | |
|---|---------|--------------------|-------------|--------------|
| + | HORIZON | EQ 20MG BASE;375MG | N022511 002 | Apr 30, 2010 |
| + | ! | EQ 20MG BASE;500MG | N022511 001 | Apr 30, 2010 |

ESOMEPRAZOLE SODIUM

INJECTABLE; INTRAVENOUS

ESOMEPRAZOLE SODIUM

| | | | | | |
|-----------|-------------------|--------------------------|--------------------------|--------------------|--------------|
| AP | ACCORD HLTHCARE | EQ 40MG BASE/VIAL | A205379 001 | Sep 25, 2015 | |
| AP | ! | AUROBINDO PHARMA LTD | EQ 40MG BASE/VIAL | A204657 002 | Aug 10, 2016 |
| AP | DEVA HOLDING AS | EQ 40MG BASE/VIAL | A207181 001 | Mar 06, 2017 | |
| AP | MYLAN LABS LTD | EQ 40MG BASE/VIAL | A202686 002 | May 17, 2017 | |
| AP | SUN PHARMA GLOBAL | EQ 40MG BASE/VIAL | A200882 002 | Mar 18, 2013 | |

NEXIUM IV

| | | | | | | |
|-----------|---|---|--------------------|--------------------------|--------------------|--------------|
| AP | + | ! | ASTRAZENECA PHARMS | EQ 40MG BASE/VIAL | N021689 002 | Mar 31, 2005 |
|-----------|---|---|--------------------|--------------------------|--------------------|--------------|

ESOMEPRAZOLE STRONTIUM

CAPSULE, DELAYED RELEASE;ORAL

ESOMEPRAZOLE STRONTIUM

| | | | | |
|---|---------------|--------|-------------|--------------|
| + | R2 PHARMA LLC | 49.3MG | N202342 002 | Aug 06, 2013 |
|---|---------------|--------|-------------|--------------|

ESTAZOLAM

TABLET;ORAL

ESTAZOLAM

| | | | | |
|-----------|--------------|------------|--------------------|--------------|
| AB | MAYNE PHARMA | 1MG | A074921 001 | Jul 10, 1997 |
| AB | ! | 2MG | A074921 002 | Jul 10, 1997 |
| AB | PAR PHARM | 1MG | A074826 001 | Jul 03, 1997 |
| AB | | 2MG | A074826 002 | Jul 03, 1997 |
| AB | WATSON LABS | 1MG | A074818 001 | Aug 19, 1997 |
| AB | | 2MG | A074818 002 | Aug 19, 1997 |

ESTRADIOL

CREAM;VAGINAL

ESTRACE

| | | | | | |
|-----------|---|--------------------|--------------|--------------------|--------------|
| AB | ! | ALLERGAN SALES LLC | 0.01% | A086069 001 | Jan 31, 1984 |
|-----------|---|--------------------|--------------|--------------------|--------------|

ESTRADIOL

| | | | | |
|-----------|--------------------|--------------|--------------------|--------------|
| AB | ALVOGEN PINE BROOK | 0.01% | A209767 001 | Mar 05, 2018 |
| AB | MYLAN PHARMS INC | 0.01% | A208788 001 | Dec 29, 2017 |
| AB | PERRIGO UK FINCO | 0.01% | A210194 001 | Jan 22, 2018 |
| AB | TEVA PHARMS USA | 0.01% | A210488 001 | Mar 30, 2018 |

FILM, EXTENDED RELEASE;TRANSDERMAL

CLIMARA

| | | | | | |
|-----------|---|----------------|--------------------|--------------------|--------------|
| AB | + | BAYER HLTHCARE | 0.06MG/24HR | N020375 006 | May 27, 2003 |
|-----------|---|----------------|--------------------|--------------------|--------------|

ESTRADIOL

| | | | | |
|------------|--------------------|----------------------|--------------------|--------------|
| AB | MYLAN TECHNOLOGIES | 0.06MG/24HR | A075182 005 | Jul 20, 2006 |
| AB1 | | 0.025MG/24HR | A201675 001 | Dec 19, 2014 |
| AB1 | | 0.0375MG/24HR | A201675 002 | Dec 19, 2014 |
| AB1 | | 0.05MG/24HR | A201675 003 | Dec 19, 2014 |
| AB1 | | 0.075MG/24HR | A201675 004 | Dec 19, 2014 |
| AB1 | | 0.1MG/24HR | A201675 005 | Dec 19, 2014 |

PRESCRIPTION DRUG PRODUCT LIST

ESTRADIOL

FILM, EXTENDED RELEASE;TRANSDERMAL

VIVELLE-DOT

| | | | | | | |
|------------|---|----------|----------------------|----------------|------------|--------------|
| <u>AB1</u> | + | NOVARTIS | <u>0.025MG/24HR</u> | <u>N020538</u> | <u>009</u> | May 03, 2002 |
| <u>AB1</u> | + | | <u>0.0375MG/24HR</u> | <u>N020538</u> | <u>005</u> | Jan 08, 1999 |
| <u>AB1</u> | + | | <u>0.05MG/24HR</u> | <u>N020538</u> | <u>006</u> | Jan 08, 1999 |
| <u>AB1</u> | + | | <u>0.075MG/24HR</u> | <u>N020538</u> | <u>007</u> | Jan 08, 1999 |
| <u>AB1</u> | + | ! | <u>0.1MG/24HR</u> | <u>N020538</u> | <u>008</u> | Jan 08, 1999 |

CLIMARA

| | | | | | | |
|------------|---|----------------|----------------------|----------------|------------|--------------|
| <u>AB2</u> | + | BAYER HLTHCARE | <u>0.025MG/24HR</u> | <u>N020375</u> | <u>004</u> | Mar 05, 1999 |
| <u>AB2</u> | + | | <u>0.0375MG/24HR</u> | <u>N020375</u> | <u>005</u> | May 27, 2003 |
| <u>AB2</u> | + | | <u>0.05MG/24HR</u> | <u>N020375</u> | <u>001</u> | Dec 22, 1994 |
| <u>AB2</u> | + | | <u>0.075MG/24HR</u> | <u>N020375</u> | <u>003</u> | Mar 23, 1998 |
| <u>AB2</u> | + | ! | <u>0.1MG/24HR</u> | <u>N020375</u> | <u>002</u> | Dec 22, 1994 |

ESTRADIOL

| | | | | | | |
|------------|--|--------------------|----------------------|----------------|------------|--------------|
| <u>AB2</u> | | MYLAN TECHNOLOGIES | <u>0.025MG/24HR</u> | <u>A075182</u> | <u>003</u> | Jan 26, 2005 |
| <u>AB2</u> | | | <u>0.0375MG/24HR</u> | <u>A075182</u> | <u>004</u> | Jul 20, 2006 |
| <u>AB2</u> | | | <u>0.05MG/24HR</u> | <u>A075182</u> | <u>006</u> | Feb 24, 2000 |
| <u>AB2</u> | | | <u>0.075MG/24HR</u> | <u>A075182</u> | <u>002</u> | Jan 26, 2005 |
| <u>AB2</u> | | | <u>0.1MG/24HR</u> | <u>A075182</u> | <u>001</u> | Feb 24, 2000 |
| <u>AB3</u> | | | <u>0.025MG/24HR</u> | <u>A206685</u> | <u>001</u> | Aug 15, 2018 |
| <u>AB3</u> | | | <u>0.0375MG/24HR</u> | <u>A206685</u> | <u>002</u> | Aug 15, 2018 |
| <u>AB3</u> | | | <u>0.05MG/24HR</u> | <u>A206685</u> | <u>003</u> | Aug 15, 2018 |
| <u>AB3</u> | | | <u>0.075MG/24HR</u> | <u>A206685</u> | <u>004</u> | Aug 15, 2018 |
| <u>AB3</u> | | | <u>0.1MG/24HR</u> | <u>A206685</u> | <u>005</u> | Aug 15, 2018 |

MINIVELLE

| | | | | | | |
|------------|---|-------|----------------------|----------------|------------|--------------|
| <u>AB3</u> | + | NOVEN | <u>0.025MG/24HR</u> | <u>N203752</u> | <u>005</u> | Sep 23, 2014 |
| <u>AB3</u> | + | | <u>0.0375MG/24HR</u> | <u>N203752</u> | <u>001</u> | Oct 29, 2012 |
| <u>AB3</u> | + | | <u>0.05MG/24HR</u> | <u>N203752</u> | <u>003</u> | Oct 29, 2012 |
| <u>AB3</u> | + | | <u>0.075MG/24HR</u> | <u>N203752</u> | <u>002</u> | Oct 29, 2012 |
| <u>AB3</u> | + | ! | <u>0.1MG/24HR</u> | <u>N203752</u> | <u>004</u> | Oct 29, 2012 |

ALORA

| | | | | | | |
|----|--|--------------------|--------------|---------|-----|--------------|
| BX | | ALLERGAN SALES LLC | 0.025MG/24HR | N020655 | 004 | Apr 05, 2002 |
| BX | | | 0.05MG/24HR | N020655 | 001 | Dec 20, 1996 |
| BX | | | 0.075MG/24HR | N020655 | 002 | Dec 20, 1996 |
| BX | | | 0.1MG/24HR | N020655 | 003 | Dec 20, 1996 |

MENOSTAR

| | | | | | | |
|---|---|----------------|--------------|---------|-----|--------------|
| + | ! | BAYER HLTHCARE | 0.014MG/24HR | N021674 | 001 | Jun 08, 2004 |
|---|---|----------------|--------------|---------|-----|--------------|

GEL;TRANSDERMAL

DIVIGEL

| | | | | | | |
|---|--|---------------------|------|---------|-----|--------------|
| + | | VERTICAL PHARMS LLC | 0.1% | N022038 | 001 | Jun 04, 2007 |
|---|--|---------------------|------|---------|-----|--------------|

GEL, METERED;TRANSDERMAL

ELESTRIN

| | | | | | | |
|---|---|---------------------|---------------------------|---------|-----|--------------|
| + | ! | MYLAN SPECIALITY LP | 0.06% (0.87GM/ACTIVATION) | N021813 | 001 | Dec 15, 2006 |
|---|---|---------------------|---------------------------|---------|-----|--------------|

ESTROGEL

| | | | | | | |
|---|---|-------------------|---------------------------|---------|-----|--------------|
| + | ! | ASCEND THERAPS US | 0.06% (1.25GM/ACTIVATION) | N021166 | 002 | Feb 09, 2004 |
|---|---|-------------------|---------------------------|---------|-----|--------------|

INSERT;VAGINAL

IMVEXXY

| | | | | | | |
|---|---|--------------------|---------|---------|-----|--------------|
| + | | THERAPEUTICSMD INC | 0.004MG | N208564 | 001 | May 29, 2018 |
| + | ! | | 0.01MG | N208564 | 002 | May 29, 2018 |

INSERT, EXTENDED RELEASE;VAGINAL

ESTRING

| | | | | | | |
|---|---|-------------------------|---------------|---------|-----|--------------|
| + | ! | PHARMACIA AND UPJOHN | 0.0075MG/24HR | N020472 | 001 | Apr 26, 1996 |
|---|---|-------------------------|---------------|---------|-----|--------------|

SPRAY;TRANSDERMAL

EVAMIST

| | | | | | | |
|---|---|---------------------|--------------|---------|-----|--------------|
| + | ! | PERRIGO PHARMA INTL | 1.53MG/SPRAY | N022014 | 001 | Jul 27, 2007 |
|---|---|---------------------|--------------|---------|-----|--------------|

TABLET;ORAL

ESTRADIOL

| | | | | | | |
|-----------|--|-----------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | BARR LABS INC | <u>0.5MG</u> | <u>A040197</u> | <u>001</u> | Oct 22, 1997 |
| <u>AB</u> | | | <u>1MG</u> | <u>A040197</u> | <u>002</u> | Oct 22, 1997 |
| <u>AB</u> | | ! | <u>2MG</u> | <u>A040197</u> | <u>003</u> | Oct 22, 1997 |
| <u>AB</u> | | EPIC PHARMA INC | <u>0.5MG</u> | <u>A040275</u> | <u>001</u> | Dec 29, 1998 |
| <u>AB</u> | | | <u>1MG</u> | <u>A040275</u> | <u>002</u> | Dec 29, 1998 |
| <u>AB</u> | | | <u>2MG</u> | <u>A040275</u> | <u>003</u> | Dec 29, 1998 |
| <u>AB</u> | | MAYNE PHARMA | <u>0.5MG</u> | <u>A040114</u> | <u>003</u> | Mar 14, 1996 |
| <u>AB</u> | | | <u>1MG</u> | <u>A040114</u> | <u>001</u> | Mar 14, 1996 |
| <u>AB</u> | | | <u>2MG</u> | <u>A040114</u> | <u>002</u> | Mar 14, 1996 |
| <u>AB</u> | | MYLAN | <u>0.5MG</u> | <u>A040326</u> | <u>001</u> | Apr 21, 1999 |
| <u>AB</u> | | | <u>1MG</u> | <u>A040326</u> | <u>002</u> | Apr 21, 1999 |
| <u>AB</u> | | | <u>2MG</u> | <u>A040326</u> | <u>003</u> | Apr 21, 1999 |

PRESCRIPTION DRUG PRODUCT LIST

ESTRADIOL

TABLET;VAGINAL

ESTRADIOL

| | | | | |
|-----------|---------------------|--------------|--------------------|--------------|
| AB | AMNEAL PHARMS | 10MCG | A205256 001 | May 29, 2015 |
| AB | GLENMARK PHARMS LTD | 10MCG | A210264 001 | Sep 14, 2018 |
| AB | TEVA PHARMS USA | 10MCG | A206388 001 | Jul 21, 2017 |

VAGIFEM

| | | | | | |
|-----------|------------|------------------|--------------|--------------------|--------------|
| AB | + ! | NOVO NORDISK INC | 10MCG | N020908 002 | Nov 25, 2009 |
|-----------|------------|------------------|--------------|--------------------|--------------|

ESTRADIOL ACETATEINSERT, EXTENDED RELEASE;VAGINAL
FEMRING

| | | | | |
|------------|-----------|---------------------|-------------|--------------|
| + | MILLICENT | EQ 0.05MG BASE/24HR | N021367 001 | Mar 20, 2003 |
| + ! | | EQ 0.1MG BASE/24HR | N021367 002 | Mar 20, 2003 |

ESTRADIOL CYPIONATE

INJECTABLE;INJECTION

DEPO-ESTRADIOL

| | | | | |
|---|-------------------------|--------|-------------|--|
| ! | PHARMACIA AND UPJOHN | 5MG/ML | A085470 003 | |
|---|-------------------------|--------|-------------|--|

ESTRADIOL VALERATE

INJECTABLE;INJECTION

DELESTROGEN

| | | | | | |
|-----------|------------|-------------------------|----------------|--------------------|--|
| AO | + ! | PAR STERILE PRODUCTS | 20MG/ML | N009402 004 | |
| AO | + ! | | 40MG/ML | N009402 003 | |

ESTRADIOL VALERATE

| | | | | | |
|-----------|------------|-------------------------|----------------|--------------------|--------------|
| AO | | LUITPOLD | 20MG/ML | A090920 001 | Jan 19, 2010 |
| AO | | | 40MG/ML | A090920 002 | Jan 19, 2010 |
| | | DELESTROGEN | | | |
| | + ! | PAR STERILE PRODUCTS | 10MG/ML | N009402 002 | |

ESTRADIOL; LEVONORGESTRELFILM, EXTENDED RELEASE;TRANSDERMAL
CLIMARA PRO

| | | | | |
|------------|----------------|---------------------------|-------------|--------------|
| + ! | BAYER HLTHCARE | 0.045MG/24HR;0.015MG/24HR | N021258 001 | Nov 21, 2003 |
|------------|----------------|---------------------------|-------------|--------------|

ESTRADIOL; NORETHINDRONE ACETATEFILM, EXTENDED RELEASE;TRANSDERMAL
COMBIPATCH

| | | | | |
|------------|------------------|-------------------------|-------------|--------------|
| + | NOVEN PHARMS INC | 0.05MG/24HR;0.14MG/24HR | N020870 001 | Aug 07, 1998 |
| + ! | | 0.05MG/24HR;0.25MG/24HR | N020870 002 | Aug 07, 1998 |

TABLET;ORAL

ACTIVELLA

| | | | | | |
|-----------|------------|-------------------|--------------------|--------------------|--------------|
| AB | + | AMNEAL PHARMS LLC | 0.5MG;0.1MG | N020907 002 | Dec 28, 2006 |
| AB | + ! | | 1MG;0.5MG | N020907 001 | Nov 18, 1998 |

AMABELZ

| | | | | | |
|-----------|--|-----------|--------------------|--------------------|--------------|
| AB | | LUPIN LTD | 0.5MG;0.1MG | A203339 001 | Jun 20, 2016 |
| AB | | | 1MG;0.5MG | A203339 002 | Jun 20, 2016 |

ESTRADIOL AND NORETHINDRONE ACETATE

| | | | | | |
|-----------|--|--------------------|--------------------|--------------------|--------------|
| AB | | ACCORD HLTHCARE | 1MG;0.5MG | A210233 001 | Feb 28, 2018 |
| AB | | BARR | 1MG;0.5MG | A079193 001 | May 11, 2010 |
| AB | | BRECKENRIDGE PHARM | 0.5MG;0.1MG | A078324 002 | Jun 09, 2011 |
| AB | | | 1MG;0.5MG | A078324 001 | Apr 17, 2008 |
| AB | | MYLAN LABS LTD | 0.5MG;0.1MG | A207261 001 | Feb 10, 2017 |
| AB | | | 1MG;0.5MG | A207261 002 | Feb 10, 2017 |
| AB | | TEVA PHARMS USA | 0.5MG;0.1MG | A200747 001 | Mar 08, 2012 |

ESTRADIOL; NORGESTIMATE

TABLET;ORAL

ESTRADIOL AND NORGESTIMATE

| | | | | |
|---|------|--------------------|-------------|--------------|
| ! | BARR | 1MG,1MG;N/A,0.09MG | A076812 001 | Apr 29, 2005 |
|---|------|--------------------|-------------|--------------|

ESTRADIOL; PROGESTERONE

CAPSULE;ORAL

BIJUVA

| | | | | |
|------------|--------------------|-----------|-------------|--------------|
| + ! | THERAPEUTICSMD INC | 1MG;100MG | N210132 001 | Oct 28, 2018 |
|------------|--------------------|-----------|-------------|--------------|

ESTRAMUSTINE PHOSPHATE SODIUM

CAPSULE;ORAL

EMCYT

| | | | | |
|------------|-------------------------|--------------------|-------------|--|
| + ! | PHARMACIA AND UPJOHN | EQ 140MG PHOSPHATE | N018045 001 | |
|------------|-------------------------|--------------------|-------------|--|

PRESCRIPTION DRUG PRODUCT LIST

ESTROGENS, CONJUGATED

CREAM; TOPICAL, VAGINAL

PREMARIN

+! WYETH PHARMS 0.625MG/GM N020216 001

INJECTABLE; INJECTION

PREMARIN

+! WYETH PHARMS 25MG/VIAL N010402 001

TABLET; ORAL

PREMARIN

+ WYETH PHARMS 0.3MG N004782 003

+ 0.45MG N004782 006 Jul 16, 2003

+! 0.625MG N004782 004

+! 0.9MG N004782 005 Jan 26, 1984

+! 1.25MG N004782 001

ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE

TABLET; ORAL-28

PREMPHASE 14/14

+! WYETH PHARMS 0.625MG, 0.625MG; N/A, 5MG N020527 002 Nov 17, 1995

PREMPRO

+! WYETH PHARMS 0.3MG; 1.5MG N020527 005 Jun 04, 2003

+! 0.45MG; 1.5MG N020527 004 Mar 12, 2003

+! 0.625MG; 2.5MG N020527 001 Nov 17, 1995

+! 0.625MG; 5MG N020527 003 Jan 09, 1998

ESTROGENS, ESTERIFIED

TABLET; ORAL

MENEST

MONARCH PHARMS 0.3MG A084951 001

0.625MG A084948 001

1.25MG A084950 001

! 2.5MG A084949 001

ESTROPIPATE

TABLET; ORAL

ESTROPIPATE

MYLAN 0.75MG A040359 001 Aug 26, 1999

1.5MG A040359 002 Aug 26, 1999

OGEN 5

PHARMACIA AND 6MG A083220 004

UPJOHN

ESZOPICLONE

TABLET; ORAL

ESZOPICLONE

| | | | | |
|-----------|----------------------|------------|--------------------|--------------|
| AB | AUROBINDO PHARMA LTD | 1MG | A208451 001 | Sep 15, 2016 |
| AB | | 2MG | A208451 002 | Sep 15, 2016 |
| AB | | 3MG | A208451 003 | Sep 15, 2016 |
| AB | DR REDDYS LABS LTD | 1MG | A091024 001 | Apr 15, 2014 |
| AB | | 2MG | A091024 002 | Apr 15, 2014 |
| AB | | 3MG | A091024 003 | Apr 15, 2014 |
| AB | GLENMARK GENERICS | 1MG | A091166 001 | Apr 15, 2014 |
| AB | | 2MG | A091166 002 | Apr 15, 2014 |
| AB | | 3MG | A091166 003 | Apr 15, 2014 |
| AB | LUPIN LTD | 1MG | A091124 001 | Sep 13, 2011 |
| AB | | 2MG | A091124 002 | Sep 13, 2011 |
| AB | | 3MG | A091124 003 | Sep 13, 2011 |
| AB | MACLEODS PHARMS LTD | 1MG | A202929 001 | Jan 30, 2015 |
| AB | | 2MG | A202929 002 | Jan 30, 2015 |
| AB | | 3MG | A202929 003 | Jan 30, 2015 |
| AB | MYLAN PHARMS INC | 1MG | A091151 001 | Mar 26, 2013 |
| AB | | 2MG | A091151 002 | Mar 26, 2013 |
| AB | | 3MG | A091151 003 | Mar 26, 2013 |
| AB | ORCHID HLTHCARE | 1MG | A091113 001 | Jun 10, 2014 |
| AB | | 2MG | A091113 002 | Jun 10, 2014 |
| AB | | 3MG | A091113 003 | Jun 10, 2014 |
| AB | SUN PHARMA GLOBAL | 1MG | A091103 001 | Apr 03, 2013 |
| AB | | 2MG | A091103 002 | Apr 03, 2013 |
| AB | | 3MG | A091103 003 | Apr 03, 2013 |
| AB | TEVA | 1MG | A091169 001 | May 23, 2011 |
| AB | | 2MG | A091169 002 | May 23, 2011 |
| AB | | 3MG | A091169 003 | May 23, 2011 |
| AB | WEST-WARD PHARMS INT | 1MG | A091153 001 | Apr 15, 2014 |

PRESCRIPTION DRUG PRODUCT LIST

ESZOPICLONE

TABLET; ORAL

ESZOPICLONE

| | | | | |
|-----------|--|------------|--------------------|--------------|
| AB | | 2MG | A091153 002 | Apr 15, 2014 |
| AB | | 3MG | A091153 003 | Apr 15, 2014 |

LUNESTA

| | | | | | |
|-----------|---|---------------------|------------|--------------------|--------------|
| AB | + | SUNOVION PHARMS INC | 1MG | N021476 001 | Dec 15, 2004 |
| AB | + | | 2MG | N021476 002 | Dec 15, 2004 |
| AB | + | ! | 3MG | N021476 003 | Dec 15, 2004 |

ETELICALCETIDE

SOLUTION; INTRAVENOUS

PARSABIV

| | | | | | | |
|--|---|---|----------------|---------------------------|-------------|--------------|
| | + | ! | KAI PHARMS INC | 2.5MG/0.5ML (2.5MG/0.5ML) | N208325 001 | Feb 07, 2017 |
| | + | ! | | 5MG/ML (5MG/ML) | N208325 002 | Feb 07, 2017 |
| | + | ! | | 10MG/2ML (5MG/ML) | N208325 003 | Feb 07, 2017 |

ETEPLIRSEN

SOLUTION; INTRAVENOUS

EXONDYS 51

| | | | | | | |
|--|---|---|---------------------|----------------------|-------------|--------------|
| | + | ! | SAREPTA THERAPS INC | 100MG/2ML (50MG/ML) | N206488 001 | Sep 19, 2016 |
| | + | ! | | 500MG/10ML (50MG/ML) | N206488 002 | Sep 19, 2016 |

ETHACRYNATE SODIUM

INJECTABLE; INJECTION

EDECRIN

| | | | | | | |
|----------------------------------|---|---|----------------------|--------------------------|--------------------|--------------|
| AP | + | ! | ATON | EQ 50MG BASE/VIAL | N016093 001 | |
| <u>ETHACRYNATE SODIUM</u> | | | | | | |
| AP | | | MYLAN INSTITUTIONAL | EQ 50MG BASE/VIAL | A204634 001 | Aug 23, 2016 |
| AP | | | PAR STERILE PRODUCTS | EQ 50MG BASE/VIAL | A205473 001 | Jul 29, 2015 |
| AP | | | ZYDUS PHARMS USA INC | EQ 50MG BASE/VIAL | A207758 001 | Nov 17, 2017 |

ETHACRYNIC ACID

TABLET; ORAL

EDECRIN

| | | | | | | |
|-------------------------------|---|---|----------------------|-------------|--------------------|--------------|
| AB | + | ! | ATON | 25MG | N016092 001 | |
| <u>ETHACRYNIC ACID</u> | | | | | | |
| AB | | | ALVOGEN | 25MG | A205709 001 | Jul 24, 2018 |
| AB | | | AMNEAL PHARMS CO | 25MG | A208805 001 | May 08, 2018 |
| AB | | | EDENBRIDGE PHARMS | 25MG | A205609 001 | Jun 30, 2016 |
| AB | | | PAR PHARM INC | 25MG | A208501 001 | Jul 21, 2017 |
| AB | | | WEST-WARD PHARMS INT | 25MG | A207262 001 | Feb 23, 2017 |

ETHAMBUTOL HYDROCHLORIDE

TABLET; ORAL

ETHAMBUTOL HYDROCHLORIDE

| | | | | | | |
|-------------------------|---|---|----------------|--------------|--------------------|--------------|
| AB | | | AKORN | 100MG | A075095 001 | Nov 30, 1999 |
| AB | | | | 400MG | A075095 002 | Nov 30, 1999 |
| AB | | | BARR | 400MG | A076057 001 | Nov 26, 2001 |
| AB | | | LUPIN | 100MG | A078939 001 | Jun 17, 2009 |
| AB | | | | 400MG | A078939 002 | Jun 17, 2009 |
| <u>MYAMBUTOL</u> | | | | | | |
| AB | + | | STI PHARMA LLC | 100MG | N016320 001 | |
| AB | + | ! | | 400MG | N016320 003 | |

ETHANOLAMINE OLEATE

INJECTABLE; INJECTION

ETHAMOLIN

| | | | | | | |
|--|---|---|-----------|---------|-------------|--------------|
| | + | ! | QOL MEDCL | 50MG/ML | N019357 001 | Dec 22, 1988 |
|--|---|---|-----------|---------|-------------|--------------|

ETHINYL ESTRADIOL; ETHYNODIOL DIACETATE

TABLET; ORAL-28

ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL

| | | | | | | |
|------------------------------|---|--|----------------|--------------------|--------------------|--------------|
| AB | | | MYLAN LABS LTD | 0.035MG;1MG | A204703 001 | Jul 28, 2016 |
| AB | | | | 0.05MG;1MG | A204704 001 | Feb 09, 2016 |
| <u>KELNOR</u> | | | | | | |
| AB | | | BARR | 0.035MG;1MG | A076785 001 | May 23, 2005 |
| <u>MALMOREDE</u> | | | | | | |
| AB | | | NOVAST LABS | 0.05MG;1MG | A209547 001 | Jul 25, 2018 |
| <u>ZOVIA 1/35E-28</u> | | | | | | |
| AB | | | MAYNE PHARMA | 0.035MG;1MG | A072721 001 | Dec 30, 1991 |
| <u>ZOVIA 1/50E-28</u> | | | | | | |
| AB | ! | | WATSON LABS | 0.05MG;1MG | A072723 001 | Dec 30, 1991 |

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; ETONOGESTREL

RING;VAGINAL

NUVARING

+! ORGANON SUB MERCK 0.015MG/24HR;0.12MG/24HR N021187 001 Oct 03, 2001

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET;ORAL

ASHLYNA

AB GLENMARK GENERICS 0.03MG,0.01MG;0.15MG,N/A A203163 001 Feb 23, 2015

DAYSEE

AB LUPIN LTD 0.03MG,0.01MG;0.15MG,N/A A091467 001 Apr 10, 2013

FAYOSIM

AB LUPIN LTD 0.02MG,0.15MG;0.025MG,0.15MG;0.03MG,0.15MG;0.01MG,N/A A205943 001 Mar 29, 2016

ICLEVIA

AB AUROBINDO PHARMA LTD 0.03MG;0.15MG A206850 001 Jun 29, 2018

INTROVALE

AB LABS LEON FARMA 0.03MG;0.15MG A079064 001 Sep 27, 2010

JAIMIESS

AB LABS LEON FARMA 0.03MG,0.01MG;0.15MG,N/A A203770 001 Dec 27, 2017

LEVONORGESTREL AND ETHINYL ESTRADIOL

AB AMNEAL PHARMS 0.03MG;0.15MG A203871 001 Nov 13, 2015

AB 0.03MG,0.01MG;0.15MG,N/A A203872 001 Dec 22, 2015

AB GLENMARK GENERICS 0.02MG;0.09MG A202791 001 Apr 09, 2015

AB GLENMARK PHARMS LTD 0.03MG;0.15MG A203164 001 Jun 12, 2015

AB LUPIN LTD 0.03MG;0.15MG A091440 001 Oct 23, 2012

AB MYLAN LABS LTD 0.03MG;0.15MG A200490 001 Apr 21, 2015

AB ! WATSON LABS 0.02MG;0.09MG A079218 001 Jun 06, 2011

LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL

AB LABS LEON FARMA 0.02MG,0.1MG;0.01MG,N/A A205131 001 Dec 14, 2017

AB LUPIN LTD 0.02MG,0.1MG;0.01MG,N/A A091674 001 Oct 26, 2011

AB MAYNE PHARMA 0.02MG,0.1MG;0.01MG,N/A A200407 001 Oct 25, 2011

AB 0.03MG,0.01MG;0.15MG,N/A A078834 001 May 31, 2011

AB MYLAN LABS LTD 0.02MG,0.15MG; A206053 001 Oct 02, 2017

0.025MG,0.15MG;0.03MG,0.15MG;

0.02MG,0.1MG;0.01MG,N/A A200493 001 Jun 17, 2015

0.03MG,0.01MG;0.15MG,N/A A200492 001 May 27, 2015

LO SIMPESSÉ

AB AUROBINDO PHARMA LTD 0.02MG,0.1MG;0.01MG,N/A A206852 001 Apr 28, 2017

LOSEASONIQUE

AB TEVA BRANDED PHARM 0.02MG,0.1MG;0.01MG,N/A N022262 001 Oct 24, 2008

QUARTETTE

AB +! TEVA BRANDED PHARM 0.02MG,0.15MG;0.025MG,0.15MG;0.03MG,0.15MG;0.01MG,N/A N204061 001 Mar 28, 2013

QUASENSE

AB WATSON LABS 0.03MG;0.15MG A077101 001 Sep 06, 2006

SEASONALE

AB +! TEVA BRANDED PHARM 0.03MG;0.15MG N021544 001 Sep 05, 2003

SEASONIQUE

AB +! TEVA BRANDED PHARM 0.03MG,0.01MG;0.15MG,N/A N021840 001 May 25, 2006

SETLAKIN

AB NOVAST LABS 0.03MG;0.15MG A090716 001 Sep 15, 2014

SIMPESSÉ

AB AUROBINDO PHARMA LTD 0.03MG,0.01MG;0.15MG,N/A A206851 001 Apr 07, 2017

BALCOLTRA

AVION PHARMS 0.02MG;0.1MG N208612 001 Jan 09, 2018

TABLET;ORAL-28

ALTAVERA

AB LABS LEON FARMA 0.03MG;0.15MG A079102 001 Aug 03, 2010

AYUNA

AB AUROBINDO PHARMA LTD 0.03MG;0.15MG A206866 001 Sep 23, 2016

ELIFEMME

AB LABS LEON FARMA 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG A202507 001 Dec 04, 2015

ENPRESSE-28

AB DURAMED PHARMS BARR 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG A075809 002 Jul 16, 2001

KURVELO

AB LUPIN LTD 0.03MG;0.15MG A091408 001 Oct 17, 2012

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL-28

LEVONEST

| | | | | |
|-----------|-----------------|---------------------------------------------------------|--------------------|--------------|
| AB | NOVAST LABS LTD | <u>0.03MG, 0.04MG, 0.03MG; 0.05MG, 0.075MG, 0.125MG</u> | <u>A090719 001</u> | Dec 29, 2010 |
|-----------|-----------------|---------------------------------------------------------|--------------------|--------------|

LEVONORGESTREL AND ETHINYL ESTRADIOL

| | | | | |
|-----------|---------------|-----------------------|--------------------|--------------|
| AB | AMNEAL PHARMS | <u>0.03MG; 0.15MG</u> | <u>A201095 001</u> | Dec 08, 2014 |
|-----------|---------------|-----------------------|--------------------|--------------|

| | | | | |
|-----------|-----------|---------------------------------------------------------|--------------------|--------------|
| AB | LUPIN LTD | <u>0.03MG, 0.04MG, 0.03MG; 0.05MG, 0.075MG, 0.125MG</u> | <u>A200248 001</u> | Nov 19, 2015 |
|-----------|-----------|---------------------------------------------------------|--------------------|--------------|

| | | | | |
|-----------|----------------|---------------------------------------------------------|--------------------|--------------|
| AB | MYLAN LABS LTD | <u>0.03MG, 0.04MG, 0.03MG; 0.05MG, 0.075MG, 0.125MG</u> | <u>A202970 001</u> | Mar 23, 2018 |
|-----------|----------------|---------------------------------------------------------|--------------------|--------------|

| | | | | |
|-----------|--|-----------------------|--------------------|--------------|
| AB | | <u>0.03MG; 0.15MG</u> | <u>A091663 001</u> | Dec 21, 2012 |
|-----------|--|-----------------------|--------------------|--------------|

LEVORA 0.15/30-28

| | | | | |
|-----------|----------------|-----------------------|--------------------|--------------|
| AB | ! MAYNE PHARMA | <u>0.03MG; 0.15MG</u> | <u>A073594 001</u> | Dec 13, 1993 |
|-----------|----------------|-----------------------|--------------------|--------------|

MARLISSA

| | | | | |
|-----------|-------------------|-----------------------|--------------------|--------------|
| AB | GLENMARK GENERICS | <u>0.03MG; 0.15MG</u> | <u>A091452 001</u> | Feb 29, 2012 |
|-----------|-------------------|-----------------------|--------------------|--------------|

MYZILRA

| | | | | |
|-----------|--------------------|---------------------------------------------------------|--------------------|--------------|
| AB | VINTAGE PHARMS LLC | <u>0.03MG, 0.04MG, 0.03MG; 0.05MG, 0.075MG, 0.125MG</u> | <u>A077502 001</u> | Nov 23, 2011 |
|-----------|--------------------|---------------------------------------------------------|--------------------|--------------|

PORTIA-28

| | | | | |
|-----------|------|-----------------------|--------------------|--------------|
| AB | BARR | <u>0.03MG; 0.15MG</u> | <u>A075866 002</u> | May 23, 2002 |
|-----------|------|-----------------------|--------------------|--------------|

TRIVORA-28

| | | | | |
|-----------|----------------|---------------------------------------------------------|--------------------|--------------|
| AB | ! MAYNE PHARMA | <u>0.03MG, 0.04MG, 0.03MG; 0.05MG, 0.075MG, 0.125MG</u> | <u>A074538 002</u> | Dec 18, 1997 |
|-----------|----------------|---------------------------------------------------------|--------------------|--------------|

AFIRMELLE

| | | | | |
|------------|----------------------|----------------------|--------------------|--------------|
| AB1 | AUROBINDO PHARMA LTD | <u>0.02MG; 0.1MG</u> | <u>A206886 001</u> | Nov 14, 2016 |
|------------|----------------------|----------------------|--------------------|--------------|

AVIANE-28

| | | | | |
|------------|---------------------|----------------------|--------------------|--------------|
| AB1 | DURAMED PHARMS BARR | <u>0.02MG; 0.1MG</u> | <u>A075796 001</u> | Apr 30, 2001 |
|------------|---------------------|----------------------|--------------------|--------------|

CERINTA

| | | | | |
|------------|--------------------|----------------------|--------------------|--------------|
| AB1 | SUN PHARM INDS LTD | <u>0.02MG; 0.1MG</u> | <u>A202817 001</u> | Jan 07, 2019 |
|------------|--------------------|----------------------|--------------------|--------------|

FALMINA

| | | | | |
|------------|-----------------|----------------------|--------------------|--------------|
| AB1 | NOVAST LABS LTD | <u>0.02MG; 0.1MG</u> | <u>A090721 001</u> | Mar 28, 2012 |
|------------|-----------------|----------------------|--------------------|--------------|

LEVONORGESTREL AND ETHINYL ESTRADIOL

| | | | | |
|------------|---------------|----------------------|--------------------|--------------|
| AB1 | AMNEAL PHARMS | <u>0.02MG; 0.1MG</u> | <u>A201108 001</u> | Feb 05, 2014 |
|------------|---------------|----------------------|--------------------|--------------|

| | | | | |
|------------|-----------|----------------------|--------------------|--------------|
| AB1 | LUPIN LTD | <u>0.02MG; 0.1MG</u> | <u>A091425 001</u> | Jan 18, 2013 |
|------------|-----------|----------------------|--------------------|--------------|

| | | | | |
|------------|----------------|----------------------|--------------------|--------------|
| AB1 | ! MAYNE PHARMA | <u>0.02MG; 0.1MG</u> | <u>A076625 001</u> | Nov 18, 2004 |
|------------|----------------|----------------------|--------------------|--------------|

| | | | | |
|------------|----------------|----------------------|--------------------|--------------|
| AB1 | MYLAN LABS LTD | <u>0.02MG; 0.1MG</u> | <u>A200245 001</u> | Oct 09, 2013 |
|------------|----------------|----------------------|--------------------|--------------|

ORSYTHIA

| | | | | |
|------------|--------------------|----------------------|--------------------|--------------|
| AB1 | VINTAGE PHARMS LLC | <u>0.02MG; 0.1MG</u> | <u>A077099 001</u> | May 11, 2011 |
|------------|--------------------|----------------------|--------------------|--------------|

VIENVA

| | | | | |
|------------|-----------------|----------------------|--------------------|--------------|
| AB1 | LABS LEON FARMA | <u>0.02MG; 0.1MG</u> | <u>A201088 001</u> | May 21, 2015 |
|------------|-----------------|----------------------|--------------------|--------------|

LESSINA-28

| | | | | |
|------------|------|----------------------|--------------------|--------------|
| AB2 | BARR | <u>0.02MG; 0.1MG</u> | <u>A075803 002</u> | Mar 20, 2002 |
|------------|------|----------------------|--------------------|--------------|

LEVONORGESTREL AND ETHINYL ESTRADIOL

| | | | | |
|------------|----------------|----------------------|--------------------|--------------|
| AB2 | ! MAYNE PHARMA | <u>0.02MG; 0.1MG</u> | <u>A077681 001</u> | May 31, 2006 |
|------------|----------------|----------------------|--------------------|--------------|

| | | | | |
|------------|----------------|----------------------|--------------------|--------------|
| AB2 | MYLAN LABS LTD | <u>0.02MG; 0.1MG</u> | <u>A202247 001</u> | Dec 08, 2014 |
|------------|----------------|----------------------|--------------------|--------------|

ETHINYL ESTRADIOL; NORELGESTROMIN

FILM, EXTENDED RELEASE; TRANSDERMAL

XULANE

| | | | | |
|---|--------------------|---------------------------|-------------|--------------|
| ! | MYLAN TECHNOLOGIES | 0.035MG/24HR; 0.15MG/24HR | A200910 001 | Apr 16, 2014 |
|---|--------------------|---------------------------|-------------|--------------|

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21

NORINYL 1+35 21-DAY

| | | | | |
|-----------|--------------------|---------------------|--------------------|--|
| AB | ALLERGAN SALES LLC | <u>0.035MG; 1MG</u> | <u>N017565 001</u> | |
|-----------|--------------------|---------------------|--------------------|--|

NORTREL 1/35-21

| | | | | |
|-----------|------|---------------------|--------------------|--------------|
| AB | BARR | <u>0.035MG; 1MG</u> | <u>A072693 001</u> | Feb 28, 1992 |
|-----------|------|---------------------|--------------------|--------------|

NORTREL 7/7/7

| | | | | |
|------|--|-----------------------------------------------|-------------|--------------|
| BARR | | 0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MG | A075478 001 | Aug 30, 2002 |
|------|--|-----------------------------------------------|-------------|--------------|

TABLET; ORAL-28

ALYACEN 1/35

| | | | | |
|-----------|-------------------|---------------------|--------------------|--------------|
| AB | GLENMARK GENERICS | <u>0.035MG; 1MG</u> | <u>A091634 001</u> | Jan 19, 2012 |
|-----------|-------------------|---------------------|--------------------|--------------|

ALYACEN 7/7/7

| | | | | |
|-----------|-------------------|------------------------------------------------------|--------------------|--------------|
| AB | GLENMARK GENERICS | <u>0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MG</u> | <u>A091636 001</u> | Jan 19, 2012 |
|-----------|-------------------|------------------------------------------------------|--------------------|--------------|

ARANELLE

| | | | | |
|-----------|------|-----------------------------------------------------|--------------------|--------------|
| AB | BARR | <u>0.035MG, 0.035MG, 0.035MG; 0.5MG, 1MG, 0.5MG</u> | <u>A076783 001</u> | Sep 29, 2004 |
|-----------|------|-----------------------------------------------------|--------------------|--------------|

BALZIVA-28

| | | | | |
|-----------|--------|-----------------------|--------------------|--------------|
| AB | ! BARR | <u>0.035MG; 0.4MG</u> | <u>A076238 001</u> | Apr 22, 2004 |
|-----------|--------|-----------------------|--------------------|--------------|

BREVICON 28-DAY

| | | | | |
|-----------|--------------------|-----------------------|--------------------|--|
| AB | ALLERGAN SALES LLC | <u>0.035MG; 0.5MG</u> | <u>N017743 001</u> | |
|-----------|--------------------|-----------------------|--------------------|--|

PRESCRIPTION DRUG PRODUCT LISTETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-28

| | | | | | | |
|------------------------------------------------------------------------|----------------------|-------------------------------------------------|----------------|------------|---------|------|
| <u>BRIELLYN</u> | | | | | | |
| AB | GLENMARK GENERICS | 0.035MG;0.4MG | A090538 | 001 | Mar 22, | 2011 |
| <u>CYCLAFEM 0.5/35</u> | | | | | | |
| AB | VINTAGE PHARMS | 0.035MG;0.5MG | A203413 | 001 | Dec 16, | 2015 |
| <u>CYCLAFEM 1/35</u> | | | | | | |
| AB | VINTAGE PHARMS LLC | 0.035MG;1MG | A076337 | 001 | Nov 12, | 2010 |
| <u>CYCLAFEM 7/7/7</u> | | | | | | |
| AB | VINTAGE PHARMS LLC | 0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG | A076338 | 001 | Nov 16, | 2010 |
| <u>CYONANZ</u> | | | | | | |
| AB | AUROBINDO PHARMA LTD | 0.035MG;0.5MG | A207055 | 001 | Oct 21, | 2016 |
| <u>DASETTA 1/35</u> | | | | | | |
| AB | NOVAST LABS LTD | 0.035MG;1MG | A090948 | 001 | Dec 22, | 2011 |
| <u>DASETTA 7/7/7</u> | | | | | | |
| AB | NOVAST LABS LTD | 0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG | A090946 | 001 | Dec 22, | 2011 |
| <u>GILDAGIA</u> | | | | | | |
| AB | VINTAGE PHARMS | 0.035MG;0.4MG | A078376 | 001 | Nov 06, | 2012 |
| <u>NORETHINDRONE AND ETHINYL ESTRADIOL</u> | | | | | | |
| AB | ACCORD HLTHCARE | 0.035MG;1MG | A206864 | 001 | Apr 28, | 2017 |
| AB | MAYNE PHARMA | 0.035MG;0.5MG | A070686 | 001 | Jan 29, | 1987 |
| AB | WATSON LABS | 0.035MG;0.4MG | A078323 | 001 | Feb 04, | 2010 |
| AB | WATSON LABS TEVA | 0.035MG;1MG | A070687 | 001 | Jan 29, | 1987 |
| <u>NORINYL 1+35 28-DAY</u> | | | | | | |
| AB | ALLERGAN SALES LLC | 0.035MG;1MG | N017565 | 002 | | |
| <u>NORTREL 0.5/35-28</u> | | | | | | |
| AB | BARR | 0.035MG;0.5MG | A072695 | 001 | Feb 28, | 1992 |
| <u>NORTREL 1/35-28</u> | | | | | | |
| AB | BARR | 0.035MG;1MG | A072696 | 001 | Feb 28, | 1992 |
| <u>NORTREL 7/7/7</u> | | | | | | |
| AB | BARR | 0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG | A075478 | 002 | Aug 30, | 2002 |
| <u>NYLIA 1/35</u> | | | | | | |
| AB | AUROBINDO PHARMA LTD | 0.035MG;1MG | A207056 | 001 | Oct 21, | 2016 |
| <u>NYLIA 7/7/7</u> | | | | | | |
| AB | AUROBINDO PHARMA LTD | 0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG | A207054 | 001 | Oct 21, | 2016 |
| <u>ORTHO-NOVUM 1/35-28</u> | | | | | | |
| AB | +! JANSSEN PHARMS | 0.035MG;1MG | N017919 | 002 | | |
| <u>ORTHO-NOVUM 7/7/7-28</u> | | | | | | |
| AB | +! JANSSEN PHARMS | 0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG | N018985 | 002 | Apr 04, | 1984 |
| <u>PHILITH</u> | | | | | | |
| AB | NOVAST LABS LTD | 0.035MG;0.4MG | A090947 | 001 | Dec 22, | 2011 |
| <u>PIRMELLA 1/35</u> | | | | | | |
| AB | LUPIN LTD | 0.035MG;1MG | A201512 | 001 | Apr 24, | 2013 |
| <u>PIRMELLA 7/7/7</u> | | | | | | |
| AB | LUPIN LTD | 0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG | A201510 | 001 | Apr 24, | 2013 |
| <u>TRI-NORINYL 28-DAY</u> | | | | | | |
| AB | +! MAYNE PHARMA | 0.035MG,0.035MG,0.035MG;0.5MG,1MG,0.5MG | N018977 | 002 | Apr 13, | 1984 |
| <u>VYFEMLA</u> | | | | | | |
| AB | LUPIN LTD | 0.035MG;0.4MG | A201886 | 001 | Sep 26, | 2013 |
| <u>WERA</u> | | | | | | |
| AB | ! NOVAST LABS LTD | 0.035MG;0.5MG | A091204 | 001 | Mar 27, | 2012 |
| <u>NORETHINDRONE AND ETHINYL ESTRADIOL</u> | | | | | | |
| AB | ! MYLAN LABS LTD | 0.05MG;1MG | A203006 | 001 | Aug 05, | 2013 |
| <u>NORETHINDRONE AND ETHINYL ESTRADIOL (10/11)</u> | | | | | | |
| | WATSON LABS TEVA | 0.035MG,0.035MG;0.5MG,1MG | A071044 | 001 | Apr 01, | 1988 |
| TABLET, CHEWABLE; ORAL | | | | | | |
| <u>FEMCON FE</u> | | | | | | |
| AB | +! APIL | 0.035MG;0.4MG | N021490 | 001 | Nov 14, | 2003 |
| <u>KAITLIB FE</u> | | | | | | |
| AB | LUPIN LTD | 0.025MG;0.8MG | A203448 | 001 | Dec 17, | 2015 |
| <u>NEXESTA FE</u> | | | | | | |
| AB | AUROBINDO PHARMA LTD | 0.035MG;0.4MG | A207535 | 001 | Feb 02, | 2017 |
| <u>NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u> | | | | | | |
| AB | ACCORD HLTHCARE | 0.035MG;0.4MG | A207066 | 001 | Mar 29, | 2017 |
| AB | AMNEAL PHARMS | 0.035MG;0.4MG | A078892 | 001 | Sep 26, | 2011 |

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET, CHEWABLE; ORAL

NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

| | | | | |
|-----------|----------------|----------------------|--------------------|--------------|
| AB | + APIL | 0.025MG;0.8MG | N022573 001 | Dec 22, 2010 |
| AB | BARR | 0.035MG;0.4MG | A078965 001 | Aug 05, 2010 |
| AB | LUPIN LTD | 0.035MG;0.4MG | A091332 001 | Mar 23, 2016 |
| AB | MYLAN LABS LTD | 0.025MG;0.8MG | A203371 001 | Apr 23, 2014 |
| AB | | 0.035MG;0.4MG | A202086 001 | Apr 01, 2015 |

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

CAPSULE; ORAL

TAYTULLA

+! APIL

0.02MG;1MG

N204426 001 Apr 19, 2013

TABLET; ORAL

AUROVELA 24 FE

| | | | | |
|-----------|----------------------|-------------------|--------------------|--------------|
| AB | AUROBINDO PHARMA LTD | 0.02MG;1MG | A207504 001 | Jun 15, 2017 |
|-----------|----------------------|-------------------|--------------------|--------------|

BLISOVI 24 FE

| | | | | |
|-----------|-----------|-------------------|--------------------|--------------|
| AB | LUPIN LTD | 0.02MG;1MG | A091398 001 | Oct 28, 2015 |
|-----------|-----------|-------------------|--------------------|--------------|

FEMHRT

| | | | | |
|-----------|------|-----------------------|--------------------|--------------|
| AB | APIL | 0.0025MG;0.5MG | N021065 001 | Jan 14, 2005 |
|-----------|------|-----------------------|--------------------|--------------|

FYAVOLV

| | | | | |
|-----------|-----------|-----------------------|--------------------|--------------|
| AB | LUPIN LTD | 0.005MG;1MG | A204213 002 | Dec 10, 2015 |
| AB | | 0.0025MG;0.5MG | A204213 001 | Dec 10, 2015 |

GILDESS 24 FE

| | | | | |
|-----------|----------------|-------------------|--------------------|--------------|
| AB | VINTAGE PHARMS | 0.02MG;1MG | A090293 001 | Dec 01, 2014 |
|-----------|----------------|-------------------|--------------------|--------------|

LARIN 24 FE

| | | | | |
|-----------|-------------|-------------------|--------------------|--------------|
| AB | NOVAST LABS | 0.02MG;1MG | A202994 001 | Feb 18, 2015 |
|-----------|-------------|-------------------|--------------------|--------------|

LERIBANE

| | | | | |
|-----------|-------------|-----------------------|--------------------|--------------|
| AB | NOVAST LABS | 0.0025MG;0.5MG | A203435 002 | Jun 03, 2016 |
| AB | | 0.005MG;1MG | A203435 001 | Jun 03, 2016 |

LOESTRIN 24 FE

| | | | | |
|-----------|---------------|-------------------|--------------------|--------------|
| AB | + APIL | 0.02MG;1MG | N021871 001 | Feb 17, 2006 |
|-----------|---------------|-------------------|--------------------|--------------|

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

| | | | | |
|-----------|------------------------|-----------------------|--------------------|--------------|
| AB | ! BARR LABS INC | 0.005MG;1MG | A076221 001 | Nov 06, 2009 |
| AB | GLENMARK GENERICS | 0.0025MG;0.5MG | A203038 001 | Apr 02, 2015 |
| AB | | 0.005MG;1MG | A203038 002 | Apr 02, 2015 |
| AB | MYLAN LABS LTD | 0.0025MG;0.5MG | A207260 001 | Feb 02, 2017 |
| AB | | 0.005MG;1MG | A207259 001 | Dec 27, 2016 |

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

| | | | | |
|-----------|------------------------|-------------------|--------------------|--------------|
| AB | ! AMNEAL PHARMS | 0.02MG;1MG | A078267 001 | Sep 01, 2009 |
| AB | BARR LABS INC | 0.02MG;1MG | A090938 001 | Dec 01, 2014 |
| AB | GLENMARK PHARMS LTD | 0.02MG;1MG | A204847 001 | Nov 17, 2017 |
| AB | MYLAN LABS LTD | 0.02MG;1MG | A202742 001 | Oct 30, 2014 |

LO LOESTRIN FE

+! APIL

0.01MG,0.01MG;1MG,N/A

N022501 001 Oct 21, 2010

TABLET; ORAL-21

AUROVELA 1.5/30

| | | | | |
|-----------|----------------------|---------------------|--------------------|--------------|
| AB | AUROBINDO PHARMA LTD | 0.03MG;1.5MG | A207581 001 | Jun 26, 2017 |
|-----------|----------------------|---------------------|--------------------|--------------|

AUROVELA 1/20

| | | | | |
|-----------|----------------------|-------------------|--------------------|--------------|
| AB | AUROBINDO PHARMA LTD | 0.02MG;1MG | A207506 001 | Jun 16, 2017 |
|-----------|----------------------|-------------------|--------------------|--------------|

GILDESS 1.5/30

| | | | | |
|-----------|--------------------|---------------------|--------------------|--------------|
| AB | VINTAGE PHARMS LLC | 0.03MG;1.5MG | A077075 002 | Jul 24, 2012 |
|-----------|--------------------|---------------------|--------------------|--------------|

GILDESS 1/20

| | | | | |
|-----------|--------------------|-------------------|--------------------|--------------|
| AB | VINTAGE PHARMS LLC | 0.02MG;1MG | A077077 002 | Jul 24, 2012 |
|-----------|--------------------|-------------------|--------------------|--------------|

JUNEL 1.5/30

| | | | | |
|-----------|------|---------------------|--------------------|--------------|
| AB | BARR | 0.03MG;1.5MG | A076381 001 | May 30, 2003 |
|-----------|------|---------------------|--------------------|--------------|

JUNEL 1/20

| | | | | |
|-----------|------|-------------------|--------------------|--------------|
| AB | BARR | 0.02MG;1MG | A076380 001 | May 30, 2003 |
|-----------|------|-------------------|--------------------|--------------|

LARIN 1.5/30

| | | | | |
|-----------|-------------|---------------------|--------------------|--------------|
| AB | NOVAST LABS | 0.03MG;1.5MG | A202996 001 | Mar 20, 2014 |
|-----------|-------------|---------------------|--------------------|--------------|

LARIN 1/20

| | | | | |
|-----------|-------------|-------------------|--------------------|--------------|
| AB | NOVAST LABS | 0.02MG;1MG | A202995 001 | Dec 04, 2013 |
|-----------|-------------|-------------------|--------------------|--------------|

LOESTRIN 21 1.5/30

| | | | | |
|-----------|------|---------------------|--------------------|--|
| AB | APIL | 0.03MG;1.5MG | N017875 001 | |
|-----------|------|---------------------|--------------------|--|

LOESTRIN 21 1/20

| | | | | |
|-----------|------|-------------------|--------------------|--|
| AB | APIL | 0.02MG;1MG | N017876 001 | |
|-----------|------|-------------------|--------------------|--|

MICROGESTIN 1.5/30

| | | | | |
|-----------|--------------|---------------------|--------------------|--------------|
| AB | MAYNE PHARMA | 0.03MG;1.5MG | A075548 002 | Jul 30, 2003 |
|-----------|--------------|---------------------|--------------------|--------------|

MICROGESTIN 1/20

| | | | | |
|-----------|--------------|-------------------|--------------------|--------------|
| AB | MAYNE PHARMA | 0.02MG;1MG | A075647 002 | Jul 30, 2003 |
|-----------|--------------|-------------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL-21

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

| | | | | |
|-----------|---------------------|---------------------|--------------------|--------------|
| <u>AB</u> | GLENMARK PHARMS LTD | <u>0.02MG;1MG</u> | <u>A206969 001</u> | Jan 20, 2016 |
| <u>AB</u> | MYLAN LABS LTD | <u>0.02MG;1MG</u> | <u>A202771 001</u> | Nov 06, 2013 |
| <u>AB</u> | | <u>0.03MG;1.5MG</u> | <u>A202770 001</u> | Feb 19, 2015 |

TRI-LEGEST 21

BARR

0.02MG, 0.03MG, 0.035MG; 1MG, 1MG, 1MG

A076405 001 Oct 26, 2007

TABLET; ORAL-28

AUROVELA FE 1.5/30

| | | | | |
|-----------|----------------------|---------------------|--------------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>0.03MG;1.5MG</u> | <u>A207580 001</u> | Jun 15, 2017 |
|-----------|----------------------|---------------------|--------------------|--------------|

AUROVELA FE 1/20

| | | | | |
|-----------|----------------------|-------------------|--------------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>0.02MG;1MG</u> | <u>A207505 001</u> | Jun 16, 2017 |
|-----------|----------------------|-------------------|--------------------|--------------|

BLISOVI FE 1.5/30

| | | | | |
|-----------|-----------|---------------------|--------------------|--------------|
| <u>AB</u> | LUPIN LTD | <u>0.03MG;1.5MG</u> | <u>A201585 001</u> | Nov 18, 2015 |
|-----------|-----------|---------------------|--------------------|--------------|

BLISOVI FE 1/20

| | | | | |
|-----------|-----------|-------------------|--------------------|--------------|
| <u>AB</u> | LUPIN LTD | <u>0.02MG;1MG</u> | <u>A201584 001</u> | Nov 18, 2015 |
|-----------|-----------|-------------------|--------------------|--------------|

ESTROSTEP FE

| | | | | |
|-----------|---------|-----------------------------------------------|--------------------|--------------|
| <u>AB</u> | +! APIL | <u>0.02MG, 0.03MG, 0.035MG; 1MG, 1MG, 1MG</u> | <u>N020130 002</u> | Oct 09, 1996 |
|-----------|---------|-----------------------------------------------|--------------------|--------------|

GILDESS FE 1.5/30

| | | | | |
|-----------|--------------------|---------------------|--------------------|--------------|
| <u>AB</u> | VINTAGE PHARMS LLC | <u>0.03MG;1.5MG</u> | <u>A077075 001</u> | Apr 28, 2005 |
|-----------|--------------------|---------------------|--------------------|--------------|

GILDESS FE 1/20

| | | | | |
|-----------|--------------------|-------------------|--------------------|--------------|
| <u>AB</u> | VINTAGE PHARMS LLC | <u>0.02MG;1MG</u> | <u>A077077 001</u> | May 20, 2005 |
|-----------|--------------------|-------------------|--------------------|--------------|

HAILEY 1.5/30

| | | | | |
|-----------|-----------------|---------------------|--------------------|--------------|
| <u>AB</u> | GLENMARK PHARMS | <u>0.03MG;1.5MG</u> | <u>A209297 001</u> | Jun 05, 2018 |
|-----------|-----------------|---------------------|--------------------|--------------|

HAILEY FE 1.5/30

| | | | | |
|-----------|-----------------|---------------------|--------------------|--------------|
| <u>AB</u> | GLENMARK PHARMS | <u>0.03MG;1.5MG</u> | <u>A209031 001</u> | Jun 05, 2018 |
|-----------|-----------------|---------------------|--------------------|--------------|

HAILEY FE 1/20

| | | | | |
|-----------|---------------------|-------------------|--------------------|--------------|
| <u>AB</u> | GLENMARK PHARMS LTD | <u>0.02MG;1MG</u> | <u>A206597 001</u> | Nov 21, 2017 |
|-----------|---------------------|-------------------|--------------------|--------------|

JUNEL FE 1.5/30

| | | | | |
|-----------|------|---------------------|--------------------|--------------|
| <u>AB</u> | BARR | <u>0.03MG;1.5MG</u> | <u>A076064 001</u> | Sep 18, 2003 |
|-----------|------|---------------------|--------------------|--------------|

JUNEL FE 1/20

| | | | | |
|-----------|------|-------------------|--------------------|--------------|
| <u>AB</u> | BARR | <u>0.02MG;1MG</u> | <u>A076081 001</u> | Sep 18, 2003 |
|-----------|------|-------------------|--------------------|--------------|

LARIN FE 1.5/30

| | | | | |
|-----------|-------------|---------------------|--------------------|--------------|
| <u>AB</u> | NOVAST LABS | <u>0.03MG;1.5MG</u> | <u>A091453 001</u> | Aug 23, 2013 |
|-----------|-------------|---------------------|--------------------|--------------|

LARIN FE 1/20

| | | | | |
|-----------|-------------|-------------------|--------------------|--------------|
| <u>AB</u> | NOVAST LABS | <u>0.02MG;1MG</u> | <u>A091454 001</u> | Aug 26, 2013 |
|-----------|-------------|-------------------|--------------------|--------------|

LOESTRIN FE 1.5/30

| | | | | |
|-----------|---------|---------------------|--------------------|--|
| <u>AB</u> | +! APIL | <u>0.03MG;1.5MG</u> | <u>N017355 001</u> | |
|-----------|---------|---------------------|--------------------|--|

LOESTRIN FE 1/20

| | | | | |
|-----------|--------|-------------------|--------------------|--|
| <u>AB</u> | + APIL | <u>0.02MG;1MG</u> | <u>N017354 001</u> | |
|-----------|--------|-------------------|--------------------|--|

MICROGESTIN FE 1.5/30

| | | | | |
|-----------|--------------|---------------------|--------------------|--------------|
| <u>AB</u> | MAYNE PHARMA | <u>0.03MG;1.5MG</u> | <u>A075548 001</u> | Feb 05, 2001 |
|-----------|--------------|---------------------|--------------------|--------------|

MICROGESTIN FE 1/20

| | | | | |
|-----------|--------------|-------------------|--------------------|--------------|
| <u>AB</u> | MAYNE PHARMA | <u>0.02MG;1MG</u> | <u>A075647 001</u> | Feb 05, 2001 |
|-----------|--------------|-------------------|--------------------|--------------|

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

| | | | | |
|-----------|--------------|-----------------------------------------------|--------------------|--------------|
| <u>AB</u> | MAYNE PHARMA | <u>0.02MG, 0.03MG, 0.035MG; 1MG, 1MG, 1MG</u> | <u>A076629 001</u> | Mar 18, 2010 |
|-----------|--------------|-----------------------------------------------|--------------------|--------------|

| | | | | |
|-----------|----------------|-------------------|--------------------|--------------|
| <u>AB</u> | MYLAN LABS LTD | <u>0.02MG;1MG</u> | <u>A202772 001</u> | Nov 14, 2013 |
|-----------|----------------|-------------------|--------------------|--------------|

| | | | | |
|-----------|--|-----------------------------------------------|--------------------|--------------|
| <u>AB</u> | | <u>0.02MG, 0.03MG, 0.035MG; 1MG, 1MG, 1MG</u> | <u>A205069 001</u> | Jun 22, 2018 |
|-----------|--|-----------------------------------------------|--------------------|--------------|

| | | | | |
|-----------|--|---------------------|--------------------|--------------|
| <u>AB</u> | | <u>0.03MG;1.5MG</u> | <u>A202741 001</u> | Feb 20, 2015 |
|-----------|--|---------------------|--------------------|--------------|

TRI-LEGEST FE

| | | | | |
|-----------|------|-----------------------------------------------|--------------------|--------------|
| <u>AB</u> | BARR | <u>0.02MG, 0.03MG, 0.035MG; 1MG, 1MG, 1MG</u> | <u>A076105 001</u> | Oct 26, 2007 |
|-----------|------|-----------------------------------------------|--------------------|--------------|

TABLET, CHEWABLE; ORAL

MIBELAS 24 FE

| | | | | |
|-----------|----------------|-------------------|--------------------|--------------|
| <u>AB</u> | LUPIN ATLANTIS | <u>0.02MG;1MG</u> | <u>A206287 001</u> | May 24, 2016 |
|-----------|----------------|-------------------|--------------------|--------------|

MINASTRIN 24 FE

| | | | | |
|-----------|---------|-------------------|--------------------|--------------|
| <u>AB</u> | +! APIL | <u>0.02MG;1MG</u> | <u>N203667 001</u> | May 08, 2013 |
|-----------|---------|-------------------|--------------------|--------------|

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

| | | | | |
|-----------|---------------|-------------------|--------------------|--------------|
| <u>AB</u> | AMNEAL PHARMS | <u>0.02MG;1MG</u> | <u>A207514 001</u> | Sep 11, 2017 |
|-----------|---------------|-------------------|--------------------|--------------|

| | | | | |
|-----------|-------------------|-------------------|--------------------|--------------|
| <u>AB</u> | CHEMO RESEARCH SL | <u>0.02MG;1MG</u> | <u>A209609 001</u> | Jul 16, 2018 |
|-----------|-------------------|-------------------|--------------------|--------------|

| | | | | |
|-----------|---------------------|-------------------|--------------------|--------------|
| <u>AB</u> | GLENMARK PHARMS LTD | <u>0.02MG;1MG</u> | <u>A210369 001</u> | Dec 26, 2017 |
|-----------|---------------------|-------------------|--------------------|--------------|

| | | | | |
|-----------|----------------|-------------------|--------------------|--------------|
| <u>AB</u> | MYLAN LABS LTD | <u>0.02MG;1MG</u> | <u>A206120 001</u> | Sep 12, 2017 |
|-----------|----------------|-------------------|--------------------|--------------|

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

ESTARYLLA

| | | | | |
|-----------|-----------------|-----------------------|--------------------|--------------|
| <u>AB</u> | LABS LEON FARMA | <u>0.035MG;0.25MG</u> | <u>A090794 001</u> | Jan 30, 2013 |
|-----------|-----------------|-----------------------|--------------------|--------------|

MILI

| | | | | |
|-----------|----------------------|-----------------------|--------------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>0.035MG;0.25MG</u> | <u>A205449 001</u> | Jul 07, 2016 |
|-----------|----------------------|-----------------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

MONO-LINYAH

| | | | | | |
|-----------|-----------------|-----------------------|----------------|------------|--------------|
| AB | NOVAST LABS LTD | <u>0.035MG;0.25MG</u> | <u>A090523</u> | <u>001</u> | May 23, 2012 |
|-----------|-----------------|-----------------------|----------------|------------|--------------|

NORGESTIMATE AND ETHINYL ESTRADIOL

| | | | | | |
|-----------|---------------|------------------------------------------------------|----------------|------------|--------------|
| AB | AMNEAL PHARMS | <u>0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG</u> | <u>A203870</u> | <u>001</u> | Nov 12, 2015 |
|-----------|---------------|------------------------------------------------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-----------------------|----------------|------------|--------------|
| AB | | <u>0.035MG;0.25MG</u> | <u>A203865</u> | <u>001</u> | Oct 27, 2015 |
|-----------|--|-----------------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|------------------------------------------------------|----------------|------------|--------------|
| AB | | <u>0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG</u> | <u>A203873</u> | <u>001</u> | May 12, 2016 |
|-----------|--|------------------------------------------------------|----------------|------------|--------------|

| | | | | | |
|-----------|-------------------|------------------------------------------------------|----------------|------------|--------------|
| AB | GLENMARK GENERICS | <u>0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG</u> | <u>A200494</u> | <u>001</u> | Jun 17, 2011 |
|-----------|-------------------|------------------------------------------------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-----------------------|----------------|------------|--------------|
| AB | | <u>0.035MG;0.25MG</u> | <u>A200538</u> | <u>001</u> | Apr 05, 2012 |
|-----------|--|-----------------------|----------------|------------|--------------|

| | | | | | |
|-----------|-----------------|------------------------------------------------------|----------------|------------|--------------|
| AB | GLENMARK PHARMS | <u>0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG</u> | <u>A204057</u> | <u>001</u> | Feb 23, 2016 |
|-----------|-----------------|------------------------------------------------------|----------------|------------|--------------|

| | | | | | |
|-----------|-----------|------------------------------------------------------|----------------|------------|--------------|
| AB | LUPIN LTD | <u>0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG</u> | <u>A205588</u> | <u>001</u> | Apr 26, 2016 |
|-----------|-----------|------------------------------------------------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-----------------------|----------------|------------|--------------|
| AB | | <u>0.035MG;0.25MG</u> | <u>A205630</u> | <u>001</u> | Oct 27, 2016 |
|-----------|--|-----------------------|----------------|------------|--------------|

| | | | | | |
|-----------|--------------|------------------------------------------------------|----------------|------------|--------------|
| AB | LUPIN PHARMS | <u>0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG</u> | <u>A200541</u> | <u>001</u> | Jun 25, 2012 |
|-----------|--------------|------------------------------------------------------|----------------|------------|--------------|

| | | | | | |
|-----------|----------------|------------------------------------------------------|----------------|------------|--------------|
| AB | MYLAN LABS LTD | <u>0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG</u> | <u>A202132</u> | <u>001</u> | Sep 09, 2015 |
|-----------|----------------|------------------------------------------------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|------------------------------------------------------|----------------|------------|--------------|
| AB | | <u>0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG</u> | <u>A201897</u> | <u>001</u> | Jan 27, 2016 |
|-----------|--|------------------------------------------------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-----------------------|----------------|------------|--------------|
| AB | | <u>0.035MG;0.25MG</u> | <u>A201896</u> | <u>001</u> | Jan 27, 2016 |
|-----------|--|-----------------------|----------------|------------|--------------|

| | | | | | |
|-----------|-----------|------------------------------------------------------|----------------|------------|--------------|
| AB | OC PHARMA | <u>0.035MG;0.035MG;0.035MG;0.18MG;0.215MG;0.25MG</u> | <u>A200383</u> | <u>001</u> | Apr 07, 2015 |
|-----------|-----------|------------------------------------------------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-----------------------|----------------|------------|--------------|
| AB | | <u>0.035MG;0.25MG</u> | <u>A200384</u> | <u>001</u> | Apr 07, 2015 |
|-----------|--|-----------------------|----------------|------------|--------------|

ORTHO CYCLEN-28

| | | | | | |
|-----------|-------------------|-----------------------|----------------|------------|--------------|
| AB | +! JANSSEN PHARMS | <u>0.035MG;0.25MG</u> | <u>N019653</u> | <u>002</u> | Dec 29, 1989 |
|-----------|-------------------|-----------------------|----------------|------------|--------------|

ORTHO TRI-CYCLEN

| | | | | | |
|-----------|-------------------|------------------------------------------------------|----------------|------------|--------------|
| AB | +! JANSSEN PHARMS | <u>0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG</u> | <u>N019697</u> | <u>001</u> | Jul 03, 1992 |
|-----------|-------------------|------------------------------------------------------|----------------|------------|--------------|

ORTHO TRI-CYCLEN LO

| | | | | | |
|-----------|-------------------|------------------------------------------------------|----------------|------------|--------------|
| AB | +! JANSSEN PHARMS | <u>0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG</u> | <u>N021241</u> | <u>001</u> | Aug 22, 2002 |
|-----------|-------------------|------------------------------------------------------|----------------|------------|--------------|

PREVIFEM

| | | | | | |
|-----------|--------------------|-----------------------|----------------|------------|--------------|
| AB | VINTAGE PHARMS LLC | <u>0.035MG;0.25MG</u> | <u>A076334</u> | <u>001</u> | Jan 09, 2004 |
|-----------|--------------------|-----------------------|----------------|------------|--------------|

SPRINTEC

| | | | | | |
|-----------|------|-----------------------|----------------|------------|--------------|
| AB | BARR | <u>0.035MG;0.25MG</u> | <u>A075804</u> | <u>001</u> | Sep 25, 2002 |
|-----------|------|-----------------------|----------------|------------|--------------|

TRI LO SPRINTEC

| | | | | | |
|-----------|---------------|------------------------------------------------------|----------------|------------|--------------|
| AB | BARR LABS INC | <u>0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG</u> | <u>A076784</u> | <u>001</u> | Jun 29, 2009 |
|-----------|---------------|------------------------------------------------------|----------------|------------|--------------|

TRI-ESTARYLLA

| | | | | | |
|-----------|-----------------|------------------------------------------------------|----------------|------------|--------------|
| AB | LABS LEON FARMA | <u>0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG</u> | <u>A090793</u> | <u>001</u> | Jan 30, 2013 |
|-----------|-----------------|------------------------------------------------------|----------------|------------|--------------|

TRI-LINYAH

| | | | | | |
|-----------|-----------------|------------------------------------------------------|----------------|------------|--------------|
| AB | NOVAST LABS LTD | <u>0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG</u> | <u>A090524</u> | <u>001</u> | May 30, 2012 |
|-----------|-----------------|------------------------------------------------------|----------------|------------|--------------|

TRI-LO-ESTARYLLA

| | | | | | |
|-----------|-----------------|------------------------------------------------------|----------------|------------|--------------|
| AB | LABS LEON FARMA | <u>0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG</u> | <u>A091232</u> | <u>001</u> | Jun 29, 2015 |
|-----------|-----------------|------------------------------------------------------|----------------|------------|--------------|

TRI-LO-MILI

| | | | | | |
|-----------|----------------------|------------------------------------------------------|----------------|------------|--------------|
| AB | AUROBINDO PHARMA LTD | <u>0.025MG,0.025MG,0.025MG;0.18MG,0.215MG,0.25MG</u> | <u>A205762</u> | <u>001</u> | Nov 04, 2016 |
|-----------|----------------------|------------------------------------------------------|----------------|------------|--------------|

TRI-MILI

| | | | | | |
|-----------|----------------------|------------------------------------------------------|----------------|------------|--------------|
| AB | AUROBINDO PHARMA LTD | <u>0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG</u> | <u>A205441</u> | <u>001</u> | Jul 06, 2016 |
|-----------|----------------------|------------------------------------------------------|----------------|------------|--------------|

TRI-PREVIFEM

| | | | | | |
|-----------|--------------------|------------------------------------------------------|----------------|------------|--------------|
| AB | VINTAGE PHARMS LLC | <u>0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG</u> | <u>A076335</u> | <u>001</u> | Mar 26, 2004 |
|-----------|--------------------|------------------------------------------------------|----------------|------------|--------------|

TRI-SPRINTEC

| | | | | | |
|-----------|------|------------------------------------------------------|----------------|------------|--------------|
| AB | BARR | <u>0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG</u> | <u>A075808</u> | <u>001</u> | Dec 29, 2003 |
|-----------|------|------------------------------------------------------|----------------|------------|--------------|

ETHINYL ESTRADIOL; NORGESTREL

TABLET; ORAL-21

CRYSELLE

| | | | | | |
|-----------|---------------------|---------------------|----------------|------------|--------------|
| AB | DURAMED PHARMS BARR | <u>0.03MG;0.3MG</u> | <u>A075840</u> | <u>001</u> | Nov 30, 2001 |
|-----------|---------------------|---------------------|----------------|------------|--------------|

TABLET; ORAL-28

CRYSELLE

| | | | | | |
|-----------|---------------------|---------------------|----------------|------------|--------------|
| AB | DURAMED PHARMS BARR | <u>0.03MG;0.3MG</u> | <u>A075840</u> | <u>002</u> | Nov 30, 2001 |
|-----------|---------------------|---------------------|----------------|------------|--------------|

ELINEST

| | | | | | |
|-----------|-----------------|---------------------|----------------|------------|--------------|
| AB | NOVAST LABS LTD | <u>0.03MG;0.3MG</u> | <u>A091105</u> | <u>001</u> | Mar 28, 2012 |
|-----------|-----------------|---------------------|----------------|------------|--------------|

LOW-OGESTREL-28

| | | | | | |
|-----------|--------------|---------------------|----------------|------------|--------------|
| AB | MAYNE PHARMA | <u>0.03MG;0.3MG</u> | <u>A075288</u> | <u>002</u> | Jul 28, 1999 |
|-----------|--------------|---------------------|----------------|------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

ETHINYL ESTRADIOL; NORGESTREL

TABLET; ORAL-28

OGESTREL 0.5/50-28

! WATSON LABS 0.05MG; 0.5MG

A075406 002 Dec 15, 1999

ETHINYL ESTRADIOL; SEGESTERONE ACETATE

RING; VAGINAL

ANNOVERA

+! THERAPEUTICSMD INC 0.013MG/24HR; 0.15MG/24HR

N209627 001 Aug 10, 2018

ETHIODIZED OIL

OIL; INTRALYMPHATIC, INTRAUTERINE

LIPIODOL

+! GUERBET EQ 4.8GM IODINE/10ML (EQ 480MG IODINE/ML)

N009190 001

ETHIONAMIDE

TABLET; ORAL

TRECATOR

+! WYETH PHARMS 250MG

N013026 002

ETHOSUXIMIDE

CAPSULE; ORAL

ETHOSUXIMIDE**AB** AKORN **250MG****A040686 001** May 28, 2008**AB** BIONPHARMA INC **250MG****A040430 001** Oct 28, 2002**AB** HERITAGE PHARMS INC **250MG****A200892 001** Sep 25, 2012ZARONTIN**AB** +! PARKE DAVIS **250MG****N012380 001**

SYRUP; ORAL

ETHOSUXIMIDE**AA** MIKART **250MG/5ML****A040506 001** Dec 22, 2003**AA** PHARM ASSOC **250MG/5ML****A040253 001** Nov 22, 2000**AA** TEVA PHARMS **250MG/5ML****A081306 001** Jul 30, 1993ZARONTIN**AA** ! PARKE-DAVIS **250MG/5ML****A080258 001**ETHOTOIN

TABLET; ORAL

PEGANONE

+! RECORDATI RARE 250MG

N010841 001

ETIDRONATE DISODIUM

TABLET; ORAL

ETIDRONATE DISODIUM

MYLAN 200MG

A075800 001 Jan 24, 2003

! 400MG

A075800 002 Jan 24, 2003

ETODOLAC

CAPSULE; ORAL

ETODOLAC**AB** ANI PHARMS INC **200MG****A075126 001** Sep 16, 1999**AB** **300MG****A075126 002** Sep 16, 1999**AB** APOTEX **200MG****A075419 001** Jul 28, 2000**AB** **300MG****A075419 002** Jul 28, 2000**AB** TARO **200MG****A075078 001** Apr 30, 1998**AB** ! **300MG****A075078 002** Apr 30, 1998

TABLET; ORAL

ETODOLAC**AB** AMNEAL PHARMS CO **400MG****A208834 001** Jun 07, 2018**AB** **500MG****A208834 002** Jun 07, 2018**AB** APOTEX INC **400MG****A076004 001** Dec 03, 2002**AB** **500MG****A076004 002** Dec 03, 2002**AB** EDENBRIDGE PHARMS **400MG****A209888 001** Nov 30, 2018**AB** **500MG****A209888 002** Nov 30, 2018**AB** SANDOZ **400MG****A074903 001** Apr 11, 1997**AB** **500MG****A074903 002** Apr 19, 1999**AB** TARO PHARM INDS **400MG****A075074 001** Mar 11, 1998**AB** ! **500MG****A075074 002** Apr 25, 2000**AB** TEVA **400MG****A075009 001** Nov 26, 1997**AB** **500MG****A075009 002** Dec 28, 1999

TABLET, EXTENDED RELEASE; ORAL

ETODOLAC**AB** TARO **400MG****A076174 001** Mar 13, 2003**AB** **500MG****A076174 002** Mar 13, 2003**AB** **600MG****A076174 003** Mar 13, 2003

PRESCRIPTION DRUG PRODUCT LIST

ETODOLAC

TABLET, EXTENDED RELEASE;ORAL

ETODOLAC

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | TEVA | <u>400MG</u> | <u>A075665</u> | <u>003</u> | Feb 05, 2001 |
| <u>AB</u> | | <u>500MG</u> | <u>A075665</u> | <u>002</u> | Jul 31, 2000 |
| <u>AB</u> | ! | <u>600MG</u> | <u>A075665</u> | <u>001</u> | Jul 31, 2000 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>400MG</u> | <u>A091134</u> | <u>001</u> | Jan 23, 2014 |
| <u>AB</u> | | <u>500MG</u> | <u>A091134</u> | <u>002</u> | Jan 23, 2014 |
| <u>AB</u> | | <u>600MG</u> | <u>A091134</u> | <u>003</u> | Jan 23, 2014 |

ETOMIDATE

INJECTABLE; INJECTION

AMIDATE

| | | | | | | |
|-----------|----|---------|---------------|----------------|------------|--------------|
| <u>AP</u> | +! | HOSPIRA | <u>2MG/ML</u> | <u>N018227</u> | <u>001</u> | Sep 07, 1982 |
|-----------|----|---------|---------------|----------------|------------|--------------|

ETOMIDATE

| | | | | | | |
|-----------|--|-------------------------|---------------|----------------|------------|--------------|
| <u>AP</u> | | AUROBINDO PHARMA LTD | <u>2MG/ML</u> | <u>A206126</u> | <u>001</u> | Feb 24, 2017 |
| <u>AP</u> | | EMCURE PHARMS LTD | <u>2MG/ML</u> | <u>A204618</u> | <u>001</u> | Aug 13, 2014 |
| <u>AP</u> | | GLAND PHARMA LTD | <u>2MG/ML</u> | <u>A209058</u> | <u>001</u> | Apr 18, 2017 |
| <u>AP</u> | | HIKMA FARMACEUTICA | <u>2MG/ML</u> | <u>A202354</u> | <u>001</u> | Feb 25, 2016 |
| <u>AP</u> | | LUITPOLD | <u>2MG/ML</u> | <u>A078867</u> | <u>001</u> | Dec 22, 2009 |
| <u>AP</u> | | MYLAN LABS LTD | <u>2MG/ML</u> | <u>A078289</u> | <u>001</u> | Jan 02, 2009 |
| <u>AP</u> | | | <u>2MG/ML</u> | <u>A201044</u> | <u>001</u> | Feb 07, 2017 |
| <u>AP</u> | | PAR STERILE PRODUCTS | <u>2MG/ML</u> | <u>A091297</u> | <u>001</u> | Jun 20, 2012 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>2MG/ML</u> | <u>A074593</u> | <u>001</u> | Nov 04, 1996 |
| <u>AP</u> | | ZYDUS PHARMS USA INC | <u>2MG/ML</u> | <u>A202360</u> | <u>001</u> | Jul 18, 2014 |

ETONOGESTREL

IMPLANT; IMPLANTATION

NEXPLANON

| | | | | | | |
|--|----|-----------------|--------------|---------|-----|--------------|
| | +! | ORGANON USA INC | 68MG/IMPLANT | N021529 | 002 | May 31, 2011 |
|--|----|-----------------|--------------|---------|-----|--------------|

ETOPOSIDE

CAPSULE; ORAL

ETOPOSIDE

| | | | | | | |
|--|---|-------|------|---------|-----|--------------|
| | ! | MYLAN | 50MG | A075635 | 001 | Sep 19, 2001 |
|--|---|-------|------|---------|-----|--------------|

INJECTABLE; INJECTION

ETOPOSIDE

| | | | | | | |
|-----------|---|-------------------------|----------------|----------------|------------|--------------|
| <u>AP</u> | | ACCORD HLTHCARE | <u>20MG/ML</u> | <u>A074513</u> | <u>001</u> | Mar 14, 1996 |
| <u>AP</u> | ! | FRESENIUS KABI USA | <u>20MG/ML</u> | <u>A074983</u> | <u>001</u> | Sep 30, 1998 |
| <u>AP</u> | | MYLAN LABS LTD | <u>20MG/ML</u> | <u>A203507</u> | <u>001</u> | Nov 20, 2017 |
| <u>AP</u> | | | <u>20MG/ML</u> | <u>A204927</u> | <u>001</u> | Oct 31, 2017 |
| <u>AP</u> | | TEVA PHARMS USA | <u>20MG/ML</u> | <u>A074529</u> | <u>001</u> | Jul 24, 1996 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>20MG/ML</u> | <u>A074290</u> | <u>001</u> | Jul 17, 1995 |

ETOPOSIDE PHOSPHATE

INJECTABLE; INJECTION

ETOPOPHOS PRESERVATIVE FREE

| | | | | | | |
|--|----|-------------------------|--------------------|---------|-----|--------------|
| | +! | BRISTOL MYERS SQUIBB | EQ 100MG BASE/VIAL | N020457 | 001 | May 17, 1996 |
|--|----|-------------------------|--------------------|---------|-----|--------------|

ETRAVIRINE

TABLET; ORAL

INTELENCE

| | | | | | | |
|--|----|-----------------|-------|---------|-----|--------------|
| | + | JANSSEN R AND D | 25MG | N022187 | 003 | Mar 26, 2012 |
| | + | | 100MG | N022187 | 001 | Jan 18, 2008 |
| | +! | | 200MG | N022187 | 002 | Dec 22, 2010 |

EVEROLIMUS

TABLET; ORAL

EVEROLIMUS

| | | | | | | |
|-----------|--|-------------------------|---------------|----------------|------------|--------------|
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>0.25MG</u> | <u>A206133</u> | <u>001</u> | Apr 12, 2018 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A206133</u> | <u>002</u> | Apr 12, 2018 |
| <u>AB</u> | | | <u>0.75MG</u> | <u>A206133</u> | <u>003</u> | Apr 12, 2018 |

ZORTRESS

| | | | | | | |
|-----------|----|----------|---------------|----------------|------------|--------------|
| <u>AB</u> | + | NOVARTIS | <u>0.25MG</u> | <u>N021560</u> | <u>001</u> | Apr 20, 2010 |
| <u>AB</u> | + | | <u>0.5MG</u> | <u>N021560</u> | <u>002</u> | Apr 20, 2010 |
| <u>AB</u> | +! | | <u>0.75MG</u> | <u>N021560</u> | <u>003</u> | Apr 20, 2010 |
| | | AFINITOR | | | | |
| | + | NOVARTIS | 2.5MG | N022334 | 003 | Jul 09, 2010 |
| | + | | 5MG | N022334 | 001 | Mar 30, 2009 |

PRESCRIPTION DRUG PRODUCT LISTEVEROLIMUS

TABLET;ORAL

AFINITOR

| | | | | | |
|----|--|-------|---------|-----|--------------|
| + | | 7.5MG | N022334 | 004 | Mar 30, 2012 |
| +! | | 10MG | N022334 | 002 | Mar 30, 2009 |

TABLET, FOR SUSPENSION;ORAL

AFINITOR DISPERZ

| | | | | | |
|----|----------------|-----|---------|-----|--------------|
| + | NOVARTIS PHARM | 2MG | N203985 | 001 | Aug 29, 2012 |
| + | | 3MG | N203985 | 002 | Aug 29, 2012 |
| +! | | 5MG | N203985 | 003 | Aug 29, 2012 |

EXEMESTANE

TABLET;ORAL

AROMASIN

| | | | | | | |
|-----------|-----------|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | <u>+!</u> | PHARMACIA AND UPJOHN | <u>25MG</u> | <u>N020753</u> | <u>001</u> | Oct 21, 1999 |
|-----------|-----------|-------------------------|-------------|----------------|------------|--------------|

EXEMESTANE

| | | | | | | |
|-----------|--|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | ALVOGEN MALTA | <u>25MG</u> | <u>A200898</u> | <u>001</u> | Jul 28, 2014 |
| <u>AB</u> | | AMNEAL PHARMS | <u>25MG</u> | <u>A206421</u> | <u>001</u> | Dec 28, 2018 |
| <u>AB</u> | | CIPLA | <u>25MG</u> | <u>A210323</u> | <u>001</u> | Apr 27, 2018 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>25MG</u> | <u>A203315</u> | <u>001</u> | Mar 10, 2017 |
| <u>AB</u> | | UPSHER SMITH LABS | <u>25MG</u> | <u>A209208</u> | <u>001</u> | Jul 26, 2017 |
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>25MG</u> | <u>A077431</u> | <u>001</u> | Apr 01, 2011 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>25MG</u> | <u>A202602</u> | <u>001</u> | Oct 03, 2018 |

EXENATIDE

SUSPENSION, EXTENDED RELEASE;SUBCUTANEOUS

BYDUREON BCISE

| | | | | | |
|----|----------------|-------------------------|---------|-----|--------------|
| +! | ASTRAZENECA AB | 2MG/0.85ML (2MG/0.85ML) | N209210 | 001 | Oct 20, 2017 |
|----|----------------|-------------------------|---------|-----|--------------|

EXENATIDE SYNTHETIC

FOR SUSPENSION, EXTENDED RELEASE;SUBCUTANEOUS

BYDUREON

| | | | | | |
|----|----------------|----------|---------|-----|--------------|
| +! | ASTRAZENECA AB | 2MG/VIAL | N022200 | 001 | Jan 27, 2012 |
|----|----------------|----------|---------|-----|--------------|

BYDUREON PEN

| | | | | | |
|----|----------------|-----|---------|-----|--------------|
| +! | ASTRAZENECA AB | 2MG | N022200 | 002 | Feb 28, 2014 |
|----|----------------|-----|---------|-----|--------------|

INJECTABLE;SUBCUTANEOUS

BYETTA

| | | | | | |
|----|----------------|--------------------------|---------|-----|--------------|
| +! | ASTRAZENECA AB | 300MCG/1.2ML (250MCG/ML) | N021773 | 001 | Apr 28, 2005 |
| +! | | 600MCG/2.4ML (250MCG/ML) | N021773 | 002 | Apr 28, 2005 |

EZETIMIBE

TABLET;ORAL

EZETIMIBE

| | | | | | | |
|-----------|--|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | ACCORD HLTHCARE | <u>10MG</u> | <u>A211550</u> | <u>001</u> | Oct 26, 2018 |
| <u>AB</u> | | ALKEM LABS LTD | <u>10MG</u> | <u>A209234</u> | <u>001</u> | Dec 21, 2017 |
| <u>AB</u> | | AMNEAL PHARMS CO | <u>10MG</u> | <u>A208803</u> | <u>001</u> | Jun 12, 2017 |
| <u>AB</u> | | APOTEX INC | <u>10MG</u> | <u>A208332</u> | <u>001</u> | Jun 12, 2017 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>10MG</u> | <u>A209838</u> | <u>001</u> | Aug 25, 2017 |
| <u>AB</u> | | GLENMARK PHARMS LTD | <u>10MG</u> | <u>A078560</u> | <u>001</u> | Jun 26, 2015 |
| <u>AB</u> | | OHM LABS INC | <u>10MG</u> | <u>A207311</u> | <u>001</u> | Jun 12, 2017 |
| <u>AB</u> | | SANDOZ INC | <u>10MG</u> | <u>A203931</u> | <u>001</u> | Jun 12, 2017 |
| <u>AB</u> | | TEVA PHARMS USA | <u>10MG</u> | <u>A078724</u> | <u>001</u> | Jun 12, 2017 |
| <u>AB</u> | | WATSON LABS INC | <u>10MG</u> | <u>A200831</u> | <u>001</u> | Jun 12, 2017 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>10MG</u> | <u>A204331</u> | <u>001</u> | Jun 12, 2017 |

ZETIA

| | | | | | | |
|-----------|-----------|---------------|-------------|----------------|------------|--------------|
| <u>AB</u> | <u>+!</u> | MSD INTL GMBH | <u>10MG</u> | <u>N021445</u> | <u>001</u> | Oct 25, 2002 |
|-----------|-----------|---------------|-------------|----------------|------------|--------------|

EZETIMIBE; SIMVASTATIN

TABLET;ORAL

EZETIMIBE AND SIMVASTATIN

| | | | | | | |
|-----------|--|------------------|------------------|----------------|------------|--------------|
| <u>AB</u> | | ALKEM LABS LTD | <u>10MG;10MG</u> | <u>A209222</u> | <u>001</u> | Dec 22, 2017 |
| <u>AB</u> | | | <u>10MG;20MG</u> | <u>A209222</u> | <u>002</u> | Dec 22, 2017 |
| <u>AB</u> | | | <u>10MG;40MG</u> | <u>A209222</u> | <u>003</u> | Dec 22, 2017 |
| <u>AB</u> | | | <u>10MG;80MG</u> | <u>A209222</u> | <u>004</u> | Dec 22, 2017 |
| <u>AB</u> | | AMNEAL PHARMS CO | <u>10MG;10MG</u> | <u>A208831</u> | <u>001</u> | Nov 21, 2017 |
| <u>AB</u> | | | <u>10MG;20MG</u> | <u>A208831</u> | <u>002</u> | Nov 21, 2017 |
| <u>AB</u> | | | <u>10MG;40MG</u> | <u>A208831</u> | <u>003</u> | Nov 21, 2017 |
| <u>AB</u> | | | <u>10MG;80MG</u> | <u>A208831</u> | <u>004</u> | Nov 21, 2017 |
| <u>AB</u> | | ANI PHARMS INC | <u>10MG;10MG</u> | <u>A201890</u> | <u>001</u> | Apr 26, 2017 |
| <u>AB</u> | | | <u>10MG;20MG</u> | <u>A201890</u> | <u>002</u> | Apr 26, 2017 |
| <u>AB</u> | | | <u>10MG;40MG</u> | <u>A201890</u> | <u>003</u> | Apr 26, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

EZETIMIBE; SIMVASTATIN

TABLET; ORAL

EZETIMIBE AND SIMVASTATIN

| | | | | |
|-----------------------|-------------------|--------------------------|---------------------------|--------------|
| AB | | <u>10MG; 80MG</u> | <u>A201890 004</u> | Apr 26, 2017 |
| AB | DR REDDYS LABS SA | <u>10MG; 10MG</u> | <u>A200909 001</u> | Apr 26, 2017 |
| AB | | <u>10MG; 20MG</u> | <u>A200909 002</u> | Apr 26, 2017 |
| AB | | <u>10MG; 40MG</u> | <u>A200909 003</u> | Apr 26, 2017 |
| AB | | <u>10MG; 80MG</u> | <u>A200909 004</u> | Apr 26, 2017 |
| AB | WATSON LABS INC | <u>10MG; 10MG</u> | <u>A202968 001</u> | Apr 26, 2017 |
| AB | | <u>10MG; 20MG</u> | <u>A202968 002</u> | Apr 26, 2017 |
| AB | | <u>10MG; 40MG</u> | <u>A202968 003</u> | Apr 26, 2017 |
| AB | | <u>10MG; 80MG</u> | <u>A202968 004</u> | Apr 26, 2017 |
| <u>VYTORIN</u> | | | | |
| AB | + MSD INTL | <u>10MG; 10MG</u> | <u>N021687 001</u> | Jul 23, 2004 |
| AB | + | <u>10MG; 20MG</u> | <u>N021687 002</u> | Jul 23, 2004 |
| AB | + | <u>10MG; 40MG</u> | <u>N021687 003</u> | Jul 23, 2004 |
| AB | + | <u>10MG; 80MG</u> | <u>N021687 004</u> | Jul 23, 2004 |

FAMCICLOVIR

TABLET; ORAL

FAMCICLOVIR

| | | | | |
|-----------|----------------------|---------------------|---------------------------|--------------|
| AB | APOTEX | <u>125MG</u> | <u>A091480 001</u> | Jul 22, 2011 |
| AB | | <u>250MG</u> | <u>A091480 002</u> | Jul 22, 2011 |
| AB | | <u>500MG</u> | <u>A091480 003</u> | Jul 22, 2011 |
| AB | AUROBINDO PHARMA LTD | <u>125MG</u> | <u>A091114 001</u> | Mar 21, 2011 |
| AB | | <u>250MG</u> | <u>A091114 002</u> | Mar 21, 2011 |
| AB | | <u>500MG</u> | <u>A091114 003</u> | Mar 21, 2011 |
| AB | CIPLA | <u>125MG</u> | <u>A078278 001</u> | Mar 21, 2011 |
| AB | | <u>250MG</u> | <u>A078278 002</u> | Mar 21, 2011 |
| AB | | <u>500MG</u> | <u>A078278 003</u> | Mar 21, 2011 |
| AB | HETERO LABS LTD V | <u>125MG</u> | <u>A202438 001</u> | Sep 10, 2014 |
| AB | | <u>250MG</u> | <u>A202438 002</u> | Sep 10, 2014 |
| AB | | <u>500MG</u> | <u>A202438 003</u> | Sep 10, 2014 |
| AB | MACLEODS PHARMS LTD | <u>125MG</u> | <u>A201022 001</u> | Jan 12, 2012 |
| AB | | <u>250MG</u> | <u>A201022 002</u> | Jan 12, 2012 |
| AB | | <u>500MG</u> | <u>A201022 003</u> | Jan 12, 2012 |
| AB | MYLAN | <u>125MG</u> | <u>A201333 001</u> | Mar 24, 2011 |
| AB | | <u>250MG</u> | <u>A201333 002</u> | Mar 24, 2011 |
| AB | | <u>500MG</u> | <u>A201333 003</u> | Mar 24, 2011 |
| AB | TEVA PHARMS | <u>125MG</u> | <u>A077487 001</u> | Aug 24, 2007 |
| AB | | <u>250MG</u> | <u>A077487 002</u> | Aug 24, 2007 |
| AB | ! | <u>500MG</u> | <u>A077487 003</u> | Aug 24, 2007 |
| AB | WEST-WARD PHARMS INT | <u>125MG</u> | <u>A090128 001</u> | Mar 21, 2011 |
| AB | | <u>250MG</u> | <u>A090128 002</u> | Mar 21, 2011 |
| AB | | <u>500MG</u> | <u>A090128 003</u> | Mar 21, 2011 |

FAMOTIDINE

FOR SUSPENSION; ORAL

FAMOTIDINE

| | | | | |
|-----------|-------------------|------------------------|---------------------------|--------------|
| AB | HI-TECH PHARMA CO | <u>40MG/5ML</u> | <u>A201995 001</u> | May 30, 2014 |
| AB | LUPIN LTD | <u>40MG/5ML</u> | <u>A090440 001</u> | Jun 29, 2010 |
| AB | NAVINTA LLC | <u>40MG/5ML</u> | <u>A091020 001</u> | May 27, 2010 |
| AB | NOVEL LABS INC | <u>40MG/5ML</u> | <u>A201695 001</u> | Dec 17, 2012 |

PEPCID

| | | | | |
|-----------|----------------|------------------------|---------------------------|--------------|
| AB | ! SALIX PHARMS | <u>40MG/5ML</u> | <u>N019527 001</u> | Feb 02, 1987 |
|-----------|----------------|------------------------|---------------------------|--------------|

INJECTABLE; INJECTION

FAMOTIDINE

| | | | | |
|-----------|------------------------|-----------------------|---------------------------|--------------|
| AP | ATHENEX INC | <u>10MG/ML</u> | <u>A075651 001</u> | Apr 16, 2001 |
| AP | | <u>10MG/ML</u> | <u>A075684 001</u> | Apr 16, 2001 |
| AP | FRESENIUS KABI USA | <u>10MG/ML</u> | <u>A075709 001</u> | Apr 16, 2001 |
| AP | MYLAN LABS LTD | <u>10MG/ML</u> | <u>A078641 001</u> | Jun 25, 2008 |
| AP | ! WEST-WARD PHARMS INT | <u>10MG/ML</u> | <u>A075488 001</u> | Apr 16, 2001 |

FAMOTIDINE PRESERVATIVE FREE

| | | | | |
|-----------|------------------------|-----------------------|---------------------------|--------------|
| AP | ATHENEX INC | <u>10MG/ML</u> | <u>A075622 001</u> | Apr 16, 2001 |
| AP | | <u>10MG/ML</u> | <u>A075825 001</u> | Apr 17, 2001 |
| AP | FRESENIUS KABI USA | <u>10MG/ML</u> | <u>A075813 001</u> | Apr 16, 2001 |
| AP | MYLAN LABS LTD | <u>10MG/ML</u> | <u>A078642 001</u> | Jun 25, 2008 |
| AP | ! WEST-WARD PHARMS INT | <u>10MG/ML</u> | <u>A075486 001</u> | Apr 16, 2001 |

FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER

| | | | | |
|----------|-----------------|------------------------|---------------------------|--------------|
| ! | BAXTER HLTHCARE | <u>0.4MG/ML</u> | <u>A075591 001</u> | May 10, 2001 |
|----------|-----------------|------------------------|---------------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

| | | | | | |
|-----------|------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | ALEMbic PHARMS LTD | <u>20MG</u> | <u>A078916</u> | <u>001</u> | May 22, 2009 |
| <u>AB</u> | | <u>40MG</u> | <u>A078916</u> | <u>002</u> | May 22, 2009 |
| <u>AB</u> | APOTEX | <u>20MG</u> | <u>A075611</u> | <u>001</u> | Jul 23, 2001 |
| <u>AB</u> | | <u>40MG</u> | <u>A075611</u> | <u>002</u> | Jul 23, 2001 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>20MG</u> | <u>A206530</u> | <u>001</u> | Dec 22, 2015 |
| <u>AB</u> | | <u>40MG</u> | <u>A206530</u> | <u>002</u> | Dec 22, 2015 |
| <u>AB</u> | CARLSBAD | <u>20MG</u> | <u>A075805</u> | <u>001</u> | Apr 16, 2001 |
| <u>AB</u> | | <u>40MG</u> | <u>A075805</u> | <u>002</u> | Apr 16, 2001 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>20MG</u> | <u>A075718</u> | <u>001</u> | Apr 16, 2001 |
| <u>AB</u> | | <u>40MG</u> | <u>A075718</u> | <u>002</u> | Apr 16, 2001 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>20MG</u> | <u>A075511</u> | <u>001</u> | Apr 16, 2001 |
| <u>AB</u> | | <u>40MG</u> | <u>A075511</u> | <u>002</u> | Apr 16, 2001 |
| <u>AB</u> | MYLAN | <u>20MG</u> | <u>A075704</u> | <u>001</u> | Apr 16, 2001 |
| <u>AB</u> | | <u>40MG</u> | <u>A075704</u> | <u>002</u> | Apr 16, 2001 |
| <u>AB</u> | PERRIGO R AND D | <u>20MG</u> | <u>A077352</u> | <u>002</u> | Jul 27, 2005 |
| <u>AB</u> | | <u>40MG</u> | <u>A077352</u> | <u>001</u> | Jul 27, 2005 |
| <u>AB</u> | TEVA | <u>20MG</u> | <u>A075311</u> | <u>001</u> | Apr 16, 2001 |
| <u>AB</u> | | <u>40MG</u> | <u>A075311</u> | <u>002</u> | Apr 16, 2001 |
| <u>AB</u> | WOCKHARDT LTD | <u>20MG</u> | <u>A075786</u> | <u>001</u> | Apr 16, 2001 |
| <u>AB</u> | | <u>40MG</u> | <u>A075786</u> | <u>002</u> | Apr 16, 2001 |
| | <u>PEPCID</u> | | | | |
| <u>AB</u> | + VALEANT PHARMS NORTH | <u>20MG</u> | <u>N019462</u> | <u>001</u> | Oct 15, 1986 |
| <u>AB</u> | +! | <u>40MG</u> | <u>N019462</u> | <u>002</u> | Oct 15, 1986 |

FAMOTIDINE; IBUPROFEN

TABLET; ORAL

DUEXIS

+! HORIZON 26.6MG; 800MG N022519 001 Apr 23, 2011

FEBUXOSTAT

TABLET; ORAL

ULORIC

+ TAKEDA PHARMS USA 40MG N021856 001 Feb 13, 2009
+! 80MG N021856 002 Feb 13, 2009FELBAMATE

SUSPENSION; ORAL

FELBAMATE

| | | | | | |
|-----------|------------------------|------------------|----------------|------------|--------------|
| <u>AB</u> | AMNEAL PHARMS | <u>600MG/5ML</u> | <u>A202385</u> | <u>001</u> | Dec 16, 2011 |
| <u>AB</u> | TARO PHARM | <u>600MG/5ML</u> | <u>A206314</u> | <u>001</u> | Jun 16, 2017 |
| | <u>FELBATOL</u> | | | | |
| <u>AB</u> | +! MYLAN SPECIALITY LP | <u>600MG/5ML</u> | <u>N020189</u> | <u>003</u> | Jul 29, 1993 |
| | <u>FELBAMATE</u> | | | | |
| <u>AB</u> | ALVOGEN MALTA | <u>400MG</u> | <u>A204595</u> | <u>001</u> | Jan 11, 2016 |
| <u>AB</u> | | <u>600MG</u> | <u>A204595</u> | <u>002</u> | Jan 11, 2016 |
| <u>AB</u> | AMNEAL PHARMS | <u>400MG</u> | <u>A201680</u> | <u>001</u> | Sep 13, 2011 |
| <u>AB</u> | | <u>600MG</u> | <u>A201680</u> | <u>002</u> | Sep 13, 2011 |
| <u>AB</u> | ANI PHARMS INC | <u>400MG</u> | <u>A202284</u> | <u>001</u> | Nov 04, 2015 |
| <u>AB</u> | | <u>600MG</u> | <u>A202284</u> | <u>002</u> | Nov 04, 2015 |
| <u>AB</u> | TARO PHARM | <u>400MG</u> | <u>A207093</u> | <u>001</u> | Apr 20, 2017 |
| <u>AB</u> | | <u>600MG</u> | <u>A207093</u> | <u>002</u> | Apr 20, 2017 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>400MG</u> | <u>A208970</u> | <u>001</u> | May 30, 2017 |
| <u>AB</u> | | <u>600MG</u> | <u>A208970</u> | <u>002</u> | May 30, 2017 |
| | <u>FELBATOL</u> | | | | |
| <u>AB</u> | + MYLAN SPECIALITY LP | <u>400MG</u> | <u>N020189</u> | <u>001</u> | Jul 29, 1993 |
| <u>AB</u> | +! | <u>600MG</u> | <u>N020189</u> | <u>002</u> | Jul 29, 1993 |

FELODIPINE

TABLET, EXTENDED RELEASE; ORAL

FELODIPINE

| | | | | | |
|-----------|----------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>2.5MG</u> | <u>A203417</u> | <u>001</u> | Jan 17, 2013 |
| <u>AB</u> | | <u>5MG</u> | <u>A203417</u> | <u>002</u> | Jan 17, 2013 |
| <u>AB</u> | | <u>10MG</u> | <u>A203417</u> | <u>003</u> | Jan 17, 2013 |
| <u>AB</u> | GLENMARK GENERICS | <u>2.5MG</u> | <u>A090365</u> | <u>001</u> | Dec 17, 2010 |
| <u>AB</u> | | <u>5MG</u> | <u>A090365</u> | <u>002</u> | Dec 17, 2010 |
| <u>AB</u> | | <u>10MG</u> | <u>A090365</u> | <u>003</u> | Dec 17, 2010 |

PRESCRIPTION DRUG PRODUCT LIST

FELODIPINE

TABLET, EXTENDED RELEASE;ORAL

FELODIPINE

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| <u>AB</u> | HERITAGE PHARMS INC | <u>2.5MG</u> | <u>A201964 001</u> | Nov 08, 2013 |
| <u>AB</u> | | <u>5MG</u> | <u>A201964 002</u> | Nov 08, 2013 |
| <u>AB</u> | | <u>10MG</u> | <u>A201964 003</u> | Nov 08, 2013 |
| <u>AB</u> | JUBILANT GENERICS | <u>2.5MG</u> | <u>A203983 001</u> | Aug 19, 2016 |
| <u>AB</u> | | <u>5MG</u> | <u>A203983 002</u> | Aug 19, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A203983 003</u> | Aug 19, 2016 |
| <u>AB</u> | MYLAN | <u>2.5MG</u> | <u>A078855 001</u> | Apr 17, 2008 |
| <u>AB</u> | | <u>5MG</u> | <u>A078855 002</u> | Apr 17, 2008 |
| <u>AB</u> | ! | <u>10MG</u> | <u>A078855 003</u> | Apr 17, 2008 |
| <u>AB</u> | ORCHID HLTHCARE | <u>2.5MG</u> | <u>A203032 001</u> | May 21, 2015 |
| <u>AB</u> | | <u>5MG</u> | <u>A203032 002</u> | May 21, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A203032 003</u> | May 21, 2015 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>2.5MG</u> | <u>A091200 001</u> | Dec 13, 2013 |
| <u>AB</u> | | <u>5MG</u> | <u>A091200 002</u> | Dec 13, 2013 |
| <u>AB</u> | | <u>10MG</u> | <u>A091200 003</u> | Dec 13, 2013 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>2.5MG</u> | <u>A075896 001</u> | Nov 02, 2004 |
| <u>AB</u> | | <u>5MG</u> | <u>A075896 002</u> | Nov 02, 2004 |
| <u>AB</u> | | <u>10MG</u> | <u>A075896 003</u> | Nov 02, 2004 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>2.5MG</u> | <u>A202170 001</u> | Nov 28, 2011 |
| <u>AB</u> | | <u>5MG</u> | <u>A202170 002</u> | Nov 28, 2011 |
| <u>AB</u> | | <u>10MG</u> | <u>A202170 003</u> | Nov 28, 2011 |
| <u>AB</u> | VINTAGE PHARMS LLC | <u>2.5MG</u> | <u>A200815 001</u> | Oct 28, 2011 |
| <u>AB</u> | | <u>5MG</u> | <u>A200815 002</u> | Oct 28, 2011 |
| <u>AB</u> | | <u>10MG</u> | <u>A200815 003</u> | Oct 28, 2011 |
| <u>AB</u> | YILING PHARM LTD | <u>2.5MG</u> | <u>A210847 001</u> | Oct 26, 2018 |
| <u>AB</u> | | <u>5MG</u> | <u>A210847 002</u> | Oct 26, 2018 |
| <u>AB</u> | | <u>10MG</u> | <u>A210847 003</u> | Oct 26, 2018 |

FENOFIBRATE

CAPSULE;ORAL

ANTARA (MICRONIZED)

| | | | | |
|-----------|------------------|--------------|--------------------|--------------|
| <u>AB</u> | + LUPIN ATLANTIS | <u>43MG</u> | <u>N021695 001</u> | Nov 30, 2004 |
| <u>AB</u> | +! | <u>130MG</u> | <u>N021695 003</u> | Nov 30, 2004 |

FENOFIBRATE

| | | | | |
|-----------|--------------------|--------------|--------------------|--------------|
| <u>AB</u> | SUN PHARM INDS LTD | <u>43MG</u> | <u>A201748 001</u> | Oct 31, 2014 |
| <u>AB</u> | | <u>130MG</u> | <u>A201748 002</u> | Oct 31, 2014 |

FENOFIBRATE (MICRONIZED)

| | | | | |
|-----------|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | AJANTA PHARMA LTD | <u>67MG</u> | <u>A210705 001</u> | Sep 10, 2018 |
| <u>AB</u> | | <u>134MG</u> | <u>A210705 002</u> | Sep 10, 2018 |
| <u>AB</u> | | <u>200MG</u> | <u>A210705 003</u> | Sep 10, 2018 |
| <u>AB</u> | AMERIGEN PHARMS LTD | <u>67MG</u> | <u>A209504 001</u> | Apr 30, 2018 |
| <u>AB</u> | | <u>134MG</u> | <u>A209504 002</u> | Apr 30, 2018 |
| <u>AB</u> | | <u>200MG</u> | <u>A209504 003</u> | Apr 30, 2018 |
| <u>AB</u> | APOTEX INC | <u>43MG</u> | <u>A202252 001</u> | Jul 26, 2013 |
| <u>AB</u> | | <u>130MG</u> | <u>A202252 002</u> | Jul 26, 2013 |
| <u>AB</u> | CNTY LINE PHARMS | <u>67MG</u> | <u>A207805 001</u> | Nov 16, 2017 |
| <u>AB</u> | | <u>134MG</u> | <u>A207805 002</u> | Nov 16, 2017 |
| <u>AB</u> | | <u>200MG</u> | <u>A207805 003</u> | Nov 16, 2017 |
| <u>AB</u> | DR REDDYS LABS SA | <u>43MG</u> | <u>A090859 001</u> | Mar 01, 2012 |
| <u>AB</u> | | <u>130MG</u> | <u>A090859 002</u> | Mar 01, 2012 |
| <u>AB</u> | GLENMARK PHARMS LTD | <u>67MG</u> | <u>A205566 001</u> | Apr 07, 2017 |
| <u>AB</u> | | <u>134MG</u> | <u>A205566 002</u> | Apr 07, 2017 |
| <u>AB</u> | | <u>200MG</u> | <u>A205566 003</u> | Apr 07, 2017 |
| <u>AB</u> | IMPAX LABS | <u>67MG</u> | <u>A075868 001</u> | Oct 27, 2003 |
| <u>AB</u> | | <u>134MG</u> | <u>A075868 002</u> | Oct 27, 2003 |
| <u>AB</u> | ! | <u>200MG</u> | <u>A075868 003</u> | Oct 27, 2003 |
| <u>AB</u> | INVAGEN PHARMS | <u>67MG</u> | <u>A207378 001</u> | Mar 28, 2017 |
| <u>AB</u> | | <u>134MG</u> | <u>A207378 002</u> | Mar 28, 2017 |
| <u>AB</u> | | <u>200MG</u> | <u>A207378 003</u> | Mar 28, 2017 |
| <u>AB</u> | MYLAN PHARMS INC | <u>43MG</u> | <u>A202579 001</u> | Jan 10, 2013 |
| <u>AB</u> | | <u>67MG</u> | <u>A202676 001</u> | Oct 23, 2012 |
| <u>AB</u> | | <u>130MG</u> | <u>A202579 002</u> | Jan 10, 2013 |
| <u>AB</u> | | <u>134MG</u> | <u>A202676 002</u> | Oct 23, 2012 |
| <u>AB</u> | | <u>200MG</u> | <u>A202676 003</u> | Oct 23, 2012 |
| <u>AB</u> | RHODES PHARMS | <u>67MG</u> | <u>A075753 001</u> | Sep 03, 2002 |
| <u>AB</u> | | <u>134MG</u> | <u>A075753 002</u> | Apr 09, 2002 |
| <u>AB</u> | | <u>200MG</u> | <u>A075753 003</u> | Apr 09, 2002 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>67MG</u> | <u>A210782 001</u> | Jun 26, 2018 |
| <u>AB</u> | | <u>134MG</u> | <u>A210782 002</u> | Jun 26, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

FENOFIBRATE

CAPSULE; ORAL

FENOFIBRATE (MICRONIZED)

| | | | | |
|-----------|---------------------|--------------|--------------------|--------------|
| AB | | 200MG | A210782 003 | Jun 26, 2018 |
| | ANTARA (MICRONIZED) | | | |
| | + LUPIN ATLANTIS | 30MG | N021695 004 | Oct 18, 2013 |
| | + LIPOFEN | 90MG | N021695 005 | Oct 18, 2013 |
| | + CIPHER PHARMS INC | 50MG | N021612 001 | Jan 11, 2006 |
| | +! | 150MG | N021612 003 | Jan 11, 2006 |

TABLET; ORAL

FENOFIBRATE

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| AB | AJANTA PHARMA LTD | 54MG | A210138 001 | Jul 23, 2018 |
| AB | | 160MG | A210138 002 | Jul 23, 2018 |
| AB | AMNEAL PHARMS LLC | 48MG | A209951 001 | Feb 09, 2018 |
| AB | | 54MG | A209950 001 | Mar 19, 2018 |
| AB | | 145MG | A209951 002 | Feb 09, 2018 |
| AB | | 160MG | A209950 002 | Mar 19, 2018 |
| AB | AUROBINDO PHARMA LTD | 48MG | A205118 001 | May 05, 2016 |
| AB | | 145MG | A205118 002 | May 05, 2016 |
| AB | CIPLA | 48MG | A208709 001 | Dec 15, 2016 |
| AB | | 145MG | A208709 002 | Dec 15, 2016 |
| AB | CNTY LINE PHARMS | 54MG | A207803 001 | Dec 19, 2017 |
| AB | | 160MG | A207803 002 | Dec 19, 2017 |
| AB | GRAVITI PHARMS | 54MG | A210606 001 | Aug 17, 2018 |
| AB | | 160MG | A210606 002 | Aug 17, 2018 |
| AB | HETERO LABS LTD III | 48MG | A204598 001 | Jul 12, 2016 |
| AB | | 145MG | A204598 002 | Jul 12, 2016 |
| AB | IMPAX LABS | 54MG | A076509 001 | Mar 26, 2008 |
| AB | ! | 160MG | A076509 002 | Mar 26, 2008 |
| AB | LUPIN LTD | 48MG | A090856 001 | Dec 23, 2011 |
| AB | | 54MG | A204019 001 | Aug 17, 2015 |
| AB | | 145MG | A090856 002 | Dec 23, 2011 |
| AB | | 160MG | A204019 002 | Aug 17, 2015 |
| AB | MYLAN | 40MG | A204475 001 | Jun 23, 2016 |
| AB | | 54MG | A076520 001 | Oct 25, 2007 |
| AB | | 120MG | A204475 002 | Jun 23, 2016 |
| AB | | 160MG | A076520 003 | Oct 25, 2007 |
| AB | MYLAN PHARMS INC | 48MG | A202856 001 | Dec 07, 2012 |
| AB | | 145MG | A202856 002 | Dec 07, 2012 |
| AB | PRINSTON INC | 48MG | A211080 001 | Aug 28, 2018 |
| AB | | 145MG | A211080 002 | Aug 28, 2018 |
| AB | RHODES PHARMS | 54MG | A076433 001 | May 13, 2005 |
| AB | | 160MG | A076433 002 | May 13, 2005 |
| AB | SUN PHARM INDS LTD | 48MG | A200884 001 | Sep 07, 2017 |
| AB | | 54MG | A076635 001 | Oct 31, 2005 |
| AB | | 145MG | A200884 002 | Sep 07, 2017 |
| AB | | 160MG | A076635 003 | Oct 31, 2005 |
| AB | VALEANT PHARMS NORTH | 48MG | A090715 001 | Apr 05, 2012 |
| AB | | 145MG | A090715 002 | Apr 05, 2012 |
| | <u>FENOGLIDE</u> | | | |
| AB | + SANTARUS INC | 40MG | N022118 001 | Aug 10, 2007 |
| AB | +! | 120MG | N022118 002 | Aug 10, 2007 |
| | <u>TRICOR</u> | | | |
| AB | + ABEVIE | 48MG | N021656 001 | Nov 05, 2004 |
| AB | +! | 145MG | N021656 002 | Nov 05, 2004 |
| | <u>TRIGLIDE</u> | | | |
| BX | +! SKYEPHARMA AG | 160MG | N021350 002 | May 07, 2005 |
| | FENOFIBRATE | | | |
| | SUN PHARM INDS LTD | 107MG | A076635 002 | Oct 31, 2005 |

FENOFIBRIC ACID

TABLET; ORAL

FIBRICOR

| | | | | |
|--|---------------------|-------|-------------|--------------|
| | + ARALEZ PHARMS INC | 35MG | N022418 001 | Aug 14, 2009 |
| | +! | 105MG | N022418 002 | Aug 14, 2009 |

PRESCRIPTION DRUG PRODUCT LIST

FENOLDOPAM MESYLATE

INJECTABLE; INJECTION

CORLOPAM

| | | | | | |
|-----------|-----------|---------|------------------------|--------------------|--------------|
| <u>AP</u> | <u>+!</u> | HOSPIRA | <u>EQ 10MG BASE/ML</u> | <u>N019922 001</u> | Sep 23, 1997 |
|-----------|-----------|---------|------------------------|--------------------|--------------|

FENOLDOPAM MESYLATE

| | | | | | |
|-----------|--|------------|------------------------|--------------------|--------------|
| <u>AP</u> | | SANDOZ INC | <u>EQ 10MG BASE/ML</u> | <u>A077155 001</u> | Feb 15, 2005 |
|-----------|--|------------|------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------------------------|------------------------|--------------------|--------------|
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>EQ 10MG BASE/ML</u> | <u>A076582 001</u> | Oct 12, 2004 |
|-----------|--|-------------------------|------------------------|--------------------|--------------|

FENOPROFEN CALCIUM

CAPSULE; ORAL

NALFON

| | | | | | |
|--|---|---------------|---------------|-------------|--|
| | + | XSPIRE PHARMA | EQ 200MG BASE | N017604 003 | |
|--|---|---------------|---------------|-------------|--|

| | | | | | |
|--|---|--|---------------|-------------|--------------|
| | + | | EQ 400MG BASE | N017604 004 | Jul 21, 2009 |
|--|---|--|---------------|-------------|--------------|

TABLET; ORAL

FENOPROFEN CALCIUM

| | | | | | |
|--|---|---------------|---------------|-------------|--------------|
| | ! | XSPIRE PHARMA | EQ 600MG BASE | A072267 001 | Aug 17, 1988 |
|--|---|---------------|---------------|-------------|--------------|

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL

DURAGESIC-100

| | | | | | |
|-----------|---|----------------|------------------|--------------------|--------------|
| <u>AB</u> | + | JANSSEN PHARMS | <u>100MCG/HR</u> | <u>N019813 001</u> | Aug 07, 1990 |
|-----------|---|----------------|------------------|--------------------|--------------|

DURAGESIC-12

| | | | | | |
|-----------|---|----------------|-------------------|--------------------|--------------|
| <u>AB</u> | + | JANSSEN PHARMS | <u>12.5MCG/HR</u> | <u>N019813 005</u> | Feb 04, 2005 |
|-----------|---|----------------|-------------------|--------------------|--------------|

DURAGESIC-25

| | | | | | |
|-----------|-----------|----------------|-----------------|--------------------|--------------|
| <u>AB</u> | <u>+!</u> | JANSSEN PHARMS | <u>25MCG/HR</u> | <u>N019813 004</u> | Aug 07, 1990 |
|-----------|-----------|----------------|-----------------|--------------------|--------------|

DURAGESIC-37

| | | | | | |
|-----------|---|----------------|-------------------|--------------------|--------------|
| <u>AB</u> | + | JANSSEN PHARMS | <u>37.5MCG/HR</u> | <u>N019813 006</u> | Jan 24, 2018 |
|-----------|---|----------------|-------------------|--------------------|--------------|

DURAGESIC-50

| | | | | | |
|-----------|---|----------------|-----------------|--------------------|--------------|
| <u>AB</u> | + | JANSSEN PHARMS | <u>50MCG/HR</u> | <u>N019813 003</u> | Aug 07, 1990 |
|-----------|---|----------------|-----------------|--------------------|--------------|

DURAGESIC-75

| | | | | | |
|-----------|---|----------------|-----------------|--------------------|--------------|
| <u>AB</u> | + | JANSSEN PHARMS | <u>75MCG/HR</u> | <u>N019813 002</u> | Aug 07, 1990 |
|-----------|---|----------------|-----------------|--------------------|--------------|

FENTANYL-100

| | | | | | |
|-----------|--|------------------|------------------|--------------------|--------------|
| <u>AB</u> | | 3M DRUG DELIVERY | <u>100MCG/HR</u> | <u>A202097 005</u> | Nov 04, 2016 |
|-----------|--|------------------|------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------|------------------|--------------------|--------------|
| <u>AB</u> | | AVEVA | <u>100MCG/HR</u> | <u>A077449 004</u> | Oct 20, 2008 |
|-----------|--|-------|------------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------|------------------|--------------------|--------------|
| <u>AB</u> | | LAVIPHARM LABS | <u>100MCG/HR</u> | <u>A077051 004</u> | Aug 04, 2006 |
|-----------|--|----------------|------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------|------------------|--------------------|--------------|
| <u>AB</u> | | MAYNE PHARMA | <u>100MCG/HR</u> | <u>A077062 004</u> | Aug 20, 2007 |
|-----------|--|--------------|------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------------|------------------|--------------------|--------------|
| <u>AB</u> | | MYLAN TECHNOLOGIES | <u>100MCG/HR</u> | <u>A076258 004</u> | Jan 28, 2005 |
|-----------|--|--------------------|------------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------|------------------|--------------------|--------------|
| <u>AB</u> | | SPECGX LLC | <u>100MCG/HR</u> | <u>A077154 004</u> | Feb 09, 2011 |
|-----------|--|------------|------------------|--------------------|--------------|

FENTANYL-12

| | | | | | |
|-----------|--|------------------|-------------------|--------------------|--------------|
| <u>AB</u> | | 3M DRUG DELIVERY | <u>12.5MCG/HR</u> | <u>A202097 001</u> | Nov 04, 2016 |
|-----------|--|------------------|-------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------|-------------------|--------------------|--------------|
| <u>AB</u> | | AVEVA | <u>12.5MCG/HR</u> | <u>A077449 005</u> | Sep 11, 2015 |
|-----------|--|-------|-------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------------|-------------------|--------------------|--------------|
| <u>AB</u> | | MYLAN TECHNOLOGIES | <u>12.5MCG/HR</u> | <u>A076258 005</u> | Jan 23, 2007 |
|-----------|--|--------------------|-------------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------|-------------------|--------------------|--------------|
| <u>AB</u> | | SPECGX LLC | <u>12.5MCG/HR</u> | <u>A077154 005</u> | Jun 11, 2015 |
|-----------|--|------------|-------------------|--------------------|--------------|

FENTANYL-25

| | | | | | |
|-----------|--|------------------|-----------------|--------------------|--------------|
| <u>AB</u> | | 3M DRUG DELIVERY | <u>25MCG/HR</u> | <u>A202097 002</u> | Nov 04, 2016 |
|-----------|--|------------------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------|-----------------|--------------------|--------------|
| <u>AB</u> | | AVEVA | <u>25MCG/HR</u> | <u>A077449 001</u> | Oct 20, 2008 |
|-----------|--|-------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------|-----------------|--------------------|--------------|
| <u>AB</u> | | LAVIPHARM LABS | <u>25MCG/HR</u> | <u>A077051 001</u> | Aug 04, 2006 |
|-----------|--|----------------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------|-----------------|--------------------|--------------|
| <u>AB</u> | | MAYNE PHARMA | <u>25MCG/HR</u> | <u>A077062 001</u> | Aug 20, 2007 |
|-----------|--|--------------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------------|-----------------|--------------------|--------------|
| <u>AB</u> | | MYLAN TECHNOLOGIES | <u>25MCG/HR</u> | <u>A076258 001</u> | Jan 28, 2005 |
|-----------|--|--------------------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------|-----------------|--------------------|--------------|
| <u>AB</u> | | SPECGX LLC | <u>25MCG/HR</u> | <u>A077154 001</u> | Feb 09, 2011 |
|-----------|--|------------|-----------------|--------------------|--------------|

FENTANYL-37

| | | | | | |
|-----------|--|-------|-------------------|--------------------|--------------|
| <u>AB</u> | | AVEVA | <u>37.5MCG/HR</u> | <u>A077449 006</u> | Dec 06, 2017 |
|-----------|--|-------|-------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------------|-------------------|--------------------|--------------|
| <u>AB</u> | | MYLAN TECHNOLOGIES | <u>37.5MCG/HR</u> | <u>A076258 006</u> | Dec 29, 2014 |
|-----------|--|--------------------|-------------------|--------------------|--------------|

FENTANYL-50

| | | | | | |
|-----------|--|------------------|-----------------|--------------------|--------------|
| <u>AB</u> | | 3M DRUG DELIVERY | <u>50MCG/HR</u> | <u>A202097 003</u> | Nov 04, 2016 |
|-----------|--|------------------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------|-----------------|--------------------|--------------|
| <u>AB</u> | | AVEVA | <u>50MCG/HR</u> | <u>A077449 002</u> | Oct 20, 2008 |
|-----------|--|-------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------|-----------------|--------------------|--------------|
| <u>AB</u> | | LAVIPHARM LABS | <u>50MCG/HR</u> | <u>A077051 002</u> | Aug 04, 2006 |
|-----------|--|----------------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------|-----------------|--------------------|--------------|
| <u>AB</u> | | MAYNE PHARMA | <u>50MCG/HR</u> | <u>A077062 002</u> | Aug 20, 2007 |
|-----------|--|--------------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------------|-----------------|--------------------|--------------|
| <u>AB</u> | | MYLAN TECHNOLOGIES | <u>50MCG/HR</u> | <u>A076258 002</u> | Jan 28, 2005 |
|-----------|--|--------------------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------|-----------------|--------------------|--------------|
| <u>AB</u> | | SPECGX LLC | <u>50MCG/HR</u> | <u>A077154 002</u> | Feb 09, 2011 |
|-----------|--|------------|-----------------|--------------------|--------------|

FENTANYL-62

| | | | | | |
|-----------|--|-------|-------------------|--------------------|--------------|
| <u>AB</u> | | AVEVA | <u>62.5MCG/HR</u> | <u>A077449 007</u> | Dec 06, 2017 |
|-----------|--|-------|-------------------|--------------------|--------------|

FENTANYL-75

| | | | | | |
|-----------|--|------------------|-----------------|--------------------|--------------|
| <u>AB</u> | | 3M DRUG DELIVERY | <u>75MCG/HR</u> | <u>A202097 004</u> | Nov 04, 2016 |
|-----------|--|------------------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------|-----------------|--------------------|--------------|
| <u>AB</u> | | AVEVA | <u>75MCG/HR</u> | <u>A077449 003</u> | Oct 20, 2008 |
|-----------|--|-------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------|-----------------|--------------------|--------------|
| <u>AB</u> | | LAVIPHARM LABS | <u>75MCG/HR</u> | <u>A077051 003</u> | Aug 04, 2006 |
|-----------|--|----------------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------|-----------------|--------------------|--------------|
| <u>AB</u> | | MAYNE PHARMA | <u>75MCG/HR</u> | <u>A077062 003</u> | Aug 20, 2007 |
|-----------|--|--------------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------------|-----------------|--------------------|--------------|
| <u>AB</u> | | MYLAN TECHNOLOGIES | <u>75MCG/HR</u> | <u>A076258 003</u> | Jan 28, 2005 |
|-----------|--|--------------------|-----------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------|-----------------|--------------------|--------------|
| <u>AB</u> | | SPECGX LLC | <u>75MCG/HR</u> | <u>A077154 003</u> | Feb 09, 2011 |
|-----------|--|------------|-----------------|--------------------|--------------|

FENTANYL-87

| | | | | | |
|-----------|--|-------|-------------------|--------------------|--------------|
| <u>AB</u> | | AVEVA | <u>87.5MCG/HR</u> | <u>A077449 008</u> | Dec 06, 2017 |
|-----------|--|-------|-------------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL

FENTANYL-62

MYLAN TECHNOLOGIES 62.5MCG/HR

A076258 007 Dec 29, 2014

FENTANYL-87

MYLAN TECHNOLOGIES 87.5MCG/HR

A076258 008 Dec 29, 2014

SPRAY; SUBLINGUAL

SUBSYS

+ INSYS DEV CO INC

0.1MG

N202788 001 Jan 04, 2012

+ 0.2MG

N202788 002 Jan 04, 2012

+! 0.4MG

N202788 003 Jan 04, 2012

+ 0.6MG

N202788 004 Jan 04, 2012

+ 0.8MG

N202788 005 Jan 04, 2012

+ 1.2MG

N202788 006 Aug 30, 2012

+ 1.6MG

N202788 007 Aug 30, 2012

FENTANYL CITRATE

INJECTABLE; INJECTION

FENTANYL CITRATEAP HOSPIRAEQ 0.05MG BASE/MLN019115 001 Jan 12, 1985FENTANYL CITRATE PRESERVATIVE FREEAP HOSPIRAEQ 0.05MG BASE/MLA072786 001 Sep 24, 1991AP +! WEST-WARD PHARMS
INTEQ 0.05MG BASE/MLN019101 001 Jul 11, 1984SUBLIMAZE PRESERVATIVE FREEAP +! AKORNEQ 0.05MG BASE/MLN016619 001

SPRAY, METERED; NASAL

LAZANDA

+ ELEFSEE PHARMS INTL

EQ 0.1MG BASE

N022569 001 Jun 30, 2011

+ EQ 0.3MG BASE

N022569 003 Dec 21, 2015

+! EQ 0.4MG BASE

N022569 002 Jun 30, 2011

TABLET; BUCCAL, SUBLINGUAL

FENTORA

+ CEPHALON

EQ 0.1MG BASE

N021947 001 Sep 25, 2006

+ EQ 0.2MG BASE

N021947 002 Sep 25, 2006

+! EQ 0.4MG BASE

N021947 003 Sep 25, 2006

+ EQ 0.6MG BASE

N021947 004 Sep 25, 2006

+ EQ 0.8MG BASE

N021947 005 Sep 25, 2006

TABLET; SUBLINGUAL

ABSTRALAB + SENTYNL THERAPS INCEQ 0.1MG BASEN022510 001 Jan 07, 2011AB +EQ 0.2MG BASEN022510 002 Jan 07, 2011AB +EQ 0.3MG BASEN022510 003 Jan 07, 2011AB +!EQ 0.4MG BASEN022510 004 Jan 07, 2011AB +EQ 0.6MG BASEN022510 005 Jan 07, 2011AB +EQ 0.8MG BASEN022510 006 Jan 07, 2011FENTANYL CITRATEAB ACTAVIS LABS FL INCEQ 0.1MG BASEA207338 001 Nov 17, 2017ABEQ 0.2MG BASEA207338 002 Nov 17, 2017ABEQ 0.3MG BASEA207338 003 Nov 17, 2017ABEQ 0.4MG BASEA207338 004 Nov 17, 2017ABEQ 0.6MG BASEA207338 005 Nov 17, 2017ABEQ 0.8MG BASEA207338 006 Nov 17, 2017

TROCHE/LOZENGE; TRANSMUCOSAL

ACTIQAB + CEPHALONEQ 0.2MG BASEN020747 001 Nov 04, 1998AB +!EQ 0.4MG BASEN020747 002 Nov 04, 1998AB +EQ 0.6MG BASEN020747 003 Nov 04, 1998AB +EQ 0.8MG BASEN020747 004 Nov 04, 1998AB +EQ 1.2MG BASEN020747 005 Nov 04, 1998AB +EQ 1.6MG BASEN020747 006 Nov 04, 1998FENTANYL CITRATEAB SPECGX LLCEQ 0.2MG BASEA078907 001 Oct 30, 2009ABEQ 0.4MG BASEA078907 002 Oct 30, 2009ABEQ 0.6MG BASEA078907 003 Oct 30, 2009ABEQ 0.8MG BASEA078907 004 Oct 30, 2009ABEQ 1.2MG BASEA078907 005 Oct 30, 2009ABEQ 1.6MG BASEA078907 006 Oct 30, 2009

PRESCRIPTION DRUG PRODUCT LIST

FERRIC CARBOXYMALTOSE

INJECTABLE; INTRAVENOUS

INJECTAFER

+! LUITPOLD 750MG IRON/15ML (50MG IRON/ML) N203565 001 Jul 25, 2013

FERRIC CITRATE

TABLET; ORAL

AURYXIA

+! KERYX BIOPHARMS EQ 210MG IRON N205874 001 Sep 05, 2014

FERRIC HEXACYANOFERRATE (II)

CAPSULE; ORAL

RADIOGARDASE (PRUSSIAN BLUE)

+! HEYL CHEMISCH 500MG N021626 001 Oct 02, 2003

FERRIC PYROPHOSPHATE CITRATE

FOR SOLUTION; INTRAVENOUS

TRIFERIC

+! ROCKWELL MEDICAL INC 272MG IRON/PACKET N208551 001 Apr 25, 2016

SOLUTION; INTRAVENOUS

TRIFERIC

+! ROCKWELL MEDICAL INC 27.2MG IRON/5ML (5.44MG IRON/ML) N206317 001 Jan 23, 2015

FERUMOXYTOL

SOLUTION; INTRAVENOUS

FERAHEME

+! AMAG PHARMS INC EQ 510MG IRON/17ML (EQ 30MG IRON/ML) N022180 001 Jun 30, 2009

FESOTERODINE FUMARATE

TABLET, EXTENDED RELEASE; ORAL

FESOTERODINE FUMARATE**AB** AUROBINDO PHARMA LTD **4MG** **A205007 001** Feb 17, 2017**AB** **8MG** **A205007 002** Feb 17, 2017**AB** ZYDUS PHARMS USA INC **4MG** **A204946 001** Oct 03, 2017**AB** **8MG** **A204946 002** Oct 03, 2017TOVIAZ**AB** + PFIZER **4MG** **N022030 001** Oct 31, 2008**AB** +! **8MG** **N022030 002** Oct 31, 2008FEXOFENADINE HYDROCHLORIDE

TABLET; ORAL

FEXOFENADINE HYDROCHLORIDE**AB** BARR **30MG** **A076191 001** Aug 31, 2005**AB** **60MG** **A076191 002** Aug 31, 2005**AB** **180MG** **A076191 003** Aug 31, 2005**AB** DR REDDYS LABS LTD **30MG** **A076502 001** Apr 11, 2006**AB** **60MG** **A076502 002** Apr 11, 2006**AB** **180MG** **A076502 003** Apr 11, 2006**AB** MYLAN **30MG** **A077081 002** Apr 11, 2008**AB** **60MG** **A077081 003** Apr 11, 2008**AB** **180MG** **A077081 001** Apr 16, 2007**AB** TEVA **30MG** **A076447 001** Sep 01, 2005**AB** **60MG** **A076447 002** Sep 01, 2005**AB** **180MG** **A076447 003** Sep 01, 2005FIDAXOMICIN

TABLET; ORAL

DIFICID

+! CUBIST PHARMS LLC 200MG N201699 001 May 27, 2011

FINAFLOXACIN

SUSPENSION/DROPS; OTIC

XTORO

+! MERLION PHARMS GMBH 0.3% N206307 001 Dec 17, 2014

FINASTERIDE

TABLET; ORAL

FINASTERIDE**AB** ACCORD HLTHCARE **1MG** **A091643 001** Nov 05, 2013**AB** **5MG** **A090121 001** Feb 23, 2010**AB** ACTAVIS TOTOWA **1MG** **A078371 001** Nov 05, 2013**AB** ACTAVIS TOTOWA TEVA **5MG** **A077914 001** Mar 28, 2007**AB** ALKEM LABS LTD **1MG** **A207750 001** Jan 06, 2017**AB** **5MG** **A204304 001** Jan 05, 2017

PRESCRIPTION DRUG PRODUCT LIST

FINASTERIDE

TABLET; ORAL

FINASTERIDE

| | | | | |
|-----------|-------------------------|------------|--------------------|--------------|
| AB | AUROBINDO PHARMA | 5MG | A078341 001 | Oct 30, 2007 |
| AB | AUROBINDO PHARMA LTD | 1MG | A203687 001 | Nov 05, 2013 |
| AB | CIPLA | 1MG | A077335 001 | Nov 20, 2014 |
| AB | DR REDDYS LABS INC | 1MG | A076436 001 | Jul 28, 2006 |
| AB | DR REDDYS LABS LTD | 5MG | A076437 001 | Feb 28, 2007 |
| AB | HETERO LABS LTD III | 1MG | A090060 001 | Jul 01, 2013 |
| AB | | 5MG | A090061 001 | Jun 07, 2010 |
| AB | MYLAN | 5MG | A077578 001 | Dec 18, 2006 |
| AB | SUN PHARMA GLOBAL | 1MG | A090508 001 | Jul 01, 2013 |
| AB | | 5MG | A090507 001 | Aug 16, 2011 |
| AB | TEVA | 1MG | A076905 001 | Nov 05, 2013 |
| AB | | 5MG | A076511 001 | Dec 15, 2006 |
| AB | ZYDUS PHARMS USA INC | 5MG | A078900 001 | Dec 28, 2009 |

PROPECIA

| | | | | |
|-----------|------------------|------------|--------------------|--------------|
| AB | + ! MERCK | 1MG | N020788 001 | Dec 19, 1997 |
|-----------|------------------|------------|--------------------|--------------|

PROSCAR

| | | | | |
|-----------|------------------|------------|--------------------|--------------|
| AB | + ! MERCK | 5MG | N020180 001 | Jun 19, 1992 |
|-----------|------------------|------------|--------------------|--------------|

FINGOLIMOD HYDROCHLORIDE

CAPSULE; ORAL

GILENYA

| | | | | |
|------------|----------|----------------|-------------|--------------|
| + | NOVARTIS | EQ 0.25MG BASE | N022527 002 | May 11, 2018 |
| + ! | | EQ 0.5MG BASE | N022527 001 | Sep 21, 2010 |

FISH OIL TRIGLYCERIDES

EMULSION; INTRAVENOUS

OMEGAVEN

| | | | | |
|------------|--------------------|-----------------------|-------------|--------------|
| + ! | FRESENIUS KABI USA | 5GM/50ML (0.1GM/ML) | N210589 001 | Jul 27, 2018 |
| + ! | | 10GM/100ML (0.1GM/ML) | N210589 002 | Jul 27, 2018 |

FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL

EMULSION; INTRAVENOUS

SMOFLIPID 20%

| | | | | |
|------------|--------------------|--------------------------------------------------------|-------------|--------------|
| + ! | FRESENIUS KABI USA | 3GM/100ML; 6GM/100ML; 5GM/100ML; 6GM/100ML (100ML) | N207648 001 | Jul 13, 2016 |
| + ! | | 3GM/100ML; 6GM/100ML; 5GM/100ML; 6GM/100ML (250ML) | N207648 002 | Jul 13, 2016 |
| + ! | | 3GM/100ML; 6GM/100ML; 5GM/100ML; 6GM/100ML (500ML) | N207648 003 | Jul 13, 2016 |
| + ! | | 3GM/100ML; 6GM/100ML; 5GM/100ML; 6GM/100ML (1000ML) | N207648 004 | Aug 10, 2018 |

FLAVOXATE HYDROCHLORIDE

TABLET; ORAL

FLAVOXATE HYDROCHLORIDE

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| AB | EPIC PHARMA | 100MG | A076835 001 | Nov 30, 2005 |
| AB | ! PADDOCK LLC | 100MG | A076831 001 | Dec 16, 2004 |

FLECAINIDE ACETATE

TABLET; ORAL

FLECAINIDE ACETATE

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| AB | AMNEAL PHARM | 50MG | A075442 001 | Jul 31, 2001 |
| AB | | 100MG | A075442 002 | Jul 31, 2001 |
| AB | | 150MG | A075442 003 | Jul 31, 2001 |
| AB | ANI PHARMS INC | 50MG | A075882 001 | Oct 28, 2002 |
| AB | | 100MG | A075882 002 | Oct 28, 2002 |
| AB | | 150MG | A075882 003 | Oct 28, 2002 |
| AB | AUROBINDO PHARMA LTD | 50MG | A202821 001 | Nov 03, 2017 |
| AB | | 100MG | A202821 002 | Nov 03, 2017 |
| AB | | 150MG | A202821 003 | Nov 03, 2017 |
| AB | SUN PHARM INDS LTD | 50MG | A076421 001 | Mar 28, 2003 |
| AB | | 100MG | A076421 002 | Mar 28, 2003 |
| AB | | 150MG | A076421 003 | Mar 28, 2003 |
| AB | WEST-WARD PHARMS INT | 50MG | A076278 001 | Jan 14, 2003 |
| AB | | 100MG | A076278 002 | Jan 14, 2003 |
| AB | ! | 150MG | A076278 003 | Jan 14, 2003 |

TAMBOCOR

| | | | | |
|-----------|---------------------------|--------------|--------------------|--------------|
| AB | + CNTY LINE PHARMS | 50MG | N018830 004 | Aug 23, 1988 |
| AB | + | 100MG | N018830 001 | Oct 31, 1985 |

PRESCRIPTION DRUG PRODUCT LIST

FLECAINIDE ACETATE

TABLET; ORAL

TAMBOCOR

AB + 150MG N018830 003 Jun 03, 1988

FLIBANSERIN

TABLET; ORAL

ADDYI

+! SPROUT PHARMS 100MG N022526 001 Aug 18, 2015

FLORBETABEN F-18

SOLUTION; INTRAVENOUS

NEURACEQ

+! LIFE MOLECULAR 30ML (1.4-135mCi/ML) N204677 001 Mar 19, 2014

FLORBETAPIR F-18

SOLUTION; INTRAVENOUS

AMYVID

+! AVID RADIOPHARMS 10-30ML (13.5-51mCi/ML) N202008 002 Apr 06, 2012

+! INC 10-50ML (13.5-51mCi/ML) N202008 003 Apr 06, 2012

FLOXURIDINE

INJECTABLE; INJECTION

FLOXURIDINE

AP FRESENIUS KABI USA 500MG/VIAL A075837 001 Feb 22, 2001

AP LUITPOLD 500MG/VIAL A203008 001 Nov 22, 2017

AP ! WEST-WARD PHARMS 500MG/VIAL A075387 001 Apr 16, 2000
INT

FLUCICLOVINE F-18

SOLUTION; INTRAVENOUS

AXUMIN

+! BLUE EARTH 9-221mCi/ML N208054 001 May 27, 2016

FLUCONAZOLE

FOR SUSPENSION; ORAL

DIFLUCAN

AB + PFIZER 50MG/5ML N020090 001 Dec 23, 1993

AB +! 200MG/5ML N020090 002 Dec 23, 1993

FLUCONAZOLE

AB AUROBINDO PHARMA 50MG/5ML A079150 001 Sep 18, 2009
LTD

AB 200MG/5ML A079150 002 Sep 18, 2009

AB IVAX SUB TEVA 50MG/5ML A077523 001 Sep 12, 2007

AB PHARMS 200MG/5ML A077523 002 Sep 12, 2007

AB WEST-WARD PHARMS 50MG/5ML A076246 001 Jul 29, 2004

AB INT 200MG/5ML A076246 002 Jul 29, 2004

INJECTABLE; INJECTION

FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER

AP HIKMA FARMACEUTICA 200MG/100ML (2MG/ML) A078764 001 Jan 30, 2012

AP 400MG/200ML (2MG/ML) A078764 002 Jan 30, 2012

AP ! HOSPIRA 200MG/100ML (2MG/ML) A076304 001 Jul 29, 2004

AP ! 400MG/200ML (2MG/ML) A076304 002 Jul 29, 2004

AP RENAISSANCE SSA LLC 200MG/100ML (2MG/ML) A077988 001 May 26, 2010

AP 400MG/200ML (2MG/ML) A077988 002 May 26, 2010

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

AP BAXTER HLTHCARE 200MG/100ML (2MG/ML) A077947 001 May 26, 2010
CORP

AP 400MG/200ML (2MG/ML) A077947 002 May 26, 2010

AP FRESENIUS KABI USA 200MG/100ML (2MG/ML) A076145 001 Jul 29, 2004

AP 400MG/200ML (2MG/ML) A076145 002 Jul 29, 2004

AP HIKMA FARMACEUTICA 200MG/100ML (2MG/ML) A076736 001 Aug 23, 2005

AP WEST-WARD PHARMS 200MG/100ML (2MG/ML) A076087 001 Jul 29, 2004

AP 400MG/200ML (2MG/ML) A076087 003 Jul 29, 2004

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 200MG/100ML (2MG/ML) A076766 001 Jul 29, 2004

AP 400MG/200ML (2MG/ML) A076766 002 Jul 29, 2004

AP HIKMA FARMACEUTICA 200MG/100ML (2MG/ML) A078698 001 Jan 30, 2012

AP 400MG/200ML (2MG/ML) A078698 002 Jan 30, 2012

AP HOSPIRA 200MG/100ML (2MG/ML) A076303 001 Jul 29, 2004

AP 400MG/200ML (2MG/ML) A076303 002 Jul 29, 2004

AP ! INFORLIFE 200MG/100ML (2MG/ML) A079104 001 Jul 30, 2009

AP ! 400MG/200ML (2MG/ML) A079104 002 Jul 30, 2009

PRESCRIPTION DRUG PRODUCT LIST

FLUCONAZOLE

INJECTABLE; INJECTION

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | |
|-----------|----------------------------------------------------------|-----------------------------|--------------------|--------------|
| <u>AP</u> | RENAISSANCE SSA LLC | <u>200MG/100ML (2MG/ML)</u> | <u>A077909 001</u> | May 26, 2010 |
| <u>AP</u> | | <u>400MG/200ML (2MG/ML)</u> | <u>A077909 002</u> | May 26, 2010 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>200MG/100ML (2MG/ML)</u> | <u>A078107 001</u> | Jul 30, 2008 |
| <u>AP</u> | | <u>400MG/200ML (2MG/ML)</u> | <u>A078107 002</u> | Jul 30, 2008 |
| | FLUCONAZOLE IN SODIUM CHLORIDE 0.9% | | | |
| | WEST-WARD PHARMS INT | 100MG/50ML (2MG/ML) | A076087 002 | Sep 26, 2008 |
| | FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | |
| | RENAISSANCE SSA LLC | 100MG/50ML (2MG/ML) | A077909 003 | Apr 20, 2015 |

TABLET; ORAL

DIFLUCAN

| | | | | | |
|-----------|---|--------|--------------|--------------------|--------------|
| <u>AB</u> | + | PFIZER | <u>50MG</u> | <u>N019949 001</u> | Jan 29, 1990 |
| <u>AB</u> | + | | <u>100MG</u> | <u>N019949 002</u> | Jan 29, 1990 |
| <u>AB</u> | + | | <u>150MG</u> | <u>N019949 004</u> | Jun 30, 1994 |
| <u>AB</u> | + | ! | <u>200MG</u> | <u>N019949 003</u> | Jan 29, 1990 |

FLUCONAZOLE

| | | | | | |
|-----------|--|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | | APOTEX | <u>50MG</u> | <u>A076665 001</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>100MG</u> | <u>A076665 002</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>150MG</u> | <u>A076665 003</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>200MG</u> | <u>A076665 004</u> | Jul 29, 2004 |
| <u>AB</u> | | AUROBINDO PHARMA | <u>50MG</u> | <u>A077731 001</u> | Oct 07, 2008 |
| <u>AB</u> | | | <u>100MG</u> | <u>A077731 002</u> | Oct 07, 2008 |
| <u>AB</u> | | | <u>150MG</u> | <u>A077731 003</u> | Oct 07, 2008 |
| <u>AB</u> | | | <u>200MG</u> | <u>A077731 004</u> | Oct 07, 2008 |
| <u>AB</u> | | DR REDDYS LABS INC | <u>50MG</u> | <u>A076658 001</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>100MG</u> | <u>A076658 002</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>150MG</u> | <u>A076658 003</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>200MG</u> | <u>A076658 004</u> | Jul 29, 2004 |
| <u>AB</u> | | GLENMARK GENERICS | <u>50MG</u> | <u>A077253 001</u> | Jan 25, 2006 |
| <u>AB</u> | | | <u>100MG</u> | <u>A077253 002</u> | Jan 25, 2006 |
| <u>AB</u> | | | <u>150MG</u> | <u>A077253 003</u> | Jan 25, 2006 |
| <u>AB</u> | | | <u>200MG</u> | <u>A077253 004</u> | Jan 25, 2006 |
| <u>AB</u> | | HARRIS PHARM | <u>50MG</u> | <u>A078423 001</u> | Mar 07, 2011 |
| <u>AB</u> | | | <u>100MG</u> | <u>A078423 002</u> | Mar 07, 2011 |
| <u>AB</u> | | | <u>150MG</u> | <u>A078423 003</u> | Mar 07, 2011 |
| <u>AB</u> | | | <u>200MG</u> | <u>A078423 004</u> | Mar 07, 2011 |
| <u>AB</u> | | IVAX SUB TEVA PHARMS | <u>50MG</u> | <u>A076077 001</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>100MG</u> | <u>A076077 002</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>150MG</u> | <u>A076077 003</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>200MG</u> | <u>A076077 004</u> | Jul 29, 2004 |
| <u>AB</u> | | MYLAN | <u>50MG</u> | <u>A076351 001</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>100MG</u> | <u>A076351 002</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>150MG</u> | <u>A076351 003</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>200MG</u> | <u>A076351 004</u> | Jul 29, 2004 |
| <u>AB</u> | | TARO | <u>50MG</u> | <u>A076507 001</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>100MG</u> | <u>A076507 002</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>150MG</u> | <u>A076507 003</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>200MG</u> | <u>A076507 004</u> | Jul 29, 2004 |
| <u>AB</u> | | TEVA | <u>50MG</u> | <u>A074681 001</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>100MG</u> | <u>A074681 002</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>150MG</u> | <u>A074681 003</u> | Jul 29, 2004 |
| <u>AB</u> | | | <u>200MG</u> | <u>A074681 004</u> | Jul 29, 2004 |
| <u>AB</u> | | UNIQUE PHARM LABS | <u>50MG</u> | <u>A076957 001</u> | Sep 28, 2005 |
| <u>AB</u> | | | <u>100MG</u> | <u>A076957 002</u> | Sep 28, 2005 |
| <u>AB</u> | | | <u>150MG</u> | <u>A076957 004</u> | Feb 27, 2017 |
| <u>AB</u> | | | <u>200MG</u> | <u>A076957 003</u> | Sep 28, 2005 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>50MG</u> | <u>A208963 001</u> | Feb 16, 2017 |
| <u>AB</u> | | | <u>100MG</u> | <u>A208963 002</u> | Feb 16, 2017 |
| <u>AB</u> | | | <u>150MG</u> | <u>A208963 003</u> | Feb 16, 2017 |
| <u>AB</u> | | | <u>200MG</u> | <u>A208963 004</u> | Feb 16, 2017 |

FLUCYTOSINE

CAPSULE; ORAL

ANCOBON

| | | | | | |
|-----------|---|---------|--------------|--------------------|--|
| <u>AB</u> | + | VALEANT | <u>250MG</u> | <u>N017001 001</u> | |
| <u>AB</u> | + | ! | <u>500MG</u> | <u>N017001 002</u> | |

PRESCRIPTION DRUG PRODUCT LIST

FLUCYTOSINE

CAPSULE; ORAL

FLUCYTOSINE

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| AB | NOVEL LABS INC | 250MG | A204652 001 | Jul 07, 2017 |
| AB | | 500MG | A204652 002 | Jul 07, 2017 |
| AB | RECIPHARM | 250MG | A207536 001 | Jun 18, 2018 |
| AB | | 500MG | A207536 002 | Jun 18, 2018 |
| AB | SIGMAPHARM LABS LLC | 250MG | A201566 001 | Jun 28, 2011 |
| AB | | 500MG | A201566 002 | Jun 28, 2011 |
| AB | WEST-WARD PHARMS INT | 250MG | A206550 001 | Oct 17, 2017 |
| AB | | 500MG | A206550 002 | Oct 17, 2017 |

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARABINE PHOSPHATE

| | | | | |
|-----------|----------------------|---------------------------|--------------------|--------------|
| AP | ACTAVIS LLC | 50MG/2ML (25MG/ML) | A203738 001 | Feb 28, 2017 |
| AP | ACTAVIS TOTOWA | 50MG/VIAL | A078610 001 | Feb 11, 2009 |
| AP | AREVA PHARMS | 50MG/2ML (25MG/ML) | A090724 001 | Sep 27, 2010 |
| AP | CUSTOPHARM INC | 50MG/VIAL | A076349 001 | Aug 28, 2003 |
| AP | ! FRESENIUS KABI USA | 50MG/2ML (25MG/ML) | A078393 001 | Oct 15, 2007 |
| AP | | 50MG/VIAL | A078544 001 | Oct 15, 2007 |
| AP | ! HOSPIRA | 50MG/VIAL | A077790 001 | Apr 06, 2007 |
| AP | MYLAN LABS LTD | 50MG/2ML (25MG/ML) | A200647 001 | Dec 21, 2011 |
| AP | | 50MG/VIAL | A200648 001 | Oct 16, 2012 |
| AP | SAGENT PHARMS | 50MG/2ML (25MG/ML) | A076661 001 | Apr 28, 2004 |
| AP | +! SANDOZ | 50MG/2ML (25MG/ML) | N022137 001 | Sep 21, 2007 |

FLUDEOXYGLUCOSE F-18

INJECTABLE; INTRAVENOUS

FLUDEOXYGLUCOSE F18

| | | | | |
|-----------|-----------------------|---------------------|--------------------|--------------|
| AP | 3D IMAGING DRUG | 20-300mCi/ML | A203778 001 | Oct 30, 2015 |
| AP | BIOMEDCL RES FDN | 20-300mCi/ML | A203710 001 | May 01, 2015 |
| AP | | 20-300mCi/ML | A203837 001 | May 01, 2015 |
| AP | BRIGHAM WOMENS | 20-300mCi/ML | A203816 001 | Oct 30, 2014 |
| AP | CARDINAL HEALTH 414 | 20-300mCi/ML | A203603 001 | Nov 13, 2015 |
| AP | CHILDRENS HOSP MI | 20-300mCi/ML | A204385 001 | Oct 29, 2014 |
| AP | CPDC | 20-300mCi/ML | A204525 001 | Oct 29, 2014 |
| AP | ESSENTIAL ISOTOPES | 20-300mCi/ML | A203946 001 | Feb 05, 2014 |
| AP | +! FEINSTEIN | 20-400mCi/ML | N021870 002 | Nov 21, 2008 |
| AP | GLOBAL ISOTOPES LLC | 20-300mCi/ML | A204463 001 | Oct 21, 2014 |
| AP | ! HOUSTON CYCLOTRON | 20-500mCi/ML | A203665 001 | Feb 14, 2013 |
| AP | JUBILANT DRAXIMAGE | 20-300mCi/ML | A203920 001 | Jun 23, 2015 |
| AP | KETTERING MEDCTR | 4-40mCi/ML | A204759 001 | Oct 27, 2015 |
| AP | KREITCHMAN PET CTR | 10-100mCi/ML | A203942 001 | Apr 11, 2016 |
| AP | LANTHEUS MEDICAL | 20-200mCi/ML | A203664 001 | Feb 04, 2014 |
| AP | MA GENERAL HOSP | 20-300mCi/ML | A204333 001 | Sep 25, 2014 |
| AP | MCPRF | 20-240mCi/ML | A203612 001 | Aug 05, 2013 |
| AP | MEM SLOAN-KETTERING | 20-300mCi/ML | A208679 001 | Dec 08, 2016 |
| AP | METHODIST HOSP RES | 20-300mCi/ML | A203904 001 | Apr 23, 2015 |
| AP | MIPS CRF | 20-300mCi/ML | A204472 001 | Sep 11, 2015 |
| AP | NCM USA BRONX LLC | 20-300mCi/ML | A204512 001 | Jan 07, 2015 |
| AP | ! PETNET | 20-200mCi/ML | A079086 001 | Feb 25, 2011 |
| AP | ! QUEEN HAMAMATSU PET | 10-100mCi/ML | A203771 001 | Aug 31, 2015 |
| AP | SHERTECH LABS LLC | 20-300mCi/ML | A204264 001 | Dec 18, 2014 |
| AP | SOFIE | 20-300mCi/ML | A203591 001 | Aug 31, 2015 |
| AP | TRUSTEES UNIV PA | 20-200mCi/ML | A203801 001 | Oct 29, 2014 |
| AP | ! UCLA BIOMEDICAL | 4-40mCi/ML | A203811 001 | Jun 27, 2013 |
| AP | UCSF RODIOPHARM | 20-300mCi/ML | A203902 001 | May 09, 2014 |
| AP | UIHC PET IMAGING | 20-300mCi/ML | A203990 001 | Aug 06, 2014 |
| AP | UNIV MICHIGAN | 20-300mCi/ML | A204531 001 | Jul 17, 2015 |
| AP | UNIV TX MD ANDERSON | 20-300mCi/ML | A203246 002 | Jan 13, 2014 |
| AP | UNIV UTAH CYCLOTRON | 20-300mCi/ML | A204498 001 | Jun 23, 2015 |
| AP | WI MEDCL CYCLOTRON | 20-500mCi/ML | A203709 001 | Oct 23, 2013 |
| AP | WUSM CYCLOTRON | 20-300mCi/ML | A203935 001 | Feb 05, 2014 |
| | HOT SHOTS NM LLC | 4-500mCi/ML | A203937 001 | Oct 30, 2014 |
| | NORTHLAND | 4-500mCi/ML | A203994 001 | Feb 04, 2015 |
| | PRECISION NUCLEAR | 20-500mCi/ML | A204546 001 | Apr 07, 2015 |
| | SPECTRON MRC LLC | 4-500mCi/ML | A203911 001 | Apr 22, 2015 |
| | UNIV TX MD ANDERSON | 20-150mCi/ML | A203246 001 | Jan 13, 2014 |

PRESCRIPTION DRUG PRODUCT LISTFLUDROCORTISONE ACETATE

TABLET; ORAL

FLUDROCORTISONE ACETATE

| | | | | | |
|-----------|--------------|--------------|----------------|------------|--------------|
| AB | BARR | <u>0.1MG</u> | <u>A040425</u> | <u>001</u> | Jan 21, 2003 |
| AB | HIKMA PHARMS | <u>0.1MG</u> | <u>A091302</u> | <u>001</u> | Jul 22, 2011 |
| AB | ! IMPAX LABS | <u>0.1MG</u> | <u>A040431</u> | <u>001</u> | Mar 18, 2002 |

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

| | | | | | |
|-----------|--------------------|-----------------------------|----------------|------------|--------------|
| AP | FRESENIUS KABI USA | <u>0.5MG/5ML (0.1MG/ML)</u> | <u>A076955</u> | <u>002</u> | Oct 12, 2004 |
| AP | | <u>1MG/10ML (0.1MG/ML)</u> | <u>A076955</u> | <u>001</u> | Oct 12, 2004 |
| AP | HIKMA FARMACEUTICA | <u>0.5MG/5ML (0.1MG/ML)</u> | <u>A078527</u> | <u>001</u> | Mar 23, 2009 |
| AP | | <u>1MG/10ML (0.1MG/ML)</u> | <u>A078527</u> | <u>002</u> | Mar 23, 2009 |
| AP | MYLAN LABS LTD | <u>0.5MG/5ML (0.1MG/ML)</u> | <u>A078595</u> | <u>001</u> | May 13, 2008 |
| AP | ! | <u>1MG/10ML (0.1MG/ML)</u> | <u>A078595</u> | <u>002</u> | May 13, 2008 |
| AP | SAGENT PHARMS | <u>0.5MG/5ML (0.1MG/ML)</u> | <u>A090584</u> | <u>001</u> | Aug 28, 2012 |
| AP | | <u>1MG/10ML (0.1MG/ML)</u> | <u>A090584</u> | <u>002</u> | Aug 28, 2012 |
| AP | SANDOZ INC | <u>0.5MG/5ML (0.1MG/ML)</u> | <u>A077071</u> | <u>001</u> | May 03, 2005 |
| AP | | <u>1MG/10ML (0.1MG/ML)</u> | <u>A077071</u> | <u>002</u> | May 03, 2005 |
| AP | WEST-WARD PHARMS | <u>0.5MG/5ML (0.1MG/ML)</u> | <u>A076256</u> | <u>002</u> | Oct 12, 2004 |
| AP | INT | | | | |
| AP | | <u>0.5MG/5ML (0.1MG/ML)</u> | <u>A076787</u> | <u>002</u> | Oct 12, 2004 |
| AP | | <u>1MG/10ML (0.1MG/ML)</u> | <u>A076256</u> | <u>001</u> | Oct 12, 2004 |
| AP | | <u>1MG/10ML (0.1MG/ML)</u> | <u>A076787</u> | <u>001</u> | Oct 12, 2004 |

FLUNISOLIDE

AEROSOL, METERED; INHALATION

AEROSPAN HFA

| | | | | | | |
|---|---|---------------------|-------------|---------|-----|--------------|
| + | ! | MYLAN SPECIALITY LP | 0.078MG/INH | N021247 | 001 | Jan 27, 2006 |
|---|---|---------------------|-------------|---------|-----|--------------|

SPRAY, METERED; NASAL

FLUNISOLIDE

| | | | | | | |
|-----------|---|-------------------|----------------------|----------------|------------|--------------|
| AB | ! | BAUSCH AND LOMB | <u>0.025MG/SPRAY</u> | <u>A074805</u> | <u>001</u> | Feb 20, 2002 |
| AB | | HI TECH PHARMA CO | <u>0.025MG/SPRAY</u> | <u>A077704</u> | <u>001</u> | Aug 03, 2006 |

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL

FLUOCINOLONE ACETONIDE

| | | | | | | |
|----------------|--------------------|---------------|--------------------|---------------|----------------|------------|
| AT | FOUGERA PHARMS INC | <u>0.01%</u> | <u>A088170</u> | <u>001</u> | Dec 16, 1982 | |
| AT | | <u>0.025%</u> | <u>A088169</u> | <u>001</u> | Dec 16, 1982 | |
| AT | G AND W LABS | <u>0.01%</u> | <u>A089526</u> | <u>001</u> | Jul 26, 1988 | |
| AT | G AND W LABS INC | <u>0.025%</u> | <u>A210747</u> | <u>001</u> | Nov 05, 2018 | |
| AT | TARO | <u>0.025%</u> | <u>A087104</u> | <u>001</u> | Apr 27, 1982 | |
| SYNALAR | | | | | | |
| AT | + | ! | MEDIMETRIKS PHARMS | <u>0.01%</u> | <u>N012787</u> | <u>004</u> |
| AT | + | ! | | <u>0.025%</u> | <u>N012787</u> | <u>002</u> |
| AT | + | ! | | <u>0.025%</u> | <u>N012787</u> | <u>005</u> |

IMPLANT; INTRAVITREAL

ILUVIEN

| | | | | | | |
|---|---|------------------|--------|---------|-----|--------------|
| + | ! | ALIMERA SCIENCES | 0.19MG | N201923 | 001 | Sep 26, 2014 |
|---|---|------------------|--------|---------|-----|--------------|

INC

RETISERT

| | | | | | | |
|---|---|-----------------|--------|---------|-----|--------------|
| + | ! | BAUSCH AND LOMB | 0.59MG | N021737 | 001 | Apr 08, 2005 |
|---|---|-----------------|--------|---------|-----|--------------|

YUTIQ

| | | | | | | |
|---|---|-----------------|--------|---------|-----|--------------|
| + | ! | EYEPOINT PHARMS | 0.18MG | N210331 | 001 | Oct 12, 2018 |
|---|---|-----------------|--------|---------|-----|--------------|

OIL; TOPICAL

DERMA-SMOOTH/FS

| | | | | | | | |
|-----------|---|---|-------------|--------------|----------------|------------|--------------|
| AT | + | ! | HILL DERMAC | <u>0.01%</u> | <u>N019452</u> | <u>001</u> | Feb 03, 1988 |
| AT | + | ! | | <u>0.01%</u> | <u>N019452</u> | <u>002</u> | Nov 09, 2005 |

FLUCINOLONE ACETONIDE

| | | | | | |
|-----------|---------------------|--------------|----------------|------------|--------------|
| AT | GLENMARK PHARMS LTD | <u>0.01%</u> | <u>A210556</u> | <u>001</u> | Oct 25, 2018 |
|-----------|---------------------|--------------|----------------|------------|--------------|

FLUOCINOLONE ACETONIDE

| | | | | | |
|-------------------------------|---------------------|--------------|----------------|------------|--------------|
| AT | AKORN | <u>0.01%</u> | <u>A091514</u> | <u>001</u> | Jun 25, 2015 |
| AT | IDENTI PHARMS INC | <u>0.01%</u> | <u>A201759</u> | <u>001</u> | Oct 17, 2011 |
| AT | | <u>0.01%</u> | <u>A201764</u> | <u>001</u> | Oct 17, 2011 |
| AT | LYNE | <u>0.01%</u> | <u>A090982</u> | <u>001</u> | Apr 25, 2016 |
| AT | | <u>0.01%</u> | <u>A203377</u> | <u>001</u> | Apr 25, 2016 |
| AT | PERRIGO ISRAEL | <u>0.01%</u> | <u>A202847</u> | <u>001</u> | Aug 09, 2013 |
| AT | | <u>0.01%</u> | <u>A202848</u> | <u>001</u> | Aug 09, 2013 |
| AT | TARO | <u>0.01%</u> | <u>A202368</u> | <u>001</u> | May 19, 2016 |
| AT | | <u>0.01%</u> | <u>A209336</u> | <u>001</u> | May 19, 2016 |
| FLUOCINONIDE ACETONIDE | | | | | |
| AT | GLENMARK PHARMS LTD | <u>0.01%</u> | <u>A210539</u> | <u>001</u> | Oct 26, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

FLUOCINOLONE ACETONIDE

OIL/DROPS;OTIC

DERMOTIC

AT +! HILL DERMAC **0.01%** **N019452 003** Nov 09, 2005

FLAC

AT PATRIN PHARMA INC **0.01%** **A210736 001** Apr 11, 2018

FLUOCINOLONE ACETONIDE

AT AKORN **0.01%** **A202705 001** Sep 09, 2016

AT IDENTI PHARMS INC **0.01%** **A091306 001** Oct 17, 2011

AT LYNE **0.01%** **A203378 001** Apr 25, 2016

AT PERRIGO ISRAEL **0.01%** **A202849 001** Jul 17, 2017

FLUOCINONIDE ACETONIDE

AT GLENMARK PHARMS LTD **0.01%** **A211815 001** Dec 14, 2018

OINTMENT;TOPICAL

FLUOCINOLONE ACETONIDE

AT FOUGERA PHARMS INC **0.025%** **A088168 001** Dec 16, 1982

AT G AND W LABS **0.025%** **A089524 001** Jul 26, 1988

AT TARO **0.025%** **A040041 001** Sep 15, 1994

SYNALAR

AT +! MEDIMETRIKS PHARMS **0.025%** **N013960 001**

SHAMPOO;TOPICAL

CAPEX

+! GALDERMA LABS LP 0.01%

N020001 001 Aug 27, 1990

SOLUTION;TOPICAL

FLUOCINOLONE ACETONIDE

AT ACTAVIS LABS UT INC **0.01%** **A208386 001** Oct 21, 2016

AT FOUGERA PHARMS INC **0.01%** **A088167 001** Dec 16, 1982

AT GAVIS PHARMS LLC **0.01%** **A206422 001** Sep 02, 2015

AT TARO **0.01%** **A089124 001** Sep 11, 1985

SYNALAR

AT +! MEDIMETRIKS PHARMS **0.01%** **N015296 001**

FLUOCINOLONE ACETONIDE; HYDROQUINONE; TRETINOIN

CREAM;TOPICAL

TRI-LUMA

+! GALDERMA LABS LP 0.01%;4%;0.05%

N021112 001 Jan 18, 2002

FLUOCINOLONE ACETONIDE; NEOMYCIN SULFATE

CREAM;TOPICAL

NEO-SYNALAR

! MEDIMETRIKS PHARMS 0.025%;EQ 3.5MG BASE/GM

A060700 001

FLUOCINONIDE

CREAM;TOPICAL

FLUOCINONIDE

AB AMNEAL PHARMS LLC **0.1%** **A211111 001** Jun 04, 2018

AB FOUGERA PHARMS INC **0.1%** **A200735 001** Jul 14, 2014

AB GLENMARK GENERICS **0.1%** **A091282 001** Jul 14, 2014

AB PERRIGO ISRAEL **0.1%** **A090256 001** Jan 14, 2014

AB TARO **0.1%** **A200734 001** Jul 14, 2014

VANOS

AB +! MEDICIS **0.1%** **N021758 001** Feb 11, 2005

FLUOCINONIDE

AB1 AMNEAL PHARMS LLC **0.05%** **A210554 001** Aug 21, 2018

AB1 FOUGERA PHARMS INC **0.05%** **A073030 001** Oct 17, 1994

AB1 G AND W LABS INC **0.05%** **A073085 001** Feb 14, 1992

AB1 TARO **0.05%** **A071500 001** Jun 10, 1987

AB1 +! **0.05%** **N019117 001** Jun 26, 1984

AB1 TELIGENT PHARMA INC **0.05%** **A211410 001** Oct 16, 2018

AB1 TEVA **0.05%** **A072488 001** Feb 06, 1989

FLUOCINONIDE EMULSIFIED BASE

AB2 FOUGERA PHARMS **0.05%** **A076586 001** Jun 23, 2004

AB2 G AND W LABS INC **0.05%** **A074204 001** Jun 13, 1995

AB2 ! TARO PHARM INDS LTD **0.05%** **A072494 001** Jan 19, 1989

AB2 TEVA **0.05%** **A072490 001** Feb 07, 1989

LIDEX-E

AB2 + CNTY LINE PHARMS **0.05%** **N016908 003**

GEL;TOPICAL

FLUOCINONIDE

AB + CNTY LINE PHARMS **0.05%** **N017373 001**

AB FOUGERA PHARMS INC **0.05%** **A072933 001** Dec 30, 1994

AB G AND W LABS INC **0.05%** **A072537 001** Feb 07, 1989

AB ! TARO **0.05%** **A074935 001** Jul 29, 1997

AB TELIGENT PHARMA INC **0.05%** **A209030 001** Jun 19, 2018

PRESCRIPTION DRUG PRODUCT LIST

FLUOCINONIDE

OINTMENT; TOPICAL

FLUOCINONIDE

| | | | | |
|-----------|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | FOUGERA PHARMS | <u>0.05%</u> | <u>A074905 001</u> | Aug 26, 1997 |
| <u>AB</u> | NOVEL LABS INC | <u>0.05%</u> | <u>A207538 001</u> | Jul 31, 2017 |
| <u>AB</u> | ! TARO | <u>0.05%</u> | <u>A075008 001</u> | Jun 30, 1999 |
| <u>AB</u> | TELIGENT PHARMA INC | <u>0.05%</u> | <u>A207680 001</u> | Sep 28, 2018 |
| <u>AB</u> | TEVA | <u>0.05%</u> | <u>A073481 001</u> | Dec 27, 1991 |

LIDEX

| | | | | |
|-----------|--------------------|--------------|--------------------|--|
| <u>AB</u> | + CNTY LINE PHARMS | <u>0.05%</u> | <u>N016909 002</u> | |
|-----------|--------------------|--------------|--------------------|--|

SOLUTION; TOPICAL

FLUOCINONIDE

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| <u>AT</u> | ENCUBE ETHICALS | <u>0.05%</u> | <u>A209699 001</u> | Nov 29, 2018 |
| <u>AT</u> | FOUGERA PHARMS INC | <u>0.05%</u> | <u>A072934 001</u> | Feb 27, 1995 |
| <u>AT</u> | G AND W LABS INC | <u>0.05%</u> | <u>A071535 001</u> | Dec 02, 1988 |
| <u>AT</u> | GLASSHOUSE PHARMS | <u>0.05%</u> | <u>A209118 001</u> | Apr 23, 2018 |
| <u>AT</u> | MACLEODS PHARMS LTD | <u>0.05%</u> | <u>A209283 001</u> | Apr 23, 2018 |
| <u>AT</u> | NOVEL LABS INC | <u>0.05%</u> | <u>A206003 001</u> | Jul 21, 2017 |
| <u>AT</u> | ! TARO | <u>0.05%</u> | <u>A074799 001</u> | Dec 31, 1996 |
| <u>AT</u> | TEVA | <u>0.05%</u> | <u>A072511 001</u> | Feb 07, 1989 |
| <u>AT</u> | ZYDUS PHARMS USA INC | <u>0.05%</u> | <u>A208948 001</u> | Jul 17, 2018 |

LIDEX

| | | | | |
|-----------|--------------------|--------------|--------------------|--------------|
| <u>AT</u> | + CNTY LINE PHARMS | <u>0.05%</u> | <u>N018849 001</u> | Apr 06, 1984 |
|-----------|--------------------|--------------|--------------------|--------------|

FLUORESCHEIN SODIUM

INJECTABLE; INTRAVENOUS

AK-FLUOR 10%

| | | | | |
|-----------|---------|---------------------------------------------|--------------------|--------------|
| <u>AP</u> | + AKORN | <u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u> | <u>N022186 001</u> | Aug 08, 2008 |
|-----------|---------|---------------------------------------------|--------------------|--------------|

FLUORESCITE

| | | | | |
|-----------|-------------------|---------------------------------------------|--------------------|--------------|
| <u>AP</u> | +! ALCON LABS INC | <u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u> | <u>N021980 001</u> | Mar 28, 2006 |
| | AK-FLUOR 25% | | | |
| | +! AKORN | EQ 500MG BASE/2ML (EQ 250MG BASE/ML) | N022186 002 | Aug 08, 2008 |

FLUOROMETHOLONE

OINTMENT; OPHTHALMIC

FML

| | | | | |
|--|-------------|------|-------------|--------------|
| | +! ALLERGAN | 0.1% | N017760 001 | Sep 04, 1985 |
|--|-------------|------|-------------|--------------|

SUSPENSION/DROPS; OPHTHALMIC

FML

| | | | | |
|--|-------------|------|-------------|--------------|
| | +! ALLERGAN | 0.1% | N016851 002 | Jul 28, 1982 |
|--|-------------|------|-------------|--------------|

FML FORTE

| | | | | |
|--|----------|-------|-------------|--------------|
| | ALLERGAN | 0.25% | N019216 001 | Apr 23, 1986 |
|--|----------|-------|-------------|--------------|

FLUOROMETHOLONE ACETATE

SUSPENSION/DROPS; OPHTHALMIC

FLAREX

| | | | | |
|--|----------------------------|------|-------------|--------------|
| | +! NOVARTIS PHARMS CORP | 0.1% | N019079 001 | Feb 11, 1986 |
|--|----------------------------|------|-------------|--------------|

FLUOROURACIL

CREAM; TOPICAL

CARAC

| | | | | |
|-----------|----------------------------|-------------|--------------------|--------------|
| <u>AB</u> | +! VALEANT PHARMS NORTH | <u>0.5%</u> | <u>N020985 001</u> | Oct 27, 2000 |
|-----------|----------------------------|-------------|--------------------|--------------|

EFUDEX

| | | | | |
|-----------|-----------------------|-----------|--------------------|--|
| <u>AB</u> | +! VALEANT PHARM INTL | <u>5%</u> | <u>N016831 003</u> | |
|-----------|-----------------------|-----------|--------------------|--|

FLUOROURACIL

| | | | | |
|-----------|------------------|-------------|--------------------|--------------|
| <u>AB</u> | MAYNE PHARMA | <u>5%</u> | <u>A077524 001</u> | Apr 11, 2008 |
| <u>AB</u> | MYLAN PHARMS INC | <u>0.5%</u> | <u>A203122 001</u> | Apr 20, 2015 |
| <u>AB</u> | TARO | <u>5%</u> | <u>A090368 001</u> | Mar 05, 2010 |

FLUOROPLEX

| | | | | |
|--|----------------|----|-------------|--|
| | +! AQUA PHARMS | 1% | N016988 001 | |
|--|----------------|----|-------------|--|

TOLAK

| | | | | |
|--|------------------------|----|-------------|--------------|
| | +! HILL DERMACEUTICALS | 4% | N022259 001 | Sep 18, 2015 |
|--|------------------------|----|-------------|--------------|

INJECTABLE; INJECTION

FLUOROURACIL

| | | | | |
|-----------|----------------------|-----------------------------|--------------------|--------------|
| <u>AP</u> | ! ACCORD HLTHCARE | <u>500MG/10ML (50MG/ML)</u> | <u>A040743 002</u> | Apr 26, 2007 |
| <u>AP</u> | ! | <u>1GM/20ML (50MG/ML)</u> | <u>A040743 001</u> | Apr 26, 2007 |
| <u>AP</u> | ! | <u>2.5GM/50ML (50MG/ML)</u> | <u>A040798 002</u> | Apr 26, 2007 |
| <u>AP</u> | ! | <u>5GM/100ML (50MG/ML)</u> | <u>A040798 001</u> | Apr 26, 2007 |
| <u>AP</u> | ! FRESENIUS KABI USA | <u>500MG/10ML (50MG/ML)</u> | <u>A040279 002</u> | Sep 30, 1998 |
| <u>AP</u> | ! | <u>1GM/20ML (50MG/ML)</u> | <u>A040279 001</u> | Sep 30, 1998 |
| <u>AP</u> | ! | <u>2.5GM/50ML (50MG/ML)</u> | <u>A040278 001</u> | Sep 30, 1998 |
| <u>AP</u> | ! | <u>5GM/100ML (50MG/ML)</u> | <u>A040278 002</u> | Sep 30, 1998 |

PRESCRIPTION DRUG PRODUCT LIST

FLUOROURACIL

INJECTABLE; INJECTION

FLUOROURACIL

| | | | | |
|-----------|-------------------|-----------------------------|--------------------|--------------|
| <u>AP</u> | GLAND PHARMA LTD | <u>500MG/10ML (50MG/ML)</u> | <u>A210123 001</u> | Oct 27, 2017 |
| <u>AP</u> | | <u>1GM/20ML (50MG/ML)</u> | <u>A210123 002</u> | Oct 27, 2017 |
| <u>AP</u> | | <u>2.5GM/50ML (50MG/ML)</u> | <u>A210124 001</u> | Dec 26, 2017 |
| <u>AP</u> | | <u>5GM/100ML (50MG/ML)</u> | <u>A210124 002</u> | Dec 26, 2017 |
| <u>AP</u> | MYLAN LABS LTD | <u>500MG/10ML (50MG/ML)</u> | <u>A202668 001</u> | Jul 17, 2012 |
| <u>AP</u> | | <u>1GM/20ML (50MG/ML)</u> | <u>A202668 002</u> | Jul 17, 2012 |
| <u>AP</u> | | <u>2.5GM/50ML (50MG/ML)</u> | <u>A202669 001</u> | Jul 17, 2012 |
| <u>AP</u> | | <u>5GM/100ML (50MG/ML)</u> | <u>A202669 002</u> | Jul 17, 2012 |
| <u>AP</u> | SAGENT PHARMS | <u>500MG/10ML (50MG/ML)</u> | <u>A203608 001</u> | May 11, 2017 |
| <u>AP</u> | | <u>1GM/20ML (50MG/ML)</u> | <u>A203608 002</u> | May 11, 2017 |
| <u>AP</u> | | <u>2.5GM/50ML (50MG/ML)</u> | <u>A203609 001</u> | Feb 17, 2016 |
| <u>AP</u> | | <u>5GM/100ML (50MG/ML)</u> | <u>A203609 002</u> | Feb 17, 2016 |
| <u>AP</u> | ! TEVA PHARMS USA | <u>500MG/10ML (50MG/ML)</u> | <u>A040333 001</u> | Jan 27, 2000 |
| <u>AP</u> | ! | <u>2.5GM/50ML (50MG/ML)</u> | <u>A040334 001</u> | Feb 25, 2000 |
| <u>AP</u> | ! | <u>5GM/100ML (50MG/ML)</u> | <u>A040334 002</u> | Feb 25, 2000 |

SOLUTION; TOPICAL

EFUDEX

| | | | | |
|-----------|-----------------------|-----------|--------------------|--|
| <u>AT</u> | +! VALEANT PHARM INTL | <u>2%</u> | <u>N016831 001</u> | |
| <u>AT</u> | +! | <u>5%</u> | <u>N016831 002</u> | |

FLUOROURACIL

| | | | | |
|-----------|------------|-----------|--------------------|--------------|
| <u>AT</u> | TARO PHARM | <u>2%</u> | <u>A076526 001</u> | Nov 05, 2003 |
| <u>AT</u> | | <u>5%</u> | <u>A076526 002</u> | Nov 05, 2003 |

FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL

FLUOXETINE HYDROCHLORIDE

| | | | | |
|-----------|----------------------|---------------------|--------------------|--------------|
| <u>AB</u> | ALEMBIC PHARMS LTD | <u>EQ 40MG BASE</u> | <u>A090223 003</u> | Mar 19, 2009 |
| <u>AB</u> | AUROBINDO PHARMA | <u>EQ 40MG BASE</u> | <u>A078619 003</u> | Jan 31, 2008 |
| <u>AB</u> | HERITAGE PHARMS INC | <u>EQ 40MG BASE</u> | <u>A201336 003</u> | Oct 01, 2012 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>EQ 40MG BASE</u> | <u>A075245 003</u> | Sep 28, 2004 |
| <u>AB</u> | MARKSANS PHARMA | <u>EQ 40MG BASE</u> | <u>A075465 003</u> | Aug 02, 2001 |
| <u>AB</u> | PAR PHARM | <u>EQ 40MG BASE</u> | <u>A076922 003</u> | Dec 16, 2004 |
| <u>AB</u> | SANDOZ | <u>EQ 40MG BASE</u> | <u>A075049 003</u> | Jan 29, 2002 |
| <u>AB</u> | SCIEGEN PHARMS INC | <u>EQ 40MG BASE</u> | <u>A204597 003</u> | Mar 16, 2015 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>EQ 40MG BASE</u> | <u>A076990 001</u> | Dec 13, 2004 |
| <u>AB</u> | TEVA | <u>EQ 40MG BASE</u> | <u>A075452 003</u> | Jan 29, 2002 |

PROZAC

| | | | | |
|-----------|---------------------|---------------------|--------------------|--------------|
| <u>AB</u> | +! ELI LILLY AND CO | <u>EQ 40MG BASE</u> | <u>N018936 003</u> | Jun 15, 1999 |
|-----------|---------------------|---------------------|--------------------|--------------|

FLUOXETINE HYDROCHLORIDE

| | | | | |
|------------|----------------------|---------------------|--------------------|--------------|
| <u>AB1</u> | ALEMBIC PHARMS LTD | <u>EQ 10MG BASE</u> | <u>A090223 001</u> | Mar 19, 2009 |
| <u>AB1</u> | | <u>EQ 20MG BASE</u> | <u>A090223 002</u> | Mar 19, 2009 |
| <u>AB1</u> | AUROBINDO PHARMA | <u>EQ 10MG BASE</u> | <u>A078619 001</u> | Jan 31, 2008 |
| <u>AB1</u> | | <u>EQ 20MG BASE</u> | <u>A078619 002</u> | Jan 31, 2008 |
| <u>AB1</u> | BARR | <u>EQ 10MG BASE</u> | <u>A074803 002</u> | Jan 30, 2002 |
| <u>AB1</u> | | <u>EQ 20MG BASE</u> | <u>A074803 001</u> | Aug 02, 2001 |
| <u>AB1</u> | HERITAGE PHARMS INC | <u>EQ 10MG BASE</u> | <u>A201336 001</u> | Oct 01, 2012 |
| <u>AB1</u> | | <u>EQ 20MG BASE</u> | <u>A201336 002</u> | Oct 01, 2012 |
| <u>AB1</u> | IVAX SUB TEVA PHARMS | <u>EQ 10MG BASE</u> | <u>A075245 002</u> | Jan 31, 2002 |
| <u>AB1</u> | | <u>EQ 20MG BASE</u> | <u>A075245 001</u> | Jan 31, 2002 |
| <u>AB1</u> | LANDELA PHARM | <u>EQ 10MG BASE</u> | <u>A075464 001</u> | Jan 30, 2002 |
| <u>AB1</u> | | <u>EQ 20MG BASE</u> | <u>A075464 002</u> | Jan 30, 2002 |
| <u>AB1</u> | MARKSANS PHARMA | <u>EQ 10MG BASE</u> | <u>A075465 001</u> | Jan 29, 2002 |
| <u>AB1</u> | | <u>EQ 20MG BASE</u> | <u>A075465 002</u> | Jan 29, 2002 |
| <u>AB1</u> | SANDOZ | <u>EQ 10MG BASE</u> | <u>A075049 001</u> | Aug 02, 2001 |
| <u>AB1</u> | | <u>EQ 20MG BASE</u> | <u>A075049 002</u> | Jan 29, 2002 |
| <u>AB1</u> | SCIEGEN PHARMS INC | <u>EQ 10MG BASE</u> | <u>A204597 001</u> | Mar 16, 2015 |
| <u>AB1</u> | | <u>EQ 20MG BASE</u> | <u>A204597 002</u> | Mar 16, 2015 |
| <u>AB1</u> | SPECGX LLC | <u>EQ 10MG BASE</u> | <u>A075658 001</u> | Jan 29, 2002 |
| <u>AB1</u> | | <u>EQ 20MG BASE</u> | <u>A075658 002</u> | Jan 29, 2002 |
| <u>AB1</u> | TEVA | <u>EQ 10MG BASE</u> | <u>A075452 001</u> | Jan 29, 2002 |
| <u>AB1</u> | | <u>EQ 20MG BASE</u> | <u>A075452 002</u> | Jan 29, 2002 |
| <u>AB1</u> | TEVA PHARMS USA | <u>EQ 10MG BASE</u> | <u>A076001 001</u> | Jan 29, 2002 |
| <u>AB1</u> | | <u>EQ 20MG BASE</u> | <u>A076001 002</u> | Jan 29, 2002 |

PROZAC

| | | | | |
|------------|--------------------|---------------------|--------------------|--------------|
| <u>AB1</u> | + ELI LILLY AND CO | <u>EQ 10MG BASE</u> | <u>N018936 006</u> | Dec 23, 1992 |
| <u>AB1</u> | + ELI LILLY AND CO | <u>EQ 20MG BASE</u> | <u>N018936 001</u> | Dec 29, 1987 |

FLUOXETINE HYDROCHLORIDE

| | | | | |
|--|-------|--------------|-------------|--------------|
| | MYLAN | EQ 10MG BASE | A078045 001 | Nov 17, 2008 |
|--|-------|--------------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

FLUOXETINE HYDROCHLORIDE

CAPSULE;ORAL

FLUOXETINE HYDROCHLORIDE

! EQ 20MG BASE

A078045 002 Nov 17, 2008

CAPSULE, DELAYED REL PELLETS;ORAL

FLUOXETINE HYDROCHLORIDE

AB BARR

EQ 90MG BASE

A076237 001 Mar 24, 2010

AB DR REDDYS LABS LTD

EQ 90MG BASE

A078572 001 Mar 22, 2010

PROZAC WEEKLY

AB +! LILLY

EQ 90MG BASE

N021235 001 Feb 26, 2001

SOLUTION;ORAL

FLUOXETINE HYDROCHLORIDE

AA LANNETT CO INC

EQ 20MG BASE/5ML

A077849 001 Feb 09, 2007

AA ! PHARM ASSOC

EQ 20MG BASE/5ML

A076015 001 Jan 30, 2002

AA SPECGX LLC

EQ 20MG BASE/5ML

A075920 001 Jan 29, 2002

AA TEVA

EQ 20MG BASE/5ML

A075506 001 Aug 02, 2001

AA WOCKHARDT BIO AG

EQ 20MG BASE/5ML

A075514 001 Aug 29, 2002

TABLET;ORAL

FLUOXETINE HYDROCHLORIDE

AB ALEMERIC PHARMS LTD

EQ 10MG BASE

A208698 001 Apr 05, 2017

AB

EQ 20MG BASE

A208698 002 Apr 05, 2017

AB +! ALVOGEN

EQ 60MG BASE

N202133 001 Oct 06, 2011

AB APPCO PHARMA LLC

EQ 60MG BASE

A211477 001 Nov 21, 2018

AB DR REDDYS LABS LTD

EQ 10MG BASE

A076006 001 Jan 30, 2002

AB

EQ 20MG BASE

A076006 002 Apr 23, 2018

AB INVENTIA HLTHCARE

EQ 60MG BASE

A209695 001 Nov 20, 2017

AB MYLAN

EQ 10MG BASE

A075755 001 Aug 02, 2001

AB !

EQ 20MG BASE

A075755 002 Aug 02, 2001

AB PAR FORM

EQ 10MG BASE

A203836 001 Aug 19, 2016

AB

EQ 20MG BASE

A203836 002 Aug 19, 2016

AB PAR PHARM INC

EQ 60MG BASE

A209419 001 Nov 16, 2017

AB SCIEGEN PHARMS INC

EQ 60MG BASE

A211282 001 Jan 10, 2019

AB TEVA

EQ 10MG BASE

A075872 001 Jan 29, 2002

AB

EQ 20MG BASE

A075872 002 Jan 04, 2019

AB TEVA PHARMS USA

EQ 60MG BASE

A211051 001 Dec 03, 2018

AB1 TORRENT PHARMS LTD

EQ 10MG BASE

A206937 001 Oct 21, 2016

AB1

EQ 20MG BASE

A206937 002 Oct 21, 2016

SARAFEM

AB1 + APIL

EQ 10MG BASE

N021860 001 May 19, 2006

AB1 +

EQ 15MG BASE

N021860 002 May 19, 2006

AB1 +!

EQ 20MG BASE

N021860 003 May 19, 2006

SELFEMRA

AB1 TEVA PHARMS USA

EQ 10MG BASE

A200151 001 Feb 03, 2014

AB1

EQ 15MG BASE

A200151 002 Feb 03, 2014

AB1

EQ 20MG BASE

A200151 003 Feb 03, 2014

FLUOXETINE HYDROCHLORIDE; OLANZAPINE

CAPSULE;ORAL

OLANZAPINE AND FLUOXETINE HYDROCHLORIDE

AB PAR PHARM

EQ 25MG BASE;EQ 3MG BASE

A077742 001 Nov 02, 2012

AB

EQ 25MG BASE;EQ 6MG BASE

A077742 002 Nov 02, 2012

AB

EQ 25MG BASE;EQ 12MG BASE

A077742 003 Nov 02, 2012

AB

EQ 50MG BASE;EQ 6MG BASE

A077742 004 Nov 02, 2012

AB

EQ 50MG BASE;EQ 12MG BASE

A077742 005 Nov 02, 2012

AB SANDOZ

EQ 25MG BASE;EQ 3MG BASE

A078901 005 Nov 16, 2012

AB

EQ 25MG BASE;EQ 6MG BASE

A078901 001 Nov 16, 2012

AB

EQ 25MG BASE;EQ 12MG BASE

A078901 003 Nov 16, 2012

AB

EQ 50MG BASE;EQ 6MG BASE

A078901 002 Nov 16, 2012

AB

EQ 50MG BASE;EQ 12MG BASE

A078901 004 Nov 16, 2012

AB TEVA PHARMS

EQ 25MG BASE;EQ 3MG BASE

A202074 001 Mar 25, 2013

AB

EQ 25MG BASE;EQ 6MG BASE

A077528 001 Jun 19, 2012

AB

EQ 25MG BASE;EQ 12MG BASE

A077528 002 Jun 19, 2012

AB

EQ 50MG BASE;EQ 6MG BASE

A077528 003 Jun 19, 2012

AB

EQ 50MG BASE;EQ 12MG BASE

A077528 004 Jun 19, 2012

SYMBYAX

AB + LILLY

EQ 25MG BASE;EQ 3MG BASE

N021520 001 Apr 09, 2007

AB +

EQ 25MG BASE;EQ 6MG BASE

N021520 002 Dec 24, 2003

AB +

EQ 25MG BASE;EQ 12MG BASE

N021520 004 Dec 24, 2003

AB +!

EQ 50MG BASE;EQ 6MG BASE

N021520 003 Dec 24, 2003

AB +

EQ 50MG BASE;EQ 12MG BASE

N021520 005 Dec 24, 2003

PRESCRIPTION DRUG PRODUCT LIST

FLUOXYMESTERONE

TABLET; ORAL

FLUOXYMESTERONE

! USL PHARMA

10MG

A088342 001 Oct 21, 1983

FLUPHENAZINE DECANOATE

INJECTABLE; INJECTION

FLUPHENAZINE DECANOATE

| | | | | |
|-----------|----------------------|----------------|--------------------|--------------|
| <u>AO</u> | AUROBINDO PHARMA LTD | <u>25MG/ML</u> | <u>A207739 001</u> | Oct 17, 2017 |
| <u>AO</u> | ! FRESENIUS KABI USA | <u>25MG/ML</u> | <u>A071413 001</u> | Jul 14, 1987 |
| <u>AO</u> | MYLAN LABS LTD | <u>25MG/ML</u> | <u>A075918 001</u> | Aug 17, 2001 |
| <u>AO</u> | PAR STERILE PRODUCTS | <u>25MG/ML</u> | <u>A203732 001</u> | Jul 03, 2014 |
| <u>AO</u> | WEST-WARD PHARMS INT | <u>25MG/ML</u> | <u>A074531 001</u> | Aug 30, 1996 |

FLUPHENAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

FLUPHENAZINE HYDROCHLORIDE

! PHARM ASSOC

5MG/ML

A074725 001 Sep 16, 1996

ELIXIR; ORAL

FLUPHENAZINE HYDROCHLORIDE

! PHARM ASSOC

2.5MG/5ML

A040146 001 Aug 21, 1996

INJECTABLE; INJECTION

FLUPHENAZINE HYDROCHLORIDE

! FRESENIUS KABI USA

2.5MG/ML

A089556 001 Apr 16, 1987

TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

| | | | | |
|-----------|----------------|--------------|--------------------|--------------|
| <u>AB</u> | LANNETT CO INC | <u>1MG</u> | <u>A089743 002</u> | Aug 25, 1988 |
| <u>AB</u> | | <u>2.5MG</u> | <u>A089743 003</u> | Aug 25, 1988 |
| <u>AB</u> | | <u>5MG</u> | <u>A089743 004</u> | Aug 25, 1988 |
| <u>AB</u> | | <u>10MG</u> | <u>A089743 001</u> | Aug 25, 1988 |
| <u>AB</u> | MYLAN | <u>1MG</u> | <u>A089804 002</u> | Aug 12, 1988 |
| <u>AB</u> | | <u>2.5MG</u> | <u>A089804 003</u> | Aug 12, 1988 |
| <u>AB</u> | | <u>5MG</u> | <u>A089804 004</u> | Aug 12, 1988 |
| <u>AB</u> | ! | <u>10MG</u> | <u>A089804 001</u> | Aug 12, 1988 |
| <u>AB</u> | SANDOZ | <u>1MG</u> | <u>A089586 002</u> | Oct 16, 1987 |
| <u>AB</u> | | <u>2.5MG</u> | <u>A089586 003</u> | Oct 16, 1987 |
| <u>AB</u> | | <u>5MG</u> | <u>A089586 004</u> | Oct 16, 1987 |
| <u>AB</u> | | <u>10MG</u> | <u>A089586 001</u> | Oct 16, 1987 |

FLURANDRENOLIDE

CREAM; TOPICAL

CORDRAN SP

| | | | | |
|-----------|------------------------|--------------|--------------------|--------------|
| <u>AT</u> | ! AQUA PHARMS | <u>0.05%</u> | <u>N012806 002</u> | |
| <u>AT</u> | <u>FLURANDRENOLIDE</u> | | | |
| <u>AT</u> | CINTEX SVCS | <u>0.05%</u> | <u>A205342 001</u> | Apr 13, 2016 |
| | CORDRAN SP | | | |
| | ! AQUA PHARMS | 0.025% | N012806 003 | |

LOTION; TOPICAL

CORDRAN

| | | | | |
|-----------|------------------------|--------------|--------------------|--------------|
| <u>AT</u> | ! AQUA PHARMS | <u>0.05%</u> | <u>N013790 001</u> | |
| <u>AT</u> | <u>FLURANDRENOLIDE</u> | | | |
| <u>AT</u> | CINTEX SVCS | <u>0.05%</u> | <u>A205343 001</u> | Dec 22, 2016 |
| <u>AT</u> | PERRIGO UK FINCO | <u>0.05%</u> | <u>A207133 001</u> | Aug 30, 2016 |

OINTMENT; TOPICAL

CORDRAN

| | | | | |
|-----------|------------------------|---------------|--------------------|--------------|
| <u>AT</u> | ! AQUA PHARMS | <u>0.05%</u> | <u>N012806 001</u> | |
| <u>AT</u> | <u>FLURANDRENOLIDE</u> | | | |
| <u>AT</u> | TELIGENT PHARMA INC | <u>0.05%</u> | <u>A207851 001</u> | Dec 30, 2016 |
| | TAPE; TOPICAL | | | |
| | CORDRAN | | | |
| | ! AQUA PHARMS LLC | 0.004MG/SQ CM | N016455 001 | |

FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL

FLURAZEPAM HYDROCHLORIDE

MYLAN PHARMS INC

15MG

A070345 002 Nov 27, 1985

!

30MG

A070345 001 Nov 27, 1985

PRESCRIPTION DRUG PRODUCT LIST

FLURBIPROFEN

TABLET; ORAL

FLURBIPROFEN

| | | | | | |
|-----------|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | MYLAN | <u>50MG</u> | <u>A074358</u> | <u>001</u> | Jun 20, 1994 |
| <u>AB</u> | ! | <u>100MG</u> | <u>A074358</u> | <u>002</u> | Jun 20, 1994 |
| <u>AB</u> | SUN PHARM INDS INC | <u>50MG</u> | <u>A075058</u> | <u>001</u> | Apr 27, 2001 |
| <u>AB</u> | | <u>100MG</u> | <u>A075058</u> | <u>002</u> | Apr 27, 2001 |
| <u>AB</u> | TEVA | <u>100MG</u> | <u>A074431</u> | <u>001</u> | May 31, 1995 |

FLURBIPROFEN SODIUM

SOLUTION/DROPS; OPHTHALMIC

FLURBIPROFEN SODIUM

| | | | | | |
|-----------|-----------------|--------------|----------------|------------|--------------|
| <u>AT</u> | BAUSCH AND LOMB | <u>0.03%</u> | <u>A074447</u> | <u>001</u> | Jan 04, 1995 |
| <u>AT</u> | +! | <u>0.03%</u> | <u>N019404</u> | <u>001</u> | Dec 31, 1986 |

OCUFENFLUTAMIDE

CAPSULE; ORAL

FLUTAMIDE

| | | | | | |
|-----------|---------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ACTAVIS LABS FL INC | <u>125MG</u> | <u>A075820</u> | <u>001</u> | Sep 18, 2001 |
| <u>AB</u> | ! | <u>125MG</u> | <u>A075780</u> | <u>001</u> | Sep 19, 2001 |
| <u>AB</u> | PAR PHARM | <u>125MG</u> | <u>A075298</u> | <u>001</u> | Sep 18, 2001 |

FLUTEMETAMOL F-18

INJECTABLE; INTRAVENOUS

VIZAMYL

| | | | | | |
|----|---------------|----------------------------|---------|-----|--------------|
| +! | GE HEALTHCARE | 121.5mCi/30ML (4.05mCi/ML) | N203137 | 002 | Oct 25, 2013 |
|----|---------------|----------------------------|---------|-----|--------------|

FLUTICASONE FUROATE

POWDER; INHALATION

ARNUITY ELLIPTA

| | | | | | |
|----|-----------------|------------|---------|-----|--------------|
| +! | GLAXOSMITHKLINE | 0.05MG/INH | N205625 | 003 | May 17, 2018 |
| +! | | 0.1MG/INH | N205625 | 001 | Aug 20, 2014 |
| +! | | 0.2MG/INH | N205625 | 002 | Aug 20, 2014 |

FLUTICASONE FUROATE; UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE

POWDER; INHALATION

TRELEGY ELLIPTA

| | | | | | |
|----|-----------------|----------------------------------------------------|---------|-----|--------------|
| +! | GLAXOSMITHKLINE | 0.1MG/INH;EQ 0.0625MG BASE/INH;EQ 0.025MG BASE/INH | N209482 | 001 | Sep 18, 2017 |
|----|-----------------|----------------------------------------------------|---------|-----|--------------|

FLUTICASONE FUROATE; VILANTEROL TRIFENATATE

POWDER; INHALATION

BREQ ELLIPTA

| | | | | | |
|----|---------------|-------------------------------|---------|-----|--------------|
| +! | GLAXO GRP LTD | 0.1MG/INH;EQ 0.025MG BASE/INH | N204275 | 001 | May 10, 2013 |
| +! | | 0.2MG/INH;EQ 0.025MG BASE/INH | N204275 | 002 | Apr 30, 2015 |

FLUTICASONE PROPIONATE

AEROSOL, METERED; INHALATION

FLOVENT HFA

| | | | | | |
|----|---------------|-------------|---------|-----|--------------|
| +! | GLAXO GRP LTD | 0.044MG/INH | N021433 | 003 | May 14, 2004 |
| +! | | 0.11MG/INH | N021433 | 002 | May 14, 2004 |
| +! | | 0.22MG/INH | N021433 | 001 | May 14, 2004 |

CREAM; TOPICAL

FLUTICASONE PROPIONATE

| | | | | | |
|-----------|-----------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ANDA REPOSITORY | <u>0.05%</u> | <u>A076633</u> | <u>001</u> | May 14, 2004 |
| <u>AB</u> | FOUGERA PHARMS | <u>0.05%</u> | <u>A076451</u> | <u>001</u> | May 14, 2004 |
| <u>AB</u> | G AND W LABS | <u>0.05%</u> | <u>A077055</u> | <u>001</u> | Jun 30, 2006 |
| <u>AB</u> | ! | <u>0.05%</u> | <u>A076793</u> | <u>001</u> | May 14, 2004 |

LOTION; TOPICAL

CUTIVATE

| | | | | | | |
|-----------|----|----------------|--------------|----------------|------------|--------------|
| <u>AB</u> | +! | FOUGERA PHARMS | <u>0.05%</u> | <u>N021152</u> | <u>001</u> | Mar 31, 2005 |
|-----------|----|----------------|--------------|----------------|------------|--------------|

FLUTICASONE PROPIONATE

| | | | | | |
|-----------|-------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | GLENMARK GENERICS | <u>0.05%</u> | <u>A090759</u> | <u>001</u> | May 02, 2011 |
| <u>AB</u> | PERRIGO ISRAEL | <u>0.05%</u> | <u>A091553</u> | <u>001</u> | Jul 30, 2013 |

OINTMENT; TOPICAL

FLUTICASONE PROPIONATE

| | | | | | |
|-----------|--------------|---------------|----------------|------------|--------------|
| <u>AB</u> | G AND W LABS | <u>0.005%</u> | <u>A077168</u> | <u>001</u> | Mar 03, 2006 |
| <u>AB</u> | ! | <u>0.005%</u> | <u>A076668</u> | <u>001</u> | May 14, 2004 |

POWDER; INHALATION

ARMONAIR RESPICLICK

| | | | | | |
|----|------------|-------------|---------|-----|--------------|
| + | TEVA PHARM | 0.055MG/INH | N208798 | 001 | Jan 27, 2017 |
| + | | 0.113MG/INH | N208798 | 002 | Jan 27, 2017 |
| +! | | 0.232MG/INH | N208798 | 003 | Jan 27, 2017 |

FLOVENT DISKUS 100

| | | | | | |
|----|---------------|-----------|---------|-----|--------------|
| +! | GLAXO GRP LTD | 0.1MG/INH | N020833 | 002 | Sep 29, 2000 |
|----|---------------|-----------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

FLUTICASONE PROPIONATE

POWDER; INHALATION

FLOVENT DISKUS 250

+! GLAXO GRP LTD 0.25MG/INH N020833 003 Sep 29, 2000

FLOVENT DISKUS 50

+! GLAXO GRP LTD 0.05MG/INH N020833 001 Sep 29, 2000

SPRAY, METERED; NASAL

FLUTICASONE PROPIONATE**AB** APOTEX INC 0.05MG/SPRAY A077538 001 Sep 12, 2007**AB** HI TECH PHARMA 0.05MG/SPRAY A077570 001 Jan 16, 2008**AB** ! WEST-WARD PHARMS 0.05MG/SPRAY A076504 001 Feb 22, 2006**AB** WOCKHARDT BIO AG 0.05MG/SPRAY A078492 001 Jan 09, 2012

KHANCE

+! OPTINOSE US INC 0.093MG N209022 001 Sep 18, 2017

FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE

AEROSOL, METERED; INHALATION

ADV AIR HFA

+! GLAXO GRP LTD 0.045MG/INH;EQ 0.021MG BASE/INH N021254 001 Jun 08, 2006

+! 0.115MG/INH;EQ 0.021MG BASE/INH N021254 002 Jun 08, 2006

+! 0.23MG/INH;EQ 0.021MG BASE/INH N021254 003 Jun 08, 2006

POWDER; INHALATION

ADV AIR DISKUS 100/50

+! GLAXO GRP LTD 0.1MG/INH;EQ 0.05MG BASE/INH N021077 001 Aug 24, 2000

ADV AIR DISKUS 250/50

+! GLAXO GRP LTD 0.25MG/INH;EQ 0.05MG BASE/INH N021077 002 Aug 24, 2000

ADV AIR DISKUS 500/50

+! GLAXO GRP LTD 0.5MG/INH;EQ 0.05MG BASE/INH N021077 003 Aug 24, 2000

AIRDUO RESPICLICK

+ TEVA PHARM 0.055MG/INH;EQ 0.014MG BASE/INH N208799 001 Jan 27, 2017

+ 0.113MG/INH;EQ 0.014MG BASE/INH N208799 002 Jan 27, 2017

+! 0.232MG;EQ 0.014MG BASE/INH N208799 003 Jan 27, 2017

FLUVASTATIN SODIUM

CAPSULE; ORAL

FLUVASTATIN SODIUM**AB** MYLAN PHARMS INC EQ 20MG BASE A090595 001 Apr 11, 2012**AB** ! EQ 40MG BASE A090595 002 Apr 11, 2012**AB** TEVA PHARMS EQ 20MG BASE A078407 001 Jun 12, 2012**AB** EQ 40MG BASE A078407 002 Jun 12, 2012

TABLET, EXTENDED RELEASE; ORAL

FLUVASTATIN SODIUM**AB** MYLAN PHARMS INC EQ 80MG BASE A202458 001 Sep 11, 2015**AB** TEVA PHARMS USA EQ 80MG BASE A079011 001 Jan 27, 2016LESCOL XL**AB** +! NOVARTIS EQ 80MG BASE N021192 001 Oct 06, 2000FLUVOXAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

FLUVOXAMINE MALEATE**AB** ACTAVIS ELIZABETH 100MG A091482 001 Apr 23, 2013**AB** ! 150MG A091482 002 Nov 18, 2013**AB** ANCHEN PHARMS 100MG A091476 001 Mar 13, 2013**AB** 150MG A091476 002 Mar 13, 2013**AB** TORRENT PHARMS LTD 100MG A203240 001 Oct 31, 2014**AB** 150MG A203240 002 Oct 31, 2014

TABLET; ORAL

FLUVOXAMINE MALEATE**AB** ANI PHARMS INC 25MG A075897 001 Jan 25, 2001**AB** 50MG A075897 002 Jan 25, 2001**AB** 100MG A075897 003 Jan 25, 2001**AB** APOTEX 25MG A075902 001 May 07, 2001**AB** 50MG A075902 002 May 07, 2001**AB** 100MG A075902 003 May 07, 2001**AB** MYLAN 25MG A075889 001 Nov 29, 2000**AB** 50MG A075889 002 Nov 29, 2000**AB** 100MG A075889 003 Nov 29, 2000**AB** TEVA 25MG A075893 001 Sep 10, 2002**AB** 50MG A075893 002 Sep 10, 2002**AB** 100MG A075893 003 Sep 10, 2002**AB** UPSHER SMITH LABS 25MG A075888 001 Nov 29, 2000**AB** 50MG A075888 002 Nov 29, 2000**AB** ! 100MG A075888 003 Nov 29, 2000

PRESCRIPTION DRUG PRODUCT LISTFLUVOXAMINE MALEATE

TABLET; ORAL

LUVOX

| | | | | | |
|-----------|------------|--------------|-----------------------|-------------------|--------------|
| AB | ANI PHARMS | 25MG | <u>N021519</u> | <u>001</u> | Dec 20, 2007 |
| AB | | 50MG | <u>N021519</u> | <u>002</u> | Dec 20, 2007 |
| AB | | 100MG | <u>N021519</u> | <u>003</u> | Dec 20, 2007 |

FOLIC ACID

INJECTABLE; INJECTION

FOLIC ACID

! FRESENIUS KABI USA 5MG/ML

A089202 001 Feb 18, 1986

TABLET; ORAL

FOLIC ACID

| | | | | | |
|-----------|---------------------|------------|-----------------------|-------------------|--------------|
| AA | ! AMNEAL PHARM | 1MG | <u>A040625</u> | <u>001</u> | Jul 21, 2005 |
| AA | ANBISON LAB | 1MG | <u>A091145</u> | <u>001</u> | Jul 12, 2013 |
| AA | CADILA PHARMS LTD | 1MG | <u>A202437</u> | <u>001</u> | Jan 27, 2014 |
| AA | CHARTWELL MOLECULAR | 1MG | <u>A090035</u> | <u>001</u> | Jun 09, 2009 |
| AA | HIKMA PHARMS | 1MG | <u>A080600</u> | <u>001</u> | |
| AA | LEADING PHARMA LLC | 1MG | <u>A040796</u> | <u>001</u> | Jan 12, 2009 |
| AA | NUVO PHARMS INC | 1MG | <u>A204418</u> | <u>001</u> | Jul 28, 2015 |
| AA | VINTAGE | 1MG | <u>A040756</u> | <u>001</u> | Jun 04, 2010 |
| AA | ! WATSON LABS | 1MG | <u>A080680</u> | <u>001</u> | |

FOLLITROPIN ALFA/BETA

INJECTABLE; SUBCUTANEOUS

FOLLISTIM AQ

+! ORGANON USA INC 300 IU/0.36ML

N021211 001 Mar 23, 2004

+! 600 IU/0.72ML

N021211 002 Mar 23, 2004

+! 900 IU/1.08ML

N021211 004 Feb 11, 2005

GONAL-F

+! EMD SERONO 450 IU/VIAL

N020378 005 Mar 26, 2004

+ 1,050 IU/VIAL

N020378 004 Feb 28, 2001

GONAL-F RFF

+! EMD SERONO 75 IU/VIAL

N021765 002 Mar 25, 2004

GONAL-F RFF REDI-JECT

+! EMD SERONO 300 IU/0.5ML

N021684 001 May 25, 2004

+! 450 IU/0.75ML

N021684 002 May 25, 2004

+! 900 IU/1.5ML

N021684 003 May 25, 2004

FOMEPIZOLE

INJECTABLE; INJECTION

ANTIZOL

| | | | | | |
|-----------|------------------|------------------------------------|-----------------------|-------------------|--------------|
| AP | +! PAR PHARM INC | <u>1.5GM/1.5ML (1GM/ML)</u> | <u>N020696</u> | <u>001</u> | Dec 04, 1997 |
|-----------|------------------|------------------------------------|-----------------------|-------------------|--------------|

FOMEPIZOLE

| | | | | | |
|-----------|---------------------|------------------------------------|-----------------------|-------------------|--------------|
| AP | LUITPOLD | <u>1.5GM/1.5ML (1GM/ML)</u> | <u>A078368</u> | <u>001</u> | Dec 14, 2007 |
| AP | MYLAN INSTITUTIONAL | <u>1.5GM/1.5ML (1GM/ML)</u> | <u>A078639</u> | <u>001</u> | Mar 03, 2008 |
| AP | NAVINTA LLC | <u>1.5GM/1.5ML (1GM/ML)</u> | <u>A078537</u> | <u>001</u> | Mar 06, 2008 |

FONDAPARINUX SODIUM

INJECTABLE; SUBCUTANEOUS

ARIXTRA

| | | | | | |
|-----------|----------------------|---------------------------|-----------------------|-------------------|--------------|
| AP | +! MYLAN IRELAND LTD | <u>2.5MG/0.5ML</u> | <u>N021345</u> | <u>001</u> | Dec 07, 2001 |
| AP | +! | <u>5MG/0.4ML</u> | <u>N021345</u> | <u>002</u> | May 28, 2004 |
| AP | +! | <u>7.5MG/0.6ML</u> | <u>N021345</u> | <u>003</u> | May 28, 2004 |
| AP | +! | <u>10MG/0.8ML</u> | <u>N021345</u> | <u>004</u> | May 28, 2004 |

FONDAPARINUX SODIUM

| | | | | | |
|-----------|----------------------|---------------------------|-----------------------|-------------------|--------------|
| AP | AUROBINDO PHARMA LTD | <u>2.5MG/0.5ML</u> | <u>A206918</u> | <u>001</u> | Dec 26, 2017 |
| AP | | <u>5MG/0.4ML</u> | <u>A206918</u> | <u>002</u> | Dec 26, 2017 |
| AP | | <u>7.5MG/0.6ML</u> | <u>A206918</u> | <u>003</u> | Dec 26, 2017 |
| AP | | <u>10MG/0.8ML</u> | <u>A206918</u> | <u>004</u> | Dec 26, 2017 |
| AP | DR REDDYS LABS LTD | <u>2.5MG/0.5ML</u> | <u>A091316</u> | <u>001</u> | Jul 11, 2011 |
| AP | | <u>5MG/0.4ML</u> | <u>A091316</u> | <u>002</u> | Jul 11, 2011 |
| AP | | <u>7.5MG/0.6ML</u> | <u>A091316</u> | <u>003</u> | Jul 11, 2011 |
| AP | | <u>10MG/0.8ML</u> | <u>A091316</u> | <u>004</u> | Jul 11, 2011 |
| AP | JIANGSU HENGRUI MED | <u>2.5MG/0.5ML</u> | <u>A206812</u> | <u>001</u> | May 15, 2018 |
| AP | | <u>5MG/0.4ML</u> | <u>A206812</u> | <u>002</u> | May 15, 2018 |
| AP | | <u>7.5MG/0.6ML</u> | <u>A206812</u> | <u>003</u> | May 15, 2018 |
| AP | | <u>10MG/0.8ML</u> | <u>A206812</u> | <u>004</u> | May 15, 2018 |
| AP | SCINOPHARM TAIWAN | <u>2.5MG/0.5ML</u> | <u>A208615</u> | <u>001</u> | Nov 14, 2018 |
| AP | | <u>5MG/0.4ML</u> | <u>A208615</u> | <u>002</u> | Nov 14, 2018 |
| AP | | <u>7.5MG/0.6ML</u> | <u>A208615</u> | <u>003</u> | Nov 14, 2018 |
| AP | | <u>10MG/0.8ML</u> | <u>A208615</u> | <u>004</u> | Nov 14, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

FORMOTEROL FUMARATE

SOLUTION; INHALATION

PERFORMIST

+! MYLAN SPECLT 0.02MG/2ML N022007 001 May 11, 2007

FORMOTEROL FUMARATE; GLYCOPYRROLATE

AEROSOL, METERED; INHALATION

BEVESPI AEROSPHERE

+! ASTRAZENECA PHARMS 0.0048MG/INH; 0.0090MG/INH N208294 001 Apr 25, 2016

FORMOTEROL FUMARATE; MOMETASONE FUROATE

AEROSOL, METERED; INHALATION

DULERA

+! MERCK SHARP DOHME 0.005MG/INH; 0.1MG/INH N022518 001 Jun 22, 2010

+! 0.005MG/INH; 0.2MG/INH N022518 002 Jun 22, 2010

FOSAMPRENAVIR CALCIUM

SUSPENSION; ORAL

LEXIVA

+! VIIV HLHCARE EQ 50MG BASE/ML N022116 001 Jun 14, 2007

TABLET; ORAL

FOSAMPRENAVIR CALCIUM**AB** MYLAN PHARMS INC **EQ 700MG BASE** **A204060 001** Apr 15, 2016LEXIVA**AB** +! VIIV HLHCARE **EQ 700MG BASE** **N021548 001** Oct 20, 2003FOSAPREPITANT DIMEGLUMINE

POWDER; INTRAVENOUS

EMEND**AP** +! MERCK AND CO INC **EQ 150MG BASE/VIAL** **N022023 002** Nov 12, 2010FOSAPREPITANT DIMEGLUMINE**AP** FRESENIUS KABI USA **EQ 150MG BASE/VIAL** **A206197 001** Jun 09, 2016FOSCARNET SODIUM

INJECTABLE; INJECTION

FOSCAVIR

+! CLINIGEN HLHCARE 2.4GM/100ML N020068 001 Sep 27, 1991

FOSFOMYCIN TROMETHAMINE

FOR SOLUTION; ORAL

MONUROL

+! ZAMBON SPA EQ 3GM BASE/PACKET N050717 001 Dec 19, 1996

FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM**AB** APOTEX INC **10MG** **A076906 001** May 17, 2005**AB** **20MG** **A076906 002** May 17, 2005**AB** **40MG** **A076906 003** May 17, 2005**AB** AUROBINDO PHARMA LTD **10MG** **A091163 001** Mar 30, 2011**AB** **20MG** **A091163 002** Mar 30, 2011**AB** **40MG** **A091163 003** Mar 30, 2011**AB** INVAGEN PHARMS **10MG** **A077222 001** Apr 20, 2005**AB** **20MG** **A077222 002** Apr 20, 2005**AB** **40MG** **A077222 003** Apr 20, 2005**AB** PRINSTON INC **10MG** **A205670 001** Aug 29, 2016**AB** **20MG** **A205670 002** Aug 29, 2016**AB** **40MG** **A205670 003** Aug 29, 2016**AB** TEVA **10MG** **A076139 001** Nov 25, 2003**AB** **20MG** **A076139 002** Nov 25, 2003**AB** ! **40MG** **A076139 003** Nov 25, 2003**AB** UPSHER SMITH LABS **10MG** **A076483 001** Apr 23, 2004**AB** **20MG** **A076483 002** Apr 23, 2004**AB** **40MG** **A076483 003** Apr 23, 2004FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE**AB** AUROBINDO PHARMA **10MG; 12.5MG** **A079245 001** Jul 09, 2009**AB** **20MG; 12.5MG** **A079245 002** Jul 09, 2009**AB** EMCURE PHARMS LTD **10MG; 12.5MG** **A079025 001** Sep 17, 2010**AB** ! **20MG; 12.5MG** **A079025 002** Sep 17, 2010**AB** INVAGEN PHARMS **10MG; 12.5MG** **A090228 001** Jul 09, 2009**AB** **20MG; 12.5MG** **A090228 002** Jul 09, 2009**AB** SANDOZ **10MG; 12.5MG** **A076961 001** Sep 28, 2005**AB** **20MG; 12.5MG** **A076961 002** Sep 28, 2005

PRESCRIPTION DRUG PRODUCT LIST

FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE

POWDER; INTRAVENOUS

AKYNZEO

+! HELSINN HLTHCARE EQ 235MG BASE;EQ 0.25MG BASE N210493 001 Apr 19, 2018

FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

CEREBYXAP +! PARKE DAVIS EQ 50MG PHENYTOIN NA/ML N020450 001 Aug 05, 1996FOSPHENYTOIN SODIUMAP AMNEAL PHARMS CO EQ 50MG PHENYTOIN NA/ML A078476 001 Mar 18, 2008AP FRESENIUS KABI USA EQ 50MG PHENYTOIN NA/ML A078052 001 Aug 06, 2007AP HIKMA FARMACEUTICA EQ 50MG PHENYTOIN NA/ML A078765 001 Dec 02, 2009AP LUITPOLD EQ 50MG PHENYTOIN NA/ML A078277 001 Aug 06, 2007AP EQ 50MG PHENYTOIN NA/ML A090099 001 May 13, 2010AP MYLAN LABS LTD EQ 50MG PHENYTOIN NA/ML A078736 001 Jun 08, 2010AP SUN PHARMA GLOBAL EQ 50MG PHENYTOIN NA/ML A078417 001 Mar 18, 2008AP WEST-WARD PHARMS EQ 50MG PHENYTOIN NA/ML A077481 001 Aug 06, 2007

INT

AP EQ 50MG PHENYTOIN NA/ML A077989 001 Aug 06, 2007AP WOCKHARDT EQ 50MG PHENYTOIN NA/ML A078137 001 Aug 06, 2007FOSTAMATINIB DISODIUM

TABLET; ORAL

TAVALISSE

+ RIGEL PHARMS INC EQ 100MG BASE N209299 001 Apr 17, 2018

+! EQ 150MG BASE N209299 002 Apr 17, 2018

FROVATRIPTAN SUCCINATE

TABLET; ORAL

FROVAAB +! ENDO PHARMS EQ 2.5MG BASE N021006 001 Nov 08, 2001FROVATRIPTAN SUCCINATEAB AMNEAL PHARMS CO EQ 2.5MG BASE A211292 001 Nov 06, 2018AB GLENMARK PHARMS LTD EQ 2.5MG BASE A204730 001 Mar 11, 2016AB MYLAN PHARMS INC EQ 2.5MG BASE A202931 001 Aug 28, 2014FULVESTRANT

INJECTABLE; INTRAMUSCULAR

FASLODEX

+! ASTRAZENECA 50MG/ML N021344 001 Apr 25, 2002

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDEAP AMNEAL PHARMS CO 10MG/ML A207552 001 Jul 20, 2016AP ! BAXTER HLTHCARE 10MG/ML A202747 001 Jan 27, 2014

CORP

AP EMCURE PHARMS LTD 10MG/ML A203428 001 Aug 26, 2014AP FRESENIUS KABI USA 10MG/ML N018902 001 May 22, 1984AP HOSPIRA 10MG/ML A075241 001 May 28, 1999AP 10MG/ML N018667 001 May 28, 1982AP WOCKHARDT 10MG/ML A077941 001 Mar 22, 2007

SOLUTION; ORAL

FUROSEMIDEAA ! WEST-WARD PHARMS 10MG/ML A070434 001 Apr 22, 1987

INT

AA WOCKHARDT BIO AG 10MG/ML A070655 001 Oct 02, 1987

INT

WEST-WARD PHARMS 40MG/5ML A070433 001 Apr 22, 1987

TABLET; ORAL

FUROSEMIDEAB IPCA LABS LTD 20MG A078010 001 Sep 18, 2006AB 40MG A078010 002 Sep 18, 2006AB 80MG A078010 003 Sep 18, 2006AB IVAX SUB TEVA 20MG N018413 001 Nov 30, 1983

PHARMS

AB 40MG N018413 002 Nov 30, 1983AB LEADING PHARMA LLC 20MG A077293 001 Nov 09, 2005AB 40MG A077293 002 Nov 09, 2005AB 80MG A077293 003 Nov 09, 2005AB MYLAN 20MG N018487 001AB 40MG N018487 002AB 80MG A070082 001 Oct 29, 1986AB PRINSTON INC 20MG A076796 001 Mar 26, 2004AB 40MG A076796 002 Mar 26, 2004

PRESCRIPTION DRUG PRODUCT LIST

FUROSEMIDE

TABLET; ORAL

FUROSEMIDE

| | | | | |
|--------------|-------------------------|-------------|--------------------|--------------|
| <u>AB</u> | | <u>80MG</u> | <u>A076796 003</u> | Mar 26, 2004 |
| <u>AB</u> | SANDOZ | <u>20MG</u> | <u>N018569 002</u> | |
| <u>AB</u> | | <u>40MG</u> | <u>N018569 001</u> | |
| <u>AB</u> | | <u>80MG</u> | <u>N018569 005</u> | Aug 14, 1984 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>20MG</u> | <u>N018823 001</u> | Nov 10, 1983 |
| <u>AB</u> | | <u>40MG</u> | <u>N018823 002</u> | Nov 10, 1983 |
| <u>AB</u> | | <u>80MG</u> | <u>A070086 001</u> | Jan 24, 1986 |
| <u>LASIX</u> | | | | |
| <u>AB</u> | + US PHARM HOLDINGS | <u>20MG</u> | <u>N016273 002</u> | |
| <u>AB</u> | + | <u>40MG</u> | <u>N016273 001</u> | |
| <u>AB</u> | +! | <u>80MG</u> | <u>N016273 003</u> | |

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

| | | | | |
|------------------|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | ACI HEALTHCARE LTD | <u>100MG</u> | <u>A206943 001</u> | May 14, 2018 |
| <u>AB</u> | | <u>300MG</u> | <u>A206943 002</u> | May 14, 2018 |
| <u>AB</u> | | <u>400MG</u> | <u>A206943 003</u> | May 14, 2018 |
| <u>AB</u> | ACTAVIS ELIZABETH | <u>100MG</u> | <u>A075350 001</u> | Sep 12, 2003 |
| <u>AB</u> | | <u>300MG</u> | <u>A075350 002</u> | Sep 12, 2003 |
| <u>AB</u> | | <u>400MG</u> | <u>A075350 003</u> | Sep 12, 2003 |
| <u>AB</u> | ALKEM | <u>100MG</u> | <u>A090858 001</u> | Dec 17, 2010 |
| <u>AB</u> | | <u>300MG</u> | <u>A090858 002</u> | Dec 17, 2010 |
| <u>AB</u> | | <u>400MG</u> | <u>A090858 003</u> | Dec 17, 2010 |
| <u>AB</u> | AMNEAL PHARMS NY | <u>100MG</u> | <u>A078428 001</u> | Jul 25, 2007 |
| <u>AB</u> | | <u>300MG</u> | <u>A078428 002</u> | Jul 25, 2007 |
| <u>AB</u> | | <u>400MG</u> | <u>A078428 003</u> | Jul 25, 2007 |
| <u>AB</u> | APOTEX INC | <u>100MG</u> | <u>A075360 001</u> | Apr 06, 2005 |
| <u>AB</u> | | <u>300MG</u> | <u>A075360 002</u> | Apr 06, 2005 |
| <u>AB</u> | | <u>400MG</u> | <u>A075360 003</u> | Apr 06, 2005 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>100MG</u> | <u>A078787 001</u> | Jan 31, 2008 |
| <u>AB</u> | | <u>300MG</u> | <u>A078787 002</u> | Jan 31, 2008 |
| <u>AB</u> | | <u>400MG</u> | <u>A078787 003</u> | Jan 31, 2008 |
| <u>AB</u> | EPIC PHARMA LLC | <u>100MG</u> | <u>A207099 001</u> | Mar 24, 2017 |
| <u>AB</u> | | <u>300MG</u> | <u>A207099 002</u> | Mar 24, 2017 |
| <u>AB</u> | | <u>400MG</u> | <u>A207099 003</u> | Mar 24, 2017 |
| <u>AB</u> | INVAGEN PHARMS | <u>100MG</u> | <u>A090705 001</u> | Dec 30, 2009 |
| <u>AB</u> | | <u>300MG</u> | <u>A090705 002</u> | Dec 30, 2009 |
| <u>AB</u> | | <u>400MG</u> | <u>A090705 003</u> | Dec 30, 2009 |
| <u>AB</u> | JIANGSU HENGRUI MED | <u>100MG</u> | <u>A091008 001</u> | Oct 26, 2017 |
| <u>AB</u> | | <u>300MG</u> | <u>A091008 002</u> | Oct 26, 2017 |
| <u>AB</u> | | <u>400MG</u> | <u>A091008 003</u> | Oct 26, 2017 |
| <u>AB</u> | MARKSANS PHARMA | <u>100MG</u> | <u>A090007 001</u> | Jul 21, 2011 |
| <u>AB</u> | | <u>300MG</u> | <u>A090007 002</u> | Jul 21, 2011 |
| <u>AB</u> | | <u>400MG</u> | <u>A090007 003</u> | Jul 21, 2011 |
| <u>AB</u> | MYLAN | <u>100MG</u> | <u>A090158 001</u> | Feb 14, 2011 |
| <u>AB</u> | | <u>300MG</u> | <u>A090158 002</u> | Feb 14, 2011 |
| <u>AB</u> | | <u>400MG</u> | <u>A090158 003</u> | Feb 14, 2011 |
| <u>AB</u> | SCIEGEN PHARMS INC | <u>100MG</u> | <u>A204989 001</u> | Feb 18, 2016 |
| <u>AB</u> | | <u>300MG</u> | <u>A204989 002</u> | Feb 18, 2016 |
| <u>AB</u> | | <u>400MG</u> | <u>A204989 003</u> | Feb 18, 2016 |
| <u>AB</u> | STRIDES PHARMA | <u>100MG</u> | <u>A211314 001</u> | Oct 16, 2018 |
| <u>AB</u> | | <u>300MG</u> | <u>A211314 002</u> | Oct 16, 2018 |
| <u>AB</u> | | <u>400MG</u> | <u>A211314 003</u> | Oct 16, 2018 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>100MG</u> | <u>A077242 001</u> | Aug 24, 2006 |
| <u>AB</u> | | <u>300MG</u> | <u>A077242 002</u> | Aug 24, 2006 |
| <u>AB</u> | | <u>400MG</u> | <u>A077242 003</u> | Aug 24, 2006 |
| <u>AB</u> | TARO PHARM | <u>100MG</u> | <u>A077261 001</u> | Aug 02, 2013 |
| <u>AB</u> | | <u>300MG</u> | <u>A077261 002</u> | Aug 02, 2013 |
| <u>AB</u> | | <u>400MG</u> | <u>A077261 003</u> | Aug 02, 2013 |
| <u>AB</u> | TEVA PHARMS | <u>100MG</u> | <u>A075435 001</u> | Oct 08, 2004 |
| <u>AB</u> | | <u>300MG</u> | <u>A075435 002</u> | Oct 08, 2004 |
| <u>AB</u> | | <u>400MG</u> | <u>A075435 003</u> | Oct 08, 2004 |
| <u>NEURONTIN</u> | | | | |
| <u>AB</u> | + PFIZER PHARMS | <u>100MG</u> | <u>N020235 001</u> | Dec 30, 1993 |
| <u>AB</u> | + | <u>300MG</u> | <u>N020235 002</u> | Dec 30, 1993 |
| <u>AB</u> | +! | <u>400MG</u> | <u>N020235 003</u> | Dec 30, 1993 |

PRESCRIPTION DRUG PRODUCT LISTGABAPENTIN

SOLUTION; ORAL

GABAPENTIN

| | | | | | |
|-----------|-------------------|------------------|----------------|------------|--------------|
| AA | ACELLA PHARMS LLC | <u>250MG/5ML</u> | <u>A076403</u> | <u>001</u> | May 01, 2012 |
| AA | AMNEAL PHARMS | <u>250MG/5ML</u> | <u>A202024</u> | <u>001</u> | Mar 23, 2012 |
| AA | HI TECH PHARMA | <u>250MG/5ML</u> | <u>A078974</u> | <u>001</u> | Feb 18, 2011 |
| AA | TARO | <u>250MG/5ML</u> | <u>A076672</u> | <u>001</u> | Jul 03, 2013 |
| AA | TRIS PHARMA INC | <u>250MG/5ML</u> | <u>A091286</u> | <u>001</u> | Mar 14, 2016 |

NEURONTIN

| | | | | | |
|-----------|----------------|------------------|----------------|------------|--------------|
| AA | +! PARKE DAVIS | <u>250MG/5ML</u> | <u>N021129</u> | <u>001</u> | Mar 02, 2000 |
|-----------|----------------|------------------|----------------|------------|--------------|

TABLET; ORAL

GABAPENTIN

| | | | | | |
|-----------|----------------------|--------------|----------------|------------|--------------|
| AB | ACI HEALTHCARE LTD | <u>600MG</u> | <u>A203244</u> | <u>002</u> | Jul 12, 2013 |
| AB | | <u>800MG</u> | <u>A203244</u> | <u>001</u> | Jul 12, 2013 |
| AB | ACTAVIS ELIZABETH | <u>600MG</u> | <u>A075694</u> | <u>001</u> | Oct 21, 2004 |
| AB | | <u>800MG</u> | <u>A075694</u> | <u>002</u> | Oct 21, 2004 |
| AB | ALKEM LABS LTD | <u>600MG</u> | <u>A206402</u> | <u>001</u> | Dec 23, 2015 |
| AB | | <u>800MG</u> | <u>A206402</u> | <u>002</u> | Dec 23, 2015 |
| AB | APOTEX INC | <u>100MG</u> | <u>A077894</u> | <u>001</u> | Oct 10, 2006 |
| AB | | <u>300MG</u> | <u>A077894</u> | <u>002</u> | Oct 10, 2006 |
| AB | | <u>400MG</u> | <u>A077894</u> | <u>003</u> | Oct 10, 2006 |
| AB | | <u>600MG</u> | <u>A077661</u> | <u>004</u> | Sep 13, 2006 |
| AB | | <u>800MG</u> | <u>A077661</u> | <u>005</u> | Sep 13, 2006 |
| AB | AUROBINDO PHARMA LTD | <u>600MG</u> | <u>A200651</u> | <u>001</u> | Oct 06, 2011 |
| AB | | <u>800MG</u> | <u>A200651</u> | <u>002</u> | Oct 06, 2011 |
| AB | CSPC OUYI PHARM CO | <u>600MG</u> | <u>A207057</u> | <u>001</u> | Oct 26, 2017 |
| AB | | <u>800MG</u> | <u>A207057</u> | <u>002</u> | Oct 26, 2017 |
| AB | GLENMARK PHARMS LTD | <u>600MG</u> | <u>A077662</u> | <u>001</u> | Aug 18, 2006 |
| AB | | <u>800MG</u> | <u>A077662</u> | <u>002</u> | Aug 18, 2006 |
| AB | INVAGEN PHARMS | <u>600MG</u> | <u>A202764</u> | <u>001</u> | Oct 16, 2012 |
| AB | | <u>800MG</u> | <u>A202764</u> | <u>002</u> | Oct 16, 2012 |
| AB | IVAX SUB TEVA PHARMS | <u>100MG</u> | <u>A076017</u> | <u>001</u> | Apr 28, 2004 |
| AB | | <u>300MG</u> | <u>A076017</u> | <u>002</u> | Apr 28, 2004 |
| AB | | <u>400MG</u> | <u>A076017</u> | <u>003</u> | Apr 28, 2004 |
| AB | | <u>600MG</u> | <u>A076017</u> | <u>004</u> | Apr 29, 2005 |
| AB | | <u>800MG</u> | <u>A076017</u> | <u>005</u> | Apr 29, 2005 |
| AB | LUPIN LTD | <u>600MG</u> | <u>A209306</u> | <u>001</u> | Aug 24, 2018 |
| AB | | <u>800MG</u> | <u>A209306</u> | <u>002</u> | Aug 24, 2018 |
| AB | MYLAN PHARMS INC | <u>600MG</u> | <u>A090335</u> | <u>001</u> | Jun 01, 2010 |
| AB | | <u>800MG</u> | <u>A090335</u> | <u>002</u> | Jun 01, 2010 |
| AB | SCIEGEN PHARMS INC | <u>600MG</u> | <u>A205101</u> | <u>001</u> | Feb 04, 2016 |
| AB | | <u>800MG</u> | <u>A205101</u> | <u>002</u> | Feb 04, 2016 |
| AB | SUN PHARM INDS LTD | <u>600MG</u> | <u>A077525</u> | <u>001</u> | Aug 24, 2006 |
| AB | | <u>800MG</u> | <u>A077525</u> | <u>002</u> | Aug 24, 2006 |
| AB | TEVA PHARMS USA | <u>600MG</u> | <u>A205807</u> | <u>001</u> | Mar 10, 2017 |
| AB | | <u>800MG</u> | <u>A205807</u> | <u>002</u> | Mar 10, 2017 |
| AB | ZYDUS PHARMS USA INC | <u>600MG</u> | <u>A078926</u> | <u>001</u> | Feb 11, 2011 |
| AB | | <u>800MG</u> | <u>A078926</u> | <u>002</u> | Feb 11, 2011 |
| | <u>NEURONTIN</u> | | | | |
| AB | + PFIZER PHARMS | <u>600MG</u> | <u>N020882</u> | <u>001</u> | Oct 09, 1998 |
| AB | +! | <u>800MG</u> | <u>N020882</u> | <u>002</u> | Oct 09, 1998 |
| | GRALISE | | | | |
| BX | +! ASSERTIO | 300MG | N022544 | 001 | Jan 28, 2011 |
| BX | +! | 600MG | N022544 | 002 | Jan 28, 2011 |

GABAPENTIN ENACARBIL

TABLET, EXTENDED RELEASE; ORAL

HORIZANT

| | | | | | |
|--|--------------------|-------|---------|-----|--------------|
| | + ARBOR PHARMS LLC | 300MG | N022399 | 002 | Dec 13, 2011 |
| | +! | 600MG | N022399 | 001 | Apr 06, 2011 |

GADOBENATE DIMEGLUMINE

INJECTABLE; INTRAVENOUS

MULTIHANCE

| | | | | | |
|--|----------------------|-------------------------|---------|-----|--------------|
| | +! BRACCO | 2.645GM/5ML (529MG/ML) | N021357 | 001 | Nov 23, 2004 |
| | +! | 5.29GM/10ML (529MG/ML) | N021357 | 002 | Nov 23, 2004 |
| | +! | 7.935GM/15ML (529MG/ML) | N021357 | 003 | Nov 23, 2004 |
| | +! | 10.58GM/20ML (529MG/ML) | N021357 | 004 | Nov 23, 2004 |
| | MULTIHANCE MULTIPACK | | | | |
| | +! BRACCO | 26.45GM/50ML (529MG/ML) | N021358 | 001 | Nov 23, 2004 |
| | +! | 52.9GM/100ML (529MG/ML) | N021358 | 002 | Nov 23, 2004 |

PRESCRIPTION DRUG PRODUCT LIST

GADOBUTROL

SOLUTION; INTRAVENOUS

GADAVIST

| | | | | | | |
|---|---|----------------|------------------------------|---------|-----|--------------|
| + | ! | BAYER HLTHCARE | 1.20944GM/2ML (604.72MG/ML) | N201277 | 006 | Dec 18, 2013 |
| + | ! | | 4.5354GM/7.5ML (604.72MG/ML) | N201277 | 001 | Mar 14, 2011 |
| + | ! | | 6.0472GM/10ML (604.72MG/ML) | N201277 | 002 | Mar 14, 2011 |
| + | ! | | 9.0708GM/15ML (604.72MG/ML) | N201277 | 003 | Mar 14, 2011 |
| + | ! | | 18.1416GM/30ML (604.72MG/ML) | N201277 | 004 | Mar 14, 2011 |
| + | ! | | 39.3068GM/65ML (604.72MG/ML) | N201277 | 005 | Mar 14, 2011 |

GADODIAMIDE

INJECTABLE; INJECTION

OMNISCAN

| | | | | | | |
|---|---|---------------|-------------------------|---------|-----|--------------|
| + | ! | GE HEALTHCARE | 287MG/ML | N020123 | 001 | Jan 08, 1993 |
| + | ! | | 28.7GM/100ML (287MG/ML) | N022066 | 002 | Sep 05, 2007 |

GADOPENTETATE DIMEGLUMINE

INJECTABLE; INJECTION

MAGNEVIST

| | | | | | | |
|---|---|----------------|-------------|---------|-----|--------------|
| + | ! | BAYER HLTHCARE | 469.01MG/ML | N019596 | 001 | Jun 02, 1988 |
| + | ! | | 469.01MG/ML | N021037 | 001 | Mar 10, 2000 |

GADOTERATE MEGLUMINE

SOLUTION; INTRAVENOUS

DOTAREM

| | | | | | | |
|---|---|---------|----------------------------|---------|-----|--------------|
| + | ! | GUERBET | 37.69GM/100ML (376.9MG/ML) | N204781 | 001 | Mar 20, 2013 |
| + | ! | | 1.8845GM/5ML (376.9MG/ML) | N204781 | 005 | Mar 31, 2017 |
| + | ! | | 3.769GM/10ML (376.9MG/ML) | N204781 | 002 | Mar 20, 2013 |
| + | ! | | 5.6535GM/15ML (376.9MG/ML) | N204781 | 003 | Mar 20, 2013 |
| + | ! | | 7.538GM/20ML (376.9MG/ML) | N204781 | 004 | Mar 20, 2013 |

GADOTERIDOL

INJECTABLE; INJECTION

PROHANCE

| | | | | | | |
|---|---|------------------|------------|---------|-----|--------------|
| + | ! | BRACCO | 279.3MG/ML | N020131 | 001 | Nov 16, 1992 |
| + | ! | BRACCO MULTIPACK | | | | |
| + | ! | BRACCO | 279.3MG/ML | N021489 | 001 | Oct 09, 2003 |

GADOXETATE DISODIUM

SOLUTION; INTRAVENOUS

EOVIST

| | | | | | | |
|---|---|----------------|------------------------------|---------|-----|--------------|
| + | ! | BAYER HLTHCARE | 1.8143GM/10ML (181.43MG/ML) | N022090 | 001 | Jul 03, 2008 |
| + | ! | | 2.72145GM/15ML (181.43MG/ML) | N022090 | 002 | Feb 04, 2013 |

GALANTAMINE HYDROBROMIDE

CAPSULE, EXTENDED RELEASE; ORAL

GALANTAMINE HYDROBROMIDE

| | | | | | | |
|-----------|---|----------------------|---------------------|----------------|------------|--------------|
| AB | | AUROBINDO PHARMA LTD | EQ 8MG BASE | A204895 | 001 | Aug 05, 2016 |
| AB | | | EQ 16MG BASE | A204895 | 002 | Aug 05, 2016 |
| AB | | | EQ 24MG BASE | A204895 | 003 | Aug 05, 2016 |
| AB | | BARR | EQ 8MG BASE | A078189 | 001 | Sep 15, 2008 |
| AB | | | EQ 16MG BASE | A078189 | 002 | Sep 15, 2008 |
| AB | | | EQ 24MG BASE | A078189 | 003 | Sep 15, 2008 |
| AB | | SUN PHARMA GLOBAL | EQ 8MG BASE | A090178 | 001 | Feb 02, 2011 |
| AB | | | EQ 16MG BASE | A090178 | 002 | Feb 02, 2011 |
| AB | | | EQ 24MG BASE | A090178 | 003 | Feb 02, 2011 |
| AB | | WATSON LABS | EQ 8MG BASE | A079028 | 001 | Dec 15, 2008 |
| AB | | | EQ 16MG BASE | A079028 | 002 | Dec 15, 2008 |
| AB | | | EQ 24MG BASE | A079028 | 003 | Dec 15, 2008 |
| | | RAZADYNE ER | | | | |
| AB | + | JANSSEN PHARMS | EQ 8MG BASE | N021615 | 001 | Apr 01, 2005 |
| AB | + | | EQ 16MG BASE | N021615 | 002 | Apr 01, 2005 |
| AB | + | | EQ 24MG BASE | N021615 | 003 | Apr 01, 2005 |

SOLUTION; ORAL

GALANTAMINE HYDROBROMIDE

| | | | | | | |
|---|---|----------------------|--------|---------|-----|--------------|
| ! | ! | WEST-WARD PHARMS INT | 4MG/ML | A078185 | 001 | Jan 30, 2009 |
|---|---|----------------------|--------|---------|-----|--------------|

TABLET; ORAL

GALANTAMINE HYDROBROMIDE

| | | | | | | |
|-----------|--|----------------------|---------------------|----------------|------------|--------------|
| AB | | APOTEX INC | EQ 4MG BASE | A077781 | 001 | Sep 27, 2011 |
| AB | | | EQ 8MG BASE | A077781 | 002 | Sep 27, 2011 |
| AB | | | EQ 12MG BASE | A077781 | 003 | Sep 27, 2011 |
| AB | | AUROBINDO PHARMA LTD | EQ 4MG BASE | A090957 | 001 | Mar 29, 2011 |
| AB | | | EQ 8MG BASE | A090957 | 002 | Mar 29, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

GALANTAMINE HYDROBROMIDE

TABLET; ORAL

GALANTAMINE HYDROBROMIDE

| | | | | |
|-----------|-------------------------|---------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 12MG BASE</u> | <u>A090957 003</u> | Mar 29, 2011 |
| <u>AB</u> | BARR | <u>EQ 4MG BASE</u> | <u>A077605 001</u> | Aug 28, 2008 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A077605 002</u> | Aug 28, 2008 |
| <u>AB</u> | | <u>EQ 12MG BASE</u> | <u>A077605 003</u> | Aug 28, 2008 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 4MG BASE</u> | <u>A077593 001</u> | Sep 11, 2008 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A077593 002</u> | Sep 11, 2008 |
| <u>AB</u> | | <u>EQ 12MG BASE</u> | <u>A077593 003</u> | Sep 11, 2008 |
| <u>AB</u> | MYLAN | <u>EQ 4MG BASE</u> | <u>A077590 001</u> | May 29, 2009 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A077590 002</u> | May 29, 2009 |
| <u>AB</u> | | <u>EQ 12MG BASE</u> | <u>A077590 003</u> | May 29, 2009 |
| <u>AB</u> | SANDOZ | <u>EQ 4MG BASE</u> | <u>A077589 001</u> | Jun 22, 2009 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A077589 002</u> | Jun 22, 2009 |
| <u>AB</u> | | <u>EQ 12MG BASE</u> | <u>A077589 003</u> | Jun 22, 2009 |
| <u>AB</u> | TEVA PHARMS | <u>EQ 4MG BASE</u> | <u>A077587 001</u> | Jul 09, 2009 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A077587 002</u> | Jul 09, 2009 |
| <u>AB</u> | | <u>EQ 12MG BASE</u> | <u>A077587 003</u> | Jul 09, 2009 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>EQ 4MG BASE</u> | <u>A077608 001</u> | Feb 11, 2009 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A077608 002</u> | Feb 11, 2009 |
| <u>AB</u> | | <u>EQ 12MG BASE</u> | <u>A077608 003</u> | Feb 11, 2009 |
| <u>AB</u> | YABAO PHARM | <u>EQ 4MG BASE</u> | <u>A077604 001</u> | Feb 06, 2009 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A077604 002</u> | Feb 06, 2009 |
| <u>AB</u> | | <u>EQ 12MG BASE</u> | <u>A077604 003</u> | Feb 06, 2009 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 4MG BASE</u> | <u>A078898 001</u> | Feb 17, 2011 |
| <u>AB</u> | | <u>EQ 8MG BASE</u> | <u>A078898 002</u> | Feb 17, 2011 |
| <u>AB</u> | | <u>EQ 12MG BASE</u> | <u>A078898 003</u> | Feb 17, 2011 |
| | <u>RAZADYNE</u> | | | |
| <u>AB</u> | +! JANSSEN PHARMS | <u>EQ 4MG BASE</u> | <u>N021169 001</u> | Feb 28, 2001 |
| <u>AB</u> | + | <u>EQ 8MG BASE</u> | <u>N021169 002</u> | Feb 28, 2001 |
| <u>AB</u> | + | <u>EQ 12MG BASE</u> | <u>N021169 003</u> | Feb 28, 2001 |

GALLIUM CITRATE GA-67

INJECTABLE; INJECTION

GALLIUM CITRATE GA 67

| | | | | |
|----|---------------------|---------|-------------|--|
| BS | LANTHEUS MEDCL | 2mCi/ML | N017478 001 | |
| BS | MALLINKRODT NUCLEAR | 2mCi/ML | N018058 001 | |

GALLIUM DOTATATE GA-68

POWDER; INTRAVENOUS

NETSPOT

| | | | | |
|---|-------------|---------------|-------------|--------------|
| + | AAA USA INC | 2.1-5.5mCi/ML | N208547 001 | Jun 01, 2016 |
|---|-------------|---------------|-------------|--------------|

GANCICLOVIR

GEL; OPHTHALMIC

ZIRGAN

| | | | | |
|---|-----------------|-------|-------------|--------------|
| + | BAUSCH AND LOMB | 0.15% | N022211 001 | Sep 15, 2009 |
|---|-----------------|-------|-------------|--------------|

GANCICLOVIR

SOLUTION; INTRAVENOUS

GANCICLOVIR

| | | | | |
|---|-------------------------|----------------------|-------------|--------------|
| + | EXELA PHARMA SCS LLC | 500MG/250ML (2MG/ML) | N209347 001 | Feb 17, 2017 |
|---|-------------------------|----------------------|-------------|--------------|

GANCICLOVIR SODIUM

INJECTABLE; INJECTION

CYTOVENE

| | | | | |
|-----------|---------------|---------------------------|--------------------|--------------|
| <u>AP</u> | +! ROCHE PALO | <u>EQ 500MG BASE/VIAL</u> | <u>N019661 001</u> | Jun 23, 1989 |
|-----------|---------------|---------------------------|--------------------|--------------|

GANCICLOVIR

| | | | | |
|-----------|-------------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 500MG BASE/VIAL</u> | <u>A090658 001</u> | Jun 21, 2010 |
| <u>AP</u> | HAINAN POLY PHARM | <u>EQ 500MG BASE/VIAL</u> | <u>A204204 001</u> | Nov 08, 2018 |
| <u>AP</u> | LUITPOLD | <u>EQ 500MG BASE/VIAL</u> | <u>A202624 001</u> | Sep 18, 2013 |
| <u>AP</u> | MYLAN LABS LTD | <u>EQ 500MG BASE/VIAL</u> | <u>A204560 001</u> | Nov 17, 2017 |
| <u>AP</u> | PAR STERILE PRODUCTS | <u>EQ 500MG BASE/VIAL</u> | <u>A204950 001</u> | Dec 06, 2016 |

GANCICLOVIR SODIUM

| | | | | |
|-----------|-------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | PHARMASCIENCE INC | <u>EQ 500MG BASE/VIAL</u> | <u>A207645 001</u> | Dec 08, 2017 |
|-----------|-------------------|---------------------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

GANIRELIX ACETATE

INJECTABLE; INJECTION

GANIRELIX ACETATE

| | | | | | |
|-----------|---|--------------------|-----------------------------|--------------------|--------------|
| <u>AP</u> | + | ORGANON USA INC | <u>EQ 250MCG BASE/0.5ML</u> | <u>N021057 001</u> | Jul 29, 1999 |
| <u>AP</u> | | SUN PHARM INDS LTD | <u>EQ 250MCG BASE/0.5ML</u> | <u>A204246 001</u> | Nov 30, 2018 |

GATIFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

GATIFLOXACIN

| | | | | | |
|----------------|---|-------------------|-------------|--------------------|--------------|
| <u>AT</u> | | HI-TECH PHARMA CO | <u>0.5%</u> | <u>A203189 001</u> | Sep 03, 2014 |
| <u>AT</u> | | LUPIN LTD | <u>0.5%</u> | <u>A202653 001</u> | Aug 28, 2013 |
| <u>AT</u> | | MYLAN PHARMS INC | <u>0.5%</u> | <u>A206446 001</u> | Jun 08, 2018 |
| <u>AT</u> | | SANDOZ INC | <u>0.5%</u> | <u>A204227 001</u> | Jul 11, 2016 |
| <u>ZYMAXID</u> | | | | | |
| <u>AT</u> | + | ALLERGAN | <u>0.5%</u> | <u>N022548 001</u> | May 18, 2010 |
| | | ZYMAR | | | |
| | + | ALLERGAN | 0.3% | N021493 001 | Mar 28, 2003 |

GEFITINIB

TABLET; ORAL

IRESSA

| | | | | |
|---|--------------------|-------|-------------|--------------|
| + | ASTRAZENECA PHARMS | 250MG | N206995 001 | Jul 13, 2015 |
|---|--------------------|-------|-------------|--------------|

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

| | | | | | |
|-----------|---|----------------------|-------------------------------|--------------------|--------------|
| <u>AP</u> | | ACCORD HLTHCARE | <u>EQ 200MG BASE/VIAL</u> | <u>A091594 001</u> | Jul 25, 2011 |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A091594 002</u> | Jul 25, 2011 |
| <u>AP</u> | | | <u>EQ 2GM BASE/VIAL</u> | <u>A091594 003</u> | Jul 25, 2011 |
| <u>AP</u> | | ACTAVIS INC | <u>200MG/5.26ML (38MG/ML)</u> | <u>A204549 001</u> | Apr 11, 2016 |
| <u>AP</u> | | | <u>1GM/26.3ML (38MG/ML)</u> | <u>A204549 002</u> | Apr 11, 2016 |
| <u>AP</u> | | | <u>2GM/52.6ML (38MG/ML)</u> | <u>A204549 003</u> | Apr 11, 2016 |
| <u>AP</u> | | ACTAVIS TOTOWA | <u>EQ 200MG BASE/VIAL</u> | <u>A079160 001</u> | Jul 25, 2011 |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A079160 002</u> | Jul 25, 2011 |
| <u>AP</u> | | | <u>EQ 2GM BASE/VIAL</u> | <u>A079160 003</u> | Jul 28, 2016 |
| <u>AP</u> | | APOTEX INC | <u>200MG/5.26ML (38MG/ML)</u> | <u>A206776 001</u> | May 23, 2017 |
| <u>AP</u> | | | <u>1GM/26.3ML (38MG/ML)</u> | <u>A206776 002</u> | May 23, 2017 |
| <u>AP</u> | | | <u>2GM/52.6ML (38MG/ML)</u> | <u>A206776 003</u> | May 23, 2017 |
| <u>AP</u> | | CIPLA | <u>EQ 200MG BASE/VIAL</u> | <u>A078759 001</u> | Jul 25, 2011 |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A078759 002</u> | Jul 25, 2011 |
| <u>AP</u> | | DR REDDYS LABS LTD | <u>EQ 200MG BASE/VIAL</u> | <u>A091365 001</u> | Jul 25, 2011 |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A091365 002</u> | Jul 25, 2011 |
| <u>AP</u> | | | <u>EQ 2GM BASE/VIAL</u> | <u>A202997 001</u> | May 07, 2013 |
| <u>AP</u> | | EMCURE PHARMS LTD | <u>EQ 200MG BASE/VIAL</u> | <u>A202063 001</u> | Sep 11, 2012 |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A202063 002</u> | Sep 11, 2012 |
| <u>AP</u> | | FRESENIUS KABI | <u>EQ 200MG BASE/VIAL</u> | <u>A090799 001</u> | Jul 25, 2011 |
| | | ONCOL | | | |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A090799 002</u> | Jul 25, 2011 |
| <u>AP</u> | | | <u>EQ 2GM BASE/VIAL</u> | <u>A090799 003</u> | May 16, 2011 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>EQ 2GM BASE/VIAL</u> | <u>A090242 003</u> | May 16, 2011 |
| <u>AP</u> | | GLAND PHARMA LTD | <u>EQ 200MG BASE/VIAL</u> | <u>A204520 001</u> | Jan 05, 2016 |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A204520 002</u> | Jan 05, 2016 |
| <u>AP</u> | | HOSPIRA | <u>EQ 200MG BASE/VIAL</u> | <u>A078339 001</u> | Jul 25, 2011 |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A078339 002</u> | Jul 25, 2011 |
| <u>AP</u> | + | HOSPIRA INC | <u>200MG/5.26ML (38MG/ML)</u> | <u>N200795 001</u> | Aug 04, 2011 |
| <u>AP</u> | + | | <u>1GM/26.3ML (38MG/ML)</u> | <u>N200795 002</u> | Aug 04, 2011 |
| <u>AP</u> | ! | | <u>EQ 2GM BASE/VIAL</u> | <u>A079183 001</u> | Nov 15, 2010 |
| <u>AP</u> | + | | <u>2GM/52.6ML (38MG/ML)</u> | <u>N200795 003</u> | Aug 04, 2011 |
| <u>AP</u> | | JIANGSU HANSON PHARM | <u>EQ 200MG BASE/VIAL</u> | <u>A202485 001</u> | May 07, 2013 |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A202485 002</u> | May 07, 2013 |
| <u>AP</u> | | LUITPOLD | <u>EQ 200MG BASE/VIAL</u> | <u>A202031 001</u> | May 07, 2013 |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A202031 002</u> | May 07, 2013 |
| <u>AP</u> | | MYLAN LABS LTD | <u>EQ 200MG BASE/VIAL</u> | <u>A200145 001</u> | Jul 25, 2011 |
| <u>AP</u> | | | <u>200MG/5.26ML (38MG/ML)</u> | <u>A205242 001</u> | Dec 06, 2017 |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A200145 002</u> | Jul 25, 2011 |
| <u>AP</u> | | | <u>1GM/26.3ML (38MG/ML)</u> | <u>A205242 002</u> | Dec 06, 2017 |
| <u>AP</u> | | | <u>EQ 2GM BASE/VIAL</u> | <u>A200145 003</u> | Jul 25, 2011 |
| <u>AP</u> | | | <u>2GM/52.6ML (38MG/ML)</u> | <u>A205242 003</u> | Dec 06, 2017 |
| <u>AP</u> | | SAGENT PHARMS | <u>200MG/5.26ML (38MG/ML)</u> | <u>A209077 001</u> | Jul 20, 2018 |
| <u>AP</u> | | | <u>1GM/26.3ML (38MG/ML)</u> | <u>A209077 002</u> | Jul 20, 2018 |
| <u>AP</u> | | | <u>2GM/52.6ML (38MG/ML)</u> | <u>A209077 003</u> | Jul 20, 2018 |
| <u>AP</u> | | SUN PHARMA GLOBAL | <u>EQ 200MG BASE/VIAL</u> | <u>A078433 001</u> | Jul 25, 2011 |
| <u>AP</u> | | | <u>EQ 1GM BASE/VIAL</u> | <u>A078433 002</u> | Jul 25, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

| | | | | | |
|-----------|-------------|---------------------------|----------------|------------|--------------|
| <u>AP</u> | TEVA PHARMS | <u>EQ 200MG BASE/VIAL</u> | <u>A077983</u> | <u>002</u> | Jan 25, 2011 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A077983</u> | <u>001</u> | Jan 25, 2011 |

GEMZAR

| | | | | | |
|-----------|----------|---------------------------|----------------|------------|--------------|
| <u>AP</u> | +! LILLY | <u>EQ 200MG BASE/VIAL</u> | <u>N020509</u> | <u>001</u> | May 15, 1996 |
| <u>AP</u> | +! | <u>EQ 1GM BASE/VIAL</u> | <u>N020509</u> | <u>002</u> | May 15, 1996 |

SOLUTION; INTRAVENOUS

GEMCITABINE HYDROCHLORIDE

| | | | | | |
|----|-----------------|-----------------------|---------|-----|--------------|
| +! | ACCORD HLTHCARE | 1GM/10ML (100MG/ML) | N209604 | 002 | Aug 03, 2017 |
| +! | | 1.5GM/15ML (100MG/ML) | N209604 | 003 | Aug 03, 2017 |
| +! | | 2GM/20ML (100MG/ML) | N209604 | 004 | Aug 03, 2017 |
| +! | | 200MG/2ML (100MG/ML) | N209604 | 001 | Aug 03, 2017 |

INFUGEM

| | | | | | |
|----|--------------------|----------------------------------------|---------|-----|--------------|
| +! | SUN PHARM INDS LTD | EQ 1200MG BASE/120ML (EQ 10MG BASE/ML) | N208313 | 001 | Jul 16, 2018 |
| +! | | EQ 1300MG BASE/130ML (EQ 10MG BASE/ML) | N208313 | 002 | Jul 16, 2018 |
| +! | | EQ 1400MG BASE/140ML (EQ 10MG BASE/ML) | N208313 | 003 | Jul 16, 2018 |
| +! | | EQ 1500MG BASE/150ML (EQ 10MG BASE/ML) | N208313 | 004 | Jul 16, 2018 |
| +! | | EQ 1600MG BASE/160ML (EQ 10MG BASE/ML) | N208313 | 005 | Jul 16, 2018 |
| +! | | EQ 1700MG BASE/170ML (EQ 10MG BASE/ML) | N208313 | 006 | Jul 16, 2018 |
| +! | | EQ 1800MG BASE/180ML (EQ 10MG BASE/ML) | N208313 | 007 | Jul 16, 2018 |
| +! | | EQ 1900MG BASE/190ML (EQ 10MG BASE/ML) | N208313 | 008 | Jul 16, 2018 |
| +! | | EQ 2000MG BASE/200ML (EQ 10MG BASE/ML) | N208313 | 009 | Jul 16, 2018 |
| +! | | EQ 2200MG BASE/220ML (EQ 10MG BASE/ML) | N208313 | 010 | Jul 16, 2018 |

GEMFIBROZIL

TABLET; ORAL

GEMFIBROZIL

| | | | | | |
|-----------|----------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | APOTEX | <u>600MG</u> | <u>A075034</u> | <u>001</u> | Jul 20, 1998 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>600MG</u> | <u>A202726</u> | <u>001</u> | Sep 16, 2015 |
| <u>AB</u> | CADILA PHARMS LTD | <u>600MG</u> | <u>A203266</u> | <u>001</u> | Jun 17, 2016 |
| <u>AB</u> | CARIBE HOLDINGS | <u>600MG</u> | <u>A078012</u> | <u>001</u> | Mar 26, 2007 |
| <u>AB</u> | CHARTWELL MOLECULES | <u>600MG</u> | <u>A074270</u> | <u>001</u> | Sep 27, 1993 |
| <u>AB</u> | HIKMA PHARMS | <u>600MG</u> | <u>A078599</u> | <u>001</u> | Aug 16, 2010 |
| <u>AB</u> | IMPAX PHARMS | <u>600MG</u> | <u>A078207</u> | <u>001</u> | Jun 01, 2007 |
| <u>AB</u> | INVAGEN PHARMS | <u>600MG</u> | <u>A077836</u> | <u>001</u> | Jul 27, 2006 |
| <u>AB</u> | NORTHSTAR HLTHCARE | <u>600MG</u> | <u>A079072</u> | <u>001</u> | Sep 13, 2010 |
| <u>AB</u> | SUN PHARM INDS INC | <u>600MG</u> | <u>A079239</u> | <u>001</u> | Dec 29, 2008 |
| <u>AB</u> | TEVA | <u>600MG</u> | <u>A074256</u> | <u>001</u> | Oct 31, 1993 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>600MG</u> | <u>A204189</u> | <u>001</u> | Aug 28, 2018 |

LOPID

| | | | | | |
|-----------|------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | +! PFIZER PHARMS | <u>600MG</u> | <u>N018422</u> | <u>003</u> | Nov 20, 1986 |
|-----------|------------------|--------------|----------------|------------|--------------|

GEMIFLOXACIN MESYLATE

TABLET; ORAL

FACTIVE

| | | | | | |
|-----------|----------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | +! LG CHEM LTD | <u>EQ 320MG BASE</u> | <u>N021158</u> | <u>001</u> | Apr 04, 2003 |
|-----------|----------------|----------------------|----------------|------------|--------------|

GEMIFLOXACIN MESYLATE

| | | | | | |
|-----------|-----------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | ORCHID HLTHCARE | <u>EQ 320MG BASE</u> | <u>A090466</u> | <u>001</u> | Jun 15, 2015 |
|-----------|-----------------|----------------------|----------------|------------|--------------|

GENTAMICIN SULFATE

CREAM; TOPICAL

GENTAMICIN SULFATE

| | | | | | |
|-----------|--------------------|---------------------|----------------|------------|--------------|
| <u>AT</u> | G AND W LABS INC | <u>EQ 0.1% BASE</u> | <u>A064056</u> | <u>001</u> | Apr 29, 1994 |
| <u>AT</u> | ! PERRIGO NEW YORK | <u>EQ 0.1% BASE</u> | <u>A062307</u> | <u>001</u> | |

INJECTABLE; INJECTION

GENTAMICIN SULFATE

| | | | | | |
|-----------|--------------------|------------------------|----------------|------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 10MG BASE/ML</u> | <u>A062366</u> | <u>002</u> | Feb 06, 1986 |
| <u>AP</u> | ! | <u>EQ 40MG BASE/ML</u> | <u>A062366</u> | <u>001</u> | Aug 04, 1983 |
| <u>AP</u> | HOSPIRA | <u>EQ 10MG BASE/ML</u> | <u>A062420</u> | <u>001</u> | Aug 15, 1983 |
| <u>AP</u> | | <u>EQ 10MG BASE/ML</u> | <u>A062612</u> | <u>004</u> | Feb 20, 1986 |
| <u>AP</u> | | <u>EQ 40MG BASE/ML</u> | <u>A062420</u> | <u>002</u> | Aug 15, 1983 |

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | | |
|-----------|-----------------|----------------------------|----------------|------------|--------------|
| <u>AP</u> | BAXTER HLTHCARE | <u>EQ 1.2MG BASE/ML</u> | <u>A062373</u> | <u>007</u> | Sep 07, 1982 |
| <u>AP</u> | | <u>EQ 1.6MG BASE/ML</u> | <u>A062373</u> | <u>008</u> | Sep 07, 1982 |
| <u>AP</u> | | <u>EQ 80MG BASE/100ML</u> | <u>A062373</u> | <u>002</u> | Sep 07, 1982 |
| <u>AP</u> | | <u>EQ 100MG BASE/100ML</u> | <u>A062373</u> | <u>005</u> | Sep 07, 1982 |
| <u>AP</u> | HOSPIRA | <u>EQ 1.2MG BASE/ML</u> | <u>A062414</u> | <u>001</u> | Aug 15, 1983 |
| <u>AP</u> | | <u>EQ 1.6MG BASE/ML</u> | <u>A062414</u> | <u>003</u> | Aug 15, 1983 |
| <u>AP</u> | | <u>EQ 80MG BASE/100ML</u> | <u>A062414</u> | <u>008</u> | Aug 15, 1983 |
| <u>AP</u> | | <u>EQ 100MG BASE/100ML</u> | <u>A062414</u> | <u>010</u> | Aug 15, 1983 |

PRESCRIPTION DRUG PRODUCT LIST

GENTAMICIN SULFATE

INJECTABLE; INJECTION

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | | |
|---|-----------------|---------------------|---------|-----|--------------|
| ! | BAXTER HLTHCARE | EQ 2MG BASE/ML | A062373 | 009 | Sep 07, 1982 |
| ! | | EQ 120MG BASE/100ML | A062373 | 006 | Sep 07, 1982 |

OINTMENT; OPHTHALMIC

GENTAMICIN SULFATE

| | | | | | | |
|-----------|---|------------|---------------------|----------------|------------|--------------|
| AT | ! | AKORN | <u>EQ 0.3% BASE</u> | <u>A064093</u> | <u>001</u> | Aug 31, 1995 |
| AT | | PERRIGO CO | <u>EQ 0.3% BASE</u> | <u>A065024</u> | <u>001</u> | Jul 30, 2004 |

TENNESSEE

OINTMENT; TOPICAL

GENTAMICIN SULFATE

| | | | | | | |
|-----------|---|---------------------|---------------------|----------------|------------|--------------|
| AT | | FOUGERA PHARMS INC | <u>EQ 0.1% BASE</u> | <u>A062533</u> | <u>001</u> | Oct 05, 1984 |
| AT | | G AND W LABS INC | <u>EQ 0.1% BASE</u> | <u>A064054</u> | <u>001</u> | Apr 29, 1994 |
| AT | ! | PERRIGO NEW YORK | <u>EQ 0.1% BASE</u> | <u>A062351</u> | <u>001</u> | Feb 18, 1982 |
| AT | | TARO | <u>EQ 0.1% BASE</u> | <u>A062477</u> | <u>001</u> | Dec 23, 1983 |
| AT | | TELIGENT PHARMA INC | <u>EQ 0.1% BASE</u> | <u>A209233</u> | <u>001</u> | Dec 31, 2018 |

SOLUTION/DROPS; OPHTHALMIC

GENOPTIC

| | | | | | | |
|-----------|---|----------|---------------------|----------------|------------|--------------|
| AT | ! | ALLERGAN | <u>EQ 0.3% BASE</u> | <u>A062452</u> | <u>001</u> | Oct 10, 1984 |
|-----------|---|----------|---------------------|----------------|------------|--------------|

GENTAK

| | | | | | | |
|-----------|--|-------|---------------------|----------------|------------|--------------|
| AT | | AKORN | <u>EQ 0.3% BASE</u> | <u>A064163</u> | <u>001</u> | Oct 12, 2001 |
|-----------|--|-------|---------------------|----------------|------------|--------------|

GENTAMICIN SULFATE

| | | | | | | |
|-----------|--|-----------------|---------------------|----------------|------------|--------------|
| AT | | AKORN | <u>EQ 0.3% BASE</u> | <u>A062635</u> | <u>001</u> | Jan 08, 1987 |
| AT | | BAUSCH AND LOMB | <u>EQ 0.3% BASE</u> | <u>A064048</u> | <u>001</u> | May 11, 1994 |
| AT | | PERRIGO CO | <u>EQ 0.3% BASE</u> | <u>A065121</u> | <u>001</u> | Jan 30, 2004 |
| AT | | TENNESSEE | | | | |
| AT | | SANDOZ INC | <u>EQ 0.3% BASE</u> | <u>A062196</u> | <u>001</u> | |

GENTAMICIN SULFATE; PREDNISOLONE ACETATE

OINTMENT; OPHTHALMIC

PRED-G

| | | | | | | |
|---|---|----------|--------------------|---------|-----|--------------|
| + | ! | ALLERGAN | EQ 0.3% BASE; 0.6% | N050612 | 001 | Dec 01, 1989 |
|---|---|----------|--------------------|---------|-----|--------------|

SUSPENSION/DROPS; OPHTHALMIC

PRED-G

| | | | | | | |
|---|---|----------|------------------|---------|-----|--------------|
| + | ! | ALLERGAN | EQ 0.3% BASE; 1% | N050586 | 001 | Jun 10, 1988 |
|---|---|----------|------------------|---------|-----|--------------|

GILTERITINIB FUMARATE

TABLET; ORAL

XOSPATA

| | | | | | | |
|---|---|----------|--------------|---------|-----|--------------|
| + | ! | ASTELLAS | EQ 40MG BASE | N211349 | 001 | Nov 28, 2018 |
|---|---|----------|--------------|---------|-----|--------------|

GLASDEGIB

TABLET; ORAL

DAURISMO

| | | | | | | |
|---|---|------------|-------|---------|-----|--------------|
| + | | PFIZER INC | 25MG | N210656 | 001 | Nov 21, 2018 |
| + | ! | | 100MG | N210656 | 002 | Nov 21, 2018 |

GLATIRAMER ACETATE

INJECTABLE; SUBCUTANEOUS

COPAXONE

| | | | | | | | |
|-----------|---|---|-----------------|----------------|----------------|------------|--------------|
| AP | + | ! | TEVA PHARMS USA | <u>20MG/ML</u> | <u>N020622</u> | <u>002</u> | Feb 12, 2002 |
| AP | + | ! | | <u>40MG/ML</u> | <u>N020622</u> | <u>003</u> | Jan 28, 2014 |

GLATIRAMER ACETATE

| | | | | | | |
|-----------|--|------------------|----------------|----------------|------------|--------------|
| AP | | MYLAN PHARMS INC | <u>20MG/ML</u> | <u>A091646</u> | <u>001</u> | Oct 03, 2017 |
| AP | | | <u>40MG/ML</u> | <u>A206936</u> | <u>001</u> | Oct 03, 2017 |

GLATOPIA

| | | | | | | |
|-----------|--|------------|----------------|----------------|------------|--------------|
| AP | | SANDOZ INC | <u>20MG/ML</u> | <u>A090218</u> | <u>001</u> | Apr 16, 2015 |
| AP | | | <u>40MG/ML</u> | <u>A206921</u> | <u>001</u> | Feb 12, 2018 |

GLECAPREVIR; PIBRENTASVIR

TABLET; ORAL

MAVYRET

| | | | | | | |
|---|---|------------|-------------|---------|-----|--------------|
| + | ! | ABBVIE INC | 100MG; 40MG | N209394 | 001 | Aug 03, 2017 |
|---|---|------------|-------------|---------|-----|--------------|

GLIMEPIRIDE

TABLET; ORAL

AMARYL

| | | | | | | | |
|-----------|---|---|-------------------|------------|----------------|------------|--------------|
| AB | + | ! | SANOFI AVENTIS US | <u>1MG</u> | <u>N020496</u> | <u>001</u> | Nov 30, 1995 |
| AB | + | | | <u>2MG</u> | <u>N020496</u> | <u>002</u> | Nov 30, 1995 |
| AB | + | | | <u>4MG</u> | <u>N020496</u> | <u>003</u> | Nov 30, 1995 |

GLIMEPIRIDE

| | | | | | | |
|-----------|--|------------------|------------|----------------|------------|--------------|
| AB | | ACCORD HLTHCARE | <u>1MG</u> | <u>A078181</u> | <u>001</u> | Aug 23, 2007 |
| AB | | | <u>2MG</u> | <u>A078181</u> | <u>002</u> | Aug 23, 2007 |
| AB | | | <u>4MG</u> | <u>A078181</u> | <u>003</u> | Aug 23, 2007 |
| AB | | AUROBINDO PHARMA | <u>1MG</u> | <u>A202759</u> | <u>001</u> | Jun 29, 2012 |

PRESCRIPTION DRUG PRODUCT LISTGLIMEPIRIDE

TABLET; ORAL

GLIMEPIRIDE

LTD

| | | | | | |
|-----------|--------------------|------------|----------------|------------|--------------|
| <u>AB</u> | | <u>2MG</u> | <u>A202759</u> | <u>002</u> | Jun 29, 2012 |
| <u>AB</u> | | <u>4MG</u> | <u>A202759</u> | <u>003</u> | Jun 29, 2012 |
| <u>AB</u> | CARLSBAD | <u>1MG</u> | <u>A077911</u> | <u>001</u> | Sep 22, 2009 |
| <u>AB</u> | | <u>2MG</u> | <u>A077911</u> | <u>002</u> | Sep 22, 2009 |
| <u>AB</u> | | <u>4MG</u> | <u>A077911</u> | <u>003</u> | Sep 22, 2009 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>1MG</u> | <u>A077091</u> | <u>001</u> | Oct 06, 2005 |
| <u>AB</u> | | <u>2MG</u> | <u>A077091</u> | <u>002</u> | Oct 06, 2005 |
| <u>AB</u> | | <u>4MG</u> | <u>A077091</u> | <u>003</u> | Oct 06, 2005 |
| <u>AB</u> | INDOCO REMEDIES | <u>1MG</u> | <u>A202112</u> | <u>001</u> | Apr 17, 2013 |
| <u>AB</u> | | <u>2MG</u> | <u>A202112</u> | <u>002</u> | Apr 17, 2013 |
| <u>AB</u> | | <u>4MG</u> | <u>A202112</u> | <u>003</u> | Apr 17, 2013 |
| <u>AB</u> | INVAGEN PHARMS | <u>1MG</u> | <u>A077295</u> | <u>001</u> | Oct 06, 2005 |
| <u>AB</u> | | <u>2MG</u> | <u>A077295</u> | <u>002</u> | Oct 06, 2005 |
| <u>AB</u> | | <u>4MG</u> | <u>A077295</u> | <u>003</u> | Oct 06, 2005 |
| <u>AB</u> | MYLAN | <u>1MG</u> | <u>A077624</u> | <u>001</u> | Nov 28, 2005 |
| <u>AB</u> | | <u>2MG</u> | <u>A077624</u> | <u>002</u> | Nov 28, 2005 |
| <u>AB</u> | | <u>4MG</u> | <u>A077624</u> | <u>003</u> | Nov 28, 2005 |
| <u>AB</u> | PRINSTON INC | <u>1MG</u> | <u>A077370</u> | <u>001</u> | Dec 23, 2005 |
| <u>AB</u> | | <u>2MG</u> | <u>A077370</u> | <u>002</u> | Dec 23, 2005 |
| <u>AB</u> | | <u>4MG</u> | <u>A077370</u> | <u>003</u> | Dec 23, 2005 |
| <u>AB</u> | | <u>8MG</u> | <u>A077370</u> | <u>004</u> | Dec 23, 2005 |
| <u>AB</u> | TEVA | <u>1MG</u> | <u>A076802</u> | <u>001</u> | Oct 06, 2005 |
| <u>AB</u> | | <u>2MG</u> | <u>A076802</u> | <u>002</u> | Oct 06, 2005 |
| <u>AB</u> | | <u>4MG</u> | <u>A076802</u> | <u>003</u> | Oct 06, 2005 |
| <u>AB</u> | VIVA HLTHCARE | <u>1MG</u> | <u>A091220</u> | <u>001</u> | Jun 29, 2012 |
| <u>AB</u> | | <u>2MG</u> | <u>A091220</u> | <u>002</u> | Jun 29, 2012 |
| <u>AB</u> | | <u>4MG</u> | <u>A091220</u> | <u>004</u> | Jun 29, 2012 |
| <u>AB</u> | | <u>8MG</u> | <u>A091220</u> | <u>006</u> | Jun 29, 2012 |
| | | 3MG | A091220 | 003 | Jun 29, 2012 |
| | | 6MG | A091220 | 005 | Jun 29, 2012 |

GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

DUETACT

| | | | | | | |
|-----------|----|-------------------|---------------------------------------------------|----------------|------------|--------------|
| <u>AB</u> | +! | TAKEDA PHARMS USA | <u>2MG;30MG</u> | <u>N021925</u> | <u>001</u> | Jul 28, 2006 |
| <u>AB</u> | + | | <u>4MG;30MG</u> | <u>N021925</u> | <u>002</u> | Jul 28, 2006 |
| | | | <u>PIOGLITAZONE HYDROCHLORIDE AND GLIMEPIRIDE</u> | | | |
| <u>AB</u> | | SANDOZ | <u>2MG;30MG</u> | <u>A201049</u> | <u>001</u> | Jan 04, 2013 |
| <u>AB</u> | | | <u>4MG;30MG</u> | <u>A201049</u> | <u>002</u> | Jan 04, 2013 |

GLIPIZIDE

TABLET; ORAL

GLIPIZIDE

| | | | | | | |
|-----------|----|--------------------|------------------|----------------|------------|--------------|
| <u>AB</u> | | ACCORD HLTHCARE | <u>5MG</u> | <u>A074550</u> | <u>001</u> | Sep 11, 1997 |
| <u>AB</u> | | | <u>10MG</u> | <u>A074550</u> | <u>002</u> | Sep 11, 1997 |
| <u>AB</u> | | ANI PHARMS INC | <u>5MG</u> | <u>A074497</u> | <u>001</u> | Aug 31, 1995 |
| <u>AB</u> | | | <u>10MG</u> | <u>A074497</u> | <u>002</u> | Aug 31, 1995 |
| <u>AB</u> | | APOTEX | <u>5MG</u> | <u>A075795</u> | <u>001</u> | Jun 13, 2001 |
| <u>AB</u> | | | <u>10MG</u> | <u>A075795</u> | <u>002</u> | Jun 13, 2001 |
| <u>AB</u> | | MYLAN | <u>5MG</u> | <u>A074226</u> | <u>001</u> | May 10, 1994 |
| <u>AB</u> | | | <u>10MG</u> | <u>A074226</u> | <u>002</u> | May 10, 1994 |
| <u>AB</u> | | SANDOZ | <u>5MG</u> | <u>A074305</u> | <u>001</u> | Apr 07, 1995 |
| <u>AB</u> | | | <u>10MG</u> | <u>A074305</u> | <u>002</u> | Apr 07, 1995 |
| <u>AB</u> | | SUN PHARM INDS INC | <u>5MG</u> | <u>A077820</u> | <u>001</u> | Jul 11, 2006 |
| <u>AB</u> | | | <u>10MG</u> | <u>A077820</u> | <u>002</u> | Jul 11, 2006 |
| <u>AB</u> | | WATSON LABS TEVA | <u>5MG</u> | <u>A074223</u> | <u>001</u> | Feb 27, 1995 |
| <u>AB</u> | | | <u>10MG</u> | <u>A074223</u> | <u>002</u> | Feb 27, 1995 |
| | | | <u>GLUCOTROL</u> | | | |
| <u>AB</u> | + | PFIZER | <u>5MG</u> | <u>N017783</u> | <u>001</u> | May 08, 1984 |
| <u>AB</u> | +! | | <u>10MG</u> | <u>N017783</u> | <u>002</u> | May 08, 1984 |

TABLET, EXTENDED RELEASE; ORAL

GLIPIZIDE

| | | | | | | |
|-----------|--|----------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>2.5MG</u> | <u>A206928</u> | <u>001</u> | May 12, 2017 |
| <u>AB</u> | | | <u>5MG</u> | <u>A206928</u> | <u>002</u> | May 12, 2017 |
| <u>AB</u> | | | <u>10MG</u> | <u>A206928</u> | <u>003</u> | May 12, 2017 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>2.5MG</u> | <u>A202298</u> | <u>001</u> | May 19, 2015 |
| <u>AB</u> | | | <u>5MG</u> | <u>A202298</u> | <u>002</u> | May 19, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A202298</u> | <u>003</u> | May 19, 2015 |
| <u>AB</u> | | PAR PHARM | <u>5MG</u> | <u>A076159</u> | <u>002</u> | Sep 20, 2013 |

PRESCRIPTION DRUG PRODUCT LIST

GLIPIZIDE

TABLET, EXTENDED RELEASE;ORAL

GLIPIZIDE

| | | | | | |
|---------------------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | <u>10MG</u> | <u>A076159</u> | <u>001</u> | Sep 20, 2013 |
| <u>AB</u> | UNIQUE PHARM LABS | <u>2.5MG</u> | <u>A204720</u> | <u>001</u> | Dec 29, 2016 |
| <u>AB</u> | | <u>5MG</u> | <u>A204720</u> | <u>002</u> | Dec 29, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A204720</u> | <u>003</u> | Dec 29, 2016 |
| <u>AB</u> | WATSON LABS | <u>2.5MG</u> | <u>A076467</u> | <u>003</u> | Mar 27, 2006 |
| <u>AB</u> | | <u>5MG</u> | <u>A076467</u> | <u>001</u> | Sep 08, 2003 |
| <u>AB</u> | | <u>10MG</u> | <u>A076467</u> | <u>002</u> | Nov 07, 2003 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>2.5MG</u> | <u>A203499</u> | <u>001</u> | Jul 16, 2018 |
| <u>AB</u> | | <u>5MG</u> | <u>A203499</u> | <u>002</u> | Jul 16, 2018 |
| <u>AB</u> | | <u>10MG</u> | <u>A203499</u> | <u>003</u> | Jul 16, 2018 |
| <u>GLUCOTROL XL</u> | | | | | |
| <u>AB</u> | + PFIZER | <u>2.5MG</u> | <u>N020329</u> | <u>003</u> | Aug 10, 1999 |
| <u>AB</u> | + | <u>5MG</u> | <u>N020329</u> | <u>001</u> | Apr 26, 1994 |
| <u>AB</u> | +! | <u>10MG</u> | <u>N020329</u> | <u>002</u> | Apr 26, 1994 |

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET;ORAL

GLIPIZIDE AND METFORMIN HYDROCHLORIDE

| | | | | | |
|-----------|-------------------------|--------------------|----------------|------------|--------------|
| <u>AB</u> | EPIC PHARMA LLC | <u>2.5MG;250MG</u> | <u>A077507</u> | <u>001</u> | Oct 27, 2005 |
| <u>AB</u> | | <u>2.5MG;500MG</u> | <u>A077507</u> | <u>002</u> | Oct 27, 2005 |
| <u>AB</u> | | <u>5MG;500MG</u> | <u>A077507</u> | <u>003</u> | Oct 27, 2005 |
| <u>AB</u> | HERITAGE PHARMS INC | <u>2.5MG;250MG</u> | <u>A078728</u> | <u>001</u> | Jun 23, 2010 |
| <u>AB</u> | | <u>2.5MG;500MG</u> | <u>A078728</u> | <u>002</u> | Jun 23, 2010 |
| <u>AB</u> | | <u>5MG;500MG</u> | <u>A078728</u> | <u>003</u> | Jun 23, 2010 |
| <u>AB</u> | MYLAN | <u>2.5MG;250MG</u> | <u>A078083</u> | <u>001</u> | Apr 12, 2007 |
| <u>AB</u> | | <u>2.5MG;500MG</u> | <u>A078083</u> | <u>002</u> | Apr 12, 2007 |
| <u>AB</u> | | <u>5MG;500MG</u> | <u>A078083</u> | <u>003</u> | Apr 12, 2007 |
| <u>AB</u> | SUN PHARM INDS INC | <u>2.5MG;250MG</u> | <u>A077620</u> | <u>001</u> | Jan 11, 2008 |
| <u>AB</u> | | <u>2.5MG;500MG</u> | <u>A077620</u> | <u>002</u> | Jan 11, 2008 |
| <u>AB</u> | | <u>5MG;500MG</u> | <u>A077620</u> | <u>003</u> | Jan 11, 2008 |
| <u>AB</u> | TEVA PHARMS | <u>2.5MG;250MG</u> | <u>A077270</u> | <u>001</u> | Oct 28, 2005 |
| <u>AB</u> | | <u>2.5MG;500MG</u> | <u>A077270</u> | <u>002</u> | Oct 28, 2005 |
| <u>AB</u> | ! | <u>5MG;500MG</u> | <u>A077270</u> | <u>003</u> | Oct 28, 2005 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>2.5MG;250MG</u> | <u>A078905</u> | <u>001</u> | Jan 31, 2011 |
| <u>AB</u> | | <u>2.5MG;500MG</u> | <u>A078905</u> | <u>002</u> | Jan 31, 2011 |
| <u>AB</u> | | <u>5MG;500MG</u> | <u>A078905</u> | <u>003</u> | Jan 31, 2011 |

GLUCAGON

INJECTABLE; INJECTION

GLUCAGON

+! LILLY 1MG/VIAL N020928 001 Sep 11, 1998

GLUCAGON HYDROCHLORIDE

INJECTABLE; INJECTION

GLUCAGEN

+! NOVO NORDISK EQ 1MG BASE/VIAL N020918 001 Jun 22, 1998

POWDER; INTRAMUSCULAR, INTRAVENOUS

GLUCAGON

+! FRESENIUS KABI USA EQ 1MG BASE/VIAL N201849 001 May 08, 2015

GLYBURIDE

TABLET;ORAL

GLYBURIDE (MICRONIZED)

| | | | | | |
|----------------|---------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | DAVA PHARMS INC | <u>1.5MG</u> | <u>A074591</u> | <u>001</u> | Dec 22, 1997 |
| <u>AB</u> | | <u>3MG</u> | <u>A074591</u> | <u>002</u> | Dec 22, 1997 |
| <u>AB</u> | | <u>4.5MG</u> | <u>A074591</u> | <u>003</u> | Dec 22, 1997 |
| <u>AB</u> | | <u>6MG</u> | <u>A074591</u> | <u>004</u> | Dec 22, 1997 |
| <u>AB</u> | HIKMA | <u>1.5MG</u> | <u>A075890</u> | <u>001</u> | Jul 31, 2003 |
| <u>AB</u> | | <u>3MG</u> | <u>A075890</u> | <u>002</u> | Jul 31, 2003 |
| <u>AB</u> | | <u>6MG</u> | <u>A075890</u> | <u>003</u> | Jul 31, 2003 |
| <u>AB</u> | MYLAN | <u>1.5MG</u> | <u>A074792</u> | <u>001</u> | Jun 26, 1998 |
| <u>AB</u> | | <u>3MG</u> | <u>A074792</u> | <u>002</u> | Jun 26, 1998 |
| <u>AB</u> | | <u>6MG</u> | <u>A074792</u> | <u>003</u> | Aug 17, 1999 |
| <u>AB</u> | TEVA | <u>1.5MG</u> | <u>A074686</u> | <u>001</u> | Apr 20, 1999 |
| <u>AB</u> | | <u>3MG</u> | <u>A074686</u> | <u>002</u> | Apr 20, 1999 |
| <u>AB</u> | | <u>4.5MG</u> | <u>A074686</u> | <u>003</u> | Apr 20, 1999 |
| <u>AB</u> | | <u>6MG</u> | <u>A074686</u> | <u>004</u> | Apr 20, 1999 |
| <u>GLYNASE</u> | | | | | |
| <u>AB</u> | + PHARMACIA AND UPJOHN | <u>1.5MG</u> | <u>N020051</u> | <u>001</u> | Mar 04, 1992 |

PRESCRIPTION DRUG PRODUCT LIST

GLYBURIDE

TABLET; ORAL

GLYNASE

| | | | | | | |
|-----------|----|--|------------|----------------|------------|--------------|
| <u>AB</u> | + | | <u>3MG</u> | <u>N020051</u> | <u>002</u> | Mar 04, 1992 |
| <u>AB</u> | +! | | <u>6MG</u> | <u>N020051</u> | <u>004</u> | Sep 24, 1993 |

GLYBURIDE

| | | | | | | |
|------------|---|-------------------------|---------------|----------------|------------|--------------|
| <u>AB1</u> | | AUROBINDO PHARMA | <u>1.25MG</u> | <u>A077537</u> | <u>001</u> | Oct 18, 2007 |
| <u>AB1</u> | | | <u>2.5MG</u> | <u>A077537</u> | <u>002</u> | Oct 18, 2007 |
| <u>AB1</u> | | | <u>5MG</u> | <u>A077537</u> | <u>003</u> | Oct 18, 2007 |
| <u>AB1</u> | | CADILA PHARMS LTD | <u>1.25MG</u> | <u>A203379</u> | <u>001</u> | Jan 04, 2019 |
| <u>AB1</u> | | | <u>2.5MG</u> | <u>A203379</u> | <u>002</u> | Jan 04, 2019 |
| <u>AB1</u> | | | <u>5MG</u> | <u>A203379</u> | <u>003</u> | Jan 04, 2019 |
| <u>AB1</u> | | EPIC PHARMA LLC | <u>1.25MG</u> | <u>A076257</u> | <u>001</u> | Jun 27, 2002 |
| <u>AB1</u> | | | <u>2.5MG</u> | <u>A076257</u> | <u>002</u> | Jun 27, 2002 |
| <u>AB1</u> | | | <u>5MG</u> | <u>A076257</u> | <u>003</u> | Jun 27, 2002 |
| <u>AB1</u> | | HERITAGE PHARMS INC | <u>1.25MG</u> | <u>A090937</u> | <u>001</u> | Feb 28, 2011 |
| <u>AB1</u> | | | <u>2.5MG</u> | <u>A090937</u> | <u>002</u> | Feb 28, 2011 |
| <u>AB1</u> | | | <u>5MG</u> | <u>A090937</u> | <u>003</u> | Feb 28, 2011 |
| <u>AB1</u> | | PHARMADAX INC | <u>1.25MG</u> | <u>A203581</u> | <u>001</u> | Apr 14, 2016 |
| <u>AB1</u> | | | <u>2.5MG</u> | <u>A203581</u> | <u>002</u> | Apr 14, 2016 |
| <u>AB1</u> | | | <u>5MG</u> | <u>A203581</u> | <u>003</u> | Apr 14, 2016 |
| <u>AB1</u> | | TEVA | <u>1.25MG</u> | <u>A074388</u> | <u>001</u> | Aug 29, 1995 |
| <u>AB1</u> | | | <u>2.5MG</u> | <u>A074388</u> | <u>002</u> | Aug 29, 1995 |
| <u>AB1</u> | ! | | <u>5MG</u> | <u>A074388</u> | <u>003</u> | Aug 29, 1995 |
| <u>AB1</u> | | ZYDUS PHARMS USA INC | <u>1.25mg</u> | <u>A206749</u> | <u>001</u> | May 10, 2016 |
| <u>AB1</u> | | | <u>2.5mg</u> | <u>A206749</u> | <u>002</u> | May 10, 2016 |
| <u>AB1</u> | | | <u>5MG</u> | <u>A206749</u> | <u>003</u> | May 10, 2016 |

DIABETA

| | | | | | | |
|------------|----|-------------------|---------------|----------------|------------|--------------|
| <u>AB2</u> | + | SANOFI AVENTIS US | <u>1.25MG</u> | <u>N017532</u> | <u>001</u> | May 01, 1984 |
| <u>AB2</u> | + | | <u>2.5MG</u> | <u>N017532</u> | <u>002</u> | May 01, 1984 |
| <u>AB2</u> | +! | | <u>5MG</u> | <u>N017532</u> | <u>003</u> | May 01, 1984 |

GLYBURIDE

| | | | | | | |
|------------|--|----------------|---------------|----------------|------------|--------------|
| <u>AB2</u> | | IMPAX LABS INC | <u>1.25MG</u> | <u>A206079</u> | <u>001</u> | Sep 30, 2015 |
| <u>AB2</u> | | | <u>2.5MG</u> | <u>A206079</u> | <u>002</u> | Sep 30, 2015 |
| <u>AB2</u> | | | <u>5MG</u> | <u>A206079</u> | <u>003</u> | Sep 30, 2015 |

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLYBURIDE AND METFORMIN HYDROCHLORIDE

| | | | | | | |
|-----------|---|-------------------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | | ACTAVIS ELIZABETH | <u>1.25MG; 250MG</u> | <u>A076716</u> | <u>001</u> | Jun 28, 2005 |
| <u>AB</u> | | | <u>2.5MG; 500MG</u> | <u>A076716</u> | <u>002</u> | Jun 28, 2005 |
| <u>AB</u> | | | <u>5MG; 500MG</u> | <u>A076716</u> | <u>003</u> | Jun 28, 2005 |
| <u>AB</u> | | AUROBINDO PHARMA | <u>1.25MG; 250MG</u> | <u>A077870</u> | <u>001</u> | Nov 14, 2007 |
| <u>AB</u> | ! | | <u>2.5MG; 500MG</u> | <u>A077870</u> | <u>002</u> | Nov 14, 2007 |
| <u>AB</u> | | | <u>5MG; 500MG</u> | <u>A077870</u> | <u>003</u> | Nov 14, 2007 |
| <u>AB</u> | | HERITAGE PHARMS INC | <u>1.25MG; 250MG</u> | <u>A079009</u> | <u>001</u> | Jun 03, 2009 |
| <u>AB</u> | | | <u>2.5MG; 500MG</u> | <u>A079009</u> | <u>002</u> | Jun 03, 2009 |
| <u>AB</u> | | | <u>5MG; 500MG</u> | <u>A079009</u> | <u>003</u> | Jun 03, 2009 |
| <u>AB</u> | | IMPAX LABS INC | <u>1.25MG; 250MG</u> | <u>A076345</u> | <u>001</u> | Feb 18, 2004 |
| <u>AB</u> | | | <u>2.5MG; 500MG</u> | <u>A076345</u> | <u>002</u> | Feb 18, 2004 |
| <u>AB</u> | | | <u>5MG; 500MG</u> | <u>A076345</u> | <u>003</u> | Feb 18, 2004 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>1.25MG; 250MG</u> | <u>A206748</u> | <u>001</u> | Feb 29, 2016 |
| <u>AB</u> | | | <u>2.5MG; 500MG</u> | <u>A206748</u> | <u>002</u> | Feb 29, 2016 |
| <u>AB</u> | | | <u>5MG; 500MG</u> | <u>A206748</u> | <u>003</u> | Feb 29, 2016 |

GLYCEROL PHENYLBUTYRATE

LIQUID; ORAL

RAVICTI

| | | | | | | |
|---|---|---------------------|----------|---------|-----|--------------|
| + | ! | HORIZON THERAPS INC | 1.1GM/ML | N203284 | 001 | Feb 01, 2013 |
|---|---|---------------------|----------|---------|-----|--------------|

GLYCINE

SOLUTION; IRRIGATION

AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER

| | | | | | | |
|-----------|---|---|-----------------|--------------------|----------------|------------|
| <u>AT</u> | + | ! | BAXTER HLTHCARE | <u>1.5GM/100ML</u> | <u>N017865</u> | <u>001</u> |
|-----------|---|---|-----------------|--------------------|----------------|------------|

GLYCINE 1.5% IN PLASTIC CONTAINER

| | | | | | | |
|-----------|--|--|-----------------|--------------------|----------------|------------|
| <u>AT</u> | | | B BRAUN | <u>1.5GM/100ML</u> | <u>N016784</u> | <u>001</u> |
| <u>AT</u> | | | ICU MEDICAL INC | <u>1.5GM/100ML</u> | <u>N018315</u> | <u>001</u> |

PRESCRIPTION DRUG PRODUCT LIST

GLYCOPYRROLATE

INJECTABLE; INJECTION

GLYCOPYRROLATE

| | | | | | |
|-----------|----------------------|-----------------|----------------|------------|--------------|
| <u>AP</u> | AMNEAL PHARMS CO | <u>0.2MG/ML</u> | <u>A208973</u> | <u>001</u> | Jun 15, 2017 |
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>0.2MG/ML</u> | <u>A210244</u> | <u>001</u> | Nov 28, 2018 |
| <u>AP</u> | FRESENIUS KABI USA | <u>0.2MG/ML</u> | <u>A209024</u> | <u>001</u> | Oct 31, 2018 |
| <u>AP</u> | | <u>0.2MG/ML</u> | <u>A209328</u> | <u>001</u> | Oct 27, 2017 |
| <u>AP</u> | ! HIKMA FARMACEUTICA | <u>0.2MG/ML</u> | <u>A090963</u> | <u>001</u> | Sep 21, 2011 |
| <u>AP</u> | LUITPOLD | <u>0.2MG/ML</u> | <u>A089335</u> | <u>001</u> | Jul 23, 1986 |
| <u>AP</u> | PIRAMAL CRITICAL | <u>0.2MG/ML</u> | <u>A210842</u> | <u>001</u> | Oct 25, 2018 |
| <u>AP</u> | PRINSTON INC | <u>0.2MG/ML</u> | <u>A210927</u> | <u>001</u> | Oct 31, 2018 |
| <u>AP</u> | SOMERSET THERAPS LLC | <u>0.2MG/ML</u> | <u>A207639</u> | <u>001</u> | Jun 23, 2017 |

POWDER; INHALATION

SEEBRI

| | | | | | |
|----|---------------------|-------------|---------|-----|--------------|
| +! | SUNOVION PHARMS INC | 15.6MCG/INH | N207923 | 001 | Oct 29, 2015 |
|----|---------------------|-------------|---------|-----|--------------|

SOLUTION; INHALATION

LONHALA MAGNAIR KIT

| | | | | | |
|----|---------------|----------|---------|-----|--------------|
| +! | SUNOVION RESP | 25MCG/ML | N208437 | 001 | Dec 05, 2017 |
|----|---------------|----------|---------|-----|--------------|

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

GLYRX-PF

| | | | | | |
|--|----------------------|---------------------|---------|-----|--------------|
| | EXELA PHARMA SCS LLC | 0.2MG/ML (0.2MG/ML) | N210997 | 001 | Jul 11, 2018 |
|--|----------------------|---------------------|---------|-----|--------------|

| | | | | | |
|--|--|----------------------|---------|-----|--------------|
| | | 0.4MG/2ML (0.2MG/ML) | N210997 | 002 | Jul 11, 2018 |
|--|--|----------------------|---------|-----|--------------|

SOLUTION; ORAL

CUVPOSA

| | | | | | |
|----|-------------|---------|---------|-----|--------------|
| +! | MERZ PHARMS | 1MG/5ML | N022571 | 001 | Jul 28, 2010 |
|----|-------------|---------|---------|-----|--------------|

TABLET; ORAL

GLYCOPYRROLATE

| | | | | | |
|-----------|---------------------|------------|----------------|------------|--------------|
| <u>AA</u> | AUROLIFE PHARMA LLC | <u>1MG</u> | <u>A202675</u> | <u>001</u> | Apr 15, 2013 |
| <u>AA</u> | BOSCOGEN | <u>1MG</u> | <u>A091182</u> | <u>001</u> | Feb 03, 2014 |
| <u>AA</u> | | <u>2MG</u> | <u>A091182</u> | <u>002</u> | Feb 03, 2014 |
| <u>AA</u> | DR REDDYS LABS LTD | <u>1MG</u> | <u>A040847</u> | <u>001</u> | Mar 21, 2008 |
| <u>AA</u> | | <u>2MG</u> | <u>A040847</u> | <u>002</u> | Mar 21, 2008 |
| <u>AA</u> | HERITAGE PHARMS INC | <u>1MG</u> | <u>A207201</u> | <u>001</u> | Jan 03, 2017 |
| <u>AA</u> | | <u>2MG</u> | <u>A207201</u> | <u>002</u> | Jan 03, 2017 |
| <u>AA</u> | LEADING PHARMA LLC | <u>1MG</u> | <u>A090195</u> | <u>001</u> | Sep 21, 2012 |
| <u>AA</u> | | <u>2MG</u> | <u>A090195</u> | <u>002</u> | Sep 21, 2012 |
| <u>AA</u> | NATCO PHARMA LTD | <u>1MG</u> | <u>A091413</u> | <u>001</u> | Jun 20, 2016 |
| <u>AA</u> | | <u>2MG</u> | <u>A091413</u> | <u>002</u> | Jun 20, 2016 |
| <u>AA</u> | ORIT LABS LLC | <u>1MG</u> | <u>A203657</u> | <u>001</u> | Nov 30, 2018 |
| <u>AA</u> | | <u>2MG</u> | <u>A203657</u> | <u>002</u> | Nov 30, 2018 |
| <u>AA</u> | OXFORD PHARMS | <u>1MG</u> | <u>A090020</u> | <u>001</u> | Oct 19, 2011 |
| <u>AA</u> | | <u>2MG</u> | <u>A090020</u> | <u>002</u> | Oct 19, 2011 |
| <u>AA</u> | ! PAR PHARM | <u>1MG</u> | <u>A040653</u> | <u>001</u> | Aug 31, 2006 |
| <u>AA</u> | ! | <u>2MG</u> | <u>A040653</u> | <u>002</u> | Aug 31, 2006 |
| <u>AA</u> | RISING PHARMS | <u>1MG</u> | <u>A040821</u> | <u>001</u> | Dec 29, 2008 |
| <u>AA</u> | | <u>2MG</u> | <u>A040821</u> | <u>002</u> | Dec 29, 2008 |
| <u>AA</u> | SUN PHARM INDS LTD | <u>1MG</u> | <u>A040844</u> | <u>001</u> | Aug 18, 2009 |
| <u>AA</u> | | <u>2MG</u> | <u>A040844</u> | <u>002</u> | Aug 18, 2009 |
| | NEXGEN PHARMA | 1.5MG | A091522 | 001 | Mar 12, 2012 |

GLYCOPYRROLATE ; INDACATEROL MALEATE

POWDER; INHALATION

UTIBRON

| | | | | | |
|----|---------------------|--------------------------|---------|-----|--------------|
| +! | SUNOVION PHARMS INC | 15.6MCG/INH; 27.5MCG/INH | N207930 | 001 | Oct 29, 2015 |
|----|---------------------|--------------------------|---------|-----|--------------|

GLYCOPYRRONIUM TOSYLATE

CLOTH; TOPICAL

QBREXZA

| | | | | | |
|----|-------------|--------------|---------|-----|--------------|
| +! | DERMIRA INC | EQ 2.4% BASE | N210361 | 001 | Jun 28, 2018 |
|----|-------------|--------------|---------|-----|--------------|

GONADOTROPIN, CHORIONIC

INJECTABLE; INJECTION

CHORIONIC GONADOTROPIN

| | | | | | |
|----------------|------------------------|--------------------------|----------------|------------|--|
| <u>AP</u> | +! FERRING | <u>10,000 UNITS/VIAL</u> | <u>N017016</u> | <u>007</u> | |
| <u>AP</u> | +! FRESENIUS KABI USA | <u>10,000 UNITS/VIAL</u> | <u>N017067</u> | <u>002</u> | |
| <u>PREGNYL</u> | | | | | |
| <u>AP</u> | +! ORGANON USA INC | <u>10,000 UNITS/VIAL</u> | <u>N017692</u> | <u>001</u> | |
| | CHORIONIC GONADOTROPIN | | | | |
| | +! FERRING | 5,000 UNITS/VIAL | N017016 | 006 | |

PRESCRIPTION DRUG PRODUCT LISTGOSERELIN ACETATE

IMPLANT; IMPLANTATION

ZOLADEX

+! TERSERA THERAPS LLC EQ 3.6MG BASE
+! EQ 10.8MG BASEN019726 001 Dec 29, 1989
N020578 001 Jan 11, 1996GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN

| | | | | | |
|-----------|-------------------|-----------------------------------------------------|----------------|------------|--------------|
| <u>AT</u> | AMRING PHARMS | <u>0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML</u> | <u>A065187</u> | <u>001</u> | Oct 28, 2005 |
| <u>AT</u> | ! BAUSCH AND LOMB | <u>0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML</u> | <u>A064047</u> | <u>001</u> | Jan 31, 1996 |

NEOSPORIN

| | | | | | |
|-----------|------------------|-----------------------------------------------------|----------------|------------|--|
| <u>AT</u> | ! MONARCH PHARMS | <u>0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML</u> | <u>A060582</u> | <u>001</u> | |
|-----------|------------------|-----------------------------------------------------|----------------|------------|--|

GRANISETRON

FILM, EXTENDED RELEASE; TRANSDERMAL

SANCUSO

+! KYOWA KIRIN 3.1MG/24HR

N022198 001 Sep 12, 2008

INJECTION, EXTENDED RELEASE; SUBCUTANEOUS

SUSTOL

+! HERON THERAPS INC 10MG/0.4ML (10MG/0.4ML)

N022445 001 Aug 09, 2016

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

GRANISETRON HYDROCHLORIDE

| | | | | | |
|-----------|----------------------|--------------------------------------------|----------------|------------|--------------|
| <u>AP</u> | AKORN INC | <u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u> | <u>A079119</u> | <u>001</u> | Sep 10, 2009 |
| <u>AP</u> | | <u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u> | <u>A079078</u> | <u>001</u> | Sep 14, 2009 |
| <u>AP</u> | | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A079078</u> | <u>002</u> | Sep 14, 2009 |
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u> | <u>A204238</u> | <u>001</u> | Jul 06, 2016 |
| <u>AP</u> | | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A204238</u> | <u>002</u> | Jul 06, 2016 |
| <u>AP</u> | BLUEPHARMA | <u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u> | <u>A078863</u> | <u>001</u> | Jun 30, 2008 |
| <u>AP</u> | | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A078880</u> | <u>001</u> | Jun 30, 2008 |
| <u>AP</u> | CIPLA LTD | <u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u> | <u>A078262</u> | <u>001</u> | Dec 31, 2007 |
| <u>AP</u> | | <u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u> | <u>A078258</u> | <u>001</u> | Jun 30, 2008 |
| <u>AP</u> | | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A078258</u> | <u>002</u> | Jun 30, 2008 |
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u> | <u>A078522</u> | <u>001</u> | Dec 31, 2007 |
| <u>AP</u> | | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A078090</u> | <u>001</u> | Jun 30, 2008 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u> | <u>A078629</u> | <u>001</u> | Dec 23, 2009 |
| <u>AP</u> | | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A078629</u> | <u>002</u> | Dec 23, 2009 |
| <u>AP</u> | LUITPOLD | <u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u> | <u>A091274</u> | <u>001</u> | Sep 22, 2010 |
| <u>AP</u> | MYLAN ASI | <u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u> | <u>A091136</u> | <u>001</u> | Apr 09, 2010 |
| <u>AP</u> | | <u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u> | <u>A091136</u> | <u>002</u> | Apr 09, 2010 |
| <u>AP</u> | | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A091137</u> | <u>002</u> | Apr 09, 2010 |
| <u>AP</u> | MYLAN LABS LTD | <u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u> | <u>A203454</u> | <u>001</u> | Apr 04, 2017 |
| <u>AP</u> | | <u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u> | <u>A203454</u> | <u>002</u> | Apr 04, 2017 |
| <u>AP</u> | | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A203453</u> | <u>001</u> | Jan 31, 2017 |
| <u>AP</u> | SANDOZ INC | <u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u> | <u>A078534</u> | <u>001</u> | Apr 30, 2009 |
| <u>AP</u> | | <u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u> | <u>A078531</u> | <u>001</u> | Apr 30, 2009 |
| <u>AP</u> | | <u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u> | <u>A078835</u> | <u>001</u> | Jun 30, 2008 |
| <u>AP</u> | | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A078531</u> | <u>002</u> | Apr 30, 2009 |
| <u>AP</u> | | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A078835</u> | <u>002</u> | Jun 30, 2008 |
| <u>AP</u> | ! TEVA PHARMS USA | <u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u> | <u>A078392</u> | <u>001</u> | Dec 31, 2007 |
| <u>AP</u> | ! | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A077297</u> | <u>001</u> | Jun 30, 2008 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u> | <u>A077913</u> | <u>001</u> | Jun 26, 2008 |
| <u>AP</u> | | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A077186</u> | <u>001</u> | Jun 30, 2008 |
| <u>AP</u> | | <u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u> | <u>A077187</u> | <u>001</u> | Jun 30, 2008 |
| <u>AP</u> | | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A077177</u> | <u>001</u> | Dec 31, 2007 |
| <u>AP</u> | WOCKHARDT USA | <u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u> | <u>A078566</u> | <u>001</u> | Feb 29, 2008 |
| <u>AP</u> | | <u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u> | <u>A078564</u> | <u>001</u> | Jun 30, 2008 |
| <u>AP</u> | | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A078565</u> | <u>001</u> | Jun 30, 2008 |

GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE

| | | | | | |
|-----------|----------------------|----------------------------------------|----------------|------------|--------------|
| <u>AP</u> | BLUEPHARMA | <u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u> | <u>A078863</u> | <u>002</u> | Jun 30, 2008 |
| <u>AP</u> | ! FRESENIUS KABI USA | <u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u> | <u>A078096</u> | <u>001</u> | Jun 30, 2008 |

TABLET; ORAL

GRANISETRON HYDROCHLORIDE

| | | | | | |
|-----------|---------------------|--------------------|----------------|------------|--------------|
| <u>AB</u> | APOTEX INC | <u>EQ 1MG BASE</u> | <u>A078843</u> | <u>001</u> | Feb 27, 2008 |
| <u>AB</u> | CHARTWELL MOLECULAR | <u>EQ 1MG BASE</u> | <u>A078037</u> | <u>001</u> | Feb 27, 2008 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 1MG BASE</u> | <u>A078846</u> | <u>001</u> | Feb 27, 2009 |
| <u>AB</u> | MYLAN | <u>EQ 1MG BASE</u> | <u>A078725</u> | <u>001</u> | Jan 30, 2008 |
| <u>AB</u> | NATCO PHARMA | <u>EQ 1MG BASE</u> | <u>A078969</u> | <u>001</u> | Jun 22, 2009 |

PRESCRIPTION DRUG PRODUCT LIST

GRANISETRON HYDROCHLORIDE

TABLET;ORAL

GRANISETRON HYDROCHLORIDE

| | | | | | |
|-----------|-------------------------|---------------------------|-----------------------|-------------------|--------------|
| AB | ORCHID HLTHCARE | <u>EQ 1MG BASE</u> | <u>A078678</u> | <u>001</u> | Feb 13, 2008 |
| AB | TARO PHARM | <u>EQ 1MG BASE</u> | <u>A090817</u> | <u>001</u> | May 28, 2010 |
| AB | ! TEVA PHARMS | <u>EQ 1MG BASE</u> | <u>A078080</u> | <u>001</u> | Dec 31, 2007 |
| AB | WEST-WARD PHARMS INT | <u>EQ 1MG BASE</u> | <u>A077842</u> | <u>001</u> | Dec 31, 2007 |

GRISEOFULVIN, MICROSIZE

SUSPENSION;ORAL

GRISEOFULVIN

| | | | | | |
|-----------|---------------------------|-------------------------|-----------------------|-------------------|--------------|
| AB | ! ACTAVIS MID ATLANTIC | <u>125MG/5ML</u> | <u>A065394</u> | <u>001</u> | Jul 06, 2007 |
| AB | CHARTWELL RX | <u>125MG/5ML</u> | <u>A065200</u> | <u>001</u> | Mar 02, 2005 |
| AB | CIPLA | <u>125MG/5ML</u> | <u>A065354</u> | <u>001</u> | Sep 10, 2007 |
| AB | VINTAGE PHARMS | <u>125MG/5ML</u> | <u>A065438</u> | <u>001</u> | Oct 08, 2010 |

TABLET;ORAL

GRISEOFULVIN

| | | | | | |
|-----------|-----------------------------------|--------------------------------------------|-----------------------|-------------------|--------------|
| AB | ! SANDOZ INC | <u>500MG</u> | <u>A091592</u> | <u>002</u> | Aug 07, 2013 |
| AB | SIGMAPHARM LABS LLC SANDOZ INC | <u>500MG</u> <u>250MG</u> | <u>A202482</u> | <u>001</u> | Oct 22, 2012 |
| | | | <u>A091592</u> | <u>001</u> | Aug 07, 2013 |

GRISEOFULVIN, ULTRAMICROSIZE

TABLET;ORAL

GRIS-PEG

| | | | | | |
|-----------|----------------------|---------------------|-----------------------|-------------------|--|
| AB | + VALEANT PHARMS INC | <u>125MG</u> | <u>N050475</u> | <u>001</u> | |
| AB | +! | <u>250MG</u> | <u>N050475</u> | <u>002</u> | |

GRISEOFULVIN, ULTRAMICROSIZE

| | | | | | |
|-----------|------------|---------------------|-----------------------|-------------------|--------------|
| AB | MOUNTAIN | <u>125MG</u> | <u>A204371</u> | <u>001</u> | Jan 09, 2014 |
| AB | | <u>250MG</u> | <u>A204371</u> | <u>002</u> | Jan 09, 2014 |
| AB | SANDOZ INC | <u>125MG</u> | <u>A202805</u> | <u>001</u> | Dec 26, 2018 |
| AB | | <u>250MG</u> | <u>A202805</u> | <u>002</u> | Dec 26, 2018 |

GRISEOFULVIN, ULTRAMICROSIZE

| | | | | | |
|-----------|---------------------|---------------------|-----------------------|-------------------|--------------|
| AB | SIGMAPHARM LABS LLC | <u>125MG</u> | <u>A202545</u> | <u>001</u> | Oct 22, 2012 |
| AB | | <u>250MG</u> | <u>A202545</u> | <u>002</u> | Oct 22, 2012 |

GUAIFENESIN; HYDROCODONE BITARTRATE

SOLUTION;ORAL

OBREDON

| | | | | | |
|----|------------------|----------------------|-----------------------|-------------------|--------------|
| +! | SOVEREIGN PHARMS | 200MG/5ML; 2.5MG/5ML | <u>N205474</u> | <u>001</u> | Nov 14, 2014 |
|----|------------------|----------------------|-----------------------|-------------------|--------------|

TABLET;ORAL

XTRELUS

| | | | | | |
|----|----------------|------------|-----------------------|-------------------|--------------|
| +! | ECI PHARMS LLC | 400MG; 5MG | <u>N208085</u> | <u>001</u> | Apr 25, 2018 |
|----|----------------|------------|-----------------------|-------------------|--------------|

GUANABENZ ACETATE

TABLET;ORAL

GUANABENZ ACETATE

| | | | | | |
|---|----------------|---------------------------|-----------------------|-------------------|--------------|
| | ANI PHARMS INC | <u>EQ 4MG BASE</u> | <u>A074149</u> | <u>001</u> | Apr 07, 1995 |
| ! | | <u>EQ 8MG BASE</u> | <u>A074149</u> | <u>002</u> | Apr 07, 1995 |

GUANFACINE HYDROCHLORIDE

TABLET;ORAL

GUANFACINE HYDROCHLORIDE

| | | | | | |
|-----------|--------------|---------------------------|-----------------------|-------------------|--------------|
| AB | AMNEAL PHARM | <u>EQ 1MG BASE</u> | <u>A075109</u> | <u>001</u> | Nov 25, 1998 |
| AB | | <u>EQ 2MG BASE</u> | <u>A075109</u> | <u>002</u> | Nov 25, 1998 |
| AB | ! MYLAN | <u>EQ 1MG BASE</u> | <u>A074796</u> | <u>001</u> | Jan 27, 1997 |
| AB | ! | <u>EQ 2MG BASE</u> | <u>A074796</u> | <u>002</u> | Jan 27, 1997 |
| AB | WATSON LABS | <u>EQ 1MG BASE</u> | <u>A074145</u> | <u>001</u> | Oct 17, 1995 |
| AB | | <u>EQ 2MG BASE</u> | <u>A074145</u> | <u>002</u> | Oct 17, 1995 |

TABLET, EXTENDED RELEASE;ORAL

GUANFACINE HYDROCHLORIDE

| | | | | | |
|-----------|-------------------|---------------------------|-----------------------|-------------------|--------------|
| AB | ACTAVIS ELIZABETH | <u>EQ 1MG BASE</u> | <u>A200881</u> | <u>001</u> | Oct 05, 2012 |
| AB | | <u>EQ 2MG BASE</u> | <u>A200881</u> | <u>002</u> | Oct 05, 2012 |
| AB | | <u>EQ 3MG BASE</u> | <u>A200881</u> | <u>003</u> | Oct 05, 2012 |
| AB | | <u>EQ 4MG BASE</u> | <u>A200881</u> | <u>004</u> | Oct 05, 2012 |
| AB | APOTEX INC | <u>EQ 1MG BASE</u> | <u>A205430</u> | <u>001</u> | Jul 25, 2018 |
| AB | | <u>EQ 2MG BASE</u> | <u>A205430</u> | <u>002</u> | Jul 25, 2018 |
| AB | | <u>EQ 3MG BASE</u> | <u>A205430</u> | <u>003</u> | Jul 25, 2018 |
| AB | | <u>EQ 4MG BASE</u> | <u>A205430</u> | <u>004</u> | Jul 25, 2018 |
| AB | MYLAN PHARMS INC | <u>EQ 1MG BASE</u> | <u>A202578</u> | <u>001</u> | Jun 02, 2015 |
| AB | | <u>EQ 2MG BASE</u> | <u>A202578</u> | <u>002</u> | Jun 02, 2015 |
| AB | | <u>EQ 3MG BASE</u> | <u>A202578</u> | <u>003</u> | Jun 02, 2015 |
| AB | | <u>EQ 4MG BASE</u> | <u>A202578</u> | <u>004</u> | Jun 02, 2015 |
| AB | SANDOZ INC | <u>EQ 1MG BASE</u> | <u>A202568</u> | <u>001</u> | Jun 03, 2015 |

PRESCRIPTION DRUG PRODUCT LIST

GUANFACINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

GUANFACINE HYDROCHLORIDE

| | | | | |
|----------------|--------------------|--------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A202568 002</u> | Jun 03, 2015 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A202568 003</u> | Jun 03, 2015 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A202568 004</u> | Jun 03, 2015 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>EQ 1MG BASE</u> | <u>A205689 001</u> | Nov 16, 2017 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A205689 002</u> | Nov 16, 2017 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A205689 003</u> | Nov 16, 2017 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A205689 004</u> | Nov 16, 2017 |
| <u>AB</u> | TEVA PHARMS USA | <u>EQ 1MG BASE</u> | <u>A201382 001</u> | Jun 02, 2015 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A201382 002</u> | Jun 02, 2015 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A201382 003</u> | Jun 02, 2015 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A201382 004</u> | Jun 02, 2015 |
| <u>AB</u> | TWI PHARMS | <u>EQ 1MG BASE</u> | <u>A201408 001</u> | Jun 02, 2015 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A201408 002</u> | Jun 02, 2015 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A201408 003</u> | Jun 02, 2015 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A201408 004</u> | Jun 02, 2015 |
| <u>INTUNIV</u> | | | | |
| <u>AB</u> | + SHIRE | <u>EQ 1MG BASE</u> | <u>N022037 001</u> | Sep 02, 2009 |
| <u>AB</u> | + SHIRE | <u>EQ 2MG BASE</u> | <u>N022037 002</u> | Sep 02, 2009 |
| <u>AB</u> | + SHIRE | <u>EQ 3MG BASE</u> | <u>N022037 003</u> | Sep 02, 2009 |
| <u>AB</u> | + SHIRE | <u>EQ 4MG BASE</u> | <u>N022037 004</u> | Sep 02, 2009 |

GUANIDINE HYDROCHLORIDE

TABLET;ORAL

GUANIDINE HYDROCHLORIDE

MERCK SHARP DOHME

125MG

N001546 001

HALCINONIDE

CREAM;TOPICAL

HALOG

+! SUN PHARM INDS INC

0.1%

N017556 001

OINTMENT;TOPICAL

HALOG

+! SUN PHARM INDS INC

0.1%

N017824 001

HALOBETASOL PROPIONATE

AEROSOL, FOAM;TOPICAL

HALOBETASOL PROPIONATE

+! MAYNE PHARMA

0.05%

N210566 001 May 24, 2018

CREAM;TOPICAL

HALOBETASOL PROPIONATE

| | | | | |
|-----------|----------------|--------------|--------------------|--------------|
| <u>AB</u> | FOUGERA PHARMS | <u>0.05%</u> | <u>A077001 001</u> | Dec 16, 2004 |
| <u>AB</u> | G AND W LABS | <u>0.05%</u> | <u>A078162 001</u> | Apr 24, 2007 |
| <u>AB</u> | PERRIGO ISRAEL | <u>0.05%</u> | <u>A077123 001</u> | Dec 16, 2004 |
| <u>AB</u> | TARO | <u>0.05%</u> | <u>A077227 001</u> | Aug 04, 2005 |

ULTRAVATE

| | | | | |
|----------------|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | + SUN PHARM INDS INC | <u>0.05%</u> | <u>N019967 001</u> | Dec 27, 1990 |
| LOTION;TOPICAL | | | | |
| BRYHALI | | | | |
| | DOW PHARM | 0.01% | N209355 001 | Nov 06, 2018 |
| ULTRAVATE | | | | |
| | +! SUN PHARM INDUSTRIES | 0.05% | N208183 001 | Nov 06, 2015 |

OINTMENT;TOPICAL

HALOBETASOL PROPIONATE

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| <u>AB</u> | ! G AND W LABS | <u>0.05%</u> | <u>A077721 001</u> | Sep 07, 2006 |
| <u>AB</u> | G AND W LABS INC | <u>0.05%</u> | <u>A077109 001</u> | Jun 14, 2005 |
| <u>AB</u> | PERRIGO ISRAEL | <u>0.05%</u> | <u>A076872 001</u> | Dec 16, 2004 |
| <u>AB</u> | TARO | <u>0.05%</u> | <u>A076994 001</u> | Dec 16, 2004 |
| <u>AB</u> | TELLIGENT PHARMA INC | <u>0.05%</u> | <u>A209978 001</u> | Mar 20, 2018 |

ULTRAVATE

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| <u>AB</u> | + SUN PHARM INDS INC | <u>0.05%</u> | <u>N019968 001</u> | Dec 17, 1990 |
|-----------|----------------------|--------------|--------------------|--------------|

HALOPERIDOL

TABLET;ORAL

HALOPERIDOL

| | | | | |
|-----------|--------|--------------|--------------------|--------------|
| <u>AB</u> | MYLAN | <u>0.5MG</u> | <u>A070278 006</u> | Jun 10, 1986 |
| <u>AB</u> | | <u>1MG</u> | <u>A070278 004</u> | Jun 10, 1986 |
| <u>AB</u> | | <u>2MG</u> | <u>A070278 001</u> | Jun 10, 1986 |
| <u>AB</u> | | <u>5MG</u> | <u>A070278 005</u> | Jun 10, 1986 |
| <u>AB</u> | | <u>10MG</u> | <u>A070278 002</u> | Jul 16, 2009 |
| <u>AB</u> | | <u>20MG</u> | <u>A070278 003</u> | Jul 16, 2009 |
| <u>AB</u> | SANDOZ | <u>0.5MG</u> | <u>A071206 001</u> | Nov 17, 1986 |

PRESCRIPTION DRUG PRODUCT LIST

HALOPERIDOL

TABLET; ORAL

HALOPERIDOL

| | | | | |
|-----------|------------------|-------------|--------------------|--------------|
| <u>AB</u> | | <u>1MG</u> | <u>A071207 001</u> | Nov 17, 1986 |
| <u>AB</u> | ! | <u>2MG</u> | <u>A071208 001</u> | Nov 17, 1986 |
| <u>AB</u> | | <u>5MG</u> | <u>A071209 001</u> | Nov 17, 1986 |
| <u>AB</u> | | <u>10MG</u> | <u>A071210 001</u> | Mar 11, 1988 |
| <u>AB</u> | | <u>20MG</u> | <u>A071211 001</u> | Mar 11, 1988 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>5MG</u> | <u>A077580 003</u> | Nov 29, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A077580 004</u> | Nov 29, 2007 |
| <u>AB</u> | | <u>20MG</u> | <u>A077580 005</u> | Nov 29, 2007 |

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

HALDOL

| | | | | | |
|-----------|---|----------------|-------------------------|--------------------|--------------|
| <u>AO</u> | + | JANSSEN PHARMS | <u>EQ 50MG BASE/ML</u> | <u>N018701 001</u> | Jan 14, 1986 |
| <u>AO</u> | + | | <u>EQ 100MG BASE/ML</u> | <u>N018701 002</u> | Jan 31, 1997 |

HALOPERIDOL DECANOATE

| | | | | | |
|-----------|--|-------------------------|-------------------------|--------------------|--------------|
| <u>AO</u> | | FRESENIUS KABI USA | <u>EQ 50MG BASE/ML</u> | <u>A074893 001</u> | Dec 19, 1997 |
| <u>AO</u> | | | <u>EQ 100MG BASE/ML</u> | <u>A074893 002</u> | Dec 19, 1997 |
| <u>AO</u> | | GLAND PHARMA LTD | <u>EQ 50MG BASE/ML</u> | <u>A205241 001</u> | May 12, 2017 |
| <u>AO</u> | | | <u>EQ 100MG BASE/ML</u> | <u>A205241 002</u> | May 12, 2017 |
| <u>AO</u> | | MYLAN LABS LTD | <u>EQ 50MG BASE/ML</u> | <u>A075440 001</u> | Feb 28, 2000 |
| <u>AO</u> | | | <u>EQ 100MG BASE/ML</u> | <u>A075440 002</u> | Feb 28, 2000 |
| <u>AO</u> | | SOMERSET THERAPS LLC | <u>EQ 50MG BASE/ML</u> | <u>A209101 001</u> | Jul 03, 2018 |
| <u>AO</u> | | | <u>EQ 100MG BASE/ML</u> | <u>A209101 002</u> | Jul 03, 2018 |
| <u>AO</u> | | TEVA PHARMS USA | <u>EQ 50MG BASE/ML</u> | <u>A075393 001</u> | May 11, 1999 |
| <u>AO</u> | | | <u>EQ 100MG BASE/ML</u> | <u>A075393 002</u> | May 11, 1999 |
| <u>AO</u> | | WEST-WARD PHARMS INT | <u>EQ 50MG BASE/ML</u> | <u>A074811 001</u> | Jan 30, 1998 |
| <u>AO</u> | | | <u>EQ 100MG BASE/ML</u> | <u>A075305 001</u> | Sep 28, 1998 |

HALOPERIDOL LACTATE

CONCENTRATE; ORAL

HALOPERIDOL

| | | | | | |
|-----------|---|----------------|-----------------------|--------------------|--------------|
| <u>AA</u> | | LANNETT CO INC | <u>EQ 2MG BASE/ML</u> | <u>A073364 001</u> | Sep 28, 1993 |
| <u>AA</u> | | PHARM ASSOC | <u>EQ 2MG BASE/ML</u> | <u>A073037 001</u> | Feb 26, 1993 |
| <u>AA</u> | ! | TEVA PHARMS | <u>EQ 2MG BASE/ML</u> | <u>A071617 001</u> | Dec 01, 1988 |

INJECTABLE; INJECTION

HALDOL

| | | | | | |
|-----------|---|----------------|-----------------------|--------------------|--|
| <u>AP</u> | + | JANSSEN PHARMS | <u>EQ 5MG BASE/ML</u> | <u>N015923 001</u> | |
|-----------|---|----------------|-----------------------|--------------------|--|

HALOPERIDOL

| | | | | | |
|-----------|--|-------------------------|-----------------------|--------------------|--------------|
| <u>AP</u> | | AKORN | <u>EQ 5MG BASE/ML</u> | <u>A204849 001</u> | Sep 06, 2017 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>EQ 5MG BASE/ML</u> | <u>A075689 001</u> | Mar 09, 2001 |
| <u>AP</u> | | GLAND PHARMA LTD | <u>EQ 5MG BASE/ML</u> | <u>A076774 001</u> | Aug 25, 2004 |
| <u>AP</u> | | MYLAN LABS LTD | <u>EQ 5MG BASE/ML</u> | <u>A078347 001</u> | Sep 14, 2009 |
| <u>AP</u> | | SAGENT PHARMS | <u>EQ 5MG BASE/ML</u> | <u>A091637 001</u> | Sep 02, 2011 |
| <u>AP</u> | | | <u>EQ 5MG BASE/ML</u> | <u>A200742 001</u> | Sep 02, 2011 |
| <u>AP</u> | | TEVA PHARMS USA | <u>EQ 5MG BASE/ML</u> | <u>A076035 001</u> | Aug 29, 2001 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>EQ 5MG BASE/ML</u> | <u>A075858 001</u> | Jun 18, 2001 |

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

| | | | | | |
|-----------|---|---------------------|------------------------|--------------------|--------------|
| <u>AP</u> | | CASI PHARMS INC | <u>5,000 UNITS/ML</u> | <u>A091659 001</u> | Jun 08, 2011 |
| <u>AP</u> | + | FRESENIUS KABI USA | <u>1,000 UNITS/ML</u> | <u>N017029 001</u> | |
| <u>AP</u> | | | <u>5,000 UNITS/ML</u> | <u>A206552 001</u> | Jun 10, 2016 |
| <u>AP</u> | + | | <u>5,000 UNITS/ML</u> | <u>N017651 006</u> | |
| <u>AP</u> | + | | <u>10,000 UNITS/ML</u> | <u>N017029 003</u> | |
| <u>AP</u> | + | | <u>20,000 UNITS/ML</u> | <u>N017029 004</u> | |
| <u>AP</u> | | GLAND PHARMA LTD | <u>5,000 UNITS/ML</u> | <u>A205323 001</u> | Feb 06, 2017 |
| <u>AP</u> | | HOSPIRA INC | <u>1,000 UNITS/ML</u> | <u>A090571 001</u> | Aug 31, 2009 |
| <u>AP</u> | | | <u>5,000 UNITS/ML</u> | <u>A090571 002</u> | Aug 31, 2009 |
| <u>AP</u> | | | <u>10,000 UNITS/ML</u> | <u>A090571 003</u> | Aug 31, 2009 |
| <u>AP</u> | | MYLAN LABS LTD | <u>1,000 UNITS/ML</u> | <u>A203851 001</u> | Nov 30, 2017 |
| <u>AP</u> | | | <u>5,000 UNITS/ML</u> | <u>A203851 002</u> | Nov 30, 2017 |
| <u>AP</u> | | | <u>10,000 UNITS/ML</u> | <u>A203851 003</u> | Nov 30, 2017 |
| <u>AP</u> | | | <u>20,000 UNITS/ML</u> | <u>A203852 001</u> | Nov 30, 2017 |
| <u>AP</u> | | NANJING KING-FRIEND | <u>1,000 UNITS/ML</u> | <u>A211005 001</u> | Dec 14, 2018 |
| <u>AP</u> | | SAGENT PHARMS | <u>1,000 UNITS/ML</u> | <u>A090808 001</u> | Jun 30, 2010 |
| <u>AP</u> | | | <u>5,000 UNITS/ML</u> | <u>A090808 002</u> | Jun 30, 2010 |
| <u>AP</u> | | | <u>10,000 UNITS/ML</u> | <u>A090808 003</u> | Jun 30, 2010 |

PRESCRIPTION DRUG PRODUCT LIST

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

| | | | | |
|-----------|----------------------------|------------------------|--------------------|--------------|
| <u>AP</u> | | <u>20,000 UNITS/ML</u> | <u>A090809 001</u> | Jun 30, 2010 |
| <u>AP</u> | SANDOZ | <u>1,000 UNITS/ML</u> | <u>A091682 001</u> | Jun 08, 2011 |
| <u>AP</u> | | <u>5,000 UNITS/ML</u> | <u>A091682 002</u> | Jun 08, 2011 |
| <u>AP</u> | | <u>10,000 UNITS/ML</u> | <u>A201002 001</u> | Jun 08, 2011 |
| <u>AP</u> | SHENZHEN TECHDOW | <u>1,000 UNITS/ML</u> | <u>A202957 001</u> | Jun 12, 2014 |
| <u>AP</u> | | <u>5,000 UNITS/ML</u> | <u>A202733 001</u> | Jun 12, 2014 |
| <u>AP</u> | | <u>5,000 UNITS/ML</u> | <u>A202957 002</u> | Jun 12, 2014 |
| <u>AP</u> | | <u>10,000 UNITS/ML</u> | <u>A203198 001</u> | Jun 12, 2014 |
| <u>AP</u> | | <u>20,000 UNITS/ML</u> | <u>A203198 002</u> | Jun 12, 2014 |
| <u>AP</u> | +! WEST-WARD PHARMS INT | <u>1,000 UNITS/ML</u> | <u>N017037 001</u> | |
| <u>AP</u> | +! | <u>5,000 UNITS/ML</u> | <u>N017037 002</u> | |
| <u>AP</u> | +! | <u>10,000 UNITS/ML</u> | <u>N017037 003</u> | |

HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|------------------------|--------------------|--------------|
| <u>AP</u> | BAXTER HLTHCARE | <u>200 UNITS/100ML</u> | <u>N018609 001</u> | Apr 28, 1982 |
|-----------|-----------------|------------------------|--------------------|--------------|

HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | |
|-----------|------------|------------------------|--------------------|--------------|
| <u>AP</u> | +! B BRAUN | <u>200 UNITS/100ML</u> | <u>N019953 001</u> | Jul 20, 1992 |
| <u>AP</u> | +! HOSPIRA | <u>200 UNITS/100ML</u> | <u>N018916 010</u> | Jun 23, 1989 |

HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|-----------|---------|---------------------------|--------------------|--------------|
| <u>AP</u> | HOSPIRA | <u>10,000 UNITS/100ML</u> | <u>N019339 003</u> | Mar 27, 1985 |
|-----------|---------|---------------------------|--------------------|--------------|

HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|------------------------|--------------------|--------------|
| <u>AP</u> | BAXTER HLTHCARE | <u>200 UNITS/100ML</u> | <u>N018609 002</u> | Apr 28, 1982 |
|-----------|-----------------|------------------------|--------------------|--------------|

HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | |
|-----------|------------|------------------------|--------------------|--------------|
| <u>AP</u> | +! HOSPIRA | <u>200 UNITS/100ML</u> | <u>N018916 011</u> | Jun 23, 1989 |
|-----------|------------|------------------------|--------------------|--------------|

HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|-----------|------------|--------------------------|--------------------|--------------|
| <u>AP</u> | +! B BRAUN | <u>4,000 UNITS/100ML</u> | <u>N019952 001</u> | Jul 20, 1992 |
| <u>AP</u> | HOSPIRA | <u>4,000 UNITS/100ML</u> | <u>N019805 001</u> | Jan 25, 1989 |

HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|-----------|------------|---------------------------|--------------------|--------------|
| <u>AP</u> | +! B BRAUN | <u>5,000 UNITS/100ML</u> | <u>N019952 004</u> | Jul 20, 1992 |
| <u>AP</u> | +! | <u>10,000 UNITS/100ML</u> | <u>N019952 005</u> | Jul 20, 1992 |
| <u>AP</u> | HOSPIRA | <u>5,000 UNITS/100ML</u> | <u>N019339 004</u> | Mar 27, 1985 |
| <u>AP</u> | | <u>5,000 UNITS/100ML</u> | <u>N019805 002</u> | Jan 25, 1989 |
| <u>AP</u> | | <u>10,000 UNITS/100ML</u> | <u>N019339 002</u> | Mar 27, 1985 |

HEPARIN SODIUM IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------------|------------------------|--------------------|--------------|
| <u>AP</u> | +! FRESENIUS KABI USA | <u>1,000 UNITS/ML</u> | <u>N017029 013</u> | Dec 05, 1985 |
| <u>AP</u> | +! | <u>5,000 UNITS/ML</u> | <u>N017029 014</u> | Dec 05, 1985 |
| <u>AP</u> | +! | <u>10,000 UNITS/ML</u> | <u>N017029 015</u> | Dec 05, 1985 |
| <u>AP</u> | +! | <u>20,000 UNITS/ML</u> | <u>N017029 016</u> | Dec 05, 1985 |

HEPARIN SODIUM PRESERVATIVE FREE

| | | | | |
|-----------|-----------------------|-----------------------|--------------------|--------------|
| <u>AP</u> | +! FRESENIUS KABI USA | <u>1,000 UNITS/ML</u> | <u>N017029 010</u> | Apr 28, 1986 |
| <u>AP</u> | SAGENT PHARMS | <u>1,000 UNITS/ML</u> | <u>A090810 001</u> | Jun 30, 2010 |
| <u>AP</u> | SHENZHEN TECHDOW | <u>1,000 UNITS/ML</u> | <u>A202732 001</u> | Jun 12, 2014 |

HEPARIN SODIUM

| | | | | |
|--|------------------------------------------------------------------------------|--------------------|-------------|--------------|
| | +! FRESENIUS KABI USA | 10,000 UNITS/ML | N017029 020 | Mar 31, 2011 |
| | ! HOSPIRA | 5,000 UNITS/ML | A088100 001 | Apr 28, 1983 |
| | +! PFIZER | 1,000 UNITS/ML | N201370 001 | Jul 21, 2011 |
| | +! | 5,000 UNITS/ML | N201370 002 | Jul 21, 2011 |
| | +! | 10,000 UNITS/ML | N201370 003 | Jul 21, 2011 |
| | STERINOVA INC | 5,000 UNITS/0.5ML | A208827 001 | Nov 19, 2018 |
| | HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| | HOSPIRA | 5,000 UNITS/100ML | N019339 001 | Mar 27, 1985 |
| | HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER | | | |
| | HOSPIRA | 5,000 UNITS/100ML | N018916 006 | Jan 31, 1984 |
| | HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER | | | |
| | HOSPIRA | 5,000 UNITS/100ML | N018916 007 | Jan 31, 1984 |
| | | 10,000 UNITS/100ML | N018916 008 | Jan 31, 1984 |
| | HEPARIN SODIUM PRESERVATIVE FREE | | | |
| | +! FRESENIUS KABI USA | 10,000 UNITS/ML | N017029 019 | Nov 22, 2010 |
| | ! HOSPIRA | 10,000 UNITS/ML | A089522 001 | May 04, 1987 |
| | +! PFIZER | 1,000 UNITS/ML | N201370 004 | Jul 21, 2011 |

HEXACHLOROPHENE

SPONGE; TOPICAL

PRE-OP

| | | | | |
|-----------|-------------------|--------------|--------------------|--|
| <u>AT</u> | +! DAVIS AND GECK | <u>480MG</u> | <u>N017433 001</u> | |
|-----------|-------------------|--------------|--------------------|--|

PRE-OP II

| | | | | |
|-----------|------------------|--------------|--------------------|--|
| <u>AT</u> | + DAVIS AND GECK | <u>480MG</u> | <u>N017433 002</u> | |
|-----------|------------------|--------------|--------------------|--|

PRESCRIPTION DRUG PRODUCT LISTHEXAMINOLEVULINATE HYDROCHLORIDEFOR SOLUTION; INTRAVESICAL
CYSVIEW KIT

+! PHOTOCURE ASA 100MG/VIAL N022555 001 May 28, 2010

HISTRELIN ACETATEIMPLANT; SUBCUTANEOUS
SUPPRELIN LA

+! ENDO PHARM 50MG N022058 001 May 03, 2007

VANTAS

+! ENDO PHARM 50MG N021732 001 Oct 12, 2004

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDEAA ABHAI LLC 1.5MG/5ML; 5MG/5ML A207487 001 Feb 21, 2017AA ACTAVIS MID 1.5MG/5ML; 5MG/5ML A088017 001 Jul 05, 1983

ATLANTIC

AA ! HI TECH PHARMA 1.5MG/5ML; 5MG/5ML A040613 001 Feb 08, 2008AA NOVEL LABS INC 1.5MG/5ML; 5MG/5ML A203535 001 Feb 13, 2017AA PADDOCK LLC 1.5MG/5ML; 5MG/5ML A205731 001 Feb 15, 2017AA WOCKHARDT BIO AG 1.5MG/5ML; 5MG/5ML A088008 001 Mar 03, 1983

TABLET; ORAL

HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATEAA AVANTHI INC 1.5MG; 5MG A207176 001 Aug 07, 2017AA NOVEL LABS INC 1.5MG; 5MG A091528 001 Apr 20, 2011TUSSIGONAA ! KING PHARMS 1.5MG; 5MG A088508 001 Jul 30, 1985HYALURONIDASE

INJECTABLE; INJECTION

AMPHADASE

+! AMPHASTAR PHARM 150 UNITS/ML N021665 001 Oct 26, 2004

VITRASE

+! BAUSCH AND LOMB 200 UNITS/VIAL N021640 002 Dec 02, 2004

HYALURONIDASE RECOMBINANT HUMAN

INJECTABLE; INJECTION

HYLENEX RECOMBINANT

+! HALOZYME THERAP 150 UNITS/ML N021859 001 Dec 02, 2005

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDRALAZINE HYDROCHLORIDEAP ! AKORN 20MG/ML A040730 001 Apr 21, 2009AP FRESENIUS KABI USA 20MG/ML A040388 001 Mar 13, 2001AP LUITPOLD 20MG/ML A040136 001 Jun 30, 1997AP MYLAN INSTITUTIONAL 20MG/ML A204680 001 Apr 28, 2016AP NAVINTA LLC 20MG/ML A202938 001 Mar 28, 2013AP X-GEN PHARMS INC 20MG/ML A203110 001 Jun 29, 2015

TABLET; ORAL

HYDRALAZINE HYDROCHLORIDEAA ALKEM LABS LTD 10MG A200737 001 Dec 07, 2012AA 25MG A200737 002 Dec 07, 2012AA 50MG A200737 003 Dec 07, 2012AA 100MG A200737 004 Dec 07, 2012AA APPCO PHARMA LLC 10MG A209251 001 Jul 09, 2018AA 25MG A209251 002 Jul 09, 2018AA 50MG A209251 003 Jul 09, 2018AA 100MG A209251 004 Jul 09, 2018AA CADILA PHARMS LTD 25MG A203845 001 Sep 18, 2014AA 50MG A203845 002 Sep 18, 2014AA 100MG A203845 003 Sep 18, 2014AA GLENMARK PHARMS LTD 10MG A090527 001 May 27, 2009AA 25MG A090527 002 May 27, 2009AA 50MG A090527 003 May 27, 2009AA 100MG A090527 004 May 27, 2009AA HERITAGE PHARMS INC 10MG A086242 001 Feb 04, 2010AA 25MG A086242 003AA 50MG A086242 002AA 100MG A086242 004 Feb 04, 2010AA HETERO LABS LTD III 10MG A040901 001 Sep 12, 2008AA 25MG A040901 002 Sep 12, 2008AA 50MG A040901 003 Sep 12, 2008AA 100MG A040901 004 Sep 12, 2008

PRESCRIPTION DRUG PRODUCT LISTHYDRALAZINE HYDROCHLORIDE

TABLET; ORAL

HYDRALAZINE HYDROCHLORIDE

| | | | | |
|-----------|--------------------|--------------|--------------------|--------------|
| <u>AA</u> | INVAGEN PHARMS | <u>10MG</u> | <u>A090255 001</u> | Dec 15, 2008 |
| <u>AA</u> | | <u>25MG</u> | <u>A090255 002</u> | Dec 15, 2008 |
| <u>AA</u> | | <u>50MG</u> | <u>A090255 003</u> | Dec 15, 2008 |
| <u>AA</u> | | <u>100MG</u> | <u>A090255 004</u> | Dec 15, 2008 |
| <u>AA</u> | PAR PHARM | <u>10MG</u> | <u>A087836 001</u> | Oct 05, 1982 |
| <u>AA</u> | | <u>25MG</u> | <u>A086961 002</u> | |
| <u>AA</u> | | <u>50MG</u> | <u>A086962 001</u> | |
| <u>AA</u> | | <u>100MG</u> | <u>A088391 001</u> | Sep 27, 1983 |
| <u>AA</u> | ! PLIVA | <u>10MG</u> | <u>A089097 001</u> | Dec 18, 1985 |
| <u>AA</u> | ! | <u>25MG</u> | <u>A088467 001</u> | May 01, 1984 |
| <u>AA</u> | ! | <u>50MG</u> | <u>A088468 001</u> | May 01, 1984 |
| <u>AA</u> | ! | <u>100MG</u> | <u>A089098 001</u> | Dec 18, 1985 |
| <u>AA</u> | SCIEGEN PHARMS INC | <u>10MG</u> | <u>A205236 001</u> | May 26, 2017 |
| <u>AA</u> | | <u>25MG</u> | <u>A205236 002</u> | May 26, 2017 |
| <u>AA</u> | | <u>50MG</u> | <u>A205236 003</u> | May 26, 2017 |
| <u>AA</u> | | <u>100MG</u> | <u>A205236 004</u> | May 26, 2017 |
| <u>AA</u> | STRIDES PHARMA | <u>25MG</u> | <u>A200770 001</u> | May 03, 2013 |
| <u>AA</u> | | <u>50MG</u> | <u>A200770 002</u> | May 03, 2013 |
| <u>AA</u> | | <u>100MG</u> | <u>A200770 003</u> | May 03, 2013 |

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDRA-ZIDE

| | | | | |
|--|-----------|------------|-------------|--------------|
| | PAR PHARM | 25MG; 25MG | A088957 001 | Oct 21, 1985 |
| | ! | 50MG; 50MG | A088946 001 | Oct 21, 1985 |

HYDRALAZINE HYDROCHLORIDE; ISOSORBIDE DINITRATE

TABLET; ORAL

BIDIL

| | | | | |
|--|---------------------|--------------|-------------|--------------|
| | +! ARBOR PHARMS LLC | 37.5MG; 20MG | N020727 001 | Jun 23, 2005 |
|--|---------------------|--------------|-------------|--------------|

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

| | | | | |
|-----------|-------------------------|---------------|--------------------|--------------|
| <u>AB</u> | ALEMBIC PHARMS LTD | <u>12.5MG</u> | <u>A200645 001</u> | Nov 30, 2010 |
| <u>AB</u> | AUROBINDO PHARMA | <u>12.5MG</u> | <u>A078164 001</u> | Sep 18, 2007 |
| <u>AB</u> | IPCA LABS LTD | <u>12.5MG</u> | <u>A079237 001</u> | Apr 02, 2009 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>12.5MG</u> | <u>A077005 001</u> | Jul 13, 2005 |
| <u>AB</u> | JUBILANT CADISTA | <u>12.5MG</u> | <u>A078391 001</u> | Feb 11, 2008 |
| <u>AB</u> | MYLAN | <u>12.5MG</u> | <u>A075640 001</u> | Jan 28, 2000 |
| <u>AB</u> | PRINSTON INC | <u>12.5MG</u> | <u>A075907 001</u> | Sep 17, 2002 |
| <u>AB</u> | SCIEGEN PHARMS INC | <u>12.5MG</u> | <u>A203561 001</u> | Jan 14, 2019 |
| <u>AB</u> | SUN PHARM INDS INC | <u>12.5MG</u> | <u>A090651 001</u> | Apr 07, 2014 |
| <u>AB</u> | UNICHEM | <u>12.5MG</u> | <u>A090510 001</u> | Jan 19, 2010 |

MICROZIDE

| | | | | |
|-----------|-----------------------|---------------|--------------------|--------------|
| <u>AB</u> | +! ALLERGAN SALES LLC | <u>12.5MG</u> | <u>N020504 001</u> | Dec 27, 1996 |
|-----------|-----------------------|---------------|--------------------|--------------|

TABLET; ORAL

HYDROCHLOROTHIAZIDE

| | | | | |
|-----------|-------------------------|---------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>12.5MG</u> | <u>A202556 001</u> | Sep 24, 2012 |
| <u>AB</u> | | <u>25MG</u> | <u>A202556 002</u> | Sep 24, 2012 |
| <u>AB</u> | | <u>50MG</u> | <u>A202556 003</u> | Sep 24, 2012 |
| <u>AB</u> | ACTAVIS ELIZABETH | <u>12.5MG</u> | <u>A040707 001</u> | Feb 27, 2007 |
| <u>AB</u> | AUROBINDO PHARMA | <u>25MG</u> | <u>A040780 001</u> | Jul 20, 2007 |
| <u>AB</u> | | <u>50MG</u> | <u>A040780 002</u> | Jul 20, 2007 |
| <u>AB</u> | HERITAGE PHARMS INC | <u>25MG</u> | <u>A085182 002</u> | |
| <u>AB</u> | | <u>50MG</u> | <u>A085182 001</u> | |
| <u>AB</u> | HIKMA INTL PHARMS | <u>50MG</u> | <u>A084878 001</u> | |
| <u>AB</u> | IPCA LABS LTD | <u>12.5MG</u> | <u>A040807 001</u> | Jul 20, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A040807 002</u> | Jul 20, 2007 |
| <u>AB</u> | | <u>50MG</u> | <u>A040807 003</u> | Jul 20, 2007 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>25MG</u> | <u>A083177 001</u> | |
| <u>AB</u> | ! | <u>50MG</u> | <u>A083177 002</u> | |
| <u>AB</u> | LEADING PHARMA LLC | <u>12.5MG</u> | <u>A040702 003</u> | May 10, 2017 |
| <u>AB</u> | | <u>25MG</u> | <u>A040702 001</u> | Mar 16, 2007 |
| <u>AB</u> | | <u>50MG</u> | <u>A040702 002</u> | Mar 16, 2007 |
| <u>AB</u> | MYLAN PHARMS INC | <u>25MG</u> | <u>A040735 002</u> | Jan 23, 2007 |
| <u>AB</u> | | <u>50MG</u> | <u>A040735 003</u> | Jan 23, 2007 |
| <u>AB</u> | OXFORD PHARMS | <u>25MG</u> | <u>A087059 001</u> | |
| <u>AB</u> | | <u>50MG</u> | <u>A087068 001</u> | |

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE

TABLET; ORAL

HYDROCHLOROTHIAZIDE

| | | | | | |
|-----------|--------------------|---------------|----------------|------------|--------------|
| <u>AB</u> | PRINSTON INC | <u>25MG</u> | <u>A040412</u> | <u>001</u> | Mar 29, 2002 |
| <u>AB</u> | | <u>50MG</u> | <u>A040412</u> | <u>002</u> | Mar 29, 2002 |
| <u>AB</u> | SCIEGEN PHARMS INC | <u>25MG</u> | <u>A203018</u> | <u>001</u> | Jul 23, 2014 |
| <u>AB</u> | | <u>50MG</u> | <u>A203018</u> | <u>002</u> | Jul 23, 2014 |
| <u>AB</u> | SUN PHARM INDS INC | <u>12.5MG</u> | <u>A040857</u> | <u>001</u> | May 30, 2008 |
| <u>AB</u> | | <u>25MG</u> | <u>A040810</u> | <u>001</u> | Mar 27, 2007 |
| <u>AB</u> | | <u>50MG</u> | <u>A040810</u> | <u>002</u> | Mar 27, 2007 |
| <u>AB</u> | UNICHEM | <u>25MG</u> | <u>A040907</u> | <u>001</u> | Aug 15, 2008 |
| <u>AB</u> | | <u>50MG</u> | <u>A040907</u> | <u>002</u> | Aug 15, 2008 |

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

AVALIDE

| | | | | | |
|-----------|------------------------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | <u>+</u> ! SANOFI AVENTIS US | <u>12.5MG; 150MG</u> | <u>N020758</u> | <u>002</u> | Sep 30, 1997 |
| <u>AB</u> | <u>+</u> ! | <u>12.5MG; 300MG</u> | <u>N020758</u> | <u>003</u> | Aug 31, 1998 |

IRBESARTAN AND HYDROCHLOROTHIAZIDE

| | | | | | |
|-----------|----------------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | ALEMBIC PHARMS LTD | <u>12.5MG; 150MG</u> | <u>A091370</u> | <u>001</u> | Oct 15, 2012 |
| <u>AB</u> | | <u>12.5MG; 300MG</u> | <u>A091370</u> | <u>002</u> | Oct 15, 2012 |
| <u>AB</u> | | <u>25MG; 300MG</u> | <u>A091370</u> | <u>003</u> | Oct 12, 2016 |
| <u>AB</u> | APOTEX INC | <u>12.5MG; 150MG</u> | <u>A201505</u> | <u>001</u> | Oct 15, 2012 |
| <u>AB</u> | | <u>12.5MG; 300MG</u> | <u>A201505</u> | <u>002</u> | Oct 15, 2012 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>12.5MG; 150MG</u> | <u>A203630</u> | <u>001</u> | Feb 22, 2013 |
| <u>AB</u> | | <u>12.5MG; 300MG</u> | <u>A203630</u> | <u>002</u> | Feb 22, 2013 |
| <u>AB</u> | | <u>25MG; 300MG</u> | <u>A203630</u> | <u>003</u> | Mar 31, 2016 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>12.5MG; 150MG</u> | <u>A203500</u> | <u>001</u> | Sep 27, 2012 |
| <u>AB</u> | | <u>12.5MG; 300MG</u> | <u>A203500</u> | <u>002</u> | Sep 27, 2012 |
| <u>AB</u> | HISUN PHARM HANGZHOU | <u>12.5MG; 150MG</u> | <u>A207896</u> | <u>001</u> | Oct 14, 2016 |
| <u>AB</u> | | <u>12.5MG; 300MG</u> | <u>A207896</u> | <u>002</u> | Oct 14, 2016 |
| <u>AB</u> | LUPIN LTD | <u>12.5MG; 150MG</u> | <u>A201524</u> | <u>001</u> | Feb 27, 2013 |
| <u>AB</u> | | <u>12.5MG; 300MG</u> | <u>A201524</u> | <u>002</u> | Feb 27, 2013 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>12.5MG; 150MG</u> | <u>A202414</u> | <u>001</u> | Sep 27, 2012 |
| <u>AB</u> | | <u>12.5MG; 300MG</u> | <u>A202414</u> | <u>002</u> | Sep 27, 2012 |
| <u>AB</u> | PRINSTON INC | <u>12.5MG; 150MG</u> | <u>A203072</u> | <u>001</u> | May 09, 2014 |
| <u>AB</u> | | <u>12.5MG; 300MG</u> | <u>A203072</u> | <u>002</u> | May 09, 2014 |
| <u>AB</u> | SANDOZ | <u>12.5MG; 150MG</u> | <u>A077446</u> | <u>001</u> | Sep 27, 2012 |
| <u>AB</u> | | <u>12.5MG; 300MG</u> | <u>A077446</u> | <u>002</u> | Sep 27, 2012 |
| <u>AB</u> | TEVA | <u>12.5MG; 150MG</u> | <u>A077369</u> | <u>001</u> | Mar 30, 2012 |
| <u>AB</u> | | <u>12.5MG; 300MG</u> | <u>A077369</u> | <u>002</u> | Mar 30, 2012 |
| <u>AB</u> | UNICHEM LABS LTD | <u>12.5MG; 150MG</u> | <u>A207018</u> | <u>001</u> | Sep 19, 2017 |
| <u>AB</u> | | <u>12.5MG; 300MG</u> | <u>A207018</u> | <u>002</u> | Sep 19, 2017 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>12.5MG; 150MG</u> | <u>A090351</u> | <u>001</u> | Oct 15, 2012 |
| <u>AB</u> | | <u>12.5MG; 300MG</u> | <u>A090351</u> | <u>002</u> | Oct 15, 2012 |
| <u>AB</u> | | <u>25MG; 300MG</u> | <u>A090351</u> | <u>003</u> | Jun 08, 2017 |

HYDROCHLOROTHIAZIDE; LISINAPRIL

TABLET; ORAL

LISINAPRIL AND HYDROCHLOROTHIAZIDE

| | | | | | |
|-----------|----------------------|---------------------|----------------|------------|--------------|
| <u>AB</u> | AUROBINDO | <u>12.5MG; 10MG</u> | <u>A077606</u> | <u>001</u> | Mar 14, 2006 |
| <u>AB</u> | | <u>12.5MG; 20MG</u> | <u>A077606</u> | <u>002</u> | Mar 14, 2006 |
| <u>AB</u> | | <u>25MG; 20MG</u> | <u>A077606</u> | <u>003</u> | Mar 14, 2006 |
| <u>AB</u> | HIKMA INTL PHARMS | <u>12.5MG; 10MG</u> | <u>A076265</u> | <u>001</u> | Jul 08, 2002 |
| <u>AB</u> | | <u>12.5MG; 20MG</u> | <u>A076265</u> | <u>002</u> | Jul 08, 2002 |
| <u>AB</u> | | <u>25MG; 20MG</u> | <u>A076265</u> | <u>003</u> | Jul 08, 2002 |
| <u>AB</u> | INVAGEN PHARMS | <u>12.5MG; 10MG</u> | <u>A204058</u> | <u>001</u> | May 23, 2017 |
| <u>AB</u> | | <u>12.5MG; 20MG</u> | <u>A204058</u> | <u>002</u> | May 23, 2017 |
| <u>AB</u> | | <u>25MG; 20MG</u> | <u>A204058</u> | <u>003</u> | May 23, 2017 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>12.5MG; 10MG</u> | <u>A075776</u> | <u>001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>12.5MG; 20MG</u> | <u>A075776</u> | <u>002</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>25MG; 20MG</u> | <u>A075776</u> | <u>003</u> | Jul 01, 2002 |
| <u>AB</u> | LUPIN | <u>12.5MG; 10MG</u> | <u>A077912</u> | <u>001</u> | Sep 27, 2006 |
| <u>AB</u> | | <u>12.5MG; 20MG</u> | <u>A077912</u> | <u>002</u> | Sep 27, 2006 |
| <u>AB</u> | | <u>25MG; 20MG</u> | <u>A077912</u> | <u>003</u> | Sep 27, 2006 |
| <u>AB</u> | MYLAN | <u>12.5MG; 10MG</u> | <u>A076113</u> | <u>001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>12.5MG; 20MG</u> | <u>A076113</u> | <u>002</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>25MG; 20MG</u> | <u>A076113</u> | <u>003</u> | Jul 01, 2002 |
| <u>AB</u> | PRINSTON INC | <u>12.5MG; 10MG</u> | <u>A076230</u> | <u>001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>12.5MG; 20MG</u> | <u>A076230</u> | <u>002</u> | Jul 01, 2002 |

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; LISINAPRIL

TABLET;ORAL

LISINAPRIL AND HYDROCHLOROTHIAZIDE

| | | | | | |
|-------------------|--------------------|--------------------|--------------------|----------------|-------------------------|
| <u>AB</u> | | <u>25MG;20MG</u> | <u>A076230</u> | <u>003</u> | Jul 01, 2002 |
| <u>AB</u> | SANDOZ | <u>12.5MG;10MG</u> | <u>A076262</u> | <u>001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>12.5MG;20MG</u> | <u>A076262</u> | <u>002</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>25MG;20MG</u> | <u>A076262</u> | <u>003</u> | Jul 01, 2002 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>12.5MG;10MG</u> | <u>A076007</u> | <u>001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>12.5MG;20MG</u> | <u>A076007</u> | <u>002</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>25MG;20MG</u> | <u>A076007</u> | <u>003</u> | Jul 01, 2002 |
| <u>AB</u> | WATSON LABS | <u>12.5MG;10MG</u> | <u>A076194</u> | <u>003</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>12.5MG;20MG</u> | <u>A076194</u> | <u>001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>25MG;20MG</u> | <u>A076194</u> | <u>002</u> | Jul 01, 2002 |
| <u>ZESTORETIC</u> | | | | | |
| <u>AB</u> | + | ALVOGEN MALTA | <u>12.5MG;10MG</u> | <u>N019888</u> | <u>003</u> Nov 18, 1993 |
| <u>AB</u> | + | ! | <u>12.5MG;20MG</u> | <u>N019888</u> | <u>001</u> Sep 20, 1990 |
| <u>AB</u> | + | ! | <u>25MG;20MG</u> | <u>N019888</u> | <u>002</u> Jul 20, 1989 |

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET;ORAL

HYZAAR

| | | | | | |
|-----------|---|-------------------|---------------------|----------------|-------------------------|
| <u>AB</u> | + | MERCK SHARP DOHME | <u>12.5MG;100MG</u> | <u>N020387</u> | <u>003</u> Oct 20, 2005 |
|-----------|---|-------------------|---------------------|----------------|-------------------------|

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE

| | | | | | |
|-----------|---|----------------------|---------------------|----------------|-------------------------|
| <u>AB</u> | | ALEMbic PHARMS LTD | <u>12.5MG;50MG</u> | <u>A091617</u> | <u>001</u> Feb 17, 2012 |
| <u>AB</u> | | | <u>12.5MG;100MG</u> | <u>A091617</u> | <u>002</u> Feb 17, 2012 |
| <u>AB</u> | | | <u>25MG;100MG</u> | <u>A091617</u> | <u>003</u> Feb 17, 2012 |
| <u>AB</u> | | AUROBINDO PHARMA | <u>12.5MG;50MG</u> | <u>A091629</u> | <u>001</u> Oct 06, 2010 |
| <u>AB</u> | | | <u>12.5MG;100MG</u> | <u>A091629</u> | <u>002</u> Oct 06, 2010 |
| <u>AB</u> | ! | | <u>25MG;100MG</u> | <u>A091629</u> | <u>003</u> Jan 06, 2010 |
| <u>AB</u> | | CADISTA PHARMS | <u>12.5MG;50MG</u> | <u>A201845</u> | <u>001</u> Sep 18, 2012 |
| <u>AB</u> | | | <u>12.5MG;100MG</u> | <u>A201845</u> | <u>002</u> Sep 18, 2012 |
| <u>AB</u> | | | <u>25MG;100MG</u> | <u>A201845</u> | <u>003</u> Sep 18, 2012 |
| <u>AB</u> | | IPCA LABS LTD | <u>12.5MG;50MG</u> | <u>A201682</u> | <u>001</u> Mar 01, 2013 |
| <u>AB</u> | | | <u>12.5MG;100MG</u> | <u>A201682</u> | <u>002</u> Mar 01, 2013 |
| <u>AB</u> | | | <u>25MG;100MG</u> | <u>A201682</u> | <u>003</u> Mar 01, 2013 |
| <u>AB</u> | | LUPIN LTD | <u>12.5MG;50MG</u> | <u>A078245</u> | <u>001</u> Oct 06, 2010 |
| <u>AB</u> | | | <u>12.5MG;100MG</u> | <u>A078245</u> | <u>002</u> May 21, 2010 |
| <u>AB</u> | | | <u>25MG;100MG</u> | <u>A078245</u> | <u>003</u> Oct 06, 2010 |
| <u>AB</u> | | MACLEODS PHARMS LTD | <u>12.5MG;50MG</u> | <u>A202289</u> | <u>001</u> Aug 09, 2012 |
| <u>AB</u> | | | <u>12.5MG;100MG</u> | <u>A202289</u> | <u>002</u> Aug 09, 2012 |
| <u>AB</u> | | | <u>25MG;100MG</u> | <u>A202289</u> | <u>003</u> Aug 09, 2012 |
| <u>AB</u> | | MYLAN | <u>12.5MG;50MG</u> | <u>A091652</u> | <u>001</u> Oct 06, 2010 |
| <u>AB</u> | | | <u>12.5MG;100MG</u> | <u>A091652</u> | <u>002</u> Apr 06, 2010 |
| <u>AB</u> | | | <u>25MG;100MG</u> | <u>A091652</u> | <u>003</u> Oct 06, 2010 |
| <u>AB</u> | | PRINSTON INC | <u>12.5MG;50MG</u> | <u>A204901</u> | <u>001</u> Nov 06, 2017 |
| <u>AB</u> | | | <u>12.5MG;100MG</u> | <u>A204901</u> | <u>002</u> Nov 06, 2017 |
| <u>AB</u> | | | <u>25MG;100MG</u> | <u>A204901</u> | <u>003</u> Nov 06, 2017 |
| <u>AB</u> | | SANDOZ | <u>12.5MG;50MG</u> | <u>A077948</u> | <u>001</u> Oct 06, 2010 |
| <u>AB</u> | | | <u>12.5MG;100MG</u> | <u>A077948</u> | <u>003</u> Aug 19, 2010 |
| <u>AB</u> | | | <u>25MG;100MG</u> | <u>A077948</u> | <u>002</u> Oct 06, 2010 |
| <u>AB</u> | | TEVA PHARMS | <u>12.5MG;50MG</u> | <u>A077157</u> | <u>001</u> Apr 06, 2010 |
| <u>AB</u> | | | <u>12.5MG;100MG</u> | <u>A077157</u> | <u>002</u> Apr 06, 2010 |
| <u>AB</u> | | | <u>25MG;100MG</u> | <u>A077157</u> | <u>003</u> Apr 06, 2010 |
| <u>AB</u> | | TORRENT PHARMS | <u>12.5MG;50MG</u> | <u>A090528</u> | <u>001</u> Oct 06, 2010 |
| <u>AB</u> | | | <u>12.5MG;100MG</u> | <u>A090528</u> | <u>003</u> Apr 06, 2010 |
| <u>AB</u> | | | <u>25MG;100MG</u> | <u>A090528</u> | <u>002</u> Oct 06, 2010 |
| <u>AB</u> | | UNICHEM LABS LTD | <u>12.5MG;50MG</u> | <u>A204832</u> | <u>001</u> Jul 21, 2017 |
| <u>AB</u> | | | <u>12.5MG;100MG</u> | <u>A204832</u> | <u>002</u> Jul 21, 2017 |
| <u>AB</u> | | | <u>25MG;100MG</u> | <u>A204832</u> | <u>003</u> Jul 21, 2017 |
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>12.5MG;50MG</u> | <u>A077732</u> | <u>002</u> Oct 06, 2010 |
| <u>AB</u> | | | <u>12.5MG;100MG</u> | <u>A077732</u> | <u>001</u> Apr 06, 2010 |
| <u>AB</u> | | | <u>25MG;100MG</u> | <u>A077732</u> | <u>003</u> Oct 06, 2010 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>12.5MG;50MG</u> | <u>A078385</u> | <u>001</u> Oct 06, 2010 |
| <u>AB</u> | | | <u>25MG;100MG</u> | <u>A078385</u> | <u>002</u> Oct 06, 2010 |

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET;ORAL

METHYLDOPA AND HYDROCHLOROTHIAZIDE

| | | | | | |
|--|---|-------|------------|---------|------------------|
| | | MYLAN | 15MG;250MG | A070265 | 002 Jan 23, 1986 |
| | ! | | 25MG;250MG | A070265 | 001 Jan 23, 1986 |

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE;ORAL

DUTOPROL

| | | | | | |
|--|---|-------------------------|--------------------------|-------------|--------------|
| | + | CONCORDIA PHARMS INC | 12.5MG;EQ 25MG TARTRATE | N021956 001 | Aug 28, 2006 |
| | + | | 12.5MG;EQ 50MG TARTRATE | N021956 002 | Aug 28, 2006 |
| | + | ! | 12.5MG;EQ 100MG TARTRATE | N021956 003 | Aug 28, 2006 |

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET;ORAL

LOPRESSOR HCT

| | | | | | |
|-----------|---|-------------------------|-------------------|--------------------|--------------|
| AB | + | US PHARMS HOLDINGS I | 25MG;50MG | N018303 001 | Dec 31, 1984 |
| AB | + | ! | 25MG;100MG | N018303 002 | Dec 31, 1984 |

METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE

| | | | | | |
|-----------|--|--------------------|-------------------|--------------------|--------------|
| AB | | ALEMBIC PHARMS LTD | 25MG;50MG | A202870 001 | Nov 06, 2013 |
| AB | | | 25MG;100MG | A202870 002 | Nov 06, 2013 |
| AB | | | 50MG;100MG | A202870 003 | Nov 06, 2013 |
| AB | | MYLAN | 25MG;50MG | A076792 001 | Aug 20, 2004 |
| AB | | | 25MG;100MG | A076792 002 | Aug 20, 2004 |
| AB | | | 50MG;100MG | A076792 003 | Aug 20, 2004 |
| AB | | SUN PHARM INDS | 25MG;50MG | A090654 001 | Jan 19, 2012 |
| AB | | | 25MG;100MG | A090654 002 | Jan 19, 2012 |
| AB | | | 50MG;100MG | A090654 003 | Jan 19, 2012 |

HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE

TABLET;ORAL

MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

| | | | | | |
|-----------|--|---------------------|---------------------|--------------------|--------------|
| AB | | GLENMARK PHARMS | 12.5MG;7.5MG | A090718 001 | Mar 17, 2010 |
| AB | | | 12.5MG;15MG | A090718 002 | Mar 17, 2010 |
| AB | | | 25MG;15MG | A090718 003 | Mar 17, 2010 |
| AB | | HERITAGE PHARMS INC | 12.5MG;7.5MG | A202150 001 | Mar 07, 2014 |
| AB | | | 12.5MG;15MG | A202150 002 | Mar 07, 2014 |
| AB | | | 25MG;15MG | A202150 003 | Mar 07, 2014 |
| AB | | TEVA | 12.5MG;7.5MG | A076980 001 | Mar 07, 2007 |
| AB | | | 12.5MG;15MG | A076980 003 | Mar 07, 2007 |
| AB | | ! | 25MG;15MG | A076980 002 | Mar 07, 2007 |

HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET;ORAL

BENICAR HCT

| | | | | | |
|-----------|---|----------------|--------------------|--------------------|--------------|
| AB | + | DAIICHI SANKYO | 12.5MG;20MG | N021532 002 | Jun 05, 2003 |
| AB | + | | 12.5MG;40MG | N021532 003 | Jun 05, 2003 |
| AB | + | ! | 25MG;40MG | N021532 005 | Jun 05, 2003 |

OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE

| | | | | | |
|-----------|--|-------------------------|--------------------|--------------------|--------------|
| AB | | ALEMBIC PHARMS LTD | 12.5MG;20MG | A204233 001 | Apr 24, 2017 |
| AB | | | 12.5MG;40MG | A204233 002 | Apr 24, 2017 |
| AB | | | 25MG;40MG | A204233 003 | Apr 24, 2017 |
| AB | | AUROBINDO PHARMA LTD | 12.5MG;20MG | A205391 001 | Apr 24, 2017 |
| AB | | | 12.5MG;40MG | A205391 002 | Apr 24, 2017 |
| AB | | | 25MG;40MG | A205391 003 | Apr 24, 2017 |
| AB | | MYLAN PHARMS INC | 12.5MG;20MG | A078827 001 | Oct 26, 2016 |
| AB | | | 12.5MG;40MG | A078827 002 | Oct 26, 2016 |
| AB | | | 25MG;40MG | A078827 003 | Oct 26, 2016 |
| AB | | PRINSTON INC | 12.5MG;20MG | A207804 001 | Apr 24, 2017 |
| AB | | | 12.5MG;40MG | A207804 002 | Apr 24, 2017 |
| AB | | | 25MG;40MG | A207804 003 | Apr 24, 2017 |
| AB | | TEVA PHARMS USA | 12.5MG;20MG | A200532 001 | Apr 24, 2017 |
| AB | | | 12.5MG;40MG | A200532 002 | Apr 24, 2017 |
| AB | | | 25MG;40MG | A200532 003 | Apr 24, 2017 |
| AB | | TORRENT PHARMS LTD | 12.5MG;20MG | A206515 001 | May 03, 2017 |
| AB | | | 12.5MG;40MG | A206515 002 | May 03, 2017 |
| AB | | | 25MG;40MG | A206515 003 | May 03, 2017 |

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

TABLET;ORAL

PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

| | | | | | |
|-----------|---|-------|------------------|--------------------|--------------|
| AB | ! | MYLAN | 25MG;80MG | A070947 001 | Apr 01, 1987 |
| | ! | | 25MG;40MG | A070947 002 | Mar 04, 1987 |

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

ACCURETIC

| | | | | | |
|-----------|----------|----------------------|----------------------------|--------------------|--------------|
| <u>AB</u> | <u>+</u> | <u>PFIZER PHARMS</u> | <u>12.5MG;EQ 10MG BASE</u> | <u>N020125 001</u> | Dec 28, 1999 |
| <u>AB</u> | <u>+</u> | | <u>12.5MG;EQ 20MG BASE</u> | <u>N020125 002</u> | Dec 28, 1999 |
| <u>AB</u> | <u>+</u> | | <u>25MG;EQ 20MG BASE</u> | <u>N020125 003</u> | Dec 28, 1999 |

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

| | | | | | |
|-----------|--|-------------------------|----------------------------|--------------------|--------------|
| <u>AB</u> | | <u>APOTEX CORP</u> | <u>12.5MG;EQ 10MG BASE</u> | <u>A091524 001</u> | Mar 12, 2013 |
| <u>AB</u> | | | <u>12.5MG;EQ 20MG BASE</u> | <u>A091524 002</u> | Mar 12, 2013 |
| <u>AB</u> | | | <u>25MG;EQ 20MG BASE</u> | <u>A091524 003</u> | Mar 12, 2013 |
| <u>AB</u> | | <u>AUROBINDO PHARMA</u> | <u>12.5MG;EQ 10MG BASE</u> | <u>A078450 001</u> | Aug 24, 2007 |
| <u>AB</u> | | | <u>12.5MG;EQ 20MG BASE</u> | <u>A078450 002</u> | Aug 24, 2007 |
| <u>AB</u> | | | <u>25MG;EQ 20MG BASE</u> | <u>A078450 003</u> | Aug 24, 2007 |
| <u>AB</u> | | <u>INVAGEN PHARMS</u> | <u>12.5MG;EQ 10MG BASE</u> | <u>A201356 001</u> | Apr 20, 2011 |
| <u>AB</u> | | | <u>12.5MG;EQ 20MG BASE</u> | <u>A201356 002</u> | Apr 20, 2011 |
| <u>AB</u> | | | <u>25MG;EQ 20MG BASE</u> | <u>A201356 003</u> | Apr 20, 2011 |
| <u>AB</u> | | <u>MYLAN</u> | <u>12.5MG;EQ 10MG BASE</u> | <u>A077093 001</u> | Mar 28, 2005 |
| <u>AB</u> | | | <u>12.5MG;EQ 20MG BASE</u> | <u>A077093 002</u> | Mar 28, 2005 |
| <u>AB</u> | | | <u>25MG;EQ 20MG BASE</u> | <u>A077093 003</u> | Mar 28, 2005 |

QUINARETIC

| | | | | | |
|-----------|--|---------------------|----------------------------|--------------------|--------------|
| <u>AB</u> | | <u>GAVIS PHARMS</u> | <u>12.5MG;EQ 10MG BASE</u> | <u>A076374 001</u> | Mar 31, 2004 |
| <u>AB</u> | | | <u>12.5MG;EQ 20MG BASE</u> | <u>A076374 002</u> | Mar 31, 2004 |
| <u>AB</u> | | | <u>25MG;EQ 20MG BASE</u> | <u>A076374 003</u> | Mar 31, 2004 |

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

TABLET; ORAL

ALDACTAZIDE

| | | | | | |
|-----------|----------|----------------------|------------------|--------------------|--------------|
| <u>AB</u> | <u>+</u> | <u>GD SEARLE LLC</u> | <u>25MG;25MG</u> | <u>N012616 004</u> | Dec 30, 1982 |
|-----------|----------|----------------------|------------------|--------------------|--------------|

SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE

| | | | | | |
|-----------|--|-------------------|------------------|--------------------|--------------|
| <u>AB</u> | | <u>MYLAN</u> | <u>25MG;25MG</u> | <u>A086513 001</u> | |
| <u>AB</u> | | <u>SUN PHARM</u> | <u>25MG;25MG</u> | <u>A089534 001</u> | Jul 02, 1987 |
| | | <u>INDUSTRIES</u> | | | |

ALDACTAZIDE

| | | | | | |
|----------|----------|----------------------|------------------|--------------------|--------------|
| <u>+</u> | <u>!</u> | <u>GD SEARLE LLC</u> | <u>50MG;50MG</u> | <u>N012616 005</u> | Dec 30, 1982 |
|----------|----------|----------------------|------------------|--------------------|--------------|

HYDROCHLOROTHIAZIDE; TELMISARTAN

TABLET; ORAL

MICARDIS HCT

| | | | | | |
|-----------|----------|-------------------|--------------------|--------------------|--------------|
| <u>AB</u> | <u>+</u> | <u>BOEHRINGER</u> | <u>12.5MG;40MG</u> | <u>N021162 001</u> | Nov 17, 2000 |
| | | <u>INGELHEIM</u> | | | |
| <u>AB</u> | <u>+</u> | | <u>12.5MG;80MG</u> | <u>N021162 002</u> | Nov 17, 2000 |
| <u>AB</u> | <u>+</u> | | <u>25MG;80MG</u> | <u>N021162 003</u> | Apr 19, 2004 |

TELMISARTAN AND HYDROCHLOROTHIAZIDE

| | | | | | |
|-----------|--|----------------------------|--------------------|--------------------|--------------|
| <u>AB</u> | | <u>ALEMBIC PHARMS LTD</u> | <u>12.5MG;40MG</u> | <u>A203010 001</u> | Feb 25, 2014 |
| <u>AB</u> | | | <u>12.5MG;80MG</u> | <u>A203010 002</u> | Feb 25, 2014 |
| <u>AB</u> | | | <u>25MG;80MG</u> | <u>A203010 003</u> | Feb 25, 2014 |
| <u>AB</u> | | <u>AUROBINDO PHARMA</u> | <u>12.5MG;40MG</u> | <u>A208727 001</u> | Dec 15, 2016 |
| | | <u>LTD</u> | | | |
| <u>AB</u> | | | <u>12.5MG;80MG</u> | <u>A208727 002</u> | Dec 15, 2016 |
| <u>AB</u> | | | <u>25MG;80MG</u> | <u>A208727 003</u> | Dec 15, 2016 |
| <u>AB</u> | | <u>LUPIN LTD</u> | <u>12.5MG;40MG</u> | <u>A091351 001</u> | Aug 07, 2014 |
| <u>AB</u> | | | <u>12.5MG;80MG</u> | <u>A091351 002</u> | Aug 07, 2014 |
| <u>AB</u> | | | <u>25MG;80MG</u> | <u>A091351 003</u> | Aug 07, 2014 |
| <u>AB</u> | | <u>MACLEODS PHARMS LTD</u> | <u>12.5MG;40MG</u> | <u>A204169 001</u> | Nov 02, 2015 |
| <u>AB</u> | | | <u>12.5MG;80MG</u> | <u>A204169 002</u> | Nov 02, 2015 |
| <u>AB</u> | | | <u>25MG;80MG</u> | <u>A204169 003</u> | Nov 02, 2015 |
| <u>AB</u> | | <u>MYLAN PHARMS INC</u> | <u>12.5MG;40MG</u> | <u>A091648 001</u> | Feb 25, 2014 |
| <u>AB</u> | | | <u>12.5MG;80MG</u> | <u>A091648 002</u> | Feb 25, 2014 |
| <u>AB</u> | | | <u>25MG;80MG</u> | <u>A091648 003</u> | Feb 25, 2014 |
| <u>AB</u> | | <u>PRINSTON INC</u> | <u>12.5MG;40MG</u> | <u>A209028 001</u> | Nov 06, 2017 |
| <u>AB</u> | | | <u>12.5MG;80MG</u> | <u>A209028 002</u> | Nov 06, 2017 |
| <u>AB</u> | | | <u>25MG;80MG</u> | <u>A209028 003</u> | Nov 06, 2017 |
| <u>AB</u> | | <u>TORRENT PHARMS LTD</u> | <u>12.5MG;40MG</u> | <u>A201192 001</u> | Feb 25, 2014 |
| <u>AB</u> | | | <u>12.5MG;80MG</u> | <u>A201192 002</u> | Feb 25, 2014 |
| <u>AB</u> | | | <u>25MG;80MG</u> | <u>A201192 003</u> | Feb 25, 2014 |
| <u>AB</u> | | <u>ZYDUS PHARMS USA</u> | <u>12.5MG;40MG</u> | <u>A204221 001</u> | Aug 15, 2017 |
| | | <u>INC</u> | | | |
| <u>AB</u> | | | <u>12.5MG;80MG</u> | <u>A204221 002</u> | Aug 15, 2017 |
| <u>AB</u> | | | <u>25MG;80MG</u> | <u>A204221 003</u> | Aug 15, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

DYAZIDE

| | | | | | |
|--------------------------------------------|------------|-------------------------|---------------------|--------------------|--------------|
| AB | + ! | GLAXOSMITHKLINE LLC | 25MG; 37.5MG | N016042 003 | Mar 03, 1994 |
| <u>TRIAMTERENE AND HYDROCHLOROTHIAZIDE</u> | | | | | |
| AB | ! | CASI PHARMS INC | 25MG; 50MG | A073191 001 | Jul 31, 1991 |
| AB | | DURAMED PHARMS BARR | 25MG; 37.5MG | A075052 001 | Jun 18, 1999 |
| AB | | IVAX SUB TEVA PHARMS | 25MG; 50MG | A074259 001 | Mar 30, 1995 |
| AB | | LANNETT CO INC | 25MG; 37.5MG | A201407 001 | Dec 09, 2011 |
| AB | | MYLAN | 25MG; 37.5MG | A074701 001 | Jun 07, 1996 |
| AB | | SANDOZ | 25MG; 37.5MG | A074821 001 | Jun 05, 1997 |

TABLET; ORAL

MAXZIDE

| | | | | | |
|--------------------------------------------|------------|-------------------------|---------------------|--------------------|--------------|
| AB | + ! | MYLAN PHARMS INC | 50MG; 75MG | N019129 001 | Oct 22, 1984 |
| <u>MAXZIDE-25</u> | | | | | |
| AB | + | MYLAN PHARMS INC | 25MG; 37.5MG | N019129 003 | May 13, 1988 |
| <u>TRIAMTERENE AND HYDROCHLOROTHIAZIDE</u> | | | | | |
| AB | | ANI PHARMS INC | 50MG; 75MG | A073467 001 | Jan 31, 1996 |
| AB | | APOTEX INC | 25MG; 37.5MG | A071251 002 | May 05, 1998 |
| AB | | | 50MG; 75MG | A071251 001 | Apr 17, 1988 |
| AB | | PLIVA | 25MG; 37.5MG | A074026 001 | Apr 26, 1996 |
| AB | | SANDOZ | 25MG; 37.5MG | A073281 001 | Apr 30, 1992 |
| AB | | | 50MG; 75MG | A072011 001 | Jun 17, 1988 |
| AB | | WATSON LABS | 25MG; 37.5MG | A073449 001 | Sep 23, 1993 |
| AB | | | 50MG; 75MG | A071851 001 | Nov 30, 1988 |
| AB | | ZYDUS PHARMS USA INC | 25MG; 37.5MG | A208360 001 | Jun 29, 2018 |
| AB | | | 50MG; 75MG | A208360 002 | Jun 29, 2018 |

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

DIOVAN HCT

| | | | | | |
|-----------|------------|----------|----------------------|--------------------|--------------|
| AB | + | NOVARTIS | 12.5MG; 80MG | N020818 001 | Mar 06, 1998 |
| AB | + | | 12.5MG; 160MG | N020818 002 | Mar 06, 1998 |
| AB | + | | 12.5MG; 320MG | N020818 004 | Apr 28, 2006 |
| AB | + | | 25MG; 160MG | N020818 003 | Jan 17, 2002 |
| AB | + ! | | 25MG; 320MG | N020818 005 | Apr 28, 2006 |

VALSARTAN AND HYDROCHLOROTHIAZIDE

| | | | | | |
|-----------|--|-------------------------|----------------------|--------------------|--------------|
| AB | | ALEMBOC PHARMS LTD | 12.5MG; 80MG | A201662 001 | Mar 21, 2013 |
| AB | | | 12.5MG; 160MG | A201662 002 | Mar 21, 2013 |
| AB | | | 12.5MG; 320MG | A201662 003 | Mar 21, 2013 |
| AB | | | 25MG; 160MG | A201662 004 | Mar 21, 2013 |
| AB | | | 25MG; 320MG | A201662 005 | Mar 21, 2013 |
| AB | | AUROBINDO PHARMA LTD | 12.5MG; 80MG | A202519 001 | Mar 21, 2013 |
| AB | | | 12.5MG; 160MG | A202519 002 | Mar 21, 2013 |
| AB | | | 12.5MG; 320MG | A202519 003 | Mar 21, 2013 |
| AB | | | 25MG; 160MG | A202519 004 | Mar 21, 2013 |
| AB | | | 25MG; 320MG | A202519 005 | Mar 21, 2013 |
| AB | | LUPIN LTD | 12.5MG; 80MG | A078946 003 | Mar 21, 2013 |
| AB | | | 12.5MG; 160MG | A078946 004 | Mar 21, 2013 |
| AB | | | 12.5MG; 320MG | A078946 001 | Mar 21, 2013 |
| AB | | | 25MG; 160MG | A078946 005 | Mar 21, 2013 |
| AB | | | 25MG; 320MG | A078946 002 | Mar 21, 2013 |
| AB | | MACLEODS PHARMS LTD | 12.5MG; 80MG | A203145 001 | Apr 19, 2013 |
| AB | | | 12.5MG; 160MG | A203145 002 | Apr 19, 2013 |
| AB | | | 12.5MG; 320MG | A203145 003 | Apr 19, 2013 |
| AB | | | 25MG; 160MG | A203145 004 | Apr 19, 2013 |
| AB | | | 25MG; 320MG | A203145 005 | Apr 19, 2013 |
| AB | | MYLAN PHARMS INC | 12.5MG; 80MG | A078020 001 | Sep 21, 2012 |
| AB | | | 12.5MG; 160MG | A078020 002 | Sep 21, 2012 |
| AB | | | 12.5MG; 320MG | A078020 004 | Sep 21, 2012 |
| AB | | | 25MG; 160MG | A078020 003 | Sep 21, 2012 |
| AB | | | 25MG; 320MG | A078020 005 | Sep 21, 2012 |
| AB | | PRINSTON INC | 12.5MG; 80MG | A206083 001 | Feb 08, 2016 |
| AB | | | 12.5MG; 160MG | A206083 002 | Feb 08, 2016 |
| AB | | | 12.5MG; 320MG | A206083 003 | Feb 08, 2016 |
| AB | | | 25MG; 160MG | A206083 004 | Feb 08, 2016 |
| AB | | | 25MG; 320MG | A206083 005 | Feb 08, 2016 |

PRESCRIPTION DRUG PRODUCT LISTHYDROCODONE BITARTRATE

CAPSULE, EXTENDED RELEASE;ORAL

ZOHYDRO ER

| | | | | | | |
|---|---|---------------------|------|---------|-----|--------------|
| + | ! | PERNIX IRELAND PAIN | 10MG | N202880 | 001 | Oct 25, 2013 |
| + | | | 15MG | N202880 | 002 | Oct 25, 2013 |
| + | | | 20MG | N202880 | 003 | Oct 25, 2013 |
| + | | | 30MG | N202880 | 004 | Oct 25, 2013 |
| + | | | 40MG | N202880 | 005 | Oct 25, 2013 |
| + | | | 50MG | N202880 | 006 | Oct 25, 2013 |

TABLET, EXTENDED RELEASE;ORAL

HYSINGLA

| | | | | | | |
|---|---|------------------|-------|---------|-----|--------------|
| + | ! | PURDUE PHARMA LP | 20MG | N206627 | 001 | Nov 20, 2014 |
| + | | | 30MG | N206627 | 002 | Nov 20, 2014 |
| + | | | 40MG | N206627 | 003 | Nov 20, 2014 |
| + | | | 60MG | N206627 | 004 | Nov 20, 2014 |
| + | | | 80MG | N206627 | 005 | Nov 20, 2014 |
| + | | | 100MG | N206627 | 006 | Nov 20, 2014 |
| + | | | 120MG | N206627 | 007 | Nov 20, 2014 |

HYDROCODONE BITARTRATE; IBUPROFEN

TABLET;ORAL

HYDROCODONE BITARTRATE AND IBUPROFEN

| | | | | | | |
|-----------|---|---------------------|--------------------|----------------|------------|--------------|
| AB | | ACTAVIS LABS FL INC | 7.5MG;200MG | A076604 | 001 | Dec 31, 2003 |
| AB | | AMNEAL PHARMS NY | 5MG;200MG | A076642 | 002 | Mar 18, 2004 |
| AB | ! | | 7.5MG;200MG | A076642 | 001 | Oct 12, 2004 |
| AB | | AUROLIFE PHARMA LLC | 7.5MG;200MG | A204575 | 001 | Jun 02, 2016 |
| AB | | SUN PHARM INDS INC | 2.5MG;200MG | A091633 | 001 | May 28, 2013 |
| AB | | | 5MG;200MG | A091633 | 002 | May 28, 2013 |
| AB | | | 7.5MG;200MG | A091633 | 003 | May 28, 2013 |
| AB | | | 10MG;200MG | A091633 | 004 | May 28, 2013 |
| AB | | TEVA | 7.5MG;200MG | A076023 | 001 | Apr 11, 2003 |
| AB | | VINTAGE PHARMS | 5MG;200MG | A077727 | 001 | Nov 06, 2006 |
| AB | | | 7.5MG;200MG | A077723 | 001 | Nov 06, 2006 |
| AB | | | 10MG;200MG | A077723 | 002 | Nov 06, 2006 |

REPREXAIN

| | | | | | | |
|-----------|--|------------------|--------------------|----------------|------------|--------------|
| AB | | AMNEAL PHARMS NY | 2.5MG;200MG | A076642 | 003 | Oct 19, 2007 |
| AB | | | 10MG;200MG | A076642 | 004 | Oct 19, 2007 |

HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION;ORAL

HYDROCODONE BITARTRATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

| | | | | | | |
|-----------|---|------------------|-------------------------|----------------|------------|--------------|
| AA | | MAYNE PHARMA INC | 5MG/5ML;60MG/5ML | A205658 | 001 | Nov 17, 2015 |
| AA | | PADDOCK LLC | 5MG/5ML;60MG/5ML | A204658 | 001 | Apr 29, 2014 |
| AA | + | ! | 5MG/5ML;60MG/5ML | N022442 | 001 | Jun 08, 2011 |

HYDROCORTISONE

CREAM;TOPICAL

ALA-CORT

| | | | | | | |
|-----------|--|------------|-------------|----------------|------------|--------------|
| AT | | CROWN LABS | 2.5% | A080706 | 007 | Jan 05, 2016 |
| AT | | | 1% | A080706 | 006 | |

ANUSOL HC

| | | | | | | |
|-----------|--|--------------|-------------|----------------|------------|--------------|
| AT | | SALIX PHARMS | 2.5% | A088250 | 001 | Jun 06, 1984 |
|-----------|--|--------------|-------------|----------------|------------|--------------|

HYDROCORTISONE

| | | | | | | |
|-----------|---|----------------------|-------------|----------------|------------|--------------|
| AT | | ACTAVIS MID ATLANTIC | 1% | A087795 | 001 | May 03, 1983 |
| AT | | | 2.5% | A089682 | 001 | Mar 10, 1988 |
| AT | ! | FOUGERA PHARMS INC | 1% | A080693 | 003 | |
| AT | ! | | 2.5% | A089414 | 001 | Dec 16, 1986 |
| AT | | LANNETT CO INC | 2.5% | A040503 | 001 | Mar 12, 2004 |
| AT | | PERRIGO NEW YORK | 2.5% | A085025 | 001 | |
| AT | | RISING PHARMS | 2.5% | A040879 | 001 | Aug 20, 2010 |
| AT | | TARO PHARM INDS LTD | 2.5% | A088799 | 001 | Nov 09, 1984 |
| AT | | TEGENT PHARMA INC | 2.5% | A203810 | 001 | Jul 23, 2018 |

ENEMA;RECTAL

COLOCORT

| | | | | | | |
|-----------|--|-------------|-------------------|----------------|------------|--------------|
| AB | | PADDOCK LLC | 100MG/60ML | A075172 | 001 | Dec 03, 1999 |
|-----------|--|-------------|-------------------|----------------|------------|--------------|

CORTENEMA

| | | | | | | |
|-----------|---|---|-------------------|----------------|------------|--|
| AB | + | ! | 100MG/60ML | N016199 | 001 | |
|-----------|---|---|-------------------|----------------|------------|--|

HYDROCORTISONE

| | | | | | | |
|-----------|--|-------------|-------------------|----------------|------------|--------------|
| AB | | TEVA PHARMS | 100MG/60ML | A074171 | 001 | May 27, 1994 |
|-----------|--|-------------|-------------------|----------------|------------|--------------|

LOTION;TOPICAL

HYDROCORTISONE

| | | | | | | |
|-----------|---|----------------|-------------|----------------|------------|--------------|
| AT | ! | FOUGERA PHARMS | 2.5% | A040351 | 001 | Jul 25, 2000 |
|-----------|---|----------------|-------------|----------------|------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

HYDROCORTISONE

LOTION; TOPICAL

HYDROCORTISONE

| | | | | | |
|-----------|---------------------|-------------|----------------|------------|--------------|
| <u>AT</u> | LANNETT CO INC | <u>2.5%</u> | <u>A040417</u> | <u>001</u> | Jul 30, 2003 |
| <u>AT</u> | TARO | <u>2.5%</u> | <u>A040247</u> | <u>001</u> | Jul 23, 1999 |
| <u>AT</u> | TELOGENT PHARMA INC | <u>2.5%</u> | <u>A203804</u> | <u>001</u> | Jul 27, 2018 |

STIE-CORT

| | | | | | |
|-----------|------------|-------------|----------------|------------|--------------|
| <u>AT</u> | PERRIGO CO | <u>2.5%</u> | <u>A089074</u> | <u>001</u> | Nov 26, 1985 |
| | ALA-SCALP | | | | |
| | CROWN LABS | 2% | A083231 | 001 | |

OINTMENT; TOPICAL

HYDROCORTISONE

| | | | | | |
|-----------|----------------------|-------------|----------------|------------|--------------|
| <u>AT</u> | ACTAVIS MID ATLANTIC | <u>1%</u> | <u>A087796</u> | <u>001</u> | Oct 13, 1982 |
| <u>AT</u> | ! FOUGERA PHARMS | <u>1%</u> | <u>A080692</u> | <u>001</u> | |
| <u>AT</u> | ! FOUGERA PHARMS INC | <u>2.5%</u> | <u>A081203</u> | <u>001</u> | May 28, 1993 |
| <u>AT</u> | PERRIGO NEW YORK | <u>2.5%</u> | <u>A085027</u> | <u>001</u> | |
| <u>AT</u> | TARO | <u>1%</u> | <u>A086257</u> | <u>001</u> | |

HYDROCORTISONE IN ABSORBASE

| | | | | | |
|-----------|-------------------|-----------|----------------|------------|--------------|
| <u>AT</u> | CMP PHARMA INC | <u>1%</u> | <u>A088138</u> | <u>001</u> | Sep 06, 1985 |
| | SOLUTION; TOPICAL | | | | |
| | TEXACORT | | | | |
| | ! MISSION PHARMA | 2.5% | A081271 | 001 | Apr 17, 1992 |

TABLET; ORAL

CORTEF

| | | | | | |
|-----------|------------------------|-------------|----------------|------------|--|
| <u>AB</u> | + PHARMACIA AND UPJOHN | <u>5MG</u> | <u>N008697</u> | <u>003</u> | |
| <u>AB</u> | + | <u>10MG</u> | <u>N008697</u> | <u>001</u> | |
| <u>AB</u> | +! | <u>20MG</u> | <u>N008697</u> | <u>002</u> | |

HYDROCORTISONE

| | | | | | |
|-----------|-------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | HIKMA INTL PHARMS | <u>5MG</u> | <u>A083365</u> | <u>002</u> | Feb 23, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A083365</u> | <u>003</u> | Feb 23, 2015 |
| <u>AB</u> | | <u>20MG</u> | <u>A083365</u> | <u>001</u> | |
| <u>AB</u> | IMPAX LABS INC | <u>5MG</u> | <u>A040646</u> | <u>001</u> | Mar 30, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A040646</u> | <u>002</u> | Mar 30, 2007 |
| <u>AB</u> | | <u>20MG</u> | <u>A040646</u> | <u>003</u> | Mar 30, 2007 |
| <u>AB</u> | PII | <u>5MG</u> | <u>A207029</u> | <u>001</u> | Apr 27, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A207029</u> | <u>002</u> | Apr 27, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A207029</u> | <u>003</u> | Apr 27, 2017 |
| <u>AB</u> | VINTAGE | <u>5MG</u> | <u>A040761</u> | <u>001</u> | Jul 16, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A040761</u> | <u>002</u> | Jul 16, 2007 |
| <u>AB</u> | | <u>20MG</u> | <u>A040761</u> | <u>003</u> | Jul 16, 2007 |

HYDROCORTISONE ACETATE

AEROSOL, METERED; RECTAL

CORTIFOAM

| | | | | | |
|----|---------------------|-----|---------|-----|--------------|
| +! | MYLAN SPECIALITY LP | 10% | N017351 | 001 | Feb 10, 1982 |
|----|---------------------|-----|---------|-----|--------------|

CREAM; TOPICAL

MICORT-HC

| | | | | | |
|--|--------------------|------|---------|-----|--------------|
| | SEBELA IRELAND LTD | 2.5% | A040396 | 001 | Feb 27, 2001 |
|--|--------------------|------|---------|-----|--------------|

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

CREAM; TOPICAL

CORTISPORIN

| | | | | | |
|----|----------------|---------------------------------------|---------|-----|--------------|
| +! | MONARCH PHARMS | 0.5%;EQ 3.5MG BASE/GM;10,000 UNITS/GM | N050218 | 001 | Aug 09, 1985 |
|----|----------------|---------------------------------------|---------|-----|--------------|

HYDROCORTISONE ACETATE; PRAMOXINE HYDROCHLORIDE

AEROSOL, METERED; TOPICAL

EPIFOAM

| | | | | | |
|----|---------------------|-------|---------|-----|--|
| BX | MYLAN SPECIALITY LP | 1%;1% | A086457 | 001 | |
|----|---------------------|-------|---------|-----|--|

PROCTOFOAM HC

| | | | | | |
|----|---------------------|-------|---------|-----|--|
| BX | MYLAN SPECIALITY LP | 1%;1% | A086195 | 001 | |
|----|---------------------|-------|---------|-----|--|

CREAM; TOPICAL

PRAMOSONE

| | | | | | |
|--|--------------------|---------|---------|-----|--|
| | SEBELA IRELAND LTD | 0.5%;1% | A083778 | 001 | |
|--|--------------------|---------|---------|-----|--|

| | | | | | |
|--|--|-------|---------|-----|--|
| | | 1%;1% | A085368 | 001 | |
|--|--|-------|---------|-----|--|

LOTION; TOPICAL

PRAMOSONE

| | | | | | |
|--|--------------------|-------|---------|-----|--|
| | SEBELA IRELAND LTD | 1%;1% | A085980 | 001 | |
|--|--------------------|-------|---------|-----|--|

| | | | | | |
|--|--|---------|---------|-----|--|
| | | 2.5%;1% | A085979 | 001 | |
|--|--|---------|---------|-----|--|

PRESCRIPTION DRUG PRODUCT LIST

HYDROCORTISONE ACETATE; UREACREAM; TOPICAL
U-CORT

TARO 1%;10% A089472 001 Jun 13, 1988

HYDROCORTISONE BUTYRATE

CREAM; TOPICAL

HYDROCORTISONE BUTYRATE**AB1** TARO PHARM INDS 0.1% **A076654 001** Aug 03, 2005LOCOID**AB1** +! PRECISION DERMAT 0.1% **N018514 001** Mar 31, 1982HYDROCORTISONE BUTYRATE**AB2** ACTAVIS MID 0.1% **A205134 001** Dec 08, 2017

ATLANTIC

AB2 GLENMARK GENERICS 0.1% **A202145 001** Sep 27, 2013LOCOID LIPOCREAM**AB2** +! PRECISION DERMAT 0.1% **N020769 001** Sep 08, 1997

LOTION; TOPICAL

HYDROCORTISONE BUTYRATE**AB** LUPIN LTD 0.1% **A210209 001** Aug 17, 2018**AB** TELIGENT PHARMA INC 0.1% **A209556 001** Nov 21, 2017LOCOID**AB** +! PRECISION DERMAT 0.1% **N022076 001** May 18, 2007

OINTMENT; TOPICAL

HYDROCORTISONE BUTYRATE**AB** TARO 0.1% **A076842 001** Dec 27, 2004LOCOID**AB** +! PRECISION DERMAT 0.1% **N018652 001** Oct 29, 1982

SOLUTION; TOPICAL

HYDROCORTISONE BUTYRATE**AT** TARO PHARM INDS 0.1% **A076364 001** Jan 14, 2004LOCOID**AT** +! PRECISION DERMAT 0.1% **N019116 001** Feb 25, 1987HYDROCORTISONE PROBUTATE

CREAM; TOPICAL

PANDEL

+! FOUGERA PHARMS 0.1% N020453 001 Feb 28, 1997

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION

SOLU-CORTEF

+! PHARMACIA AND EQ 100MG BASE/VIAL N009866 001
UPJOHN

+! EQ 250MG BASE/VIAL N009866 002

+! EQ 500MG BASE/VIAL N009866 003

+! EQ 1GM BASE/VIAL N009866 004

HYDROCORTISONE VALERATE

CREAM; TOPICAL

HYDROCORTISONE VALERATE**AB** GLENMARK PHARMS LTD 0.2% **A211129 001** Oct 12, 2018**AB** PERRIGO NEW YORK 0.2% **A075666 001** May 24, 2000**AB** ! TARO 0.2% **A075042 001** Aug 25, 1998

OINTMENT; TOPICAL

HYDROCORTISONE VALERATE**AB** GLENMARK PHARMS LTD 0.2% **A211750 001** Dec 14, 2018**AB** ! TARO 0.2% **A075043 001** Aug 25, 1998HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OTIC

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE**AT** ! BAUSCH AND LOMB 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML **A064053 001** Dec 29, 1995**AT** SANDOZ INC 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML **A062423 001** Aug 25, 1983

SUSPENSION/DROPS; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

! SANDOZ INC 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML A062874 001 May 11, 1988

SUSPENSION/DROPS; OTIC

CASPORYN HC**AT** +! CASPER PHARMA LLC 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML **N060613 001**NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE**AT** AMRING PHARMS 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML **A065219 001** May 01, 2006**AT** SANDOZ INC 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML **A062488 001** Nov 06, 1985OTICAIR**AT** BAUSCH AND LOMB 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML **A064065 001** Aug 28, 1996

PRESCRIPTION DRUG PRODUCT LISTHYDROGEN PEROXIDE

SOLUTION; TOPICAL

ESKATA

+! ACLARIS

40%

N209305 001 Dec 14, 2017

HYDROMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION

DILAUDIDAP +! FRESENIUS KABI USA1MG/MLN019034 003 Apr 30, 2009AP +!2MG/MLN019034 004 Apr 30, 2009HYDROMORPHONE HYDROCHLORIDEAP AKORN10MG/MLA078228 001 Apr 14, 2010AP10MG/MLA078261 001 Apr 14, 2010AP BARR10MG/MLA076444 001 Apr 25, 2003AP EUROHLTH INTL SARL2MG/MLA202159 001 Apr 27, 2018AP HOSPIRA INC1MG/MLN200403 001 Dec 01, 2011AP2MG/MLN200403 002 Dec 01, 2011AP4MG/MLN200403 003 Dec 01, 2011AP10MG/MLA078591 001 Jun 17, 2008

SOLUTION; ORAL

DILAUDIDAA +! RHODES PHARMS5MG/5MLN019891 001 Dec 07, 1992HYDROMORPHONE HYDROCHLORIDEAA ASCENT PHARMS INC5MG/5MLA210176 001 Oct 27, 2017AA WEST-WARD PHARMS5MG/5MLA074653 001 Jul 29, 1998

TABLET; ORAL

DILAUDIDAB + RHODES PHARMS2MGN019892 003 Nov 09, 2007AB +4MGN019892 002 Nov 09, 2007AB +!8MGN019892 001 Dec 07, 1992HYDROMORPHONE HYDROCHLORIDEAB ASCENT PHARMS INC2MGA210506 001 Jan 17, 2018AB4MGA210506 002 Jan 17, 2018AB8MGA210506 003 Jan 17, 2018AB AUROLIFE PHARMA LLC2MGA205814 001 May 13, 2016AB4MGA205814 002 May 13, 2016AB8MGA205814 003 May 13, 2016AB ELITE LABS8MGA076723 001 Oct 18, 2005AB LANNETT CO INC2MGA077471 002 Dec 09, 2009AB4MGA077471 003 Dec 09, 2009AB8MGA077471 001 Dec 09, 2009AB SPECGX LLC2MGA076855 002 Sep 19, 2007AB4MGA076855 003 Sep 19, 2007AB8MGA076855 001 Dec 23, 2004AB WEST-WARD PHARMS4MGA074597 003 May 29, 2009

TABLET, EXTENDED RELEASE; ORAL

EXALGOAB + SPECGX LLC8MGN021217 001 Mar 01, 2010AB +12MGN021217 002 Mar 01, 2010AB +16MGN021217 003 Mar 01, 2010AB +!32MGN021217 004 Aug 24, 2012HYDROMORPHONE HYDROCHLORIDEAB OSMOTICA8MGA205629 001 Jul 07, 2016AB12MGA205629 002 Jul 07, 2016AB16MGA205629 003 Jul 07, 2016AB32MGA205629 004 Jul 07, 2016AB PADDOCK LLC8MGA204278 001 Apr 06, 2015AB12MGA204278 002 Apr 06, 2015AB16MGA204278 003 Apr 06, 2015AB32MGA204278 004 Sep 20, 2017HYDROXOCOBALAMIN

INJECTABLE; INJECTION

CYANOKIT

+! SERB SA

5GM/VIAL (5GM/KIT)

N022041 001 Apr 08, 2011

HYDROXOCOBALAMIN

! ACTAVIS LLC

1MG/ML

A085998 001

PRESCRIPTION DRUG PRODUCT LIST

HYDROXYAMPHETAMINE HYDROBROMIDE; TROPICAMIDE

SOLUTION/DROPS;OPHTHALMIC

PAREMYD

+! AKORN

1%;0.25%

N019261 001 Jan 30, 1992

HYDROXYCHLOROQUINE SULFATE

TABLET;ORAL

HYDROXYCHLOROQUINE SULFATE

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| AB | ALKALOIDA ZRT | 200MG | A201691 001 | May 08, 2018 |
| AB | AMNEAL PHARMS CO | 200MG | A210577 001 | May 15, 2018 |
| AB | HIKMA PHARMS | 200MG | A040760 001 | Aug 15, 2007 |
| AB | IPCA LABS LTD | 200MG | A040766 001 | Jun 14, 2007 |
| AB | LUPIN LTD | 200MG | A210543 001 | Jul 06, 2018 |
| AB | MYLAN | 200MG | A040274 001 | May 29, 1998 |
| AB | RISING PHARMS | 200MG | A210959 001 | Jan 15, 2019 |
| AB | SANDOZ | 200MG | A040104 001 | Nov 30, 1995 |
| AB | TEVA PHARMS | 200MG | A040081 001 | Sep 30, 1994 |
| AB | TWI PHARMS | 200MG | A210441 001 | May 01, 2018 |
| AB | WATSON LABS | 200MG | A040133 001 | Nov 30, 1995 |
| AB | ZYDUS PHARMS USA INC | 200MG | A040657 001 | Sep 21, 2007 |

PLAQUENIL

| | | | | |
|-----------|----------------------------|--------------|--------------------|--|
| AB | +! CONCORDIA PHARMS INC | 200MG | N009768 001 | |
|-----------|----------------------------|--------------|--------------------|--|

HYDROXYPROGESTERONE CAPROATE

SOLUTION;INTRAMUSCULAR

HYDROXYPROGESTERONE CAPROATE

| | | | | |
|-----------|---------------------|------------------------------|--------------------|--------------|
| AP | LUITPOLD | 250MG/ML (250MG/ML) | A210723 001 | Jun 21, 2018 |
| AP | SLAYBACK PHARMA LLC | 1250MG/5ML (250MG/ML) | A210618 001 | Dec 28, 2018 |

MAKENA

| | | | | |
|-----------|--------------------|------------------------------|--------------------|--------------|
| AP | +! AMAG PHARMA USA | 1250MG/5ML (250MG/ML) | N021945 001 | Feb 03, 2011 |
|-----------|--------------------|------------------------------|--------------------|--------------|

MAKENA PRESERVATIVE FREE

| | | | | |
|-----------|--------------------|----------------------------|--------------------|--------------|
| AP | +! AMAG PHARMA USA | 250MG/ML (250MG/ML) | N021945 002 | Feb 19, 2016 |
|-----------|--------------------|----------------------------|--------------------|--------------|

HYDROXYPROGESTERONE CAPROATE

! ASPEN GLOBAL INC 250MG/ML (250MG/ML)

A200271 001 Aug 24, 2015

SOLUTION;SUBCUTANEOUS

MAKENA (AUTOINJECTOR)

+! AMAG PHARMA USA 275MG/1.1ML (250MG/ML)

N021945 004 Feb 14, 2018

HYDROXYPROPYL CELLULOSE

INSERT;OPHTHALMIC

LACRISERT

+! ATON

5MG

N018771 001

HYDROXYUREA

CAPSULE;ORAL

HYDREA

| | | | | |
|-----------|----------------------------|--------------|--------------------|--|
| AB | +! BRISTOL MYERS SQUIBB | 500MG | N016295 001 | |
|-----------|----------------------------|--------------|--------------------|--|

HYDROXYUREA

| | | | | |
|-----------|------|--------------|--------------------|--------------|
| AB | BARR | 500MG | A075143 001 | Oct 16, 1998 |
|-----------|------|--------------|--------------------|--------------|

| | | | | |
|-----------|-----------|--------------|--------------------|--------------|
| AB | PAR PHARM | 500MG | A075340 001 | Feb 24, 1999 |
|-----------|-----------|--------------|--------------------|--------------|

DROXIA

+ BRISTOL MYERS
SQUIBB 200MG

N016295 002 Feb 25, 1998

+ 300MG

N016295 003 Feb 25, 1998

+ 400MG

N016295 004 Feb 25, 1998

TABLET;ORAL

SIKLOS

+ ADDMEDICA SAS 100MG
+ 1GM

N208843 001 Dec 21, 2017

N208843 002 Dec 21, 2017

HYDROXYZINE HYDROCHLORIDE

INJECTABLE;INJECTION

HYDROXYZINE HYDROCHLORIDE

! LUITPOLD 25MG/ML

A087408 001

! 50MG/ML

A087408 002

SYRUP;ORAL

HYDROXYZINE HYDROCHLORIDE

| | | | | |
|-----------|--------------------|-----------------|--------------------|--------------|
| AA | ! HI TECH PHARMA | 10MG/5ML | A040010 001 | Oct 28, 1994 |
| AA | ! LANNETT CO INC | 10MG/5ML | A201674 001 | Aug 21, 2013 |
| AA | ! VINTAGE PHARMS | 10MG/5ML | A040391 001 | Apr 10, 2002 |
| AA | ! WOCKHARDT BIO AG | 10MG/5ML | A087294 001 | Apr 12, 1982 |

PRESCRIPTION DRUG PRODUCT LIST

HYDROXYZINE HYDROCHLORIDE

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE

| | | | | |
|-----------|---------------------|-------------|--------------------|--------------|
| <u>AB</u> | AMNEAL PHARM | <u>10MG</u> | <u>A040808 001</u> | Sep 24, 2008 |
| <u>AB</u> | | <u>25MG</u> | <u>A040808 002</u> | Sep 24, 2008 |
| <u>AB</u> | | <u>50MG</u> | <u>A040808 003</u> | Sep 24, 2008 |
| <u>AB</u> | ECI PHARMS LLC | <u>10MG</u> | <u>A040804 001</u> | Jun 30, 2008 |
| <u>AB</u> | | <u>25MG</u> | <u>A040804 002</u> | Jun 30, 2008 |
| <u>AB</u> | | <u>50MG</u> | <u>A040804 003</u> | Jun 30, 2008 |
| <u>AB</u> | ELITE LABS INC | <u>10MG</u> | <u>A040604 002</u> | Dec 28, 2004 |
| <u>AB</u> | | <u>25MG</u> | <u>A040604 003</u> | Dec 28, 2004 |
| <u>AB</u> | | <u>50MG</u> | <u>A040604 001</u> | Dec 28, 2004 |
| <u>AB</u> | HERITAGE PHARMA | <u>10MG</u> | <u>A204279 001</u> | Aug 20, 2014 |
| <u>AB</u> | | <u>25MG</u> | <u>A204279 002</u> | Aug 20, 2014 |
| <u>AB</u> | | <u>50MG</u> | <u>A204279 003</u> | Aug 20, 2014 |
| <u>AB</u> | HETERO LABS LTD III | <u>10MG</u> | <u>A040805 001</u> | May 29, 2008 |
| <u>AB</u> | | <u>25MG</u> | <u>A040805 002</u> | May 29, 2008 |
| <u>AB</u> | | <u>50MG</u> | <u>A040805 003</u> | May 29, 2008 |
| <u>AB</u> | INVAGEN PHARMS | <u>10MG</u> | <u>A040812 001</u> | Mar 12, 2008 |
| <u>AB</u> | | <u>25MG</u> | <u>A040812 002</u> | Mar 12, 2008 |
| <u>AB</u> | | <u>50MG</u> | <u>A040812 003</u> | Mar 12, 2008 |
| <u>AB</u> | KVK TECH | <u>10MG</u> | <u>A040786 001</u> | Mar 20, 2007 |
| <u>AB</u> | | <u>25MG</u> | <u>A040787 001</u> | Mar 20, 2007 |
| <u>AB</u> | | <u>50MG</u> | <u>A040788 001</u> | Mar 20, 2007 |
| <u>AB</u> | MYLAN | <u>10MG</u> | <u>A091176 001</u> | Jun 07, 2010 |
| <u>AB</u> | | <u>25MG</u> | <u>A091176 002</u> | Jun 07, 2010 |
| <u>AB</u> | | <u>50MG</u> | <u>A091176 003</u> | Jun 07, 2010 |
| <u>AB</u> | NORTHSTAR HLTHCARE | <u>10MG</u> | <u>A040840 002</u> | Mar 31, 2008 |
| <u>AB</u> | | <u>25MG</u> | <u>A040840 003</u> | Mar 31, 2008 |
| <u>AB</u> | | <u>50MG</u> | <u>A040840 001</u> | Mar 31, 2008 |
| <u>AB</u> | NUVO PHARM | <u>10MG</u> | <u>A207120 001</u> | Mar 29, 2017 |
| <u>AB</u> | | <u>50MG</u> | <u>A207122 001</u> | Mar 29, 2017 |
| <u>AB</u> | NUVO PHARMS INC | <u>25MG</u> | <u>A207121 001</u> | Mar 29, 2017 |
| <u>AB</u> | ! PLIVA | <u>10MG</u> | <u>A088617 001</u> | Jan 10, 1986 |
| <u>AB</u> | ! | <u>25MG</u> | <u>A088618 001</u> | Jan 10, 1986 |
| <u>AB</u> | ! | <u>50MG</u> | <u>A088619 001</u> | Jan 10, 1986 |
| <u>AB</u> | PRINSTON INC | <u>10MG</u> | <u>A040579 001</u> | May 27, 2005 |
| <u>AB</u> | | <u>25MG</u> | <u>A040574 001</u> | May 27, 2005 |
| <u>AB</u> | | <u>50MG</u> | <u>A040580 001</u> | May 27, 2005 |

HYDROXYZINE PAMOATE

CAPSULE; ORAL

HYDROXYZINE PAMOATE

| | | | | |
|-----------------|---------------------|-------------------------------|--------------------|--------------|
| <u>AB</u> | BARR | <u>EQ 25MG HYDROCHLORIDE</u> | <u>A088496 001</u> | Jun 15, 1984 |
| <u>AB</u> | | <u>EQ 50MG HYDROCHLORIDE</u> | <u>A088487 001</u> | Jun 15, 1984 |
| <u>AB</u> | HERITAGE PHARMA | <u>EQ 25MG HYDROCHLORIDE</u> | <u>A201507 001</u> | Jun 03, 2013 |
| <u>AB</u> | | <u>EQ 50MG HYDROCHLORIDE</u> | <u>A201507 002</u> | Jun 03, 2013 |
| <u>AB</u> | IMPAX LABS INC | <u>EQ 25MG HYDROCHLORIDE</u> | <u>A040156 001</u> | Jul 15, 1996 |
| <u>AB</u> | | <u>EQ 50MG HYDROCHLORIDE</u> | <u>A040156 002</u> | Jul 15, 1996 |
| <u>AB</u> | SANDOZ | <u>EQ 25MG HYDROCHLORIDE</u> | <u>A087479 001</u> | |
| <u>AB</u> | | <u>EQ 50MG HYDROCHLORIDE</u> | <u>A086183 001</u> | |
| <u>VISTARIL</u> | | | | |
| <u>AB</u> | + PFIZER | <u>EQ 25MG HYDROCHLORIDE</u> | <u>N011459 002</u> | |
| <u>AB</u> | +! | <u>EQ 50MG HYDROCHLORIDE</u> | <u>N011459 004</u> | |
| | HYDROXYZINE PAMOATE | | | |
| | BARR | <u>EQ 100MG HYDROCHLORIDE</u> | <u>A088488 001</u> | Jun 15, 1984 |

IBANDRONATE SODIUM

INJECTABLE; INTRAVENOUS

BONIVA

| | | | | |
|---------------------------|----------------------|------------------------|--------------------|--------------|
| <u>AP</u> | +! ROCHE | <u>EQ 3MG BASE/3ML</u> | <u>N021858 001</u> | Jan 06, 2006 |
| <u>IBANDRONATE SODIUM</u> | | | | |
| <u>AP</u> | ACCORD HLTHCARE | <u>EQ 3MG BASE/3ML</u> | <u>A206058 001</u> | Feb 05, 2016 |
| <u>AP</u> | APOTEX INC | <u>EQ 3MG BASE/3ML</u> | <u>A204222 001</u> | Oct 16, 2015 |
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>EQ 3MG BASE/3ML</u> | <u>A205332 001</u> | Aug 19, 2015 |
| <u>AP</u> | EMCURE PHARMS LTD | <u>EQ 3MG BASE/3ML</u> | <u>A203987 001</u> | Sep 02, 2014 |
| <u>AP</u> | MYLAN LABS LTD | <u>EQ 3MG BASE/3ML</u> | <u>A202671 001</u> | Sep 02, 2014 |
| <u>AP</u> | SAGENT PHARMS | <u>EQ 3MG BASE/3ML</u> | <u>A202235 001</u> | Sep 02, 2014 |
| <u>AP</u> | SUN PHARM INDS LTD | <u>EQ 3MG BASE/3ML</u> | <u>A090853 001</u> | Feb 14, 2014 |
| TABLET; ORAL | | | | |
| <u>BONIVA</u> | | | | |
| <u>AB</u> | +! HOFFMANN LA ROCHE | <u>EQ 150MG BASE</u> | <u>N021455 002</u> | Mar 24, 2005 |

PRESCRIPTION DRUG PRODUCT LIST

IBANDRONATE SODIUM

TABLET; ORAL

IBANDRONATE SODIUM

| | | | | |
|-----------|----------------------|----------------------|--------------------|--------------|
| <u>AB</u> | APOTEX INC | <u>EQ 150MG BASE</u> | <u>A078948 001</u> | Mar 19, 2012 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 150MG BASE</u> | <u>A204502 001</u> | Mar 11, 2016 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 150MG BASE</u> | <u>A078997 001</u> | Apr 30, 2012 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>EQ 150MG BASE</u> | <u>A206887 001</u> | Oct 31, 2017 |
| <u>AB</u> | ORCHID HLTHCARE | <u>EQ 150MG BASE</u> | <u>A078998 001</u> | Mar 19, 2012 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>EQ 150MG BASE</u> | <u>A078996 001</u> | Aug 15, 2012 |
| <u>AB</u> | WATSON LABS TEVA | <u>EQ 150MG BASE</u> | <u>A079003 001</u> | Mar 20, 2012 |

IBRUTINIB

CAPSULE; ORAL

IMBRUVICA

| | | | | |
|---|-------------------|-------|-------------|--------------|
| + | PHARMACYCLICS INC | 70MG | N205552 002 | Dec 20, 2017 |
| + | ! | 140MG | N205552 001 | Nov 13, 2013 |

TABLET; ORAL

IMBRUVICA

| | | | | |
|---|-------------------|-------|-------------|--------------|
| + | PHARMACYCLICS INC | 140MG | N210563 001 | Feb 16, 2018 |
| + | | 280MG | N210563 002 | Feb 16, 2018 |
| + | | 420MG | N210563 003 | Feb 16, 2018 |
| + | ! | 560MG | N210563 004 | Feb 16, 2018 |

IBUPROFEN

SOLUTION; INTRAVENOUS

CALDOLOR

| | | | | | |
|---|---|-------------------|----------------------|-------------|--------------|
| + | ! | CUMBERLAND PHARMS | 800MG/8ML (100MG/ML) | N022348 002 | Jun 11, 2009 |
|---|---|-------------------|----------------------|-------------|--------------|

SUSPENSION; ORAL

IBUPROFEN

| | | | | | |
|-----------|---|----------------------|------------------|--------------------|--------------|
| <u>AB</u> | ! | ACTAVIS MID ATLANTIC | <u>100MG/5ML</u> | <u>A074978 001</u> | Mar 25, 1998 |
| <u>AB</u> | | HI-TECH PHARMACAL | <u>100MG/5ML</u> | <u>A205647 001</u> | Nov 03, 2016 |
| <u>AB</u> | | PERRIGO R AND D | <u>100MG/5ML</u> | <u>A076925 001</u> | Sep 23, 2004 |
| <u>AB</u> | | TARO | <u>100MG/5ML</u> | <u>A209204 001</u> | Jun 23, 2017 |
| | | AUROBINDO PHARMA LTD | 100MG/5ML | A209178 001 | Feb 16, 2018 |

TABLET; ORAL

IBU-TAB

| | | | | | |
|-----------|--|------|--------------|--------------------|--------------|
| <u>AB</u> | | ALRA | <u>400MG</u> | <u>A071058 001</u> | Aug 11, 1988 |
| <u>AB</u> | | | <u>600MG</u> | <u>A071059 001</u> | Aug 11, 1988 |

IBUPROFEN

| | | | | | |
|-----------|---|--------------------|--------------|--------------------|--------------|
| <u>AB</u> | | AMNEAL PHARMS NY | <u>400MG</u> | <u>A071334 001</u> | Nov 25, 1986 |
| <u>AB</u> | | | <u>400MG</u> | <u>A078558 001</u> | Jun 18, 2007 |
| <u>AB</u> | | | <u>600MG</u> | <u>A071335 001</u> | Nov 25, 1986 |
| <u>AB</u> | | | <u>600MG</u> | <u>A078558 002</u> | Jun 18, 2007 |
| <u>AB</u> | | | <u>800MG</u> | <u>A071935 001</u> | Oct 13, 1987 |
| <u>AB</u> | | | <u>800MG</u> | <u>A078558 003</u> | Jun 18, 2007 |
| <u>AB</u> | | CONTRACT PHARMACAL | <u>400MG</u> | <u>A071268 002</u> | Oct 15, 1986 |
| <u>AB</u> | | | <u>600MG</u> | <u>A071268 001</u> | Oct 15, 1986 |
| <u>AB</u> | | | <u>800MG</u> | <u>A071268 003</u> | Jul 01, 1988 |
| <u>AB</u> | | DR REDDYS LA | <u>400MG</u> | <u>A075682 001</u> | Nov 14, 2001 |
| <u>AB</u> | | | <u>600MG</u> | <u>A075682 002</u> | Nov 14, 2001 |
| <u>AB</u> | ! | | <u>800MG</u> | <u>A075682 003</u> | Nov 14, 2001 |
| <u>AB</u> | | DR REDDYS LABS INC | <u>400MG</u> | <u>A076112 001</u> | Oct 31, 2001 |
| <u>AB</u> | | | <u>600MG</u> | <u>A076112 002</u> | Oct 31, 2001 |
| <u>AB</u> | | | <u>800MG</u> | <u>A076112 003</u> | Oct 31, 2001 |
| <u>AB</u> | | GRANULES INDIA LTD | <u>400MG</u> | <u>A091625 001</u> | Sep 15, 2015 |
| <u>AB</u> | | | <u>600MG</u> | <u>A091625 002</u> | Sep 15, 2015 |
| <u>AB</u> | | | <u>800MG</u> | <u>A091625 003</u> | Sep 15, 2015 |
| <u>AB</u> | | HEC PHARM | <u>400MG</u> | <u>A204062 001</u> | Sep 10, 2018 |
| <u>AB</u> | | | <u>600MG</u> | <u>A204062 002</u> | Sep 10, 2018 |
| <u>AB</u> | | | <u>800MG</u> | <u>A204062 003</u> | Sep 10, 2018 |
| <u>AB</u> | | MARKSANS PHARMA | <u>400MG</u> | <u>A090796 001</u> | Dec 21, 2010 |
| <u>AB</u> | | | <u>600MG</u> | <u>A090796 002</u> | Dec 21, 2010 |
| <u>AB</u> | | | <u>800MG</u> | <u>A090796 003</u> | Dec 21, 2010 |
| <u>AB</u> | | PERRIGO R AND D | <u>400MG</u> | <u>A077114 001</u> | Jul 18, 2005 |
| <u>AB</u> | | | <u>600MG</u> | <u>A077114 002</u> | Jul 18, 2005 |
| <u>AB</u> | | | <u>800MG</u> | <u>A077114 003</u> | Jul 18, 2005 |
| <u>AB</u> | | SHANDONG XINHUA | <u>400MG</u> | <u>A202413 001</u> | Nov 23, 2016 |
| <u>AB</u> | | | <u>600MG</u> | <u>A202413 002</u> | Nov 23, 2016 |
| <u>AB</u> | | | <u>800MG</u> | <u>A202413 003</u> | Nov 23, 2016 |
| <u>AB</u> | | STRIDES PHARMA | <u>400MG</u> | <u>A078329 001</u> | Feb 05, 2009 |

PRESCRIPTION DRUG PRODUCT LIST

IBUPROFEN

TABLET; ORAL

IBUPROFEN

| | | | | |
|-----------|----------------|--------------|--------------------|--------------|
| <u>AB</u> | | <u>600MG</u> | <u>A078329 002</u> | Feb 05, 2009 |
| <u>AB</u> | | <u>800MG</u> | <u>A078329 003</u> | Feb 05, 2009 |
| <u>AB</u> | VINTAGE PHARMS | <u>400MG</u> | <u>A071644 001</u> | Feb 01, 1988 |

IBUPROFEN LYSINE

INJECTABLE; INTRAVENOUS

IBUPROFEN LYSINE

| | | | | |
|-----------|------------------|-------------------------------------------|--------------------|--------------|
| <u>AP</u> | X-GEN PHARMS INC | <u>EQ 20MG BASE/2ML (EQ 10MG BASE/ML)</u> | <u>A202402 001</u> | Mar 30, 2016 |
|-----------|------------------|-------------------------------------------|--------------------|--------------|

NEOPROFEN

| | | | | |
|-----------|-------------------|-------------------------------------------|--------------------|--------------|
| <u>AP</u> | +! RECORDATI RARE | <u>EQ 20MG BASE/2ML (EQ 10MG BASE/ML)</u> | <u>N021903 001</u> | Apr 13, 2006 |
|-----------|-------------------|-------------------------------------------|--------------------|--------------|

IBUPROFEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE HYDROCHLORIDE AND IBUPROFEN

| | | | | |
|-----------|-------------------|-------------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS ELIZABETH | <u>400MG; 5MG</u> | <u>A078769 001</u> | Jan 04, 2008 |
| <u>AB</u> | ! BARR LABS INC | <u>400MG; 5MG</u> | <u>A078316 001</u> | Nov 29, 2007 |

IBUTILIDE FUMARATE

INJECTABLE; INJECTION

CORVERT

| | | | | |
|-----------|-------------------------|-----------------|--------------------|--------------|
| <u>AP</u> | +! PHARMACIA AND UPJOHN | <u>0.1MG/ML</u> | <u>N020491 001</u> | Dec 28, 1995 |
|-----------|-------------------------|-----------------|--------------------|--------------|

IBUTILIDE FUMARATE

| | | | | |
|-----------|---------------------|-----------------|--------------------|--------------|
| <u>AP</u> | LUITPOLD | <u>0.1MG/ML</u> | <u>A090240 001</u> | Jan 11, 2010 |
| <u>AP</u> | MYLAN INSTITUTIONAL | <u>0.1MG/ML</u> | <u>A090643 001</u> | Jan 11, 2010 |

ICATIBANT ACETATE

INJECTABLE; SUBCUTANEOUS

FIRAZYR

| | | | | |
|----|---------------------|-------------------------------------------|--------------------|--------------|
| +! | SHIRE ORPHAN THERAP | <u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u> | <u>N022150 001</u> | Aug 25, 2011 |
|----|---------------------|-------------------------------------------|--------------------|--------------|

ICODEXTRINSOLUTION; INTRAPERITONEAL
EXTRANEAL

| | | | | |
|----|-----------------|--------------------|--------------------|--------------|
| +! | BAXTER HLTHCARE | <u>7.5GM/100ML</u> | <u>N021321 001</u> | Dec 20, 2002 |
|----|-----------------|--------------------|--------------------|--------------|

ICOSAPENT ETHYL

CAPSULE; ORAL

VASCEPA

| | | | | |
|----|---------------|--------------|--------------------|--------------|
| + | AMARIN PHARMS | <u>500MG</u> | <u>N202057 002</u> | Feb 16, 2017 |
| +! | | <u>1GM</u> | <u>N202057 001</u> | Jul 26, 2012 |

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDAMYCIN PFS

| | | | | |
|-----------|-------------------------|---------------|--------------------|--------------|
| <u>AP</u> | +! PHARMACIA AND UPJOHN | <u>1MG/ML</u> | <u>N050734 001</u> | Feb 17, 1997 |
|-----------|-------------------------|---------------|--------------------|--------------|

IDARUBICIN HYDROCHLORIDE

| | | | | |
|-----------|----------------------|---------------|--------------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>1MG/ML</u> | <u>A065440 001</u> | Aug 04, 2009 |
| <u>AP</u> | MYLAN LABS LTD | <u>1MG/ML</u> | <u>A200144 001</u> | Oct 11, 2012 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>1MG/ML</u> | <u>A065275 001</u> | Dec 14, 2006 |

| | | | | |
|-----------|--|---------------|--------------------|--------------|
| <u>AP</u> | | <u>1MG/ML</u> | <u>A065288 001</u> | May 15, 2007 |
|-----------|--|---------------|--------------------|--------------|

IDARUBICIN HYDROCHLORIDE PFS

| | | | | |
|-----------|-----------------|---------------|--------------------|--------------|
| <u>AP</u> | TEVA PHARMS USA | <u>1MG/ML</u> | <u>A065036 001</u> | May 01, 2002 |
|-----------|-----------------|---------------|--------------------|--------------|

IDELALISIB

TABLET; ORAL

ZYDELIG

| | | | | |
|----|---------------------|--------------|--------------------|--------------|
| + | GILEAD SCIENCES INC | <u>100MG</u> | <u>N205858 001</u> | Jul 23, 2014 |
| +! | | <u>150MG</u> | <u>N205858 002</u> | Jul 23, 2014 |

IFOSFAMIDE

INJECTABLE; INJECTION

IFEX

| | | | | | |
|-----------|---|-----------------|-----------------|--------------------|--------------|
| <u>AP</u> | + | BAXTER HLTHCARE | <u>1GM/VIAL</u> | <u>N019763 001</u> | Dec 30, 1988 |
| <u>AP</u> | + | | <u>3GM/VIAL</u> | <u>N019763 002</u> | Dec 30, 1988 |

IFOSFAMIDE

| | | | | | |
|-----------|---|--------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | ! | FRESENIUS KABI USA | <u>1GM/VIAL</u> | <u>A076078 001</u> | May 28, 2002 |
| <u>AP</u> | ! | | <u>3GM/VIAL</u> | <u>A076078 002</u> | May 28, 2002 |
| <u>AP</u> | | MYLAN LABS LTD | <u>1GM/20ML (50MG/ML)</u> | <u>A201689 001</u> | Nov 26, 2012 |
| <u>AP</u> | | | <u>3GM/60ML (50MG/ML)</u> | <u>A201689 002</u> | Nov 26, 2012 |
| <u>AP</u> | ! | TEVA PHARMS USA | <u>1GM/20ML (50MG/ML)</u> | <u>A076657 001</u> | Apr 04, 2007 |
| <u>AP</u> | ! | | <u>3GM/60ML (50MG/ML)</u> | <u>A076657 002</u> | Apr 04, 2007 |

PRESCRIPTION DRUG PRODUCT LISTIFOSFAMIDE

INJECTABLE; INJECTION

IFOSFAMIDE

| | | | | |
|-----------|-------------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | WEST-WARD PHARMS INT | <u>1GM/20ML (50MG/ML)</u> | <u>A076619 001</u> | Jun 29, 2011 |
| <u>AP</u> | | <u>3GM/60ML (50MG/ML)</u> | <u>A076619 002</u> | Jun 29, 2011 |

ILOPERIDONE

TABLET; ORAL

FANAPT

| | | | | | |
|-----------|----|------------------|-------------|--------------------|--------------|
| <u>AB</u> | +! | VANDA PHARMS INC | <u>1MG</u> | <u>N022192 001</u> | May 06, 2009 |
| <u>AB</u> | + | | <u>2MG</u> | <u>N022192 002</u> | May 06, 2009 |
| <u>AB</u> | + | | <u>4MG</u> | <u>N022192 003</u> | May 06, 2009 |
| <u>AB</u> | + | | <u>6MG</u> | <u>N022192 004</u> | May 06, 2009 |
| <u>AB</u> | + | | <u>8MG</u> | <u>N022192 005</u> | May 06, 2009 |
| <u>AB</u> | + | | <u>10MG</u> | <u>N022192 006</u> | May 06, 2009 |
| <u>AB</u> | + | | <u>12MG</u> | <u>N022192 007</u> | May 06, 2009 |

ILOPERIDONE

| | | | | | |
|-----------|--|-------------------|-------------|--------------------|--------------|
| <u>AB</u> | | INVENTIA HLTHCARE | <u>1MG</u> | <u>A207231 001</u> | Nov 28, 2016 |
| <u>AB</u> | | | <u>2MG</u> | <u>A207231 002</u> | Nov 28, 2016 |
| <u>AB</u> | | | <u>4MG</u> | <u>A207231 003</u> | Nov 28, 2016 |
| <u>AB</u> | | | <u>6MG</u> | <u>A207231 004</u> | Nov 28, 2016 |
| <u>AB</u> | | | <u>8MG</u> | <u>A207231 005</u> | Nov 28, 2016 |
| <u>AB</u> | | | <u>10MG</u> | <u>A207231 006</u> | Nov 28, 2016 |
| <u>AB</u> | | | <u>12MG</u> | <u>A207231 007</u> | Nov 28, 2016 |

ILOPROST

SOLUTION; INHALATION

VENTAVIS

| | | | | | |
|--|----|---------------------|---------------------|-------------|--------------|
| | +! | ACTELION PHARMS LTD | 10MCG/ML (10MCG/ML) | N021779 002 | Dec 08, 2005 |
| | +! | | 20MCG/ML (20MCG/ML) | N021779 003 | Aug 07, 2009 |

IMATINIB MESYLATE

TABLET; ORAL

GLEEVEC

| | | | | | |
|-----------|----|----------|----------------------|--------------------|--------------|
| <u>AB</u> | + | NOVARTIS | <u>EQ 100MG BASE</u> | <u>N021588 001</u> | Apr 18, 2003 |
| <u>AB</u> | +! | | <u>EQ 400MG BASE</u> | <u>N021588 002</u> | Apr 18, 2003 |

IMATINIB MESYLATE

| | | | | | |
|-----------|--|-------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | APOTEX INC | <u>EQ 100MG BASE</u> | <u>A079179 001</u> | Aug 05, 2016 |
| <u>AB</u> | | | <u>EQ 400MG BASE</u> | <u>A079179 002</u> | Aug 05, 2016 |
| <u>AB</u> | | DR REDDYS LABS LTD | <u>EQ 100MG BASE</u> | <u>A206547 001</u> | Aug 13, 2018 |
| <u>AB</u> | | | <u>EQ 400MG BASE</u> | <u>A206547 002</u> | Aug 13, 2018 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>EQ 100MG BASE</u> | <u>A204644 001</u> | Jun 21, 2017 |
| <u>AB</u> | | | <u>EQ 400MG BASE</u> | <u>A204644 002</u> | Jun 21, 2017 |
| <u>AB</u> | | SUN PHARMA GLOBAL | <u>EQ 100MG BASE</u> | <u>A078340 001</u> | Dec 03, 2015 |
| <u>AB</u> | | | <u>EQ 400MG BASE</u> | <u>A078340 002</u> | Dec 03, 2015 |
| <u>AB</u> | | TEVA PHARMS USA | <u>EQ 100MG BASE</u> | <u>A204285 001</u> | Aug 04, 2016 |
| <u>AB</u> | | | <u>EQ 400MG BASE</u> | <u>A204285 002</u> | Aug 04, 2016 |
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>EQ 100MG BASE</u> | <u>A207586 001</u> | Jul 13, 2018 |
| <u>AB</u> | | | <u>EQ 400MG BASE</u> | <u>A207586 002</u> | Jul 13, 2018 |

IMIGLUCERASE

INJECTABLE; INJECTION

CEREZYME

| | | | | | |
|--|----|---------|----------------|-------------|--------------|
| | + | GENZYME | 200 UNITS/VIAL | N020367 001 | May 23, 1994 |
| | +! | | 400 UNITS/VIAL | N020367 002 | Sep 22, 1999 |

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

| | | | | | |
|-----------|--|--------------------|-------------|--------------------|--------------|
| <u>AB</u> | | LEADING PHARMA LLC | <u>10MG</u> | <u>A040903 001</u> | Oct 24, 2012 |
| <u>AB</u> | | | <u>25MG</u> | <u>A040903 002</u> | Oct 24, 2012 |
| <u>AB</u> | | | <u>50MG</u> | <u>A040903 003</u> | Oct 24, 2012 |
| <u>AB</u> | | LUPIN LTD | <u>10MG</u> | <u>A090441 002</u> | Mar 11, 2010 |
| <u>AB</u> | | | <u>25MG</u> | <u>A090441 003</u> | Mar 11, 2010 |
| <u>AB</u> | | | <u>50MG</u> | <u>A090441 001</u> | Mar 11, 2010 |
| <u>AB</u> | | PAR PHARM | <u>10MG</u> | <u>A088292 001</u> | Oct 21, 1983 |
| <u>AB</u> | | | <u>25MG</u> | <u>A088262 001</u> | Oct 21, 1983 |
| <u>AB</u> | | | <u>50MG</u> | <u>A088276 001</u> | Oct 21, 1983 |
| <u>AB</u> | | SANDOZ | <u>10MG</u> | <u>A084936 002</u> | |
| <u>AB</u> | | | <u>25MG</u> | <u>A083745 001</u> | |
| <u>AB</u> | | | <u>50MG</u> | <u>A084937 001</u> | |
| <u>AB</u> | | SPECGX LLC | <u>10MG</u> | <u>A087846 002</u> | May 22, 1984 |
| <u>AB</u> | | | <u>25MG</u> | <u>A087846 003</u> | May 22, 1984 |

PRESCRIPTION DRUG PRODUCT LIST

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

| | | | | |
|-----------------|--------------------------|-------------|--------------------|--------------|
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>10MG</u> | <u>A081048 001</u> | Jun 05, 1990 |
| <u>AB</u> | | <u>25MG</u> | <u>A081049 001</u> | Jun 05, 1990 |
| <u>AB</u> | | <u>50MG</u> | <u>A081050 001</u> | Jun 05, 1990 |
| <u>TOFRANIL</u> | | | | |
| <u>AB</u> | ! SPECGX LLC | <u>50MG</u> | <u>A087846 001</u> | May 22, 1984 |
| | IMIPRAMINE HYDROCHLORIDE | | | |
| | OXFORD PHARMS | 10MG | A040751 003 | Feb 28, 2008 |
| | | 25MG | A040751 002 | Feb 28, 2008 |

IMIPRAMINE PAMOATE

CAPSULE; ORAL

IMIPRAMINE PAMOATE

| | | | | |
|-----------|---------------------------|-------------------------------|--------------------|--------------|
| <u>AB</u> | LUPIN LTD | <u>EQ 75MG HYDROCHLORIDE</u> | <u>A090444 001</u> | Apr 16, 2010 |
| <u>AB</u> | | <u>EQ 100MG HYDROCHLORIDE</u> | <u>A090444 002</u> | Apr 16, 2010 |
| <u>AB</u> | | <u>EQ 125MG HYDROCHLORIDE</u> | <u>A090444 003</u> | Apr 16, 2010 |
| <u>AB</u> | | <u>EQ 150MG HYDROCHLORIDE</u> | <u>A090444 004</u> | Apr 16, 2010 |
| <u>AB</u> | ! WEST-WARD PHARMS INT | <u>EQ 75MG HYDROCHLORIDE</u> | <u>A091099 001</u> | Apr 16, 2010 |
| <u>AB</u> | | <u>EQ 100MG HYDROCHLORIDE</u> | <u>A091099 002</u> | Apr 16, 2010 |
| <u>AB</u> | | <u>EQ 125MG HYDROCHLORIDE</u> | <u>A091099 003</u> | Apr 16, 2010 |
| <u>AB</u> | | <u>EQ 150MG HYDROCHLORIDE</u> | <u>A091099 004</u> | Apr 16, 2010 |

IMIQUIMOD

CREAM; TOPICAL

ALDARA

| | | | | |
|-----------|------------|-----------|--------------------|--------------|
| <u>AB</u> | +! MEDICIS | <u>5%</u> | <u>N020723 001</u> | Feb 27, 1997 |
|-----------|------------|-----------|--------------------|--------------|

IMIQUIMOD

| | | | | |
|-----------|-------------------|-----------|--------------------|--------------|
| <u>AB</u> | ANDA REPOSITORY | <u>5%</u> | <u>A091044 001</u> | Feb 28, 2011 |
| <u>AB</u> | APOTEX INC | <u>5%</u> | <u>A091308 001</u> | Apr 06, 2012 |
| <u>AB</u> | FOUGERA PHARMS | <u>5%</u> | <u>A078548 001</u> | Feb 25, 2010 |
| <u>AB</u> | GLENMARK GENERICS | <u>5%</u> | <u>A201994 001</u> | Mar 06, 2012 |
| <u>AB</u> | PERRIGO ISRAEL | <u>5%</u> | <u>A078837 001</u> | Sep 07, 2010 |
| <u>AB</u> | TARO | <u>5%</u> | <u>A200173 001</u> | Apr 15, 2011 |
| ZYCLARA | | | | |
| | +! MEDICIS | 2.5% | N022483 002 | Jul 15, 2011 |
| | +! | 3.75% | N022483 001 | Mar 25, 2010 |

INAMRINONE LACTATE

INJECTABLE; INJECTION

AMRINONE LACTATE

| | | | | |
|---|-------------------------|----------------|-------------|--------------|
| ! | WEST-WARD PHARMS INT | EQ 5MG BASE/ML | A075513 001 | May 09, 2000 |
|---|-------------------------|----------------|-------------|--------------|

INDACATEROL MALEATE

POWDER; INHALATION

ARCAPTA NEOHALER

| | | | | |
|----|---------------------|---------------|-------------|--------------|
| +! | SUNOVION PHARMS INC | EQ 75MCG BASE | N022383 001 | Jul 01, 2011 |
|----|---------------------|---------------|-------------|--------------|

INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

| | | | | |
|-----------|---------------------|---------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS ELIZABETH | <u>1.25MG</u> | <u>A074722 001</u> | Jun 17, 1996 |
| <u>AB</u> | | <u>2.5MG</u> | <u>A074722 002</u> | Jun 17, 1996 |
| <u>AB</u> | AMERIGEN PHARMS LTD | <u>1.25MG</u> | <u>A075201 001</u> | Dec 04, 1998 |
| <u>AB</u> | | <u>2.5MG</u> | <u>A075201 002</u> | Dec 04, 1998 |
| <u>AB</u> | ANI PHARMS INC | <u>1.25MG</u> | <u>A074299 002</u> | Apr 29, 1996 |
| <u>AB</u> | | <u>2.5MG</u> | <u>A074299 001</u> | Jul 27, 1995 |
| <u>AB</u> | MYLAN | <u>1.25MG</u> | <u>A074461 002</u> | Mar 26, 1997 |
| <u>AB</u> | ! | <u>2.5MG</u> | <u>A074461 001</u> | Mar 27, 1996 |

INDINAVIR SULFATE

CAPSULE; ORAL

CRIXIVAN

| | | | | |
|----|-------------------|---------------|-------------|--------------|
| + | MERCK SHARP DOHME | EQ 200MG BASE | N020685 003 | Mar 13, 1996 |
| +! | | EQ 400MG BASE | N020685 001 | Mar 13, 1996 |

PRESCRIPTION DRUG PRODUCT LIST

INDIUM IN-111 CHLORIDE

INJECTABLE; INJECTION

INDICLOR

+! GE HEALTHCARE 2mCi/0.2ML

N019862 001 Dec 29, 1992

INDIUM IN 111 CHLORIDE

+! MALLINKRODT NUCLEAR 5mCi/0.5ML

N019841 001 Sep 27, 1994

INDIUM IN-111 OXYQUINOLINE

INJECTABLE; INJECTION

INDIUM IN 111 OXYQUINOLINE

+! GE HEALTHCARE 1mCi/ML

N019044 001 Dec 24, 1985

INDIUM IN-111 PENTETATE DISODIUM

INJECTABLE; INTRATHECAL

MPI INDIUM DTPA IN 111

+! GE HEALTHCARE 1mCi/ML

N017707 001 Feb 18, 1982

INDIUM IN-111 PENTETREOTIDE KIT

INJECTABLE; INJECTION

OCTREOSCAN

+! MALLINKRODT NUCLEAR 3mCi/ML

N020314 001 Jun 02, 1994

INDOCYANINE GREEN

INJECTABLE; INJECTION

IC-GREEN**AP** +! AKORN **25MG/VIAL** **N011525 001**INDOCYANINE GREEN**AP** DIAGNOSTIC GREEN **25MG/VIAL** **A040811 001** Nov 21, 2007

POWDER; INTRAVENOUS, INTERSTITIAL

SPY AGENT GREEN KIT

+! NOVADAQ TECH 25MG/VIAL

N211580 001 Nov 21, 2018

INDOMETHACIN

CAPSULE; ORAL

INDOMETHACIN**AB** GLENMARK GENERICS **25MG** **A091276 001** Dec 22, 2010**AB** **50MG** **A091276 002** Dec 22, 2010**AB** HERITAGE PHARMS INC **25MG** **N018851 001** May 18, 1984**AB** **50MG** **N018851 002** May 18, 1984**AB** HETERO LABS LTD III **25MG** **A091240 001** Apr 12, 2011**AB** **50MG** **A091240 002** Apr 12, 2011**AB** IVAX SUB TEVA **25MG** **A070719 001** Feb 12, 1986

PHARMS

AB **50MG** **A070756 001** Feb 12, 1986**AB** JUBILANT GENERICS **25MG** **A205215 001** Aug 25, 2017**AB** **50MG** **A205215 002** Aug 25, 2017**AB** MYLAN **25MG** **N018858 001** Apr 20, 1984**AB** ! **50MG** **A070624 001** Sep 04, 1985**AB** SANDOZ **25MG** **A070673 001** Apr 29, 1987**AB** **50MG** **A070674 001** Apr 29, 1987**AB** SUN PHARM INDS INC **25MG** **A091401 001** Mar 28, 2013**AB** **50MG** **A091401 002** Mar 28, 2013**AB** ZYDUS PHARMS USA **25MG** **A090403 001** Nov 15, 2010

INC

AB **50MG** **A090403 002** Nov 15, 2010

TIVORBEX

+ IROKO PHARMS LLC 20MG

N204768 001 Feb 24, 2014

+! 40MG

N204768 002 Feb 24, 2014

CAPSULE, EXTENDED RELEASE; ORAL

INDOMETHACIN**AB** AMNEAL PHARMS **75MG** **A091549 001** Dec 01, 2010**AB** AUROBINDO PHARMA **75MG** **A204243 001** Dec 27, 2016

LTD

AB AVANTHI INC **75MG** **A079175 001** Mar 06, 2009**AB** CHARTWELL RX **75MG** **A200529 001** Nov 30, 2010**AB** GLENMARK PHARMS LTD **75MG** **A203501 001** Jun 22, 2017**AB** HETERO LABS LTD III **75MG** **A201807 001** Sep 28, 2012**AB** JUBILANT GENERICS **75MG** **A202706 001** Oct 05, 2015**AB** MYLAN PHARMS INC **75MG** **A202139 001** Mar 20, 2014**AB** NOVAST LABS **75MG** **A204853 001** May 08, 2017**AB** ! SANDOZ **75MG** **A074464 001** May 28, 1998**AB** ZYDUS PHARMS USA **75MG** **A202711 001** Sep 25, 2017

INC

PRESCRIPTION DRUG PRODUCT LIST

INDOMETHACIN

INJECTABLE; INJECTION

INDOMETHACIN

+! FRESENIUS KABI USA EQ 1MG BASE/VIAL N022536 001 Mar 17, 2010

SUPPOSITORY; RECTAL

INDOMETHACIN

! G AND W LABS 50MG A073314 001 Aug 31, 1992

SUSPENSION; ORAL

INDOCIN

+! IROKO PHARMS 25MG/5ML N018332 001 Oct 10, 1985

INDOMETHACIN SODIUM

INJECTABLE; INJECTION

INDOCINAP +! RECORDATI RARE EQ 1MG BASE/VIAL N018878 001 Jan 30, 1985INDOMETHACIN SODIUMAP HOSPIRA INC EQ 1MG BASE/VIAL A204118 001 Apr 19, 2016AP NAVINTA LLC EQ 1MG BASE/VIAL A206561 001 Jul 19, 2017AP WEST-WARD PHARMS EQ 1MG BASE/VIAL A078713 001 Jul 16, 2008
INTINGENOL MEBUTATE

GEL; TOPICAL

INGENOL MEBUTATEAB PERRIGO UK FINCO 0.015% A209018 001 Jan 07, 2019AB 0.05% A209019 001 Jan 09, 2019PICATOAB +! LEO LABS 0.015% N202833 001 Jan 23, 2012AB +! 0.05% N202833 002 Jan 23, 2012INOTERSEN SODIUM

SOLUTION; SUBCUTANEOUS

TEGSEDI

+! AKCEA THERAPS EQ 284MG BASE/1.5ML (EQ 189.3MG N211172 001 Oct 05, 2018

BASE/ML)

INSULIN ASPART

SOLUTION; INTRAVENOUS, SUBCUTANEOUS

FIASP

+! NOVO 1000 UNITS/10ML (100 UNITS/ML) N208751 001 Sep 29, 2017

SOLUTION; SUBCUTANEOUS

FIASP FLEXTOUCH

+! NOVO 300 UNITS/3ML (100 UNITS/ML) N208751 002 Sep 29, 2017

INSULIN ASPART PROTAMINE RECOMBINANT; INSULIN ASPART RECOMBINANT

INJECTABLE; SUBCUTANEOUS

NOVOLOG MIX 70/30

+! NOVO NORDISK INC 700 UNITS/10ML; 300 UNITS/10ML (70 N021172 001 Nov 01, 2001

UNITS/ML; 30 UNITS/ML)

NOVOLOG MIX 70/30 FLEXPEN

+! NOVO NORDISK INC 210 UNITS/3ML; 90 UNITS/3ML (70 N021172 004 May 03, 2002

UNITS/ML; 30 UNITS/ML)

INSULIN ASPART RECOMBINANT

INJECTABLE; SUBCUTANEOUS

NOVOLOG

+! NOVO NORDISK INC 1000 UNITS/10ML (100 UNITS/ML) N020986 001 Jun 07, 2000

NOVOLOG FLEXPEN

+! NOVO NORDISK INC 300 UNITS/3ML (100 UNITS/ML) N020986 003 Jan 19, 2001

NOVOLOG PENFILL

+! NOVO NORDISK INC 300 UNITS/3ML (100 UNITS/ML) N020986 002 Jun 07, 2000

INSULIN ASPART; INSULIN DEGLUDEC

SOLUTION; SUBCUTANEOUS

RYZODEG 70/30

+! NOVO 90 UNITS/3ML; 210 UNITS/3ML (30 N203313 001 Sep 25, 2015

UNITS/ML; 70 UNITS/ML)

INSULIN DEGLUDEC

SOLUTION; SUBCUTANEOUS

TRESIBA

+ NOVO 300 UNITS/3ML (100 UNITS/ML) N203314 001 Sep 25, 2015

+! 600 UNITS/3ML (200 UNITS/ML) N203314 002 Sep 25, 2015

PRESCRIPTION DRUG PRODUCT LIST

INSULIN DEGLUDEC; LIRAGLUTIDE

SOLUTION;SUBCUTANEOUS

XULTOPHY 100/3.6

+! NOVO

300 UNITS/3ML;10.8MG/3ML (100
UNITS/ML;3.6MG/ML)

N208583 001 Nov 21, 2016

INSULIN DETEMIR RECOMBINANT

INJECTABLE;SUBCUTANEOUS

LEVEMIR

+! NOVO NORDISK INC

1000 UNITS/10ML (100 UNITS/ML)

N021536 001 Jun 16, 2005

LEVEMIR FLEXTOUCH

+! NOVO NORDISK INC

300 UNITS/3ML (100 UNITS/ML)

N021536 005 Oct 31, 2013

INSULIN GLARGINE

SOLUTION;SUBCUTANEOUS

BASAGLAR

ELI LILLY AND CO

300 UNITS/3ML (100 UNITS/ML)

N205692 001 Dec 16, 2015

INSULIN GLARGINE RECOMBINANT

INJECTABLE;INJECTION

LANTUS

+! SANOFI AVENTIS US

100 UNITS/ML

N021081 001 Apr 20, 2000

LANTUS SOLOSTAR

+! SANOFI AVENTIS US

300 UNITS/3ML (100 UNITS/ML)

N021081 002 Apr 27, 2007

SOLUTION;SUBCUTANEOUS

TOUJEO MAX SOLOSTAR

+! SANOFI US SERVICES

900 UNITS/3ML (300 UNITS/ML)

N206538 002 Mar 26, 2018

TOUJEO SOLOSTAR

+! SANOFI US SERVICES

450 UNITS/1.5ML (300 UNITS/ML)

N206538 001 Feb 25, 2015

INSULIN GLARGINE; LIXISENATIDE

SOLUTION;SUBCUTANEOUS

SOLIQUA 100/33

+! SANOFI-AVENTIS US

300 UNITS/3ML;99MCG/3ML (100
UNITS/ML;33MCG/ML)

N208673 001 Nov 21, 2016

INSULIN GLULISINE RECOMBINANT

INJECTABLE;INTRAVENOUS, SUBCUTANEOUS

APIDRA

+! SANOFI AVENTIS US

1000 UNITS/10ML (100 UNITS/ML)

N021629 001 Apr 16, 2004

INJECTABLE;SUBCUTANEOUS

APIDRA SOLOSTAR

+ SANOFI AVENTIS US

300 UNITS/3ML

N021629 003 Feb 24, 2009

INSULIN HUMAN

SOLUTION;SUBCUTANEOUS

HUMULIN R

+! LILLY

10000 UNITS/20ML (500 UNITS/ML)

N018780 004 Mar 31, 1994

HUMULIN R KWIKPEN

+! LILLY

1500 UNITS/3ML (500 UNITS/ML)

N018780 002 Dec 29, 2015

INSULIN LISPRO

SOLUTION;INTRAVENOUS, SUBCUTANEOUS

ADMELOG

+ SANOFI-AVENTIS US

300 UNITS/3ML (100 UNITS/ML)

N209196 003 Oct 19, 2018

+ 1000 UNITS/10ML (100 UNITS/ML)

N209196 001 Dec 11, 2017

ADMELOG SOLOSTAR

+ SANOFI-AVENTIS US

300 UNITS/3ML (100 UNITS/ML)

N209196 002 Dec 11, 2017

INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT

INJECTABLE;INJECTION

HUMALOG MIX 50/50

+! LILLY

50 UNITS/ML;50 UNITS/ML

N021018 001 Dec 22, 1999

HUMALOG MIX 50/50 KWIKPEN

+! LILLY

50 UNITS/ML;50 UNITS/ML

N021018 002 Sep 06, 2007

HUMALOG MIX 75/25

+! LILLY

75 UNITS/ML;25 UNITS/ML

N021017 001 Dec 22, 1999

HUMALOG MIX 75/25 KWIKPEN

+! LILLY

75 UNITS/ML;25 UNITS/ML

N021017 002 Sep 06, 2007

INSULIN LISPRO RECOMBINANT

INJECTABLE;INJECTION

HUMALOG

+! LILLY

100 UNITS/ML

N020563 001 Jun 14, 1996

HUMALOG KWIKPEN

+! LILLY

100 UNITS/ML

N020563 003 Sep 06, 2007

PRESCRIPTION DRUG PRODUCT LIST

INSULIN LISPRO RECOMBINANT

SOLUTION; SUBCUTANEOUS

HUMALOG KWIKPEN

+! ELI LILLY AND CO 200 UNITS/ML N205747 001 May 26, 2015

INSULIN RECOMBINANT HUMAN

POWDER; INHALATION

AFREZZA

+ MANNKIND 4 UNITS/INH N022472 001 Jun 27, 2014

+! 8 UNITS/INH N022472 002 Jun 27, 2014

+ 12 UNITS/INH N022472 003 Apr 17, 2015

IOBENGUANE I-131

SOLUTION; INTRAVENOUS

AZEDRA

+! PROGENICS PHARMS 15mCi/ML N209607 001 Jul 30, 2018
INCIOBENGUANE SULFATE I-123

SOLUTION; INTRAVENOUS

ADREVIEW

+! GE HEALTHCARE 10mCi/5ML (2mCi/ML) N022290 001 Sep 19, 2008

IODIXANOL

INJECTABLE; INJECTION

VISIPAQUE 270

+! GE HEALTHCARE 55% N020351 001 Mar 22, 1996

VISIPAQUE 320

+! GE HEALTHCARE 65.2% N020351 002 Mar 22, 1996

65.2% N020808 002 Aug 29, 1997

IOFLUPANE I-123

SOLUTION; INTRAVENOUS

DATSCAN

+! GE HLTHCARE INC 5mCi/2.5ML (2mCi/ML) N022454 001 Jan 14, 2011

IOHEXOL

FOR SOLUTION; ORAL

ORALTAG

INTERPHARMA PRAHA 9.7GM/BOT

N205383 001 Mar 26, 2015

AS

INJECTABLE; INJECTION

OMNIPAQUE 140

+! GE HEALTHCARE 30.2% N018956 005 Nov 30, 1988

SOLUTION; INJECTION, ORAL

OMNIPAQUE 350

+! GE HEALTHCARE 75.5% N018956 004 Dec 26, 1985

75.5% N020608 003 Oct 24, 1995

SOLUTION; INJECTION, ORAL, RECTAL

OMNIPAQUE 180

+! GE HEALTHCARE 38.8% N018956 001 Dec 26, 1985

OMNIPAQUE 240

+! GE HEALTHCARE 51.8% N018956 002 Dec 26, 1985

OMNIPAQUE 300

+! GE HEALTHCARE 64.7% N018956 003 Dec 26, 1985

64.7% N020608 002 Oct 24, 1995

SOLUTION; ORAL

OMNIPAQUE 12

+! GE HEALTHCARE 2.6% N018956 009 Apr 17, 2018

OMNIPAQUE 9

+! GE HEALTHCARE 1.9% N018956 008 Apr 17, 2018

IOPAMIDOL

INJECTABLE; INJECTION

ISOVUE-300**AP +! BRACCO 61% N018735 002 Dec 31, 1985****ISOVUE-370****AP +! BRACCO 76% N018735 003 Dec 31, 1985****SCANLUX-300****AP SANOCHEMIA CORP USA 61% A090394 001 Jun 18, 2010****SCANLUX-370****AP SANOCHEMIA CORP USA 76% A090394 002 Jun 18, 2010**

ISOVUE-200

+! BRACCO 41% N018735 006 Jul 07, 1987

ISOVUE-250

+! BRACCO 51% N018735 007 Jul 06, 1992

PRESCRIPTION DRUG PRODUCT LIST

IOPAMIDOL

INJECTABLE; INJECTION

ISOVUE-250

+! 51% N020327 002 Oct 12, 1994

ISOVUE-300

+! BRACCO 61% N020327 003 Oct 12, 1994

ISOVUE-370

+! BRACCO 76% N020327 004 Oct 12, 1994

ISOVUE-M 200

+! BRACCO 41% N018735 001 Dec 31, 1985

ISOVUE-M 300

+! BRACCO 61% N018735 004 Dec 31, 1985

IOPROMIDE

INJECTABLE; INJECTION

ULTRAVIST (PHARMACY BULK)

+! BAYER HLTHCARE 49.9% N021425 003 Mar 12, 2004

+! 62.3% N021425 001 Sep 20, 2002

+! 76.9% N021425 002 Sep 20, 2002

ULTRAVIST 240

+! BAYER HLTHCARE 49.9% N020220 003 May 10, 1995

ULTRAVIST 300

+! BAYER HLTHCARE 62.3% N020220 002 May 10, 1995

ULTRAVIST 370

+! BAYER HLTHCARE 76.9% N020220 001 May 10, 1995

IOTHALAMATE MEGLUMINE

INJECTABLE; INJECTION

CONRAY

+! LIEBEL-FLARSHEIM 60% N013295 001

CONRAY 43

+! LIEBEL-FLARSHEIM 43% N013295 002

SOLUTION; INTRAVESICAL

CYSTO-CONRAY II

LIEBEL-FLARSHEIM 17.2% N017057 002

IOTHALAMATE SODIUM I-125

INJECTABLE; INJECTION

GLOFIL-125

ISOTEX 250-300uCi/ML N017279 001

IOVERSOL

INJECTABLE; INJECTION

OPTIRAY 240

+! LIEBEL-FLARSHEIM 51% N019710 002 Dec 30, 1988

OPTIRAY 300

+! LIEBEL-FLARSHEIM 64% N019710 004 Jan 22, 1992

+! 64% N020923 004 May 13, 1999

OPTIRAY 320

+! LIEBEL-FLARSHEIM 68% N019710 001 Dec 30, 1988

+! 68% N020923 002 May 29, 1998

OPTIRAY 350

+! LIEBEL-FLARSHEIM 74% N019710 005 Jan 22, 1992

+! 74% N020923 003 May 28, 1998

IPRATROPIUM BROMIDE

AEROSOL, METERED; INHALATION

ATROVENT HFA

+! BOEHRINGER 0.021MG/INH N021527 001 Nov 27, 2004

INGELHEIM

SOLUTION; INHALATION

IPRATROPIUM BROMIDE**AN** AUROBINDO PHARMA LTD **0.02%** **A206543 001** Oct 27, 2016**AN** LANDELA PHARM **0.02%** **A077072 001** Jul 19, 2005**AN** NEPHRON **0.02%** **A075562 001** Sep 27, 2001**AN** ! RITEDOSE CORP **0.02%** **A075693 001** Jan 26, 2001**AN** SUN PHARMA GLOBAL **0.02%** **A207903 001** Jan 03, 2017**AN** WATSON LABS **0.02%** **A076291 001** May 09, 2005

SPRAY, METERED; NASAL

IPRATROPIUM BROMIDE**AB** APOTEX INC **0.042MG/SPRAY** **A076155 001** Apr 18, 2003**AB** BAUSCH AND LOMB **0.021MG/SPRAY** **A076025 001** Mar 31, 2003**AB** **0.042MG/SPRAY** **A076103 001** Mar 31, 2003**AB** MYLAN SPECIALITY LP **0.021MG/SPRAY** **A075552 001** Mar 31, 2003

PRESCRIPTION DRUG PRODUCT LIST

IPRATROPIUM BROMIDE

SPRAY, METERED;NASAL

IPRATROPIUM BROMIDE

| | | | | | |
|-----------|---|-------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | <u>0.042MG/SPRAY</u> | <u>A075553 001</u> | Mar 31, 2003 | |
| <u>AB</u> | ! | WEST-WARD PHARMS INT | <u>0.021MG/SPRAY</u> | <u>A076664 001</u> | Nov 05, 2003 |
| <u>AB</u> | ! | | <u>0.042MG/SPRAY</u> | <u>A076598 001</u> | Nov 05, 2003 |

IRBESARTAN

TABLET;ORAL

AVAPRO

| | | | | | |
|-----------|---|-------------------|--------------|--------------------|--------------|
| <u>AB</u> | + | SANOFI AVENTIS US | <u>75MG</u> | <u>N020757 001</u> | Sep 30, 1997 |
| <u>AB</u> | + | | <u>150MG</u> | <u>N020757 002</u> | Sep 30, 1997 |
| <u>AB</u> | + | | <u>300MG</u> | <u>N020757 003</u> | Sep 30, 1997 |

IRBESARTAN

| | | | | | |
|-----------|--|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | | ALEMBIC PHARMS LTD | <u>75MG</u> | <u>A091236 001</u> | Oct 15, 2012 |
| <u>AB</u> | | | <u>150MG</u> | <u>A091236 002</u> | Oct 15, 2012 |
| <u>AB</u> | | | <u>300MG</u> | <u>A091236 003</u> | Oct 15, 2012 |
| <u>AB</u> | | AMNEAL PHARMS | <u>75MG</u> | <u>A204740 001</u> | Apr 17, 2018 |
| <u>AB</u> | | | <u>150MG</u> | <u>A204740 002</u> | Apr 17, 2018 |
| <u>AB</u> | | | <u>300MG</u> | <u>A204740 003</u> | Apr 17, 2018 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>75MG</u> | <u>A203081 001</u> | Sep 27, 2012 |
| <u>AB</u> | | | <u>150MG</u> | <u>A203081 002</u> | Sep 27, 2012 |
| <u>AB</u> | | | <u>300MG</u> | <u>A203081 003</u> | Sep 27, 2012 |
| <u>AB</u> | | CHARTWELL MOLECULAR | <u>75MG</u> | <u>A077205 001</u> | Nov 14, 2012 |
| <u>AB</u> | | | <u>150MG</u> | <u>A077205 002</u> | Nov 14, 2012 |
| <u>AB</u> | | | <u>300MG</u> | <u>A077205 003</u> | Nov 14, 2012 |
| <u>AB</u> | | HETERO LABS LTD V | <u>75MG</u> | <u>A202910 001</u> | Sep 27, 2012 |
| <u>AB</u> | | | <u>150MG</u> | <u>A202910 002</u> | Sep 27, 2012 |
| <u>AB</u> | | | <u>300MG</u> | <u>A202910 003</u> | Sep 27, 2012 |
| <u>AB</u> | | HISUN PHARM HANGZHOU | <u>75MG</u> | <u>A206194 001</u> | Jun 14, 2016 |
| <u>AB</u> | | | <u>150MG</u> | <u>A206194 002</u> | Jun 14, 2016 |
| <u>AB</u> | | | <u>300MG</u> | <u>A206194 003</u> | Jun 14, 2016 |
| <u>AB</u> | | JUBILANT GENERICS | <u>75MG</u> | <u>A203534 001</u> | Feb 23, 2015 |
| <u>AB</u> | | | <u>150MG</u> | <u>A203534 002</u> | Feb 23, 2015 |
| <u>AB</u> | | | <u>300MG</u> | <u>A203534 003</u> | Feb 23, 2015 |
| <u>AB</u> | | LUPIN LTD | <u>75MG</u> | <u>A201531 001</u> | Oct 15, 2012 |
| <u>AB</u> | | | <u>150MG</u> | <u>A201531 002</u> | Oct 15, 2012 |
| <u>AB</u> | | | <u>300MG</u> | <u>A201531 003</u> | Oct 15, 2012 |
| <u>AB</u> | | MACLEODS PHARMS LTD | <u>75MG</u> | <u>A202254 001</u> | Oct 03, 2012 |
| <u>AB</u> | | | <u>150MG</u> | <u>A202254 002</u> | Oct 03, 2012 |
| <u>AB</u> | | | <u>300MG</u> | <u>A202254 003</u> | Oct 03, 2012 |
| <u>AB</u> | | NEOPHARMA | <u>75MG</u> | <u>A203161 001</u> | Sep 27, 2012 |
| <u>AB</u> | | | <u>150MG</u> | <u>A203161 002</u> | Sep 27, 2012 |
| <u>AB</u> | | | <u>300MG</u> | <u>A203161 003</u> | Sep 27, 2012 |
| <u>AB</u> | | PRINSTON INC | <u>75MG</u> | <u>A203071 001</u> | Sep 27, 2012 |
| <u>AB</u> | | | <u>150MG</u> | <u>A203071 002</u> | Sep 27, 2012 |
| <u>AB</u> | | | <u>300MG</u> | <u>A203071 003</u> | Sep 27, 2012 |
| <u>AB</u> | | SANDOZ | <u>75MG</u> | <u>A077466 001</u> | Sep 27, 2012 |
| <u>AB</u> | | | <u>150MG</u> | <u>A077466 002</u> | Sep 27, 2012 |
| <u>AB</u> | | | <u>300MG</u> | <u>A077466 003</u> | Sep 27, 2012 |
| <u>AB</u> | | SCIEGEN PHARMS INC | <u>75MG</u> | <u>A204774 001</u> | Dec 07, 2015 |
| <u>AB</u> | | | <u>150MG</u> | <u>A204774 002</u> | Dec 07, 2015 |
| <u>AB</u> | | | <u>300MG</u> | <u>A204774 003</u> | Dec 07, 2015 |
| <u>AB</u> | | TEVA PHARMS | <u>75MG</u> | <u>A077159 001</u> | Mar 30, 2012 |
| <u>AB</u> | | | <u>150MG</u> | <u>A077159 002</u> | Mar 30, 2012 |
| <u>AB</u> | | | <u>300MG</u> | <u>A077159 003</u> | Mar 30, 2012 |
| <u>AB</u> | | UNICHEM LABS LTD | <u>75MG</u> | <u>A203020 001</u> | Dec 07, 2015 |
| <u>AB</u> | | | <u>150MG</u> | <u>A203020 002</u> | Dec 07, 2015 |
| <u>AB</u> | | | <u>300MG</u> | <u>A203020 003</u> | Dec 07, 2015 |
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>75MG</u> | <u>A090201 001</u> | Oct 15, 2012 |
| <u>AB</u> | | | <u>150MG</u> | <u>A090201 002</u> | Oct 15, 2012 |
| <u>AB</u> | | | <u>300MG</u> | <u>A090201 003</u> | Oct 15, 2012 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>75MG</u> | <u>A079213 001</u> | Sep 27, 2012 |
| <u>AB</u> | | | <u>150MG</u> | <u>A079213 002</u> | Sep 27, 2012 |
| <u>AB</u> | | | <u>300MG</u> | <u>A079213 003</u> | Sep 27, 2012 |

PRESCRIPTION DRUG PRODUCT LIST

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

CAMPTOSAR

| | | | | | |
|-----------|------------|-------------------|-----------------------------|--------------------|--------------|
| <u>AP</u> | <u>+</u> ! | <u>PFIZER INC</u> | <u>40MG/2ML (20MG/ML)</u> | <u>N020571 001</u> | Jun 14, 1996 |
| <u>AP</u> | <u>+</u> ! | | <u>100MG/5ML (20MG/ML)</u> | <u>N020571 002</u> | Jun 14, 1996 |
| <u>AP</u> | <u>+</u> ! | | <u>300MG/15ML (20MG/ML)</u> | <u>N020571 003</u> | Aug 05, 2010 |

IRINOTECAN HYDROCHLORIDE

| | | | | | |
|-----------|----------|-----------------------------|-----------------------------|--------------------|--------------|
| <u>AP</u> | | <u>ACCORD HLTHCARE</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A079068 001</u> | Nov 21, 2008 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A079068 002</u> | Nov 21, 2008 |
| <u>AP</u> | | <u>ACTAVIS TOTOWA</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A078589 001</u> | Feb 27, 2008 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A078589 002</u> | Feb 27, 2008 |
| <u>AP</u> | | | <u>500MG/25ML (20MG/ML)</u> | <u>A078589 003</u> | Nov 18, 2015 |
| <u>AP</u> | | <u>AKORN</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A090726 001</u> | Sep 16, 2009 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A090726 002</u> | Sep 16, 2009 |
| <u>AP</u> | | <u>CIPLA LTD</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A077219 001</u> | Feb 20, 2008 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A077219 002</u> | Feb 20, 2008 |
| <u>AP</u> | | <u>EMCURE PHARMS LTD</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A200771 001</u> | Feb 14, 2012 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A200771 002</u> | Feb 14, 2012 |
| <u>AP</u> | | <u>FRESENIUS KABI ONCOL</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A078188 001</u> | Feb 27, 2008 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A078188 002</u> | Feb 27, 2008 |
| <u>AP</u> | | <u>FRESENIUS KABI USA</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A077776 001</u> | Feb 27, 2008 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A077776 002</u> | Feb 27, 2008 |
| <u>AP</u> | | <u>HIKMA FARMACEUTICA</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A091032 001</u> | Dec 20, 2010 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A091032 002</u> | Dec 20, 2010 |
| <u>AP</u> | | <u>HISUN PHARM HANGZHOU</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A090016 001</u> | Jan 28, 2009 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A090016 002</u> | Jan 28, 2009 |
| <u>AP</u> | | <u>HOSPIRA</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A077915 001</u> | Feb 27, 2008 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A077915 002</u> | Feb 27, 2008 |
| <u>AP</u> | <u>!</u> | | <u>500MG/25ML (20MG/ML)</u> | <u>A078796 001</u> | Feb 27, 2008 |
| <u>AP</u> | | <u>INGENUS PHARMS LLC</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A206935 001</u> | May 26, 2017 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A206935 002</u> | May 26, 2017 |
| <u>AP</u> | | <u>INTAS PHARMS USA</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A203054 001</u> | Aug 31, 2017 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A203054 002</u> | Aug 31, 2017 |
| <u>AP</u> | | <u>JIANGSU HENGRUI MED</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A090675 002</u> | Dec 16, 2011 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A090675 001</u> | Dec 16, 2011 |
| <u>AP</u> | | <u>MUSTAFA NEVZAT ILAC</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A090393 002</u> | May 13, 2011 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A090393 003</u> | May 13, 2011 |
| <u>AP</u> | | <u>NEOPHARMA</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A078953 001</u> | Apr 15, 2010 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A078953 002</u> | Apr 15, 2010 |
| <u>AP</u> | | <u>PLIVA LACHEMA</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A078122 001</u> | Oct 31, 2008 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A078122 002</u> | Oct 31, 2008 |
| <u>AP</u> | | <u>QILU PHARM CO LTD</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A203380 001</u> | May 03, 2016 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A203380 002</u> | May 03, 2016 |
| <u>AP</u> | | | <u>300MG/15ML (20MG/ML)</u> | <u>A203380 003</u> | May 03, 2016 |
| <u>AP</u> | | <u>SHILPA MEDICARE LTD</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A208718 001</u> | Dec 28, 2018 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A208718 002</u> | Dec 28, 2018 |
| <u>AP</u> | | <u>TEVA PHARMS USA</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A090101 002</u> | Feb 27, 2008 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A090101 003</u> | Feb 27, 2008 |
| <u>AP</u> | | | <u>500MG/25ML (20MG/ML)</u> | <u>A090101 001</u> | Nov 26, 2008 |
| <u>AP</u> | | <u>WEST-WARD PHARMS INT</u> | <u>40MG/2ML (20MG/ML)</u> | <u>A078753 001</u> | Dec 24, 2008 |
| <u>AP</u> | | | <u>100MG/5ML (20MG/ML)</u> | <u>A078753 002</u> | Dec 24, 2008 |

INJECTABLE, LIPOSOMAL; INTRAVENOUS

ONIVYDE

| | | | | |
|------------|------------------|---------------------------------------------|--------------------|--------------|
| <u>+</u> ! | <u>IPSEN INC</u> | <u>EQ 43MG BASE/10ML (EQ 4.3MG BASE/ML)</u> | <u>N207793 001</u> | Oct 22, 2015 |
|------------|------------------|---------------------------------------------|--------------------|--------------|

IRON DEXTRAN

INJECTABLE; INJECTION

DEXFERRUM

| | | | | | |
|-----------|------------|---------------------------|------------------------|--------------------|--------------|
| <u>BP</u> | | <u>LUITPOLD</u> | <u>EQ 50MG IRON/ML</u> | <u>N040024 001</u> | Feb 23, 1996 |
| | | <u>INFED</u> | | | |
| <u>BP</u> | <u>+</u> ! | <u>ALLERGAN SALES LLC</u> | <u>EQ 50MG IRON/ML</u> | <u>N017441 001</u> | |
| | | <u>PROFERDEX</u> | | | |
| <u>BP</u> | | <u>NEW RIVER</u> | <u>EQ 50MG IRON/ML</u> | <u>N017807 001</u> | |

IRON SUCROSE

INJECTABLE; INTRAVENOUS

VENOFER

| | | | | |
|------------|-----------------|---------------------------------------------|--------------------|--------------|
| <u>+</u> | <u>LUITPOLD</u> | <u>EQ 50MG BASE/2.5ML (EQ 20MG BASE/ML)</u> | <u>N021135 002</u> | Mar 20, 2005 |
| <u>+</u> ! | | <u>EQ 100MG BASE/5ML (EQ 20MG BASE/ML)</u> | <u>N021135 001</u> | Nov 06, 2000 |
| <u>+</u> | | <u>EQ 200MG BASE/10ML (EQ 20MG BASE/ML)</u> | <u>N021135 004</u> | Feb 09, 2007 |

PRESCRIPTION DRUG PRODUCT LIST

ISAVUCONAZONIUM SULFATE

CAPSULE; ORAL

CRESEMBA

+! ASTELLAS 186MG N207500 001 Mar 06, 2015

POWDER; INTRAVENOUS

CRESEMBA

+! ASTELLAS 372MG N207501 001 Mar 06, 2015

ISOCARBOAZID

TABLET; ORAL

MARPLAN

+! VALIDUS PHARMS INC 10MG N011961 001

ISOFLURANE

LIQUID; INHALATION

FORANEAN +! BAXTER HLTHCARE 99.9% N017624 001ISOFLURANEAN HALOCARBON PRODS 99.9% A075225 001 Oct 20, 1999AN PIRAMAL CRITICAL 99.9% A074416 001 Sep 30, 1994AN PIRAMAL ENT 99.9% A074502 001 Jun 27, 1995ISONIAZID

INJECTABLE; INJECTION

ISONIAZID

! SANDOZ INC 100MG/ML A040648 001 Jul 05, 2005

SYRUP; ORAL

ISONIAZID

! CMP PHARMA INC 50MG/5ML A088235 001 Nov 10, 1983

TABLET; ORAL

ISONIAZIDAA BARR 100MG A080936 001AA 300MG A080937 002AA MIKART 100MG A040090 001 Jun 26, 1997AA 300MG A040090 002 Jun 26, 1997AA +! SANDOZ 100MG N008678 002AA +! 300MG N008678 003AA THEPHARMANETWORK 100MG A202610 001 Oct 29, 2014

LLC

AA 300MG A202610 002 Oct 29, 2014LANIAZIDAA LANNETT 300MG A089776 001 Jun 13, 1988ISONIAZID; PYRAZINAMIDE; RIFAMPIN

TABLET; ORAL

RIFATER

+! SANOFI AVENTIS US 50MG; 300MG; 120MG N050705 001 May 31, 1994

ISONIAZID; RIFAMPIN

CAPSULE; ORAL

RIFAMATE

! SANOFI AVENTIS US 150MG; 300MG A061884 001

ISOPROTERENOL HYDROCHLORIDE

INJECTABLE; INJECTION

ISOPROTERENOL HYDROCHLORIDEAP AMNEAL PHARMS CO 0.2MG/ML A210576 001 Oct 17, 2018AP AMPHASTAR PHARMS 0.2MG/ML A210106 001 Jun 18, 2018

INC

AP CIPLA 0.2MG/ML A210322 001 Jun 12, 2018AP NEXUS PHARMS 0.2MG/ML A206961 001 Aug 02, 2017ISUPRELAP +! VALEANT PHARMS 0.2MG/ML N010515 001

NORTH

ISOSORBIDE DINITRATE

CAPSULE, EXTENDED RELEASE; ORAL

DILATRATE-SR

+! AUXILIUM PHARMS LLC 40MG N019790 001 Sep 02, 1988

TABLET; ORAL

ISORDILAB + VALEANT PHARMS 5MG N012093 007 Jul 29, 1988

NORTH

ISOSORBIDE DINITRATEAB HIKMA INTL PHARMS 5MG A086067 001 Oct 29, 1987AB 10MG A086066 001 Oct 29, 1987AB 20MG A088088 001 Nov 02, 1987

PRESCRIPTION DRUG PRODUCT LIST

ISOSORBIDE DINITRATE

TABLET; ORAL

ISOSORBIDE DINITRATE

| | | | | | |
|--------------------------------|-----------|-------------------------|----------------|------------|------------------|
| <u>AB</u> | | <u>30MG</u> | <u>A040591</u> | <u>001</u> | Jan 10, 2007 |
| <u>AB</u> | PAR PHARM | <u>5MG</u> | <u>A086923</u> | <u>001</u> | Mar 12, 1987 |
| <u>AB</u> | | <u>10MG</u> | <u>A086925</u> | <u>001</u> | Mar 12, 1987 |
| <u>AB</u> | | <u>20MG</u> | <u>A087537</u> | <u>001</u> | Oct 02, 1987 |
| <u>AB</u> | ! | <u>30MG</u> | <u>A087946</u> | <u>001</u> | Jan 12, 1988 |
| <u>AB</u> | SANDOZ | <u>5MG</u> | <u>A086221</u> | <u>001</u> | Jan 07, 1988 |
| <u>AB</u> | | <u>10MG</u> | <u>A086223</u> | <u>001</u> | Jan 07, 1988 |
| <u>AB</u> | | <u>20MG</u> | <u>A089367</u> | <u>001</u> | Apr 07, 1988 |
| ISORDIL | | | | | |
| | +! | VALEANT PHARMS NORTH | 40MG | N012093 | 001 Jul 29, 1988 |
| TABLET, EXTENDED RELEASE; ORAL | | | | | |
| <u>ISOSORBIDE DINITRATE</u> | | | | | |
| | ! | SUN PHARM INDS INC | 40MG | A040009 | 001 Dec 30, 1998 |

ISOSORBIDE MONONITRATE

TABLET; ORAL

ISOSORBIDE MONONITRATE

| | | | | | |
|----------------|-------------------|----------------|----------------|----------------|-------------------------|
| <u>AB</u> | ACTAVIS ELIZABETH | <u>10MG</u> | <u>A075037</u> | <u>002</u> | Oct 30, 1998 |
| <u>AB</u> | | <u>20MG</u> | <u>A075037</u> | <u>001</u> | Oct 30, 1998 |
| <u>AB</u> | ANI PHARMS INC | <u>20MG</u> | <u>A075147</u> | <u>001</u> | Nov 27, 1998 |
| <u>AB</u> | HIKMA PHARMS | <u>20MG</u> | <u>A075361</u> | <u>001</u> | Oct 05, 2000 |
| <u>MONOKET</u> | | | | | |
| <u>AB</u> | + | LANNETT CO INC | <u>10MG</u> | <u>N020215</u> | <u>002</u> Jun 30, 1993 |
| <u>AB</u> | +! | | <u>20MG</u> | <u>N020215</u> | <u>001</u> Jun 30, 1993 |

TABLET, EXTENDED RELEASE; ORAL

ISOSORBIDE MONONITRATE

| | | | | | |
|-----------|-----------------|--------------|----------------|------------|--------------|
| <u>AB</u> | DEXCEL LTD | <u>30MG</u> | <u>A075522</u> | <u>002</u> | Sep 20, 2016 |
| <u>AB</u> | | <u>60MG</u> | <u>A075522</u> | <u>001</u> | Apr 17, 2000 |
| <u>AB</u> | | <u>120MG</u> | <u>A210822</u> | <u>001</u> | Aug 29, 2018 |
| <u>AB</u> | HIKMA PHARMS | <u>30MG</u> | <u>A076813</u> | <u>002</u> | Mar 30, 2006 |
| <u>AB</u> | | <u>60MG</u> | <u>A076813</u> | <u>001</u> | Jan 07, 2005 |
| <u>AB</u> | LANNETT CO INC | <u>30MG</u> | <u>A075155</u> | <u>002</u> | Jan 13, 2000 |
| <u>AB</u> | | <u>60MG</u> | <u>A075155</u> | <u>001</u> | Oct 30, 1998 |
| <u>AB</u> | ! | <u>120MG</u> | <u>A075155</u> | <u>003</u> | Aug 04, 2000 |
| <u>AB</u> | NESHER PHARMS | <u>30MG</u> | <u>A075395</u> | <u>001</u> | Mar 16, 2000 |
| <u>AB</u> | | <u>60MG</u> | <u>A075395</u> | <u>002</u> | Mar 16, 2000 |
| <u>AB</u> | | <u>120MG</u> | <u>A075395</u> | <u>003</u> | Mar 16, 2000 |
| <u>AB</u> | RICONPHARMA LLC | <u>30MG</u> | <u>A210918</u> | <u>001</u> | Nov 05, 2018 |
| <u>AB</u> | | <u>60MG</u> | <u>A210918</u> | <u>002</u> | Nov 05, 2018 |
| <u>AB</u> | | <u>120MG</u> | <u>A210918</u> | <u>003</u> | Nov 05, 2018 |
| <u>AB</u> | TORRENT PHARMS | <u>30MG</u> | <u>A200270</u> | <u>001</u> | Jun 03, 2011 |
| <u>AB</u> | | <u>60MG</u> | <u>A200495</u> | <u>001</u> | Jun 03, 2011 |
| <u>AB</u> | | <u>120MG</u> | <u>A200495</u> | <u>002</u> | Jun 03, 2011 |
| <u>AB</u> | VINTAGE PHARMS | <u>30MG</u> | <u>A090598</u> | <u>001</u> | Aug 11, 2010 |
| <u>AB</u> | | <u>60MG</u> | <u>A090598</u> | <u>002</u> | Aug 11, 2010 |
| <u>AB</u> | | <u>120MG</u> | <u>A090598</u> | <u>003</u> | Aug 11, 2010 |

ISOSULFAN BLUE

INJECTABLE; INJECTION

ISOSULFAN BLUE

| | | | | | |
|-----------|-------------------------|---------------------|----------------|----------------|-------------------------|
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>1%</u> | <u>A206831</u> | <u>001</u> | Feb 02, 2016 |
| <u>AP</u> | BELOTECA INC | <u>1%</u> | <u>A210714</u> | <u>001</u> | Jan 16, 2019 |
| <u>AP</u> | ! | MYLAN INSTITUTIONAL | <u>1%</u> | <u>A090874</u> | <u>001</u> Jul 20, 2010 |

ISOTRETINOIN

CAPSULE; ORAL

AMNESTEEM

| | | | | | |
|-----------|------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | MYLAN PHARMS INC | <u>10MG</u> | <u>A075945</u> | <u>001</u> | Nov 08, 2002 |
| <u>AB</u> | | <u>20MG</u> | <u>A075945</u> | <u>002</u> | Nov 08, 2002 |
| <u>AB</u> | | <u>40MG</u> | <u>A075945</u> | <u>003</u> | Nov 08, 2002 |

CLARAVIS

| | | | | | |
|-----------|-----------------|-------------|----------------|------------|--------------|
| <u>AB</u> | TEVA PHARMS USA | <u>10MG</u> | <u>A076356</u> | <u>001</u> | Apr 11, 2003 |
| <u>AB</u> | | <u>20MG</u> | <u>A076135</u> | <u>002</u> | Apr 11, 2003 |
| <u>AB</u> | | <u>30MG</u> | <u>A076135</u> | <u>003</u> | May 11, 2006 |
| <u>AB</u> | ! | <u>40MG</u> | <u>A076135</u> | <u>001</u> | Apr 11, 2003 |

ISOTRETINOIN

| | | | | | |
|-----------|------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | AMNEAL PHARMS NY | <u>10MG</u> | <u>A207792</u> | <u>001</u> | Sep 29, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A207792</u> | <u>002</u> | Sep 29, 2017 |
| <u>AB</u> | | <u>30MG</u> | <u>A207792</u> | <u>003</u> | Sep 29, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

ISOTRETINOIN

CAPSULE;ORAL

ISOTRETINOIN

AB 40MG A207792 004 Sep 29, 2017

MYORISAN

AB DOUGLAS PHARMS 10MG A076485 001 Jan 19, 2012
AB 20MG A076485 002 Jan 19, 2012
AB 30MG A076485 004 Aug 25, 2015
AB 40MG A076485 003 Jan 19, 2012

ZENATANE

AB DR REDDYS LABS LTD 10MG A202099 001 Mar 25, 2013
AB 20MG A202099 002 Mar 25, 2013
AB 30MG A202099 004 Feb 23, 2015
AB 40MG A202099 003 Mar 25, 2013

ABSORICA

BX + SUN PHARM INDS INC 10MG N021951 001 May 25, 2012
BX + 20MG N021951 002 May 25, 2012
BX + 30MG N021951 003 May 25, 2012
BX +! 40MG N021951 004 May 25, 2012
+ 25MG N021951 005 Aug 15, 2014
+ 35MG N021951 006 Aug 15, 2014

ISRADIPINE

CAPSULE;ORAL

ISRADIPINE

AB ELITE LABS INC 2.5MG A077169 001 Apr 24, 2006
AB 5MG A077169 002 Apr 24, 2006
AB WATSON LABS TEVA 2.5MG A077317 001 Jan 05, 2006
AB ! 5MG A077317 002 Jan 05, 2006

ITRACONAZOLE

CAPSULE;ORAL

ITRACONAZOLE

AB ACCORD HLTHCARE 100MG A205991 001 May 26, 2016
AB ALEMBIC PHARMS LTD 100MG A206741 001 Dec 13, 2016
AB ALKEM LABS LTD 100MG A208591 001 Jun 12, 2017
AB AMNEAL PHARMS 100MG A205080 001 Sep 26, 2016
AB JUBILANT GENERICS 100MG A203445 001 Feb 23, 2017
AB MYLAN PHARMS INC 100MG A200463 001 Jul 20, 2012
AB PAR PHARM INC 100MG A205724 001 Dec 13, 2016
AB SANDOZ 100MG A076104 001 May 28, 2004
AB TORRENT PHARMS LTD 100MG A209460 001 Aug 24, 2018
AB ZYDUS PHARMS USA 100MG A204672 001 Sep 19, 2017
INC

SPORANOX

AB +! JANSSEN PHARMS 100MG N020083 001 Sep 11, 1992
TOLSURA
+! MAYNE PHARMA INTL 65MG N208901 001 Dec 11, 2018
SOLUTION;ORAL

ITRACONAZOLE

AA AMNEAL PHARMS 10MG/ML A205573 001 Oct 30, 2015

SPORANOX

AA +! JANSSEN PHARMS 10MG/ML N020657 001 Feb 21, 1997
TABLET;ORAL
ONMEL
+! SEBELA IRELAND LTD 200MG N022484 001 Apr 29, 2010

IVABRADINE HYDROCHLORIDE

TABLET;ORAL

CORLANOR

+ AMGEN INC EQ 5MG BASE N206143 001 Apr 15, 2015
+! EQ 7.5MG BASE N206143 002 Apr 15, 2015

IVACAFTOR

GRANULE;ORAL

KALYDECO

+ VERTEX PHARMS INC 50MG/PACKET N207925 001 Mar 17, 2015
+! 75MG/PACKET N207925 002 Mar 17, 2015

TABLET;ORAL

KALYDECO

+! VERTEX PHARMS 150MG N203188 001 Jan 31, 2012

PRESCRIPTION DRUG PRODUCT LIST

IVACAFTOR; IVACAFTOR, TEZACAFTOR

TABLET, TABLET;ORAL

SYMDEKO (COPACKAGED)

+! VERTEX PHARMS INC 150MG,N/A;150MG, 100MG N210491 001 Feb 12, 2018

IVACAFTOR; LUMACAFTOR

GRANULE;ORAL

ORKAMBI

+ VERTEX PHARMS INC 125MG/PACKET;100MG/PACKET N211358 001 Aug 07, 2018

+! 188MG/PACKET;150MG/PACKET N211358 002 Aug 07, 2018

TABLET;ORAL

ORKAMBI

+ VERTEX PHARMS INC 125MG;100MG N206038 002 Sep 28, 2016

+! 125MG;200MG N206038 001 Jul 02, 2015

IVERMECTIN

CREAM;TOPICAL

SOOLANTRA

+! GALDERMA LABS LP 1% N206255 001 Dec 19, 2014

LOTION;TOPICAL

SKLICE

+! ARBOR PHARMS LLC 0.5% N202736 001 Feb 07, 2012

TABLET;ORAL

IVERMECTIN**AB** EDENBRIDGE PHARMS **3MG** **A204154 001** Oct 24, 2014STROMEKTOL**AB** +! MERCK SHARP DOHME **3MG** **N050742 002** Oct 08, 1998IVOSIDENIB

TABLET;ORAL

TIBSOVO

+! AGIOS PHARMS INC 250MG N211192 001 Jul 20, 2018

IXABEPILONE

INJECTABLE; INTRAVENOUS

IXEMPRA KIT

+! R-PHARM US LLC 15MG/VIAL N022065 001 Oct 16, 2007

+! 45MG/VIAL N022065 002 Oct 16, 2007

IXAZOMIB CITRATE

CAPSULE;ORAL

NINLARO

+ MILLENNIUM PHARMS EQ 2.3MG BASE N208462 001 Nov 20, 2015

+ EQ 3MG BASE N208462 002 Nov 20, 2015

+! EQ 4MG BASE N208462 003 Nov 20, 2015

KETAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

KETALAR**AP** +! PAR STERILE **EQ 10MG BASE/ML** **N016812 001**

PRODUCTS

AP +! **EQ 50MG BASE/ML** **N016812 002****AP** +! **EQ 100MG BASE/ML** **N016812 003**KETAMINE HYDROCHLORIDE**AP** HOSPIRA **EQ 50MG BASE/ML** **A074549 001** Jun 27, 1996**AP** **EQ 100MG BASE/ML** **A074549 002** Jun 27, 1996**AP** MYLAN INSTITUTIONAL **EQ 10MG BASE/ML** **A076092 001** Sep 30, 2008**AP** **EQ 50MG BASE/ML** **A076092 002** Dec 28, 2001**AP** **EQ 100MG BASE/ML** **A076092 003** Oct 25, 2002**AP** WEST-WARD PHARMS **EQ 50MG BASE/ML** **A074524 001** Mar 22, 1996

INT

AP **EQ 100MG BASE/ML** **A074524 002** Mar 22, 1996KETOCONAZOLE

AEROSOL, FOAM;TOPICAL

EXTINA**AT** +! MYLAN PHARMS INC **2%** **N021738 001** Jun 12, 2007KETOCONAZOLE**AT** PERRIGO ISRAEL **2%** **A091550 001** Aug 25, 2011

CREAM;TOPICAL

KETOCONAZOLE**AB** FOUGERA PHARMS **2%** **A076294 001** Apr 28, 2004**AB** ! TEVA **2%** **A075581 001** Apr 25, 2000KETOZOILE**AB** TARO **2%** **A075638 001** Dec 18, 2002

PRESCRIPTION DRUG PRODUCT LIST

| | | | | | | |
|---------------------------------|---|----------------------|----------------|----------------|------------|--------------|
| <u>KETOCONAZOLE</u> | | | | | | |
| GEL; TOPICAL | | | | | | |
| XOLEGEL | | | | | | |
| | + | AQUA PHARMS | 2% | N021946 | 001 | Jul 28, 2006 |
| SHAMPOO; TOPICAL | | | | | | |
| <u>KETOCONAZOLE</u> | | | | | | |
| <u>AB</u> | | PERRIGO NEW YORK | <u>2%</u> | <u>A076419</u> | <u>001</u> | Jan 07, 2004 |
| <u>AB</u> | | TOLMAR | <u>2%</u> | <u>A076942</u> | <u>001</u> | Apr 11, 2005 |
| <u>NIZORAL</u> | | | | | | |
| <u>AB</u> | + | JANSSEN PHARMS | <u>2%</u> | <u>N019927</u> | <u>001</u> | Aug 31, 1990 |
| TABLET; ORAL | | | | | | |
| <u>KETOCONAZOLE</u> | | | | | | |
| <u>AB</u> | | MYLAN | <u>200MG</u> | <u>A075597</u> | <u>001</u> | Dec 23, 1999 |
| <u>AB</u> | | STRIDES PHARMA | <u>200MG</u> | <u>A210457</u> | <u>001</u> | Jun 18, 2018 |
| <u>AB</u> | | TARO | <u>200MG</u> | <u>A075319</u> | <u>001</u> | Jun 15, 1999 |
| <u>AB</u> | ! | TEVA | <u>200MG</u> | <u>A075273</u> | <u>001</u> | Jun 15, 1999 |
| <u>KETOPROFEN</u> | | | | | | |
| CAPSULE; ORAL | | | | | | |
| <u>KETOPROFEN</u> | | | | | | |
| <u>AB</u> | | HERITAGE PHARMS INC | <u>50MG</u> | <u>A074014</u> | <u>002</u> | Jan 29, 1993 |
| <u>AB</u> | | | <u>75MG</u> | <u>A074014</u> | <u>003</u> | Jan 29, 1993 |
| <u>AB</u> | | TEVA | <u>50MG</u> | <u>A073516</u> | <u>001</u> | Dec 22, 1992 |
| <u>AB</u> | ! | | <u>75MG</u> | <u>A073517</u> | <u>001</u> | Dec 22, 1992 |
| | | HERITAGE PHARMS INC | 25MG | A074014 | 001 | Jan 29, 1993 |
| CAPSULE, EXTENDED RELEASE; ORAL | | | | | | |
| <u>KETOPROFEN</u> | | | | | | |
| | ! | MYLAN | 200MG | A075679 | 001 | Feb 20, 2002 |
| <u>KETOROLAC TROMETHAMINE</u> | | | | | | |
| INJECTABLE; INJECTION | | | | | | |
| <u>KETOROLAC TROMETHAMINE</u> | | | | | | |
| <u>AP</u> | | AMPHASTAR PHARM | <u>15MG/ML</u> | <u>A076209</u> | <u>001</u> | Jul 21, 2004 |
| <u>AP</u> | | | <u>30MG/ML</u> | <u>A076209</u> | <u>002</u> | Jul 21, 2004 |
| <u>AP</u> | | BAXTER HLTHCARE CORP | <u>15MG/ML</u> | <u>A209900</u> | <u>002</u> | Jul 25, 2018 |
| <u>AP</u> | | | <u>30MG/ML</u> | <u>A209900</u> | <u>001</u> | Sep 15, 2017 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>15MG/ML</u> | <u>A075784</u> | <u>001</u> | Jan 11, 2002 |
| <u>AP</u> | | | <u>15MG/ML</u> | <u>A203242</u> | <u>001</u> | Oct 07, 2015 |
| <u>AP</u> | | | <u>30MG/ML</u> | <u>A075784</u> | <u>002</u> | Jan 11, 2002 |
| <u>AP</u> | | | <u>30MG/ML</u> | <u>A203242</u> | <u>002</u> | Oct 07, 2015 |
| <u>AP</u> | | GLAND PHARMA LTD | <u>15MG/ML</u> | <u>A204216</u> | <u>001</u> | Nov 01, 2016 |
| <u>AP</u> | | | <u>30MG/ML</u> | <u>A204216</u> | <u>002</u> | Nov 01, 2016 |
| <u>AP</u> | ! | HOSPIRA | <u>15MG/ML</u> | <u>A074802</u> | <u>001</u> | Jun 05, 1997 |
| <u>AP</u> | | | <u>15MG/ML</u> | <u>A074993</u> | <u>001</u> | Jan 27, 1999 |
| <u>AP</u> | ! | | <u>30MG/ML</u> | <u>A074802</u> | <u>002</u> | Jun 05, 1997 |
| <u>AP</u> | | | <u>30MG/ML</u> | <u>A074993</u> | <u>002</u> | Jan 27, 1999 |
| <u>AP</u> | | SAGENT PHARMS | <u>15MG/ML</u> | <u>A091065</u> | <u>001</u> | Nov 27, 2013 |
| <u>AP</u> | | | <u>30MG/ML</u> | <u>A091065</u> | <u>002</u> | Nov 27, 2013 |
| <u>AP</u> | | SANDOZ INC | <u>30MG/ML</u> | <u>A076271</u> | <u>002</u> | Oct 06, 2004 |
| <u>AP</u> | | WOCKHARDT | <u>15MG/ML</u> | <u>A077942</u> | <u>001</u> | Mar 27, 2007 |
| <u>AP</u> | | | <u>30MG/ML</u> | <u>A077942</u> | <u>002</u> | Mar 27, 2007 |
| SOLUTION/DROPS; OPHTHALMIC | | | | | | |
| <u>ACULAR</u> | | | | | | |
| <u>AT</u> | + | ALLERGAN | <u>0.5%</u> | <u>N019700</u> | <u>001</u> | Nov 09, 1992 |
| <u>ACULAR LS</u> | | | | | | |
| <u>AT</u> | + | ALLERGAN | <u>0.4%</u> | <u>N021528</u> | <u>001</u> | May 30, 2003 |
| <u>KETOROLAC TROMETHAMINE</u> | | | | | | |
| <u>AT</u> | | AKORN | <u>0.4%</u> | <u>A078399</u> | <u>001</u> | Nov 05, 2009 |
| <u>AT</u> | | | <u>0.5%</u> | <u>A078434</u> | <u>001</u> | Nov 05, 2009 |
| <u>AT</u> | | APOTEX INC | <u>0.4%</u> | <u>A077308</u> | <u>001</u> | Nov 05, 2009 |
| <u>AT</u> | | | <u>0.5%</u> | <u>A076109</u> | <u>001</u> | Nov 05, 2009 |
| <u>AT</u> | | AUROBINDO PHARMA LTD | <u>0.4%</u> | <u>A205191</u> | <u>001</u> | Nov 15, 2018 |
| <u>AT</u> | | SANDOZ INC | <u>0.4%</u> | <u>A078721</u> | <u>001</u> | Nov 05, 2009 |
| <u>AT</u> | | | <u>0.5%</u> | <u>A076583</u> | <u>001</u> | Nov 05, 2009 |
| <u>AT</u> | | SUN PHARMA GLOBAL | <u>0.5%</u> | <u>A090017</u> | <u>001</u> | Nov 05, 2009 |
| ACUVAIL | | | | | | |
| | + | ALLERGAN | 0.45% | N022427 | 001 | Jul 22, 2009 |
| SPRAY, METERED; NASAL | | | | | | |
| SPRIX | | | | | | |
| | + | EGALET US INC | 15.75MG/SPRAY | N022382 | 001 | May 14, 2010 |

PRESCRIPTION DRUG PRODUCT LISTKETOROLAC TROMETHAMINE

TABLET; ORAL

KETOROLAC TROMETHAMINE

| | | | | |
|-----------|------------------|-------------|--------------------|--------------|
| AB | CYCLE PHARMS LTD | 10MG | A210616 001 | Aug 16, 2018 |
| AB | ! | 10MG | A074761 001 | May 16, 1997 |
| AB | PLIVA | 10MG | A075284 001 | Jun 23, 1999 |
| AB | TEVA | 10MG | A074754 001 | May 16, 1997 |

KETOROLAC TROMETHAMINE; PHENYLEPHRINE HYDROCHLORIDE

SOLUTION; IRRIGATION

OMIDRIA

| | | | | |
|---|--------|-------------------------|-------------|--------------|
| + | OMEROS | EQ 0.3% BASE;EQ 1% BASE | N205388 001 | May 30, 2014 |
|---|--------|-------------------------|-------------|--------------|

L-GLUTAMINE

FOR SOLUTION; ORAL

ENDARI

| | | | | |
|---|--------------|------------|-------------|--------------|
| + | EMMAUS MEDCL | 5GM/PACKET | N208587 001 | Jul 07, 2017 |
|---|--------------|------------|-------------|--------------|

NUTRESTORE

| | | | | |
|---|--------------|------------|-------------|--------------|
| + | EMMAUS MEDCL | 5GM/PACKET | N021667 001 | Jun 10, 2004 |
|---|--------------|------------|-------------|--------------|

LABETALOL HYDROCHLORIDE

INJECTABLE; INJECTION

LABETALOL HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|---------------|--------------------|--------------|
| AP | AKORN INC | 5MG/ML | A075431 001 | Nov 29, 1999 |
| AP | GLAND PHARMA LTD | 5MG/ML | A090699 001 | Apr 03, 2012 |
| AP | ! | 5MG/ML | A075239 001 | Nov 29, 1999 |
| AP | ! | 5MG/ML | A075240 001 | Nov 29, 1999 |
| AP | MYLAN ASI | 5MG/ML | A079134 001 | Feb 03, 2010 |
| AP | WEST-WARD PHARMS INT | 5MG/ML | A075303 001 | May 28, 1999 |

TABLET; ORAL

LABETALOL HYDROCHLORIDE

| | | | | |
|-----------------|-------------------------|--------------|--------------------|--------------|
| AB | INNOGENIX | 100MG | A075215 001 | Jul 29, 1999 |
| AB | | 200MG | A075215 002 | Jul 29, 1999 |
| AB | | 300MG | A075215 003 | Jul 29, 1999 |
| AB | IVAX SUB TEVA PHARMS | 100MG | A074787 001 | Aug 03, 1998 |
| AB | | 200MG | A074787 002 | Aug 03, 1998 |
| AB | | 300MG | A074787 003 | Aug 03, 1998 |
| AB | PAR FORM | 100MG | A200908 001 | Jul 10, 2012 |
| AB | | 200MG | A200908 002 | Jul 10, 2012 |
| AB | | 300MG | A200908 003 | Jul 10, 2012 |
| AB | SANDOZ | 100MG | A075113 001 | Aug 04, 1998 |
| AB | ! | 200MG | A075113 002 | Aug 04, 1998 |
| AB | | 300MG | A075113 003 | Aug 04, 1998 |
| AB | TWI PHARMS | 100MG | A209603 001 | Jun 20, 2018 |
| AB | | 200MG | A209603 002 | Jun 20, 2018 |
| AB | | 300MG | A209603 003 | Jun 20, 2018 |
| AB | WATSON LABS | 100MG | A075133 001 | Aug 03, 1998 |
| AB | | 200MG | A075133 002 | Aug 03, 1998 |
| AB | | 300MG | A075133 003 | Aug 03, 1998 |
| AB | ZYDUS PHARMS USA INC | 100MG | A207743 001 | Sep 19, 2017 |
| AB | | 200MG | A207743 002 | Sep 19, 2017 |
| AB | | 300MG | A207743 003 | Sep 19, 2017 |
| TRANDATE | | | | |
| AB | CNTY LINE PHARMS | 100MG | N018716 001 | May 24, 1985 |
| AB | | 200MG | N018716 002 | Aug 01, 1984 |
| AB | | 300MG | N018716 003 | Aug 01, 1984 |

LACOSAMIDE

SOLUTION; INTRAVENOUS

VIMPAT

| | | | | |
|---|---------|----------------------|-------------|--------------|
| + | UCB INC | 200MG/20ML (10MG/ML) | N022254 001 | Oct 28, 2008 |
|---|---------|----------------------|-------------|--------------|

SOLUTION; ORAL

VIMPAT

| | | | | |
|---|---------|---------|-------------|--------------|
| + | UCB INC | 10MG/ML | N022255 001 | Apr 20, 2010 |
|---|---------|---------|-------------|--------------|

TABLET; ORAL

VIMPAT

| | | | | |
|---|---------|-------|-------------|--------------|
| + | UCB INC | 50MG | N022253 001 | Oct 28, 2008 |
| + | | 100MG | N022253 002 | Oct 28, 2008 |
| + | | 150MG | N022253 003 | Oct 28, 2008 |
| + | | 200MG | N022253 004 | Oct 28, 2008 |

PRESCRIPTION DRUG PRODUCT LIST

LACTULOSE

FOR SOLUTION; ORAL

LACTULOSE

| | | | | | |
|---|-------------------|-------------|---------|-----|--------------|
| ! | CUMBERLAND PHARMS | 10GM/PACKET | A074712 | 001 | Dec 10, 1997 |
| ! | | 20GM/PACKET | A074712 | 002 | Dec 10, 1997 |

SOLUTION; ORAL

CONSTILAC

| | | | | | |
|-----------|------|------------------|----------------|------------|--------------|
| AA | ALRA | <u>10GM/15ML</u> | A071054 | 001 | Jul 26, 1988 |
|-----------|------|------------------|----------------|------------|--------------|

LACTULOSE

| | | | | | |
|-----------|-------------------------|------------------|----------------|------------|--------------|
| AA | ANI PHARMS | <u>10GM/15ML</u> | A078430 | 001 | Nov 28, 2007 |
| AA | BIO-PHARM INC | <u>10GM/15ML</u> | A207786 | 001 | Jun 11, 2018 |
| AA | FRESENIUS KABI | <u>10GM/15ML</u> | A090503 | 001 | Jan 25, 2012 |
| AA | ! HI TECH PHARMA | <u>10GM/15ML</u> | A074076 | 001 | Jul 03, 1995 |
| AA | LANNETT CO INC | <u>10GM/15ML</u> | A075993 | 001 | Jul 26, 2001 |
| AA | LIFEPHARMA | <u>10GM/15ML</u> | A209517 | 001 | Nov 23, 2018 |
| AA | PHARM ASSOC | <u>10GM/15ML</u> | A074623 | 001 | Jul 30, 1996 |
| AA | VISTAPHARM | <u>10GM/15ML</u> | A074138 | 001 | Sep 30, 1992 |
| AA | WEST-WARD PHARMS INT | <u>10GM/15ML</u> | A073591 | 001 | May 29, 1992 |
| AA | WOCKHARDT BIO AG | <u>10GM/15ML</u> | A074602 | 001 | Nov 14, 1996 |

SOLUTION; ORAL, RECTAL

CHOLAC

| | | | | | |
|-----------|------|------------------|----------------|------------|--------------|
| AA | ALRA | <u>10GM/15ML</u> | A071331 | 001 | Jul 26, 1988 |
|-----------|------|------------------|----------------|------------|--------------|

ENULOSE

| | | | | | |
|-----------|---------------------------|------------------|----------------|------------|--------------|
| AA | ! ACTAVIS MID ATLANTIC | <u>10GM/15ML</u> | A071548 | 001 | Aug 15, 1988 |
|-----------|---------------------------|------------------|----------------|------------|--------------|

GENERLAC

| | | | | | |
|-----------|------------------|------------------|----------------|------------|--------------|
| AA | WOCKHARDT BIO AG | <u>10GM/15ML</u> | A074603 | 001 | Oct 31, 1996 |
|-----------|------------------|------------------|----------------|------------|--------------|

LACTULOSE

| | | | | | |
|-----------|----------------|------------------|----------------|------------|--------------|
| AA | ANI PHARMS | <u>10GM/15ML</u> | A090426 | 001 | Nov 21, 2008 |
| AA | BIO-PHARM INC | <u>10GM/15ML</u> | A203762 | 001 | Mar 27, 2015 |
| AA | FRESENIUS KABI | <u>10GM/15ML</u> | A090502 | 001 | Jan 25, 2012 |
| AA | HI TECH PHARMA | <u>10GM/15ML</u> | A074077 | 001 | Jul 03, 1995 |

LAMIVUDINE

SOLUTION; ORAL

EPIVIR

| | | | | | |
|-----------|------------------|----------------|----------------|------------|--------------|
| AA | +! VIIV HLTHCARE | <u>10MG/ML</u> | N020596 | 001 | Nov 17, 1995 |
|-----------|------------------|----------------|----------------|------------|--------------|

LAMIVUDINE

| | | | | | |
|-----------|-------------------------|----------------|----------------|------------|--------------|
| AA | AUROBINDO PHARMA LTD | <u>10MG/ML</u> | A077695 | 001 | Nov 21, 2016 |
| AA | LANNETT CO INC | <u>10MG/ML</u> | A203564 | 001 | Oct 31, 2014 |
| | EPIVIR-HBV | | | | |
| | +! GLAXOSMITHKLINE | 5MG/ML | N021004 | 001 | Dec 08, 1998 |

TABLET; ORAL

EPIVIR

| | | | | | |
|-----------|-----------------|--------------|----------------|------------|--------------|
| AB | + VIIV HLTHCARE | <u>150MG</u> | N020564 | 001 | Nov 17, 1995 |
| AB | +! | <u>300MG</u> | N020564 | 003 | Jun 24, 2002 |

EPIVIR-HBV

| | | | | | |
|-----------|--------------------|--------------|----------------|------------|--------------|
| AB | +! GLAXOSMITHKLINE | <u>100MG</u> | N021003 | 001 | Dec 08, 1998 |
|-----------|--------------------|--------------|----------------|------------|--------------|

LAMIVUDINE

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| AB | APOTEX | <u>150MG</u> | A091606 | 001 | Dec 02, 2011 |
| AB | | <u>300MG</u> | A091606 | 002 | Dec 02, 2011 |
| AB | APOTEX INC | <u>100MG</u> | A202941 | 001 | Jan 02, 2014 |
| AB | ARISE PHARMS | <u>150MG</u> | A206974 | 001 | Nov 21, 2016 |
| AB | | <u>300MG</u> | A206974 | 002 | Nov 21, 2016 |
| AB | AUROBINDO PHARMA LTD | <u>150MG</u> | A077464 | 001 | Nov 21, 2016 |
| AB | | <u>150MG</u> | A202032 | 001 | Nov 17, 2011 |
| AB | | <u>300MG</u> | A077464 | 002 | Nov 21, 2016 |
| AB | | <u>300MG</u> | A202032 | 002 | Nov 17, 2011 |
| AB | CIPLA | <u>150MG</u> | A077221 | 001 | Mar 03, 2017 |
| AB | | <u>300MG</u> | A077221 | 002 | Mar 03, 2017 |
| AB | ECI PHARMS LLC | <u>150MG</u> | A203586 | 001 | Nov 21, 2016 |
| AB | HETERO LABS LTD V | <u>100MG</u> | A203260 | 001 | Jan 02, 2014 |
| AB | | <u>150MG</u> | A203277 | 001 | Jan 06, 2014 |
| AB | | <u>300MG</u> | A203277 | 002 | Jan 06, 2014 |
| AB | LUPIN LTD | <u>150MG</u> | A205217 | 001 | Dec 18, 2014 |
| AB | | <u>300MG</u> | A205217 | 002 | Dec 18, 2014 |
| AB | MYLAN PHARMS INC | <u>100MG</u> | A204002 | 001 | Dec 31, 2014 |
| AB | | <u>150MG</u> | A204528 | 001 | Mar 04, 2016 |
| AB | | <u>300MG</u> | A204528 | 002 | Mar 04, 2016 |
| AB | STRIDES PHARMA | <u>150MG</u> | A090457 | 001 | Apr 19, 2018 |
| AB | | <u>300MG</u> | A090457 | 002 | Apr 19, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

CIMDUO

+! MYLAN LABS LTD 300MG;300MG N022141 001 Feb 28, 2018

TEMIXYS

CELLTRION 300MG;300MG N211284 001 Nov 16, 2018

LAMIVUDINE; ZIDOVUDINE

TABLET; ORAL

COMBIVIR**AB +! VIIV HLTHCARE 150MG;300MG N020857 001 Sep 26, 1997****LAMIVUDINE AND ZIDOVUDINE****AB AUROBINDO PHARMA LTD 150MG;300MG A077558 001 May 05, 2017****AB 150MG;300MG A202418 001 May 15, 2012****AB CIPLA 150MG;300MG A077411 001 Sep 07, 2018****AB HETERO LABS LTD III 150MG;300MG A079124 001 Sep 17, 2015****AB HETERO LABS LTD V 150MG;300MG A203259 001 Feb 03, 2014****AB LUPIN LTD 150MG;300MG A090246 001 May 15, 2012****AB MACLEODS PHARMS LTD 150MG;300MG A090679 001 Aug 29, 2018****AB MYLAN PHARMS INC 150MG;300MG A204005 001 Aug 28, 2014****AB SHANGHAI DESANO 150MG;300MG A206375 001 Apr 10, 2018****AB STRIDES PHARMA 150MG;300MG A079128 001 May 13, 2015**LAMOTRIGINE

TABLET; ORAL

LAMICTAL**AB +! GLAXOSMITHKLINE LLC 25MG N020241 005 Dec 27, 1994****AB + 100MG N020241 001 Dec 27, 1994****AB + 150MG N020241 002 Dec 27, 1994****AB + 200MG N020241 003 Dec 27, 1994****LAMOTRIGINE****AB ALEMBIC PHARMS LTD 25MG A090607 001 Jan 13, 2011****AB 100MG A090607 002 Jan 13, 2011****AB 150MG A090607 003 Jan 13, 2011****AB 200MG A090607 004 Jan 13, 2011****AB ALKEM LABS LTD 25MG A200694 001 Jun 14, 2013****AB 100MG A200694 002 Jun 14, 2013****AB 150MG A200694 003 Jun 14, 2013****AB 200MG A200694 004 Jun 14, 2013****AB APOTEX INC 25MG A078625 001 Jan 27, 2009****AB 100MG A078625 002 Jan 27, 2009****AB 150MG A078625 003 Jan 27, 2009****AB 200MG A078625 004 Jan 27, 2009****AB AUROBINDO PHARMA 25MG A078956 001 Jan 27, 2009****AB 100MG A078956 002 Jan 27, 2009****AB 150MG A078956 003 Jan 27, 2009****AB 200MG A078956 004 Jan 27, 2009****AB CIPLA 25MG A077783 001 Nov 01, 2010****AB 100MG A077783 002 Nov 01, 2010****AB 150MG A077783 003 Nov 01, 2010****AB 200MG A077783 004 Nov 01, 2010****AB DR REDDYS LABS LTD 25MG A076708 001 Jan 27, 2009****AB 100MG A076708 002 Jan 27, 2009****AB 150MG A076708 003 Jan 27, 2009****AB 200MG A076708 004 Jan 27, 2009****AB GLENMARK GENERICS 25MG A090169 001 May 12, 2012****AB 100MG A090169 002 May 12, 2012****AB 150MG A090169 003 May 12, 2012****AB 200MG A090169 004 May 12, 2012****AB JUBILANT CADISTA 25MG A079132 001 Jan 27, 2009****AB 100MG A079132 002 Jan 27, 2009****AB 150MG A079132 003 Jan 27, 2009****AB 200MG A079132 004 Jan 27, 2009****AB LUPIN LTD 25MG A078691 001 Jun 01, 2010****AB 100MG A078691 002 Jun 01, 2010****AB 150MG A078691 003 Jun 01, 2010****AB 200MG A078691 004 Jun 01, 2010****AB MYLAN 25MG A077420 001 Jan 27, 2009****AB 100MG A077420 002 Jan 27, 2009****AB 150MG A077420 003 Jan 27, 2009****AB 200MG A077420 004 Jan 27, 2009****AB TARO PHARM INDS 25MG A078525 001 Jan 27, 2009****AB 100MG A078525 002 Jan 27, 2009**

PRESCRIPTION DRUG PRODUCT LIST

LAMOTRIGINE

TABLET; ORAL

LAMOTRIGINE

| | | | | |
|-----------|------------------|--------------|--------------------|--------------|
| <u>AB</u> | | <u>150MG</u> | <u>A078525 003</u> | Jan 27, 2009 |
| <u>AB</u> | | <u>200MG</u> | <u>A078525 004</u> | Jan 27, 2009 |
| <u>AB</u> | TEVA | <u>25MG</u> | <u>A076388 001</u> | Aug 30, 2006 |
| <u>AB</u> | | <u>100MG</u> | <u>A076388 002</u> | Aug 30, 2006 |
| <u>AB</u> | | <u>150MG</u> | <u>A076388 003</u> | Aug 30, 2006 |
| <u>AB</u> | | <u>200MG</u> | <u>A076388 004</u> | Aug 30, 2006 |
| <u>AB</u> | TORRENT PHARMS | <u>25MG</u> | <u>A078947 001</u> | Jan 27, 2009 |
| <u>AB</u> | | <u>100MG</u> | <u>A078947 002</u> | Jan 27, 2009 |
| <u>AB</u> | | <u>150MG</u> | <u>A078947 003</u> | Jan 27, 2009 |
| <u>AB</u> | | <u>200MG</u> | <u>A078947 004</u> | Jan 27, 2009 |
| <u>AB</u> | UNICHEM LABS LTD | <u>25MG</u> | <u>A090170 001</u> | Oct 06, 2011 |
| <u>AB</u> | | <u>100MG</u> | <u>A090170 002</u> | Oct 06, 2011 |
| <u>AB</u> | | <u>150MG</u> | <u>A090170 003</u> | Oct 06, 2011 |
| <u>AB</u> | | <u>200MG</u> | <u>A090170 004</u> | Oct 06, 2011 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>25MG</u> | <u>A077633 001</u> | Jan 27, 2009 |
| <u>AB</u> | | <u>100MG</u> | <u>A077633 003</u> | Jan 27, 2009 |
| <u>AB</u> | | <u>150MG</u> | <u>A077633 004</u> | Jan 27, 2009 |
| <u>AB</u> | | <u>200MG</u> | <u>A077633 005</u> | Jan 27, 2009 |
| | | 50MG | A077633 002 | Jan 27, 2009 |
| | | 250MG | A077633 006 | Jan 27, 2009 |

TABLET, CHEWABLE; ORAL

LAMICTAL CD

| | | | | | |
|-----------|---|---------------------|-------------|--------------------|--------------|
| <u>AB</u> | + | GLAXOSMITHKLINE LLC | <u>2MG</u> | <u>N020764 004</u> | Sep 08, 2000 |
| <u>AB</u> | + | | <u>5MG</u> | <u>N020764 001</u> | Aug 24, 1998 |
| <u>AB</u> | + | ! | <u>25MG</u> | <u>N020764 002</u> | Aug 24, 1998 |

LAMOTRIGINE

| | | | | |
|-----------|-------------------------|-------------|--------------------|--------------|
| <u>AB</u> | ALEMBIC PHARMS LTD | <u>5MG</u> | <u>A201168 001</u> | Jun 12, 2014 |
| <u>AB</u> | | <u>25MG</u> | <u>A201168 002</u> | Jun 12, 2014 |
| <u>AB</u> | AUROBINDO PHARMA | <u>5MG</u> | <u>A090401 002</u> | Nov 04, 2009 |
| <u>AB</u> | | <u>25MG</u> | <u>A090401 003</u> | Nov 04, 2009 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>5MG</u> | <u>A076701 001</u> | Jan 22, 2009 |
| <u>AB</u> | | <u>25MG</u> | <u>A076701 002</u> | Jan 22, 2009 |
| <u>AB</u> | GLENMARK PHARMS LTD | <u>5MG</u> | <u>A079099 001</u> | Feb 19, 2009 |
| <u>AB</u> | | <u>25MG</u> | <u>A079099 002</u> | Feb 19, 2009 |
| <u>AB</u> | JUBILANT GENERICS | <u>5MG</u> | <u>A200220 001</u> | Feb 28, 2011 |
| <u>AB</u> | | <u>25MG</u> | <u>A200220 002</u> | Feb 28, 2011 |
| <u>AB</u> | TARO | <u>5MG</u> | <u>A079204 001</u> | Feb 04, 2009 |
| <u>AB</u> | | <u>25MG</u> | <u>A079204 002</u> | Feb 04, 2009 |
| <u>AB</u> | TEVA | <u>5MG</u> | <u>A076420 001</u> | Jun 21, 2006 |
| <u>AB</u> | | <u>25MG</u> | <u>A076420 002</u> | Jun 21, 2006 |
| <u>AB</u> | WATSON LABS | <u>2MG</u> | <u>A076928 001</u> | Jan 22, 2009 |
| <u>AB</u> | | <u>5MG</u> | <u>A076928 002</u> | Jan 22, 2009 |
| <u>AB</u> | | <u>25MG</u> | <u>A076928 003</u> | Jan 22, 2009 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>5MG</u> | <u>A078009 002</u> | Jan 22, 2009 |
| <u>AB</u> | | <u>25MG</u> | <u>A078009 003</u> | Jan 22, 2009 |

TABLET, EXTENDED RELEASE; ORAL

LAMICTAL XR

| | | | | | |
|-----------|---|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | + | GLAXOSMITHKLINE LLC | <u>25MG</u> | <u>N022115 001</u> | May 29, 2009 |
| <u>AB</u> | + | ! | <u>50MG</u> | <u>N022115 002</u> | May 29, 2009 |
| <u>AB</u> | + | | <u>100MG</u> | <u>N022115 003</u> | May 29, 2009 |
| <u>AB</u> | + | ! | <u>200MG</u> | <u>N022115 004</u> | May 29, 2009 |
| <u>AB</u> | + | | <u>250MG</u> | <u>N022115 006</u> | Jun 21, 2011 |
| <u>AB</u> | + | | <u>300MG</u> | <u>N022115 005</u> | Apr 14, 2010 |

LAMOTRIGINE

| | | | | |
|-----------|-------------------|--------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS ELIZABETH | <u>100MG</u> | <u>A200672 003</u> | Oct 17, 2013 |
| <u>AB</u> | | <u>200MG</u> | <u>A200672 004</u> | Oct 17, 2013 |
| <u>AB</u> | | <u>25MG</u> | <u>A200672 001</u> | Oct 17, 2013 |
| <u>AB</u> | | <u>50MG</u> | <u>A200672 002</u> | Oct 17, 2013 |
| <u>AB</u> | | <u>250MG</u> | <u>A203733 001</u> | Nov 13, 2013 |
| <u>AB</u> | | <u>300MG</u> | <u>A200672 005</u> | Oct 17, 2013 |
| <u>AB</u> | AMNEAL PHARMS | <u>25MG</u> | <u>A207497 001</u> | Nov 30, 2018 |
| <u>AB</u> | | <u>50MG</u> | <u>A207497 002</u> | Nov 30, 2018 |
| <u>AB</u> | | <u>100MG</u> | <u>A207497 003</u> | Nov 30, 2018 |
| <u>AB</u> | | <u>200MG</u> | <u>A207497 004</u> | Nov 30, 2018 |
| <u>AB</u> | | <u>250MG</u> | <u>A207497 005</u> | Nov 30, 2018 |
| <u>AB</u> | | <u>300MG</u> | <u>A207497 006</u> | Nov 30, 2018 |
| <u>AB</u> | ANCHEN PHARMS | <u>25MG</u> | <u>A201374 001</u> | Dec 26, 2012 |
| <u>AB</u> | | <u>50MG</u> | <u>A201374 002</u> | Dec 26, 2012 |
| <u>AB</u> | | <u>100MG</u> | <u>A201374 003</u> | Dec 26, 2012 |

PRESCRIPTION DRUG PRODUCT LIST

LAMOTRIGINE

TABLET, EXTENDED RELEASE;ORAL

LAMOTRIGINE

| | | | | | |
|-----------|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | <u>200MG</u> | <u>A201374</u> | <u>004</u> | Dec 26, 2012 |
| <u>AB</u> | | <u>250MG</u> | <u>A201374</u> | <u>005</u> | Dec 26, 2012 |
| <u>AB</u> | | <u>300MG</u> | <u>A201374</u> | <u>006</u> | Dec 26, 2012 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>25MG</u> | <u>A202383</u> | <u>001</u> | Jun 19, 2013 |
| <u>AB</u> | | <u>50MG</u> | <u>A202383</u> | <u>002</u> | Jun 19, 2013 |
| <u>AB</u> | | <u>100MG</u> | <u>A202383</u> | <u>003</u> | Jun 19, 2013 |
| <u>AB</u> | | <u>200MG</u> | <u>A202383</u> | <u>004</u> | Jun 19, 2013 |
| <u>AB</u> | | <u>300MG</u> | <u>A202383</u> | <u>005</u> | Jun 19, 2013 |
| <u>AB</u> | HANDA PHARMS LLC | <u>100MG</u> | <u>A202887</u> | <u>003</u> | Jun 17, 2013 |
| <u>AB</u> | | <u>200MG</u> | <u>A202887</u> | <u>004</u> | Jun 17, 2013 |
| <u>AB</u> | PAR PHARM | <u>25MG</u> | <u>A201791</u> | <u>001</u> | Jan 18, 2013 |
| <u>AB</u> | | <u>50MG</u> | <u>A201791</u> | <u>002</u> | Jan 18, 2013 |
| <u>AB</u> | | <u>100MG</u> | <u>A201791</u> | <u>003</u> | Jan 18, 2013 |
| <u>AB</u> | | <u>200MG</u> | <u>A201791</u> | <u>004</u> | Jan 18, 2013 |
| <u>AB</u> | | <u>250MG</u> | <u>A201791</u> | <u>005</u> | Jan 18, 2013 |
| <u>AB</u> | | <u>300MG</u> | <u>A201791</u> | <u>006</u> | Jan 18, 2013 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>25MG</u> | <u>A203370</u> | <u>001</u> | Dec 23, 2013 |
| <u>AB</u> | | <u>50MG</u> | <u>A203370</u> | <u>002</u> | Dec 23, 2013 |
| <u>AB</u> | | <u>100MG</u> | <u>A203370</u> | <u>003</u> | Dec 23, 2013 |
| <u>AB</u> | | <u>200MG</u> | <u>A203370</u> | <u>004</u> | Dec 23, 2013 |
| <u>AB</u> | WOCKHARDT LTD | <u>25MG</u> | <u>A202498</u> | <u>001</u> | Jan 04, 2013 |
| <u>AB</u> | | <u>50MG</u> | <u>A202498</u> | <u>002</u> | Jan 04, 2013 |
| <u>AB</u> | | <u>100MG</u> | <u>A202498</u> | <u>003</u> | Jan 04, 2013 |
| <u>AB</u> | | <u>200MG</u> | <u>A202498</u> | <u>004</u> | Jan 04, 2013 |
| <u>AB</u> | | <u>300MG</u> | <u>A202498</u> | <u>005</u> | Jan 04, 2013 |

TABLET, ORALLY DISINTEGRATING;ORAL

LAMICTAL ODT

| | | | | | | |
|-----------|---|---------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | GLAXOSMITHKLINE LLC | <u>25MG</u> | <u>N022251</u> | <u>001</u> | May 08, 2009 |
| <u>AB</u> | + | ! | <u>50MG</u> | <u>N022251</u> | <u>002</u> | May 08, 2009 |
| <u>AB</u> | + | | <u>100MG</u> | <u>N022251</u> | <u>003</u> | May 08, 2009 |
| <u>AB</u> | + | | <u>200MG</u> | <u>N022251</u> | <u>004</u> | May 08, 2009 |

LAMOTRIGINE

| | | | | | |
|-----------|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | IMPAX LABS INC | <u>25MG</u> | <u>A200828</u> | <u>001</u> | Jul 15, 2013 |
| <u>AB</u> | | <u>50MG</u> | <u>A200828</u> | <u>002</u> | Jul 15, 2013 |
| <u>AB</u> | | <u>100MG</u> | <u>A200828</u> | <u>003</u> | Jul 15, 2013 |
| <u>AB</u> | | <u>200MG</u> | <u>A200828</u> | <u>004</u> | Jul 15, 2013 |
| <u>AB</u> | PAR PHARM | <u>25MG</u> | <u>A204158</u> | <u>001</u> | Oct 27, 2015 |
| <u>AB</u> | | <u>50MG</u> | <u>A204158</u> | <u>002</u> | Oct 27, 2015 |
| <u>AB</u> | | <u>100MG</u> | <u>A204158</u> | <u>003</u> | Oct 27, 2015 |
| <u>AB</u> | | <u>200MG</u> | <u>A204158</u> | <u>004</u> | Oct 27, 2015 |
| <u>AB</u> | SCIEGEN PHARMS INC | <u>25MG</u> | <u>A206382</u> | <u>001</u> | Jun 17, 2016 |
| <u>AB</u> | | <u>50MG</u> | <u>A206382</u> | <u>002</u> | Jun 17, 2016 |
| <u>AB</u> | | <u>100MG</u> | <u>A206382</u> | <u>003</u> | Jun 17, 2016 |
| <u>AB</u> | | <u>200MG</u> | <u>A206382</u> | <u>004</u> | Jun 17, 2016 |

LANREOTIDE ACETATE

SOLUTION;SUBCUTANEOUS

SOMATULINE DEPOT

| | | | | | | |
|---|---|--------------|-------------------------------------------|---------|-----|--------------|
| + | ! | IPSEN PHARMA | EQ 60MG BASE/0.2ML (EQ 60MG BASE/0.2ML) | N022074 | 001 | Aug 30, 2007 |
| + | ! | | EQ 90MG BASE/0.3ML (EQ 90MG BASE/0.3ML) | N022074 | 002 | Aug 30, 2007 |
| + | ! | | EQ 120MG BASE/0.5ML (EQ 120MG BASE/0.5ML) | N022074 | 003 | Aug 30, 2007 |

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

| | | | | | |
|-----------|----------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | AJANTA PHARMA LTD | <u>15MG</u> | <u>A203957</u> | <u>001</u> | Oct 14, 2016 |
| <u>AB</u> | | <u>30MG</u> | <u>A203957</u> | <u>002</u> | Oct 14, 2016 |
| <u>AB</u> | BRECKENRIDGE PHARM | <u>15MG</u> | <u>A203964</u> | <u>001</u> | Oct 17, 2018 |
| <u>AB</u> | | <u>30MG</u> | <u>A203964</u> | <u>002</u> | Oct 17, 2018 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>15MG</u> | <u>A091269</u> | <u>001</u> | Oct 15, 2010 |
| <u>AB</u> | | <u>30MG</u> | <u>A091269</u> | <u>002</u> | Oct 15, 2010 |
| <u>AB</u> | INVENTIA HLTHCARE | <u>15MG</u> | <u>A205868</u> | <u>001</u> | Aug 30, 2017 |
| <u>AB</u> | | <u>30MG</u> | <u>A205868</u> | <u>002</u> | Aug 30, 2017 |
| <u>AB</u> | KRKA TOVARNA ZDRAVIL | <u>15MG</u> | <u>A091212</u> | <u>001</u> | Sep 16, 2013 |
| <u>AB</u> | | <u>30MG</u> | <u>A091212</u> | <u>002</u> | Sep 16, 2013 |
| <u>AB</u> | LABS LICONSA | <u>15MG</u> | <u>A203203</u> | <u>001</u> | Jul 25, 2016 |
| <u>AB</u> | | <u>30MG</u> | <u>A203203</u> | <u>002</u> | Jul 25, 2016 |
| <u>AB</u> | LANNETT CO INC | <u>15MG</u> | <u>A207156</u> | <u>001</u> | Sep 28, 2017 |
| <u>AB</u> | | <u>30MG</u> | <u>A207156</u> | <u>002</u> | Sep 28, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

| | | | | | |
|-----------|--------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | MYLAN PHARMS INC | <u>15MG</u> | <u>A090763</u> | <u>001</u> | Nov 10, 2009 |
| <u>AB</u> | | <u>30MG</u> | <u>A090763</u> | <u>002</u> | Nov 10, 2009 |
| <u>AB</u> | NATCO PHARMA LTD | <u>15MG</u> | <u>A201921</u> | <u>001</u> | Dec 18, 2012 |
| <u>AB</u> | | <u>30MG</u> | <u>A201921</u> | <u>002</u> | Dec 18, 2012 |
| <u>AB</u> | SANDOZ | <u>15MG</u> | <u>A090331</u> | <u>001</u> | Apr 23, 2010 |
| <u>AB</u> | | <u>30MG</u> | <u>A090331</u> | <u>002</u> | Apr 23, 2010 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>15MG</u> | <u>A202637</u> | <u>001</u> | Sep 13, 2013 |
| <u>AB</u> | | <u>30MG</u> | <u>A091509</u> | <u>001</u> | Sep 13, 2013 |
| <u>AB</u> | TEVA PHARMS | <u>15MG</u> | <u>A077255</u> | <u>001</u> | Nov 10, 2009 |
| <u>AB</u> | | <u>30MG</u> | <u>A077255</u> | <u>002</u> | Nov 10, 2009 |
| <u>AB</u> | WOCKHARDT USA | <u>15MG</u> | <u>A202176</u> | <u>001</u> | Sep 14, 2012 |
| <u>AB</u> | | <u>30MG</u> | <u>A202176</u> | <u>002</u> | Sep 14, 2012 |
| <u>AB</u> | ZYDUS HLTHCARE | <u>15MG</u> | <u>A202366</u> | <u>001</u> | Aug 19, 2013 |
| <u>AB</u> | | <u>30MG</u> | <u>A202366</u> | <u>002</u> | Aug 19, 2013 |

PREVACID

| | | | | | |
|-----------|---------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | + TAKEDA PHARMS USA | <u>15MG</u> | <u>N020406</u> | <u>001</u> | May 10, 1995 |
| <u>AB</u> | +! | <u>30MG</u> | <u>N020406</u> | <u>002</u> | May 10, 1995 |

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

LANSOPRAZOLE

| | | | | | |
|-----------|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | MYLAN PHARMS INC | <u>15MG</u> | <u>A202396</u> | <u>001</u> | Nov 28, 2018 |
| <u>AB</u> | | <u>30MG</u> | <u>A202396</u> | <u>002</u> | Nov 28, 2018 |
| <u>AB</u> | TEVA PHARMS USA | <u>15MG</u> | <u>A208784</u> | <u>001</u> | Sep 21, 2017 |
| <u>AB</u> | | <u>30MG</u> | <u>A208784</u> | <u>002</u> | Sep 21, 2017 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>15MG</u> | <u>A200816</u> | <u>001</u> | Nov 27, 2018 |
| <u>AB</u> | | <u>30MG</u> | <u>A200816</u> | <u>002</u> | Nov 27, 2018 |

PREVACID

| | | | | | |
|-----------|---------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | + TAKEDA PHARMS USA | <u>15MG</u> | <u>N021428</u> | <u>001</u> | Aug 30, 2002 |
| <u>AB</u> | +! | <u>30MG</u> | <u>N021428</u> | <u>002</u> | Aug 30, 2002 |

LANTHANUM CARBONATE

POWDER;ORAL

FOSRENOL

| | | | | | |
|--|-----------------|---------------|---------|-----|--------------|
| | + SHIRE DEV LLC | EQ 750MG BASE | N204734 | 001 | Sep 24, 2014 |
| | +! | EQ 1GM BASE | N204734 | 002 | Sep 24, 2014 |

TABLET, CHEWABLE;ORAL

FOSRENOL

| | | | | | |
|-----------|-------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | + SHIRE LLC | <u>EQ 500MG BASE</u> | <u>N021468</u> | <u>002</u> | Oct 26, 2004 |
| <u>AB</u> | +! | <u>EQ 750MG BASE</u> | <u>N021468</u> | <u>003</u> | Nov 23, 2005 |
| <u>AB</u> | +! | <u>EQ 1GM BASE</u> | <u>N021468</u> | <u>004</u> | Nov 23, 2005 |

LANTHANUM CARBONATE

| | | | | | |
|-----------|------------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | NATCO PHARMA LTD | <u>EQ 500MG BASE</u> | <u>A090978</u> | <u>001</u> | Aug 11, 2017 |
| <u>AB</u> | | <u>EQ 750MG BASE</u> | <u>A090978</u> | <u>002</u> | Aug 11, 2017 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A090978</u> | <u>003</u> | Aug 11, 2017 |

LAPATINIB DITOSYLATE

TABLET;ORAL

TYKERB

| | | | | | |
|--|----------------------------|---------------|---------|-----|--------------|
| | +! NOVARTIS PHARMS CORP | EQ 250MG BASE | N022059 | 001 | Mar 13, 2007 |
|--|----------------------------|---------------|---------|-----|--------------|

LAROTRECTINIB

CAPSULE;ORAL

VITRAKVI

| | | | | | |
|--|---------------------|-------|---------|-----|--------------|
| | + LOXO ONCOLOGY INC | 25MG | N210861 | 001 | Nov 26, 2018 |
| | +! | 100MG | N210861 | 002 | Nov 26, 2018 |

SOLUTION;ORAL

VITRAKVI

| | | | | | |
|--|----------------------|---------|---------|-----|--------------|
| | +! LOXO ONCOLOGY INC | 20MG/ML | N211710 | 001 | Nov 26, 2018 |
|--|----------------------|---------|---------|-----|--------------|

LATANOPROST

EMULSION;OPHTHALMIC

XELPROS

| | | | | | |
|--|----------------------|--------|---------|-----|--------------|
| | +! SUN PHARMA GLOBAL | 0.005% | N206185 | 001 | Sep 12, 2018 |
|--|----------------------|--------|---------|-----|--------------|

SOLUTION/DROPS;OPHTHALMIC

LATANOPROST

| | | | | | |
|-----------|--------------------|---------------|----------------|------------|--------------|
| <u>AT</u> | AKORN | <u>0.005%</u> | <u>A090887</u> | <u>001</u> | Jul 19, 2011 |
| <u>AT</u> | AMRING PHARMS | <u>0.005%</u> | <u>A200925</u> | <u>001</u> | Mar 22, 2011 |
| <u>AT</u> | BAUSCH AND LOMB | <u>0.005%</u> | <u>A201006</u> | <u>001</u> | Mar 22, 2011 |
| <u>AT</u> | DR REDDYS LABS LTD | <u>0.005%</u> | <u>A202077</u> | <u>001</u> | Feb 11, 2013 |
| <u>AT</u> | FDC LTD | <u>0.005%</u> | <u>A202442</u> | <u>001</u> | Apr 22, 2016 |
| <u>AT</u> | MYLAN | <u>0.005%</u> | <u>A201786</u> | <u>001</u> | Mar 22, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

LATANOPROST

SOLUTION/DROPS;OPHTHALMIC

LATANOPROST

| | | | | | |
|-----------|---|------------|---------------|--------------------|--------------|
| AT | + | SANDOZ INC | 0.005% | A091449 001 | Mar 22, 2011 |
|-----------|---|------------|---------------|--------------------|--------------|

XALATAN

| | | | | | |
|-----------|---|-------------------------|---------------|--------------------|--------------|
| AT | + | PHARMACIA AND UPJOHN | 0.005% | N020597 001 | Jun 05, 1996 |
|-----------|---|-------------------------|---------------|--------------------|--------------|

LATANOPROSTENE BUNOD

SOLUTION/DROPS;OPHTHALMIC

VYZULTA

| | | | | |
|---|-----------------|--------|-------------|--------------|
| + | BAUSCH AND LOMB | 0.024% | N207795 001 | Nov 02, 2017 |
|---|-----------------|--------|-------------|--------------|

LEDIPASVIR; SOFOSBUVIR

TABLET;ORAL

HARVONI

| | | | | |
|---|---------------------|------------|-------------|--------------|
| + | GILEAD SCIENCES INC | 90MG;400MG | N205834 001 | Oct 10, 2014 |
|---|---------------------|------------|-------------|--------------|

LEFLUNOMIDE

TABLET;ORAL

ARAVA

| | | | | | |
|-----------|---|-------------------|-------------|--------------------|--------------|
| AB | + | SANOFI AVENTIS US | 10MG | N020905 001 | Sep 10, 1998 |
| AB | + | | 20MG | N020905 002 | Sep 10, 1998 |

LEFLUNOMIDE

| | | | | | |
|-----------|--|---------------------|-------------|--------------------|--------------|
| AB | | ALEMBIC PHARMS LTD | 10MG | A091369 001 | Nov 21, 2011 |
| AB | | | 20MG | A091369 002 | Nov 21, 2011 |
| AB | | APOTEX INC | 10MG | A077090 001 | Sep 13, 2005 |
| AB | | | 20MG | A077090 002 | Sep 13, 2005 |
| AB | | BARR | 10MG | A077083 001 | Sep 13, 2005 |
| AB | | | 20MG | A077083 002 | Sep 13, 2005 |
| AB | | HERITAGE PHARMS INC | 10MG | A077086 001 | Sep 13, 2005 |
| AB | | | 20MG | A077086 002 | Sep 13, 2005 |
| AB | | TEVA PHARMS | 10MG | A077084 001 | Sep 13, 2005 |
| AB | | | 20MG | A077084 002 | Sep 13, 2005 |

ARAVA

| | | | | |
|---|-------------------|-------|-------------|--------------|
| + | SANOFI AVENTIS US | 100MG | N020905 003 | Sep 10, 1998 |
|---|-------------------|-------|-------------|--------------|

LENALIDOMIDE

CAPSULE;ORAL

REVLIMID

| | | | | |
|---|---------|-------|-------------|--------------|
| + | CELGENE | 2.5MG | N021880 005 | Dec 21, 2011 |
| + | | 5MG | N021880 001 | Dec 27, 2005 |
| + | | 10MG | N021880 002 | Dec 27, 2005 |
| + | | 15MG | N021880 003 | Jun 29, 2006 |
| + | | 20MG | N021880 006 | Jun 05, 2013 |
| + | | 25MG | N021880 004 | Jun 29, 2006 |

LENVATINIB MESYLATE

CAPSULE;ORAL

LENVIMA

| | | | | |
|---|-----------|--------------|-------------|--------------|
| + | EISAI INC | EQ 4MG BASE | N206947 001 | Feb 13, 2015 |
| + | | EQ 10MG BASE | N206947 002 | Feb 13, 2015 |

LESINURAD

TABLET;ORAL

ZURAMPIC

| | | | | |
|---|---------------------|-------|-------------|--------------|
| + | IRONWOOD PHARMS INC | 200MG | N207988 001 | Dec 22, 2015 |
|---|---------------------|-------|-------------|--------------|

LETERMOVIR

SOLUTION;INTRAVENOUS

PREVMIS

| | | | | |
|---|-------------------|----------------------|-------------|--------------|
| + | MERCK SHARP DOHME | 240MG/12ML (20MG/ML) | N209940 001 | Nov 08, 2017 |
| + | | 480MG/24ML (20MG/ML) | N209940 002 | Nov 08, 2017 |

TABLET;ORAL

PREVMIS

| | | | | |
|---|-------------------|-------|-------------|--------------|
| + | MERCK SHARP DOHME | 240MG | N209939 001 | Nov 08, 2017 |
| + | | 480MG | N209939 002 | Nov 08, 2017 |

LETROZOLE

TABLET;ORAL

FEMARA

| | | | | | |
|-----------|---|-----------------|--------------|--------------------|--------------|
| AB | + | NOVARTIS PHARMS | 2.5MG | N020726 001 | Jul 25, 1997 |
|-----------|---|-----------------|--------------|--------------------|--------------|

LETROZOLE

| | | | | | |
|-----------|--|--------------------|--------------|--------------------|--------------|
| AB | | ACCORD HLTHCARE | 2.5MG | A090934 001 | Jun 03, 2011 |
| AB | | APOTEX INC | 2.5MG | A091303 001 | Apr 19, 2012 |
| AB | | BEIJING YILING | 2.5MG | A205869 001 | Nov 14, 2018 |
| AB | | DR REDDYS LABS LTD | 2.5MG | A091191 001 | Jun 03, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

LETROZOLE

TABLET; ORAL

LETROZOLE

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | EUGIA PHARMA | <u>2.5MG</u> | <u>A211717</u> | <u>001</u> | Jan 11, 2019 |
| <u>AB</u> | FRESENIUS KABI ONCOL | <u>2.5MG</u> | <u>A090491</u> | <u>001</u> | Jun 03, 2011 |
| <u>AB</u> | HIKMA PHARMS | <u>2.5MG</u> | <u>A203796</u> | <u>001</u> | Jun 03, 2016 |
| <u>AB</u> | INDICUS PHARMA | <u>2.5MG</u> | <u>A201804</u> | <u>001</u> | Jun 03, 2011 |
| <u>AB</u> | JIANGSU HENGRUI MED | <u>2.5MG</u> | <u>A202716</u> | <u>001</u> | May 16, 2013 |
| <u>AB</u> | NATCO PHARMA LTD | <u>2.5MG</u> | <u>A200161</u> | <u>001</u> | Jun 03, 2011 |
| <u>AB</u> | TEVA PHARMS | <u>2.5MG</u> | <u>A090289</u> | <u>001</u> | Jun 03, 2011 |
| <u>AB</u> | VINTAGE PHARMS LLC | <u>2.5MG</u> | <u>A090789</u> | <u>001</u> | Jun 03, 2011 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>2.5MG</u> | <u>A090838</u> | <u>001</u> | Jun 03, 2011 |

LETROZOLE; RIBOCICLIB SUCCINATE

TABLET, TABLET; ORAL

KISQALI FEMARA CO-PACK (COPACKAGED)

| | | | | | | |
|---|---|-------------------------|-----------------------------|---------|-----|--------------|
| + | ! | NOVARTIS PHARMS CORP | 2.5MG,N/A;N/A,EQ 200MG BASE | N209935 | 001 | May 04, 2017 |
|---|---|-------------------------|-----------------------------|---------|-----|--------------|

LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

| | | | | | | |
|-----------|--------------------|---------------------------|---------------------------|----------------|--------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 10MG BASE/ML</u> | <u>A207226</u> | <u>001</u> | Jul 27, 2018 | |
| <u>AP</u> | INGENUS PHARMS LLC | <u>EQ 10MG BASE/VIAL</u> | <u>A210917</u> | <u>001</u> | Nov 23, 2018 | |
| <u>AP</u> | TEVA PHARMS USA | <u>EQ 100MG BASE/VIAL</u> | <u>A081277</u> | <u>001</u> | Sep 28, 1993 | |
| <u>AP</u> | | <u>EQ 350MG BASE/VIAL</u> | <u>A040174</u> | <u>001</u> | Jun 12, 1997 | |
| <u>AP</u> | ! | WEST-WARD PHARMS INT | <u>EQ 50MG BASE/VIAL</u> | <u>A089384</u> | <u>001</u> | Sep 14, 1987 |
| <u>AP</u> | ! | | <u>EQ 100MG BASE/VIAL</u> | <u>A089717</u> | <u>001</u> | Mar 28, 1988 |

LEUCOVORIN CALCIUM PRESERVATIVE FREE

| | | | | | | |
|-----------|--------------------|---------------------------|---------------------------|----------------|--------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 200MG BASE/VIAL</u> | <u>A040258</u> | <u>001</u> | Feb 26, 1999 | |
| <u>AP</u> | ! | | <u>EQ 500MG BASE/VIAL</u> | <u>A040286</u> | <u>001</u> | Feb 26, 1999 |
| <u>AP</u> | MYLAN LABS LTD | <u>EQ 100MG BASE/VIAL</u> | <u>A203800</u> | <u>001</u> | May 19, 2017 | |
| <u>AP</u> | | <u>EQ 200MG BASE/VIAL</u> | <u>A203800</u> | <u>002</u> | May 19, 2017 | |
| <u>AP</u> | | <u>EQ 350MG BASE/VIAL</u> | <u>A203800</u> | <u>003</u> | May 19, 2017 | |
| <u>AP</u> | SAGENT PHARMS | <u>EQ 50MG BASE/VIAL</u> | <u>A200753</u> | <u>001</u> | Sep 06, 2012 | |
| <u>AP</u> | | <u>EQ 100MG BASE/VIAL</u> | <u>A200753</u> | <u>002</u> | Sep 06, 2012 | |
| <u>AP</u> | | <u>EQ 200MG BASE/VIAL</u> | <u>A200753</u> | <u>003</u> | Sep 06, 2012 | |
| <u>AP</u> | | <u>EQ 350MG BASE/VIAL</u> | <u>A200855</u> | <u>001</u> | Sep 06, 2012 | |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A209110</u> | <u>001</u> | Oct 26, 2017 | |
| <u>AP</u> | ! | WEST-WARD PHARMS INT | <u>EQ 10MG BASE/ML</u> | <u>A040347</u> | <u>001</u> | Apr 25, 2000 |
| <u>AP</u> | ! | | <u>EQ 200MG BASE/VIAL</u> | <u>A040056</u> | <u>001</u> | May 23, 1995 |
| <u>AP</u> | ! | | <u>EQ 350MG BASE/VIAL</u> | <u>A040335</u> | <u>001</u> | Apr 20, 2000 |

LEUCOVORIN CALCIUM

FRESENIUS KABI USA EQ 10MG BASE/ML

A207241 001 Mar 14, 2018

TABLET; ORAL

LEUCOVORIN CALCIUM

| | | | | | |
|-----------|-------------------------|---------------------|----------------|------------|--------------|
| <u>AB</u> | BARR | <u>EQ 5MG BASE</u> | <u>A071198</u> | <u>001</u> | Sep 24, 1987 |
| <u>AB</u> | | <u>EQ 25MG BASE</u> | <u>A071199</u> | <u>001</u> | Sep 24, 1987 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>EQ 5MG BASE</u> | <u>A072733</u> | <u>001</u> | Feb 22, 1993 |
| <u>AB</u> | ! | <u>EQ 25MG BASE</u> | <u>A072736</u> | <u>001</u> | Feb 22, 1993 |
| | | EQ 10MG BASE | A072734 | 001 | Feb 22, 1993 |
| | | EQ 15MG BASE | A072735 | 001 | Feb 22, 1993 |

LEUPROLIDE ACETATE

FOR SUSPENSION; INTRAMUSCULAR

LUTRATE DEPOT KIT

+! GP-PHARM SA 22.5MG/VIAL

N205054 001 Aug 28, 2018

INJECTABLE; INJECTION

LEUPROLIDE ACETATE

| | | | | | | |
|-----------|---|-------------------|-------------------------|----------------|------------|------------------|
| <u>AP</u> | ! | SANDOZ | <u>1MG/0.2ML</u> | <u>A074728</u> | <u>001</u> | Aug 04, 1998 |
| <u>AP</u> | | SUN PHARMA GLOBAL | <u>1MG/0.2ML</u> | <u>A078885</u> | <u>001</u> | Mar 09, 2009 |
| <u>AP</u> | | TEVA PHARMS USA | <u>1MG/0.2ML</u> | <u>A075471</u> | <u>001</u> | Oct 25, 2000 |
| | | LUPRON DEPOT | | | | |
| | + | ! | ABBVIE ENDOCRINE INC | 3.75MG | N020011 | 002 Oct 26, 1995 |
| | + | ! | | 7.5MG/VIAL | N019732 | 001 Jan 26, 1989 |
| | + | ! | | 11.25MG/VIAL | N020708 | 001 Mar 07, 1997 |
| | + | | | 22.5MG/VIAL | N020517 | 001 Dec 22, 1995 |
| | + | ! | | 30MG/VIAL | N020517 | 002 May 30, 1997 |
| | + | ! | | 45MG/VIAL | N020517 | 003 Jun 17, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

LEUPROLIDE ACETATE

INJECTABLE; INJECTION

LUPRON DEPOT-PED

| | | | | | | |
|---|---|-------------------------|--------------|---------|-----|--------------|
| + | ! | ABBVIE ENDOCRINE INC | 7.5MG/VIAL | N020263 | 002 | Apr 16, 1993 |
| + | ! | | 11.25MG/VIAL | N020263 | 005 | Jan 21, 1994 |
| + | ! | | 11.25MG/VIAL | N020263 | 007 | Aug 15, 2011 |
| + | ! | | 15MG/VIAL | N020263 | 006 | Jan 21, 1994 |
| + | ! | | 30MG/VIAL | N020263 | 008 | Aug 15, 2011 |

INJECTABLE; SUBCUTANEOUS

ELIGARD

| | | | | | | |
|---|---|---------------|-------------|---------|-----|--------------|
| + | ! | TOLMAR THERAP | 7.5MG/VIAL | N021343 | 001 | Jan 23, 2002 |
| + | ! | | 22.5MG/VIAL | N021379 | 001 | Jul 24, 2002 |
| + | ! | | 30MG/VIAL | N021488 | 001 | Feb 13, 2003 |
| + | ! | | 45MG/VIAL | N021731 | 001 | Dec 14, 2004 |

LEUPROLIDE ACETATE; NORETHINDRONE ACETATE

INJECTABLE, TABLET; INTRAMUSCULAR, ORAL

LUPANETA PACK

| | | | | | | |
|---|---|------------------|-----------------------------|---------|-----|--------------|
| + | | ABBVIE ENDOCRINE | 3.75MG/VIAL, N/A; N/A, 5MG | N203696 | 001 | Dec 14, 2012 |
| + | ! | | 11.25MG/VIAL, N/A; N/A, 5MG | N203696 | 002 | Dec 14, 2012 |

LEVALBUTEROL HYDROCHLORIDE

SOLUTION; INHALATION

LEVALBUTEROL HYDROCHLORIDE

| | | | | | | |
|-----------|---|-------------------------|------------------------|----------------|------------|--------------|
| <u>AN</u> | | AUROBINDO PHARMA LTD | <u>EQ 0.25% BASE</u> | <u>A207628</u> | <u>001</u> | Jan 31, 2017 |
| <u>AN</u> | | | <u>EQ 0.0103% BASE</u> | <u>A207625</u> | <u>001</u> | Dec 30, 2016 |
| <u>AN</u> | | | <u>EQ 0.021% BASE</u> | <u>A207625</u> | <u>002</u> | Dec 30, 2016 |
| <u>AN</u> | | | <u>EQ 0.042% BASE</u> | <u>A207625</u> | <u>003</u> | Dec 30, 2016 |
| <u>AN</u> | | CIPLA | <u>EQ 0.021% BASE</u> | <u>A078171</u> | <u>002</u> | Dec 13, 2013 |
| <u>AN</u> | | | <u>EQ 0.042% BASE</u> | <u>A078171</u> | <u>003</u> | Dec 13, 2013 |
| <u>AN</u> | | | <u>EQ 0.0103% BASE</u> | <u>A078171</u> | <u>001</u> | Dec 13, 2013 |
| <u>AN</u> | | IMPAX LABS INC | <u>EQ 0.0103% BASE</u> | <u>A077756</u> | <u>003</u> | Apr 09, 2008 |
| <u>AN</u> | | | <u>EQ 0.021% BASE</u> | <u>A077756</u> | <u>001</u> | Apr 09, 2008 |
| <u>AN</u> | | | <u>EQ 0.042% BASE</u> | <u>A077756</u> | <u>002</u> | Apr 09, 2008 |
| <u>AN</u> | | MYLAN SPECIALITY LP | <u>EQ 0.0103% BASE</u> | <u>A077800</u> | <u>001</u> | Mar 15, 2013 |
| <u>AN</u> | | | <u>EQ 0.021% BASE</u> | <u>A077800</u> | <u>002</u> | Mar 15, 2013 |
| <u>AN</u> | | | <u>EQ 0.042% BASE</u> | <u>A077800</u> | <u>003</u> | Mar 15, 2013 |
| <u>AN</u> | | | <u>EQ 0.25% BASE</u> | <u>A078309</u> | <u>001</u> | Mar 20, 2009 |
| <u>AN</u> | | RITEDOSE CORP | <u>EQ 0.0103% BASE</u> | <u>A203653</u> | <u>001</u> | Mar 22, 2016 |
| <u>AN</u> | | | <u>EQ 0.021% BASE</u> | <u>A203653</u> | <u>002</u> | Mar 22, 2016 |
| <u>AN</u> | | | <u>EQ 0.042% BASE</u> | <u>A203653</u> | <u>003</u> | Mar 22, 2016 |
| <u>AN</u> | | SUN PHARMA GLOBAL | <u>EQ 0.0103% BASE</u> | <u>A207820</u> | <u>001</u> | Nov 05, 2018 |
| <u>AN</u> | | | <u>EQ 0.021% BASE</u> | <u>A207820</u> | <u>002</u> | Nov 05, 2018 |
| <u>AN</u> | | | <u>EQ 0.042% BASE</u> | <u>A207820</u> | <u>003</u> | Nov 05, 2018 |
| <u>AN</u> | | TEVA PARENTERAL | <u>EQ 0.25% BASE</u> | <u>A200875</u> | <u>001</u> | Sep 11, 2014 |
| <u>AN</u> | | TEVA PHARMS USA | <u>EQ 0.0103% BASE</u> | <u>A090297</u> | <u>001</u> | Apr 26, 2013 |
| <u>AN</u> | | | <u>EQ 0.021% BASE</u> | <u>A090297</u> | <u>002</u> | Apr 26, 2013 |
| <u>AN</u> | | | <u>EQ 0.042% BASE</u> | <u>A090297</u> | <u>003</u> | Apr 26, 2013 |
| | | <u>XOPENEX</u> | | | | |
| <u>AN</u> | + | OAK PHARMS INC | <u>EQ 0.0103% BASE</u> | <u>N020837</u> | <u>003</u> | Jan 30, 2002 |
| <u>AN</u> | + | | <u>EQ 0.021% BASE</u> | <u>N020837</u> | <u>001</u> | Mar 25, 1999 |
| <u>AN</u> | + | | <u>EQ 0.042% BASE</u> | <u>N020837</u> | <u>002</u> | Mar 25, 1999 |
| <u>AN</u> | + | | <u>EQ 0.25% BASE</u> | <u>N020837</u> | <u>004</u> | Jul 18, 2003 |

LEVALBUTEROL TARTRATE

AEROSOL, METERED; INHALATION

XOPENEX HFA

| | | | | | | |
|---|---|----------|---------------------|---------|-----|--------------|
| + | ! | SUNOVION | EQ 0.045MG BASE/INH | N021730 | 001 | Mar 11, 2005 |
|---|---|----------|---------------------|---------|-----|--------------|

LEVETIRACETAM

INJECTABLE; IV (INFUSION)

KEPPRA

| | | | | | | |
|-----------|---|-------------------------|-----------------------------|----------------|------------|--------------|
| <u>AP</u> | + | UCB INC | <u>500MG/5ML (100MG/ML)</u> | <u>N021872</u> | <u>001</u> | Jul 31, 2006 |
| | | <u>LEVETIRACETAM</u> | | | | |
| <u>AP</u> | | AKORN | <u>500MG/5ML (100MG/ML)</u> | <u>A209934</u> | <u>001</u> | May 04, 2018 |
| <u>AP</u> | | AUROBINDO PHARMA LTD | <u>500MG/5ML (100MG/ML)</u> | <u>A204312</u> | <u>001</u> | Feb 01, 2016 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>500MG/5ML (100MG/ML)</u> | <u>A090813</u> | <u>001</u> | May 26, 2010 |
| <u>AP</u> | | | <u>500MG/5ML (100MG/ML)</u> | <u>A090876</u> | <u>001</u> | Aug 13, 2015 |
| <u>AP</u> | | HAINAN POLY PHARM | <u>500MG/5ML (100MG/ML)</u> | <u>A209781</u> | <u>001</u> | Mar 20, 2018 |
| <u>AP</u> | | HIKMA FARMACEUTICA | <u>500MG/5ML (100MG/ML)</u> | <u>A090981</u> | <u>001</u> | Oct 13, 2011 |
| <u>AP</u> | | HOSPIRA INC | <u>500MG/5ML (100MG/ML)</u> | <u>A202869</u> | <u>001</u> | Apr 06, 2012 |
| <u>AP</u> | | JUBILANT GENERICS | <u>500MG/5ML (100MG/ML)</u> | <u>A206838</u> | <u>001</u> | Jun 02, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

LEVETIRACETAM

INJECTABLE; IV (INFUSION)

LEVETIRACETAM

| | | | | |
|-----------|--------------------|-----------------------------|--------------------|--------------|
| <u>AP</u> | LUITPOLD | <u>500MG/5ML (100MG/ML)</u> | <u>A202143 001</u> | Jan 31, 2012 |
| <u>AP</u> | MYLAN LABS LTD | <u>500MG/5ML (100MG/ML)</u> | <u>A203308 001</u> | Sep 16, 2016 |
| <u>AP</u> | SAGENT PHARMS | <u>500MG/5ML (100MG/ML)</u> | <u>A091627 001</u> | Jun 26, 2013 |
| <u>AP</u> | SUN PHARM INDS LTD | <u>500MG/5ML (100MG/ML)</u> | <u>A090754 001</u> | Jun 16, 2010 |
| <u>AP</u> | X GEN PHARMS | <u>500MG/5ML (100MG/ML)</u> | <u>A091485 001</u> | Aug 05, 2011 |

LEVETIRACETAM IN SODIUM CHLORIDE

| | | | | |
|-----------|-------------------------|-------------------------------|--------------------|--------------|
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>500MG/100ML (5MG/ML)</u> | <u>A207160 001</u> | Jan 04, 2017 |
| <u>AP</u> | | <u>1000MG/100ML (10MG/ML)</u> | <u>A207160 002</u> | Jan 04, 2017 |
| <u>AP</u> | | <u>1500MG/100ML (15MG/ML)</u> | <u>A207160 003</u> | Jan 04, 2017 |
| <u>AP</u> | GLAND PHARMA LTD | <u>500MG/100ML (5MG/ML)</u> | <u>A206880 001</u> | Oct 25, 2017 |
| <u>AP</u> | | <u>1000MG/100ML (10MG/ML)</u> | <u>A206880 002</u> | Oct 25, 2017 |
| <u>AP</u> | | <u>1500MG/100ML (15MG/ML)</u> | <u>A206880 003</u> | Oct 25, 2017 |
| <u>AP</u> | +! HQ SPECIALITY PHARMA | <u>500MG/100ML (5MG/ML)</u> | <u>N202543 001</u> | Nov 09, 2011 |
| <u>AP</u> | +! | <u>1000MG/100ML (10MG/ML)</u> | <u>N202543 002</u> | Nov 09, 2011 |
| <u>AP</u> | +! | <u>1500MG/100ML (15MG/ML)</u> | <u>N202543 003</u> | Nov 09, 2011 |

SOLUTION; ORAL

KEPPRA

| | | | | |
|-----------|------------|-----------------|--------------------|--------------|
| <u>AA</u> | +! UCB INC | <u>100MG/ML</u> | <u>N021505 001</u> | Jul 15, 2003 |
|-----------|------------|-----------------|--------------------|--------------|

LEVETIRACETAM

| | | | | |
|-----------|----------------------|-----------------|--------------------|--------------|
| <u>AA</u> | ACTAVIS MID ATLANTIC | <u>100MG/ML</u> | <u>A078976 001</u> | Jan 15, 2009 |
| <u>AA</u> | AMNEAL PHARMS | <u>100MG/ML</u> | <u>A090992 001</u> | Oct 27, 2009 |
| <u>AA</u> | AUROBINDO PHARMA LTD | <u>100MG/ML</u> | <u>A079063 001</u> | Jan 15, 2009 |
| <u>AA</u> | BRECKENRIDGE PHARM | <u>100MG/ML</u> | <u>A079120 001</u> | Jan 16, 2009 |
| <u>AA</u> | HETERO LABS LTD III | <u>100MG/ML</u> | <u>A203052 001</u> | Feb 28, 2013 |
| <u>AA</u> | HI-TECH PHARMACAL | <u>100MG/ML</u> | <u>A090601 001</u> | Feb 28, 2012 |
| <u>AA</u> | LANNETT CO INC | <u>100MG/ML</u> | <u>A090079 001</u> | Apr 11, 2012 |
| <u>AA</u> | | <u>100MG/ML</u> | <u>A090263 001</u> | Apr 03, 2009 |
| <u>AA</u> | LUPIN LTD | <u>100MG/ML</u> | <u>A090893 001</u> | Oct 17, 2011 |
| <u>AA</u> | ORIT LABS LLC | <u>100MG/ML</u> | <u>A203067 001</u> | May 09, 2013 |
| <u>AA</u> | PHARM ASSOC | <u>100MG/ML</u> | <u>A201157 001</u> | Jun 04, 2015 |
| <u>AA</u> | TARO | <u>100MG/ML</u> | <u>A078774 001</u> | Feb 10, 2009 |
| <u>AA</u> | TOLMAR | <u>100MG/ML</u> | <u>A079107 001</u> | Jan 15, 2009 |
| <u>AA</u> | TRIS PHARMA INC | <u>100MG/ML</u> | <u>A090461 001</u> | Sep 30, 2010 |
| <u>AA</u> | WOCKHARDT BIO AG | <u>100MG/ML</u> | <u>A090028 001</u> | Mar 03, 2010 |

TABLET; ORAL

KEPPRA

| | | | | |
|-----------|-----------|--------------|--------------------|--------------|
| <u>AB</u> | + UCB INC | <u>250MG</u> | <u>N021035 001</u> | Nov 30, 1999 |
| <u>AB</u> | + | <u>500MG</u> | <u>N021035 002</u> | Nov 30, 1999 |
| <u>AB</u> | + | <u>750MG</u> | <u>N021035 003</u> | Nov 30, 1999 |
| <u>AB</u> | +! | <u>1GM</u> | <u>N021035 004</u> | Jan 06, 2006 |

LEVETIRACETAM

| | | | | |
|-----------|--------------------|--------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>250MG</u> | <u>A090843 001</u> | Feb 14, 2011 |
| <u>AB</u> | | <u>500MG</u> | <u>A090843 002</u> | Feb 14, 2011 |
| <u>AB</u> | | <u>750MG</u> | <u>A090843 003</u> | Feb 14, 2011 |
| <u>AB</u> | | <u>1GM</u> | <u>A090843 004</u> | Feb 14, 2011 |
| <u>AB</u> | ACI HEALTHCARE LTD | <u>250MG</u> | <u>A078042 001</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>500MG</u> | <u>A078042 002</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>750MG</u> | <u>A078042 003</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>1GM</u> | <u>A078042 004</u> | Jan 15, 2009 |
| <u>AB</u> | ACIC PHARMS | <u>250MG</u> | <u>A090767 001</u> | Jul 28, 2010 |
| <u>AB</u> | | <u>500MG</u> | <u>A090767 002</u> | Jul 28, 2010 |
| <u>AB</u> | | <u>750MG</u> | <u>A090767 003</u> | Jul 28, 2010 |
| <u>AB</u> | | <u>1GM</u> | <u>A090767 004</u> | Jul 28, 2010 |
| <u>AB</u> | APOTEX INC | <u>250MG</u> | <u>A078869 001</u> | Mar 13, 2009 |
| <u>AB</u> | | <u>500MG</u> | <u>A078869 002</u> | Mar 13, 2009 |
| <u>AB</u> | | <u>750MG</u> | <u>A078869 003</u> | Mar 13, 2009 |
| <u>AB</u> | | <u>1GM</u> | <u>A078869 004</u> | Mar 13, 2009 |
| <u>AB</u> | AUROBINDO PHARMA | <u>250MG</u> | <u>A078993 001</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>500MG</u> | <u>A078993 002</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>750MG</u> | <u>A078993 003</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>1GM</u> | <u>A078993 004</u> | Jan 15, 2009 |
| <u>AB</u> | BRECKENRIDGE PHARM | <u>250MG</u> | <u>A090511 001</u> | Aug 18, 2011 |
| <u>AB</u> | | <u>500MG</u> | <u>A090511 002</u> | Aug 18, 2011 |
| <u>AB</u> | | <u>750MG</u> | <u>A090511 003</u> | Aug 18, 2011 |
| <u>AB</u> | | <u>1GM</u> | <u>A090511 004</u> | Aug 18, 2011 |
| <u>AB</u> | CHARTWELL RX | <u>250MG</u> | <u>A201293 001</u> | Jun 14, 2011 |

PRESCRIPTION DRUG PRODUCT LISTLEVETIRACETAM

TABLET; ORAL

LEVETIRACETAM

| | | | | |
|-----------|--------------------------------|--------------|--------------------|--------------|
| <u>AB</u> | | <u>500MG</u> | <u>A201293 002</u> | Jun 14, 2011 |
| <u>AB</u> | | <u>750MG</u> | <u>A201293 003</u> | Jun 14, 2011 |
| <u>AB</u> | | <u>1GM</u> | <u>A201293 004</u> | Jun 14, 2011 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>250MG</u> | <u>A076920 001</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>500MG</u> | <u>A076920 002</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>750MG</u> | <u>A076920 003</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>1GM</u> | <u>A078904 001</u> | Jan 15, 2009 |
| <u>AB</u> | HETERO LABS LTD III | <u>250MG</u> | <u>A090515 001</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>500MG</u> | <u>A090515 002</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>750MG</u> | <u>A090515 003</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>1GM</u> | <u>A090515 004</u> | Oct 08, 2010 |
| <u>AB</u> | INVAGEN PHARMS | <u>250MG</u> | <u>A078234 001</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>500MG</u> | <u>A078234 002</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>750MG</u> | <u>A078234 003</u> | Jan 15, 2009 |
| <u>AB</u> | LUPIN | <u>250MG</u> | <u>A078154 001</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>500MG</u> | <u>A078154 002</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>750MG</u> | <u>A078154 003</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>1GM</u> | <u>A090025 001</u> | Jan 15, 2009 |
| <u>AB</u> | MYLAN | <u>250MG</u> | <u>A076919 001</u> | Nov 04, 2008 |
| <u>AB</u> | | <u>500MG</u> | <u>A076919 002</u> | Nov 04, 2008 |
| <u>AB</u> | | <u>750MG</u> | <u>A076919 003</u> | Nov 04, 2008 |
| <u>AB</u> | | <u>1GM</u> | <u>A090261 001</u> | Dec 08, 2009 |
| <u>AB</u> | ORCHID HLTHCARE | <u>250MG</u> | <u>A078526 001</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>500MG</u> | <u>A078526 002</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>750MG</u> | <u>A078526 003</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>1GM</u> | <u>A090484 001</u> | Aug 05, 2010 |
| <u>AB</u> | OXFORD PHARMS | <u>250MG</u> | <u>A077319 001</u> | Mar 20, 2009 |
| <u>AB</u> | | <u>500MG</u> | <u>A077319 002</u> | Mar 20, 2009 |
| <u>AB</u> | | <u>750MG</u> | <u>A077319 003</u> | Mar 20, 2009 |
| <u>AB</u> | PRINSTON INC | <u>250MG</u> | <u>A078106 001</u> | Feb 10, 2009 |
| <u>AB</u> | | <u>500MG</u> | <u>A078106 002</u> | Feb 10, 2009 |
| <u>AB</u> | | <u>750MG</u> | <u>A078106 003</u> | Feb 10, 2009 |
| <u>AB</u> | | <u>1GM</u> | <u>A078106 004</u> | Feb 10, 2009 |
| <u>AB</u> | SECAN PHARMS | <u>500MG</u> | <u>A205102 004</u> | Dec 16, 2015 |
| <u>AB</u> | | <u>1GM</u> | <u>A205102 003</u> | Dec 16, 2015 |
| <u>AB</u> | TARO | <u>250MG</u> | <u>A078960 004</u> | Feb 01, 2010 |
| <u>AB</u> | | <u>500MG</u> | <u>A078960 003</u> | Feb 01, 2010 |
| <u>AB</u> | | <u>750MG</u> | <u>A078960 002</u> | Feb 01, 2010 |
| <u>AB</u> | | <u>1GM</u> | <u>A078960 001</u> | Feb 01, 2010 |
| <u>AB</u> | TEVA PHARMS | <u>250MG</u> | <u>A078101 001</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>500MG</u> | <u>A078101 002</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>750MG</u> | <u>A078101 003</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>1GM</u> | <u>A078101 004</u> | Jan 15, 2009 |
| <u>AB</u> | TORRENT PHARMS | <u>250MG</u> | <u>A078858 001</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>500MG</u> | <u>A078858 002</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>750MG</u> | <u>A078858 003</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>1GM</u> | <u>A078858 004</u> | Jan 15, 2009 |
| <u>AB</u> | VINTAGE PHARMS | <u>250MG</u> | <u>A091491 001</u> | Dec 14, 2010 |
| <u>AB</u> | | <u>500MG</u> | <u>A091491 002</u> | Dec 14, 2010 |
| <u>AB</u> | | <u>750MG</u> | <u>A091491 003</u> | Dec 14, 2010 |
| <u>AB</u> | | <u>1GM</u> | <u>A091491 004</u> | Dec 14, 2010 |
| <u>AB</u> | WOCKHARDT | <u>250MG</u> | <u>A079042 001</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>500MG</u> | <u>A079042 002</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>750MG</u> | <u>A079042 003</u> | Jan 15, 2009 |
| <u>AB</u> | | <u>1GM</u> | <u>A079042 004</u> | Jan 15, 2009 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>250MG</u> | <u>A078918 001</u> | Apr 29, 2009 |
| <u>AB</u> | | <u>1GM</u> | <u>A078918 002</u> | Apr 29, 2009 |
| | <u>ROWEEPPRA</u> | | | |
| <u>AB</u> | LOTUS PHARM CO LTD | <u>250MG</u> | <u>A090906 002</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>500MG</u> | <u>A090906 001</u> | Nov 05, 2010 |
| <u>AB</u> | | <u>750MG</u> | <u>A090906 003</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>1GM</u> | <u>A090906 004</u> | Oct 31, 2016 |
| | TABLET, EXTENDED RELEASE; ORAL | | | |
| | <u>KEPPRA XR</u> | | | |
| <u>AB</u> | + UCB INC | <u>500MG</u> | <u>N022285 001</u> | Sep 12, 2008 |
| <u>AB</u> | +! | <u>750MG</u> | <u>N022285 002</u> | Feb 12, 2009 |
| | <u>LEVETIRACETAM</u> | | | |
| <u>AB</u> | ACTAVIS ELIZABETH | <u>500MG</u> | <u>A091557 001</u> | Sep 12, 2011 |
| <u>AB</u> | | <u>750MG</u> | <u>A091557 002</u> | Sep 12, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

LEVETIRACETAM

TABLET, EXTENDED RELEASE;ORAL

LEVETIRACETAM

| | | | | | |
|-----------|-----------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ACTAVIS LABS FL INC | <u>500MG</u> | <u>A091093</u> | <u>001</u> | Sep 12, 2011 |
| <u>AB</u> | | <u>750MG</u> | <u>A091093</u> | <u>002</u> | Sep 12, 2011 |
| <u>AB</u> | ANCHEN PHARMS | <u>500MG</u> | <u>A091360</u> | <u>001</u> | Oct 04, 2011 |
| <u>AB</u> | | <u>750MG</u> | <u>A091360</u> | <u>002</u> | Oct 04, 2011 |
| <u>AB</u> | APOTEX INC | <u>500MG</u> | <u>A091261</u> | <u>001</u> | Sep 12, 2011 |
| <u>AB</u> | | <u>750MG</u> | <u>A091261</u> | <u>002</u> | Sep 12, 2011 |
| <u>AB</u> | DEXCEL PHARMA | <u>500MG</u> | <u>A202167</u> | <u>001</u> | Sep 04, 2015 |
| <u>AB</u> | | <u>750MG</u> | <u>A202167</u> | <u>002</u> | Sep 04, 2015 |
| <u>AB</u> | ECI PHARMS LLC | <u>500MG</u> | <u>A204754</u> | <u>001</u> | Aug 26, 2016 |
| <u>AB</u> | | <u>750MG</u> | <u>A204754</u> | <u>002</u> | Aug 26, 2016 |
| <u>AB</u> | INTELLIPHARMACEUTIC S | <u>500MG</u> | <u>A204511</u> | <u>001</u> | Feb 23, 2016 |
| <u>AB</u> | | <u>750MG</u> | <u>A204511</u> | <u>002</u> | Feb 23, 2016 |
| <u>AB</u> | LOTUS PHARM CO LTD | <u>500MG</u> | <u>A202095</u> | <u>002</u> | Jun 06, 2016 |
| <u>AB</u> | | <u>750MG</u> | <u>A202095</u> | <u>001</u> | Jun 06, 2016 |
| <u>AB</u> | LUPIN LTD | <u>500MG</u> | <u>A091399</u> | <u>001</u> | Sep 12, 2011 |
| <u>AB</u> | | <u>750MG</u> | <u>A091399</u> | <u>002</u> | Sep 12, 2011 |
| <u>AB</u> | PHARMADAX INC | <u>500MG</u> | <u>A201464</u> | <u>001</u> | May 25, 2012 |
| <u>AB</u> | | <u>750MG</u> | <u>A201464</u> | <u>002</u> | May 25, 2012 |
| <u>AB</u> | PHARMTAK INC | <u>500MG</u> | <u>A207175</u> | <u>001</u> | Sep 28, 2017 |
| <u>AB</u> | | <u>750MG</u> | <u>A207175</u> | <u>002</u> | Sep 28, 2017 |
| <u>AB</u> | PRINSTON INC | <u>500MG</u> | <u>A202533</u> | <u>001</u> | Jul 20, 2012 |
| <u>AB</u> | | <u>500MG</u> | <u>A203468</u> | <u>001</u> | May 21, 2015 |
| <u>AB</u> | | <u>750MG</u> | <u>A202533</u> | <u>002</u> | Jul 20, 2012 |
| <u>AB</u> | | <u>750MG</u> | <u>A203468</u> | <u>002</u> | May 21, 2015 |
| <u>AB</u> | ROUSES POINT PHARMS | <u>500MG</u> | <u>A202524</u> | <u>001</u> | Aug 27, 2012 |
| <u>AB</u> | | <u>750MG</u> | <u>A202524</u> | <u>002</u> | Aug 27, 2012 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>500MG</u> | <u>A091285</u> | <u>001</u> | Sep 12, 2011 |
| <u>AB</u> | | <u>750MG</u> | <u>A091285</u> | <u>002</u> | Sep 12, 2011 |
| <u>AB</u> | SUN PHARMA GLOBAL | <u>500MG</u> | <u>A203059</u> | <u>001</u> | Sep 09, 2013 |
| <u>AB</u> | | <u>750MG</u> | <u>A203059</u> | <u>002</u> | Sep 09, 2013 |
| <u>AB</u> | TEVA PHARMS | <u>500MG</u> | <u>A091430</u> | <u>001</u> | Sep 12, 2011 |
| <u>AB</u> | | <u>750MG</u> | <u>A091430</u> | <u>002</u> | Sep 12, 2011 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>500MG</u> | <u>A091338</u> | <u>001</u> | May 29, 2012 |
| <u>AB</u> | | <u>750MG</u> | <u>A091338</u> | <u>002</u> | May 29, 2012 |
| | ELEPSIA XR | | | | |
| | + SPARC | 1GM | N204417 | 001 | Dec 20, 2018 |
| | +! | 1.5GM | N204417 | 002 | Dec 20, 2018 |
| | LEVETIRACETAM | | | | |
| | APOTEX INC | 1GM | A202958 | 001 | Feb 25, 2015 |
| | TABLET, FOR SUSPENSION;ORAL | | | | |
| | SPRITAM | | | | |
| | + APRECIA PHARMS | 250MG | N207958 | 001 | Jul 31, 2015 |
| | + | 500MG | N207958 | 002 | Jul 31, 2015 |
| | + | 750MG | N207958 | 003 | Jul 31, 2015 |
| | +! | 1GM | N207958 | 004 | Jul 31, 2015 |

LEVOBUNOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

AKBETA

| | | | | | |
|-----------|-------|--------------|----------------|------------|--------------|
| <u>AT</u> | AKORN | <u>0.25%</u> | <u>A074779</u> | <u>001</u> | Oct 29, 1996 |
| <u>AT</u> | | <u>0.5%</u> | <u>A074780</u> | <u>001</u> | Oct 29, 1996 |

BETAGAN

| | | | | | |
|-----------|-------------|--------------|----------------|------------|--------------|
| <u>AT</u> | +! ALLERGAN | <u>0.25%</u> | <u>N019814</u> | <u>001</u> | Jun 28, 1989 |
| <u>AT</u> | +! | <u>0.5%</u> | <u>N019219</u> | <u>002</u> | Dec 19, 1985 |

LEVOBUNOLOL HYDROCHLORIDE

| | | | | | |
|-----------|-----------------|-------------|----------------|------------|--------------|
| <u>AT</u> | BAUSCH AND LOMB | <u>0.5%</u> | <u>A074326</u> | <u>001</u> | Mar 04, 1994 |
| <u>AT</u> | SANDOZ INC | <u>0.5%</u> | <u>A074850</u> | <u>001</u> | Oct 28, 1996 |

LEVOCARNITINE

INJECTABLE;INJECTION

CARNITOR

| | | | | | |
|-----------|------------------------|-----------------|----------------|------------|--------------|
| <u>AP</u> | +! LEADIANT BIOSCI INC | <u>200MG/ML</u> | <u>N020182</u> | <u>001</u> | Dec 16, 1992 |
|-----------|------------------------|-----------------|----------------|------------|--------------|

LEVOCARNITINE

| | | | | | |
|-----------|-------------------------|-----------------|----------------|------------|--------------|
| <u>AP</u> | LUITPOLD | <u>200MG/ML</u> | <u>A075861</u> | <u>001</u> | Jun 22, 2001 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>200MG/ML</u> | <u>A075567</u> | <u>001</u> | Mar 29, 2001 |

PRESCRIPTION DRUG PRODUCT LIST

LEVOCARNITINE

SOLUTION;ORAL

CARNITOR

AA **+** LEADIANT BIOSCI INC **1GM/10ML** **N019257 001** Apr 10, 1986

CARNITOR_SF

AA **+** LEADIANT BIOSCI INC **1GM/10ML** **N019257 002** Mar 28, 2007

LEVOCARNITINE

AA HI TECH PHARMA **1GM/10ML** **A077399 001** Oct 25, 2007

AA LYNE **1GM/10ML** **A076851 001** Aug 10, 2004

TABLET;ORAL

CARNITOR

AB **+** LEADIANT BIOSCI INC **330MG** **N018948 001** Dec 27, 1985

LEVOCARNITINE

AB ANDA REPOSITORY **330MG** **A076858 001** Sep 20, 2004

LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION;ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

AA APOTEX INC **2.5MG/5ML** **A202915 001** Aug 21, 2014

AA **!** L PERRIGO CO **2.5MG/5ML** **A091263 001** Nov 07, 2011

AA LANNETT CO INC **2.5MG/5ML** **A204599 001** May 15, 2017

AA TARO PHARM INDS LTD **2.5MG/5ML** **A202673 001** Jul 26, 2013

XYZAL

AA **+** SANOFI AVENTIS US **2.5MG/5ML** **N022157 001** Jan 28, 2008

TABLET;ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

AB APOTEX INC **5MG** **A203027 001** Feb 13, 2015

AB DR REDDYS LABS LTD **5MG** **A090392 001** Feb 24, 2011

AB GLENMARK GENERICS **5MG** **A090385 001** Feb 24, 2011

AB HETERO LABS LTD III **5MG** **A091264 001** Jun 29, 2012

AB MACLEODS PHARMS LTD **5MG** **A205564 001** Jan 11, 2016

AB MICRO LABS LTD **5MG** **A202046 001** Sep 17, 2013

INDIA

AB NEOPHARMA **5MG** **A204323 001** Dec 20, 2016

AB SCIEGEN PHARMS INC **5MG** **A203646 001** Sep 09, 2014

AB SUN PHARM INDS LTD **5MG** **A201653 001** Jun 26, 2015

AB SUN PHARMA GLOBAL **5MG** **A090362 001** Jan 31, 2013

AB SYNTHON PHARMS **5MG** **A090229 001** Nov 26, 2010

AB TEVA PHARMS **5MG** **A090199 001** Aug 22, 2011

XYZAL

AB **+** SANOFI AVENTIS US **5MG** **N022064 001** May 25, 2007

LEVODOPA

POWDER; INHALATION

INBRIJA

+ ACORDA **42MG** **N209184 001** Dec 21, 2018

LEVOFLOXACIN

INJECTABLE; INJECTION

LEVOFLOXACIN

AP **!** AUROBINDO PHARMA LTD **EQ 500MG/20ML (EQ 25MG/ML)** **A202328 001** Jan 24, 2013

AP **!** **EQ 750MG/30ML (EQ 25MG/ML)** **A202328 002** Jan 24, 2013

AP BAXTER HLTHCARE CORP **EQ 500MG/20ML (EQ 25MG/ML)** **A091436 001** Jun 05, 2013

LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

AP AUROBINDO PHARMA LTD **EQ 250MG/50ML (EQ 5MG/ML)** **A206919 001** Feb 10, 2016

AP **EQ 500MG/100ML (EQ 5MG/ML)** **A206919 002** Feb 10, 2016

AP **EQ 750MG/150ML (EQ 5MG/ML)** **A206919 003** Feb 10, 2016

AP BAXTER HLTHCARE CORP **EQ 250MG/50ML (EQ 5MG/ML)** **A091397 001** Aug 08, 2013

AP **EQ 500MG/100ML (EQ 5MG/ML)** **A091397 002** Aug 08, 2013

AP **EQ 750MG/150ML (EQ 5MG/ML)** **A091397 003** Aug 08, 2013

AP FRESENIUS KABI USA **EQ 250MG/50ML (EQ 5MG/ML)** **A200674 001** Jun 19, 2013

AP **EQ 500MG/100ML (EQ 5MG/ML)** **A200674 002** Jun 19, 2013

AP **EQ 750MG/150ML (EQ 5MG/ML)** **A200674 003** Jun 19, 2013

AP HIKMA FARMACEUTICA **EQ 250MG/50ML (EQ 5MG/ML)** **A091375 001** Sep 16, 2011

AP **EQ 500MG/100ML (EQ 5MG/ML)** **A091375 002** Sep 16, 2011

AP **EQ 750MG/150ML (EQ 5MG/ML)** **A091375 003** Sep 16, 2011

AP HOSPIRA INC **EQ 250MG/50ML (EQ 5MG/ML)** **A078579 001** Sep 03, 2015

AP **EQ 500MG/100ML (EQ 5MG/ML)** **A078579 002** Sep 03, 2015

AP **EQ 750MG/150ML (EQ 5MG/ML)** **A078579 003** Sep 03, 2015

AP **!** INFORLIFE **EQ 250MG/50ML (EQ 5MG/ML)** **A090343 001** Jul 07, 2011

AP **!** **EQ 500MG/100ML (EQ 5MG/ML)** **A090343 002** Jul 07, 2011

PRESCRIPTION DRUG PRODUCT LIST

LEVOFLOXACIN

INJECTABLE; INJECTION

LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

AP ! EQ 750MG/150ML (EQ 5MG/ML) A090343 003 Jul 07, 2011
SOLUTION; ORAL

LEVOFLOXACIN

AA ! HI TECH PHARMA 250MG/10ML A091678 001 Jun 20, 2011
AA LANNETT CO INC 250MG/10ML A205222 001 May 25, 2018

SOLUTION/DROPS; OPHTHALMIC

LEVOFLOXACIN

AT AKORN 0.5% A090268 001 Dec 20, 2010
AT MYLAN LABS LTD 0.5% A204899 001 Dec 08, 2017
AT ! RISING PHARMS 0.5% A077700 001 Dec 20, 2010
AT WATSON LABS TEVA 0.5% A076826 001 Feb 10, 2011

TABLET; ORAL

LEVOFLOXACIN

AB APOTEX INC 250MG A090787 001 Sep 29, 2011
AB 500MG A090787 002 Sep 29, 2011
AB 750MG A090787 003 Sep 29, 2011
AB AUROBINDO PHARMA LTD 250MG A201043 001 Jun 20, 2011
AB 500MG A201043 002 Jun 20, 2011
AB ! 750MG A201043 003 Jun 20, 2011
AB CIPLA LTD 250MG A076890 001 Mar 30, 2012
AB 500MG A076890 002 Mar 30, 2012
AB 750MG A076890 003 Mar 30, 2012
AB DR REDDYS LABS INC 250MG A076710 001 Jun 20, 2011
AB 500MG A076710 002 Jun 20, 2011
AB 750MG A076710 003 Jun 20, 2011
AB GLENMARK GENERICS 250MG A200250 001 Jun 20, 2011
AB 500MG A200250 002 Jun 20, 2011
AB 750MG A200250 003 Jun 20, 2011
AB HETERO LABS LTD V 250MG A202801 001 Jan 08, 2015
AB 500MG A202801 002 Jan 08, 2015
AB 750MG A202801 003 Jan 08, 2015
AB JUBILANT GENERICS 250MG A203613 001 Jun 19, 2015
AB 500MG A203613 002 Jun 19, 2015
AB LUPIN 250MG A078424 001 Jun 20, 2011
AB 500MG A078424 002 Jun 20, 2011
AB 750MG A078424 003 Jun 20, 2011
AB MACLEODS PHARMS LTD 250MG A200839 001 Mar 22, 2012
AB 500MG A200839 002 Mar 22, 2012
AB 750MG A200839 003 Mar 22, 2012
AB ORCHID HLTHCARE 250MG A202200 001 Jan 30, 2012
AB 500MG A202200 002 Jan 30, 2012
AB 750MG A202200 003 Jan 30, 2012
AB SANDOZ 250MG A077438 001 Jun 20, 2011
AB 500MG A077438 002 Jun 20, 2011
AB 750MG A077438 003 Jun 20, 2011
AB TEVA 250MG A076361 001 Jun 20, 2011
AB 500MG A076361 002 Jun 20, 2011
AB 750MG A076361 003 Jun 20, 2011
AB TORRENT PHARMS 250MG A090722 001 Jun 20, 2011
AB 500MG A090722 002 Jun 20, 2011
AB 750MG A090722 003 Jun 20, 2011
AB WOCKHARDT 250MG A090367 001 Jun 20, 2011
AB 500MG A090367 002 Jun 20, 2011
AB 750MG A090367 003 Jun 20, 2011
AB ZYDUS PHARMS USA INC 250MG A077652 001 Sep 07, 2012
AB 500MG A077652 002 Sep 07, 2012
AB 750MG A077652 003 Sep 07, 2012

LEVOLEUCOVORIN

POWDER; INTRAVENOUS

KHAPZORY

+! SPECTRUM PHARMS 175MG/VIAL N211226 001 Oct 19, 2018
+! 300MG/VIAL N211226 002 Oct 19, 2018

PRESCRIPTION DRUG PRODUCT LIST

LEVOLEUCOVORIN CALCIUM

POWDER; INTRAVENOUS

FUSILEV

| | | | | | |
|-----------|-----------|-----------------|---------------------------------|---------------------------|--------------|
| AP | +! | SPECTRUM PHARMS | <u>EQ 50MG BASE/VIAL</u> | <u>N020140 001</u> | Mar 07, 2008 |
|-----------|-----------|-----------------|---------------------------------|---------------------------|--------------|

LEVOLEUCOVORIN CALCIUM

| | | | | | |
|-----------|--|-------------|---------------------------------|---------------------------|--------------|
| AP | | ACTAVIS LLC | <u>EQ 50MG BASE/VIAL</u> | <u>A206516 001</u> | Feb 13, 2017 |
|-----------|--|-------------|---------------------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|------------------|---------------------------------|---------------------------|--------------|
| AP | | AMNEAL PHARMS CO | <u>EQ 50MG BASE/VIAL</u> | <u>A207547 001</u> | Feb 13, 2017 |
|-----------|--|------------------|---------------------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|-------------------------|---------------------------------|---------------------------|--------------|
| AP | | WEST-WARD PHARMS INT | <u>EQ 50MG BASE/VIAL</u> | <u>A206263 001</u> | Jun 16, 2016 |
|-----------|--|-------------------------|---------------------------------|---------------------------|--------------|

+! ACTAVIS LLC

SOLUTION; INTRAVENOUS

| | | |
|---------|-----|--------------|
| N208723 | 001 | Sep 29, 2016 |
|---------|-----|--------------|

LEVOLEUCOVORIN CALCIUM

| | | | | | |
|-----------|--|------------------|------------------------------------------------------|---------------------------|--------------|
| AP | | AMNEAL PHARMS CO | <u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u> | <u>A207548 001</u> | Sep 08, 2017 |
|-----------|--|------------------|------------------------------------------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|------------------|------------------------------------------------------|---------------------------|--------------|
| AP | | GLAND PHARMA LTD | <u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u> | <u>A210892 001</u> | Sep 14, 2018 |
|-----------|--|------------------|------------------------------------------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|--|----------------------------------------------------|---------------------------|--------------|
| AP | | | <u>EQ 250MG BASE/25ML (EQ 10MG BASE/ML)</u> | <u>A210892 002</u> | Sep 14, 2018 |
|-----------|--|--|----------------------------------------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|--------------------|------------------------------------------------------|---------------------------|--------------|
| AP | | INGENUS PHARMS LLC | <u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u> | <u>A210623 001</u> | May 03, 2018 |
|-----------|--|--------------------|------------------------------------------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|--|----------------------------------------------------|---------------------------|--------------|
| AP | | | <u>EQ 250MG BASE/25ML (EQ 10MG BASE/ML)</u> | <u>A210623 002</u> | May 03, 2018 |
|-----------|--|--|----------------------------------------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|----------------|------------------------------------------------------|---------------------------|--------------|
| AP | | MYLAN TEORANTA | <u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u> | <u>A203576 001</u> | Oct 20, 2015 |
|-----------|--|----------------|------------------------------------------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|--|----------------------------------------------------|---------------------------|--------------|
| AP | | | <u>EQ 250MG BASE/25ML (EQ 10MG BASE/ML)</u> | <u>A203576 002</u> | Oct 20, 2015 |
|-----------|--|--|----------------------------------------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|------------|------------------------------------------------------|---------------------------|--------------|
| AP | | SANDOZ INC | <u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u> | <u>A203563 001</u> | Mar 09, 2015 |
|-----------|--|------------|------------------------------------------------------|---------------------------|--------------|

| | | | | | |
|-----------|----------|--|----------------------------------------------------|---------------------------|--------------|
| AP | ! | | <u>EQ 250MG BASE/25ML (EQ 10MG BASE/ML)</u> | <u>A203563 002</u> | Mar 09, 2015 |
|-----------|----------|--|----------------------------------------------------|---------------------------|--------------|

LEVOMILNACIPRAN HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

FETZIMA

| | | | | | | |
|---|--|--------------------|--------------|---------|-----|--------------|
| + | | ALLERGAN SALES LLC | EQ 20MG BASE | N204168 | 001 | Jul 25, 2013 |
|---|--|--------------------|--------------|---------|-----|--------------|

| | | | | | | |
|---|--|--|--------------|---------|-----|--------------|
| + | | | EQ 40MG BASE | N204168 | 002 | Jul 25, 2013 |
|---|--|--|--------------|---------|-----|--------------|

| | | | | | | |
|---|--|--|--------------|---------|-----|--------------|
| + | | | EQ 80MG BASE | N204168 | 003 | Jul 25, 2013 |
|---|--|--|--------------|---------|-----|--------------|

| | | | | | | |
|----|--|--|---------------|---------|-----|--------------|
| +! | | | EQ 120MG BASE | N204168 | 004 | Jul 25, 2013 |
|----|--|--|---------------|---------|-----|--------------|

LEVONORDEFIN; MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

SCANDONEST L

| | | | | | | |
|---|--|---------|---------------|---------|-----|--------------|
| ! | | DEPROCO | 0.05MG/ML; 2% | A088388 | 001 | Oct 10, 1984 |
|---|--|---------|---------------|---------|-----|--------------|

LEVONORGESTREL

INTRAUTERINE DEVICE; INTRAUTERINE

KYLEENA

| | | | | | | |
|---|---|----------------|--------|---------|-----|--------------|
| + | ! | BAYER HLTHCARE | 19.5MG | N208224 | 001 | Sep 16, 2016 |
|---|---|----------------|--------|---------|-----|--------------|

LILETTA

| | | | | | | |
|--|--|--------------|------|---------|-----|--------------|
| | | MEDICINES360 | 52MG | N206229 | 001 | Feb 26, 2015 |
|--|--|--------------|------|---------|-----|--------------|

MIRENA

| | | | | | | |
|---|---|----------------|------|---------|-----|--------------|
| + | ! | BAYER HLTHCARE | 52MG | N021225 | 001 | Dec 06, 2000 |
|---|---|----------------|------|---------|-----|--------------|

SKYLA

| | | | | | | |
|---|---|----------------|--------|---------|-----|--------------|
| + | ! | BAYER HLTHCARE | 13.5MG | N203159 | 001 | Jan 09, 2013 |
|---|---|----------------|--------|---------|-----|--------------|

TABLET; ORAL

LEVONORGESTREL

| | | | | | |
|-----------|--|--------------------|----------------------|---------------------------|--------------|
| AB | | LOTUS PHARM CO LTD | <u>0.75MG</u> | <u>A202684 001</u> | Sep 02, 2016 |
|-----------|--|--------------------|----------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|----------------|----------------------|---------------------------|--------------|
| AB | | MYLAN LABS LTD | <u>0.75MG</u> | <u>A202740 001</u> | Sep 02, 2016 |
|-----------|--|----------------|----------------------|---------------------------|--------------|

| | | | | | |
|-----------|----------|-----------------|----------------------|---------------------------|--------------|
| AB | ! | PERRIGO R AND D | <u>0.75MG</u> | <u>A090740 001</u> | Dec 30, 2010 |
|-----------|----------|-----------------|----------------------|---------------------------|--------------|

LEVORPHANOL TARTRATE

TABLET; ORAL

LEVORPHANOL TARTRATE

| | | | | | |
|-----------|----------|---------------------|-------------------|---------------------------|--------------|
| AB | ! | SENTYNL THERAPS INC | <u>2MG</u> | <u>A074278 001</u> | Mar 31, 2000 |
|-----------|----------|---------------------|-------------------|---------------------------|--------------|

| | | | | | |
|-----------|--|---------------|-------------------|---------------------------|--------------|
| AB | | VIRTUS PHARMS | <u>2MG</u> | <u>A211484 001</u> | Dec 13, 2018 |
|-----------|--|---------------|-------------------|---------------------------|--------------|

| | | | | | | |
|--|--|---------------------|-----|---------|-----|--------------|
| | | SENTYNL THERAPS INC | 1MG | A074278 | 002 | Jun 18, 2018 |
|--|--|---------------------|-----|---------|-----|--------------|

| | | | | | | |
|--|--|--|-----|---------|-----|--------------|
| | | | 3MG | A074278 | 003 | Jun 18, 2018 |
|--|--|--|-----|---------|-----|--------------|

LEVOTHYROXINE SODIUM

CAPSULE; ORAL

TIROSINT

| | | | | | | |
|---|--|-------------------------|---------|---------|-----|--------------|
| + | | INSTITUT BIOCHIMIQUE | 0.013MG | N021924 | 013 | Aug 01, 2007 |
|---|--|-------------------------|---------|---------|-----|--------------|

| | | | | | | |
|---|--|--|---------|---------|-----|--------------|
| + | | | 0.025MG | N021924 | 002 | Oct 13, 2006 |
|---|--|--|---------|---------|-----|--------------|

| | | | | | | |
|---|--|--|--------|---------|-----|--------------|
| + | | | 0.05MG | N021924 | 003 | Oct 13, 2006 |
|---|--|--|--------|---------|-----|--------------|

| | | | | | | |
|---|--|--|---------|---------|-----|--------------|
| + | | | 0.075MG | N021924 | 004 | Oct 13, 2006 |
|---|--|--|---------|---------|-----|--------------|

| | | | | | | |
|---|--|--|---------|---------|-----|--------------|
| + | | | 0.088MG | N021924 | 010 | Oct 02, 2009 |
|---|--|--|---------|---------|-----|--------------|

| | | | | | | |
|---|--|--|-------|---------|-----|--------------|
| + | | | 0.1MG | N021924 | 005 | Oct 13, 2006 |
|---|--|--|-------|---------|-----|--------------|

| | | | | | | |
|---|--|--|---------|---------|-----|--------------|
| + | | | 0.112MG | N021924 | 008 | Oct 02, 2009 |
|---|--|--|---------|---------|-----|--------------|

| | | | | | | |
|---|--|--|---------|---------|-----|--------------|
| + | | | 0.125MG | N021924 | 006 | Oct 13, 2006 |
|---|--|--|---------|---------|-----|--------------|

| | | | | | | |
|---|--|--|---------|---------|-----|--------------|
| + | | | 0.137MG | N021924 | 009 | Oct 02, 2009 |
|---|--|--|---------|---------|-----|--------------|

| | | | | | | |
|---|--|--|--------|---------|-----|--------------|
| + | | | 0.15MG | N021924 | 007 | Oct 13, 2006 |
|---|--|--|--------|---------|-----|--------------|

| | | | | | | |
|---|--|--|---------|---------|-----|--------------|
| + | | | 0.175MG | N021924 | 011 | Apr 25, 2017 |
|---|--|--|---------|---------|-----|--------------|

| | | | | | | |
|---|---|--|---------|---------|-----|--------------|
| + | ! | | 0.200MG | N021924 | 012 | Apr 25, 2017 |
|---|---|--|---------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM

POWDER; INTRAVENOUS

LEVOTHYROXINE SODIUM

| | | | | | |
|-----------|-----------|----------------------|--------------------|--------------------|--------------|
| <u>AP</u> | <u>+!</u> | FRESENIUS KABI USA | <u>100MCG/VIAL</u> | <u>N202231 001</u> | Jun 24, 2011 |
| <u>AP</u> | <u>+!</u> | | <u>200MCG/VIAL</u> | <u>N202231 002</u> | Jun 24, 2011 |
| <u>AP</u> | <u>+!</u> | | <u>500MCG/VIAL</u> | <u>N202231 003</u> | Jun 24, 2011 |
| <u>AP</u> | | MAIA PHARMS INC | <u>100MCG/VIAL</u> | <u>A208749 001</u> | Dec 21, 2018 |
| <u>AP</u> | | | <u>200MCG/VIAL</u> | <u>A208749 002</u> | Dec 21, 2018 |
| <u>AP</u> | | | <u>500MCG/VIAL</u> | <u>A208749 003</u> | Dec 21, 2018 |
| <u>AP</u> | | PAR STERILE PRODUCTS | <u>200MCG/VIAL</u> | <u>A205366 001</u> | Dec 07, 2015 |
| <u>AP</u> | | PIRAMAL CRITICAL | <u>100MCG/VIAL</u> | <u>A206163 001</u> | Jun 29, 2016 |
| <u>AP</u> | | | <u>500MCG/VIAL</u> | <u>A206163 002</u> | Jun 29, 2016 |

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET; ORAL

SYNTHROID

| | | | | | | |
|---------------|-----------|--------|------------------------|----------------|-------------|--------------|
| <u>--></u> | <u>+</u> | ABBVIE | <u>--> AB1, AB2</u> | <u>0.025MG</u> | N021402 001 | Jul 24, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2</u> | <u>0.05MG</u> | N021402 002 | Jul 24, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2</u> | <u>0.075MG</u> | N021402 003 | Jul 24, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2</u> | <u>0.088MG</u> | N021402 004 | Jul 24, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2</u> | <u>0.1MG</u> | N021402 005 | Jul 24, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2</u> | <u>0.112MG</u> | N021402 006 | Jul 24, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2</u> | <u>0.125MG</u> | N021402 007 | Jul 24, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2</u> | <u>0.137MG</u> | N021402 008 | Jul 24, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2</u> | <u>0.15MG</u> | N021402 009 | Jul 24, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2</u> | <u>0.175MG</u> | N021402 010 | Jul 24, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2</u> | <u>0.2MG</u> | N021402 012 | Jul 24, 2002 |
| <u>--></u> | <u>+!</u> | | <u>--> AB1, AB2</u> | <u>0.3MG</u> | N021402 011 | Jul 24, 2002 |

LEVO-T

| | | | | | | |
|---------------|-----------|--------------|-----------------------------|----------------|-------------|--------------|
| <u>--></u> | <u>+</u> | CEDIPROF INC | <u>--> AB1, AB2, AB3</u> | <u>0.025MG</u> | N021342 001 | Mar 01, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.05MG</u> | N021342 002 | Mar 01, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.075MG</u> | N021342 003 | Mar 01, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.088MG</u> | N021342 004 | Mar 01, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.1MG</u> | N021342 005 | Mar 01, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.112MG</u> | N021342 006 | Mar 01, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.125MG</u> | N021342 007 | Mar 01, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.137MG</u> | N021342 012 | Dec 08, 2003 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.15MG</u> | N021342 008 | Mar 01, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.175MG</u> | N021342 009 | Mar 01, 2002 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.2MG</u> | N021342 010 | Mar 01, 2002 |
| <u>--></u> | <u>+!</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.3MG</u> | N021342 011 | Mar 01, 2002 |

UNITHROID

| | | | | | | |
|---------------|----------|-----------|-----------------------------|----------------|-------------|--------------|
| <u>--></u> | <u>+</u> | STEVENS J | <u>--> AB1, AB2, AB3</u> | <u>0.025MG</u> | N021210 001 | Aug 21, 2000 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.05MG</u> | N021210 002 | Aug 21, 2000 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.075MG</u> | N021210 003 | Aug 21, 2000 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.088MG</u> | N021210 004 | Aug 21, 2000 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.1MG</u> | N021210 005 | Aug 21, 2000 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.112MG</u> | N021210 006 | Aug 21, 2000 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.125MG</u> | N021210 007 | Aug 21, 2000 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.137MG</u> | N021210 012 | Feb 08, 2008 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.15MG</u> | N021210 008 | Aug 21, 2000 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.175MG</u> | N021210 009 | Aug 21, 2000 |
| <u>--></u> | <u>+</u> | | <u>--> AB1, AB2, AB3</u> | <u>0.2MG</u> | N021210 010 | Aug 21, 2000 |

PRESCRIPTION DRUG PRODUCT LIST

LEVOTHYROXINE SODIUM **

**See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET;ORAL

UNITHROID

--> +! --> AB1,AB2,AB3 0.3MG N021210 011 Aug 21, 2000

LEVOTHYROXINE SODIUM

--> MYLAN --> 0.025MG A076187 001 Jun 05, 2002
 --> AB1,AB2,AB3,AB4
 --> 0.05MG A076187 002 Jun 05, 2002
 --> AB1,AB2,AB3,AB4
 --> 0.075MG A076187 003 Jun 05, 2002
 --> AB1,AB2,AB3,AB4
 --> 0.088MG A076187 004 Jun 05, 2002
 --> AB1,AB2,AB3,AB4
 --> 0.1MG A076187 005 Jun 05, 2002
 --> AB1,AB2,AB3,AB4
 --> 0.112MG A076187 006 Jun 05, 2002
 --> AB1,AB2,AB3,AB4
 --> 0.125MG A076187 007 Jun 05, 2002
 --> AB1,AB2,AB3,AB4
 --> 0.137MG A076187 012 Dec 13, 2006
 --> AB1,AB2,AB3,AB4
 --> 0.15MG A076187 008 Jun 05, 2002
 --> AB1,AB2,AB3,AB4
 --> 0.175MG A076187 009 Jun 05, 2002
 --> AB1,AB2,AB3,AB4
 --> 0.2MG A076187 010 Jun 05, 2002
 --> ! --> 0.3MG A076187 011 Jun 05, 2002
 --> AB1,AB2,AB3,AB4

LEVOXYL

--> + KING PHARMS --> AB1,AB3 0.025MG N021301 001 May 25, 2001
 --> + --> AB1,AB3 0.05MG N021301 002 May 25, 2001
 --> + --> AB1,AB3 0.075MG N021301 003 May 25, 2001
 --> + --> AB1,AB3 0.088MG N021301 004 May 25, 2001
 --> + --> AB1,AB3 0.1MG N021301 005 May 25, 2001
 --> + --> AB1,AB3 0.112MG N021301 006 May 25, 2001
 --> + --> AB1,AB3 0.125MG N021301 007 May 25, 2001
 --> + --> AB1,AB3 0.137MG N021301 008 May 25, 2001
 --> + --> AB1,AB3 0.15MG N021301 009 May 25, 2001
 --> + --> AB1,AB3 0.175MG N021301 010 May 25, 2001
 --> +! --> AB1,AB3 0.2MG N021301 011 May 25, 2001

EUTHYROX

AB2 PROVELL 0.025MG N021292 001 May 31, 2002
AB2 0.05MG N021292 002 May 31, 2002
AB2 0.075MG N021292 003 May 31, 2002
AB2 0.088MG N021292 004 May 31, 2002
AB2 0.1MG N021292 005 May 31, 2002
AB2 0.112MG N021292 006 May 31, 2002
AB2 0.125MG N021292 007 May 31, 2002
AB2 0.137MG N021292 008 May 31, 2002
AB2 0.15MG N021292 009 May 31, 2002
AB2 0.175MG N021292 010 May 31, 2002
AB2 0.2MG N021292 011 May 31, 2002

LIDOCAINE

OINTMENT;TOPICAL

LIDOCAINE

AT ALEOR 5% A211469 001 Nov 23, 2018
 DERMACEUTICALS
AT ALKEM LABS LTD 5% A207810 001 Mar 10, 2017
AT AMNEAL PHARMS 5% A206297 001 Aug 07, 2015
AT ! FOUGERA PHARMS INC 5% A080198 001
AT G AND W LABS INC 5% A211019 001 Dec 12, 2018
AT GLENMARK PHARMS LTD 5% A206498 001 Sep 09, 2016
AT RICONPHARMA LLC 5% A208604 001 Sep 20, 2017
AT SEPTODONT INC 5% A040911 001 May 23, 2011
AT STRIDES PHARMA 5% A210958 001 Dec 11, 2018
AT TARO 5% A086724 001
AT TELIGENT PHARMA INC 5% A205318 001 Feb 01, 2016
AT TEVA PHARMS USA 5% A210256 001 Jan 16, 2018
AT VITRUVIAS THERAP 5% A208822 001 Sep 25, 2017

PRESCRIPTION DRUG PRODUCT LIST

LIDOCAINE

PATCH; TOPICAL

LIDOCAINE

| | | | | | |
|-----------|---------------------|-----------|----------------|------------|--------------|
| AB | ACTAVIS LABS UT INC | 5% | A200675 | 001 | Aug 23, 2012 |
| AB | MYLAN TECHNOLOGIES | 5% | A202346 | 001 | Aug 07, 2015 |

LIDODERM

| | | | | | |
|-----------|-----------------------|-----------|----------------|------------|--------------|
| AB | +! TEIKOKU PHARMA USA | 5% | N020612 | 001 | Mar 19, 1999 |
| | ZTLIDO | | | | |
| | +! SCILEX PHARMS INC | 1.8% | N207962 | 001 | Feb 28, 2018 |

LIDOCAINE HYDROCHLORIDE

GEL; OPHTHALMIC

AKTEN

| | | | | | |
|--|----------|------|---------|-----|--------------|
| | +! AKORN | 3.5% | N022221 | 001 | Oct 07, 2008 |
|--|----------|------|---------|-----|--------------|

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE

| | | | | | |
|-----------|----------------------|-------------|----------------|------------|--------------|
| AP | AUROBINDO PHARMA LTD | 1% | A207182 | 001 | Oct 30, 2017 |
| AP | | 2% | A207182 | 002 | Oct 30, 2017 |
| AP | B BRAUN MEDICAL INC | 1% | A208474 | 001 | Aug 03, 2018 |
| AP | HOSPIRA | 0.5% | A088328 | 001 | May 17, 1984 |
| AP | | 1% | A083158 | 001 | |
| AP | | 1% | A088329 | 001 | May 17, 1984 |
| AP | | 2% | A040078 | 001 | Jun 23, 1995 |
| AP | | 2% | A083158 | 002 | |
| AP | | 2% | A088294 | 001 | May 17, 1984 |
| AP | | 20% | A083158 | 003 | |
| AP | INTL MEDICATION | 1% | A083173 | 001 | |
| AP | | 2% | A083173 | 002 | |
| AP | LUITPOLD | 1% | A080850 | 001 | |
| AP | | 1% | A091564 | 001 | Aug 14, 2015 |
| AP | SPECTRA MDCL DEVICES | 1% | A208017 | 001 | Apr 18, 2018 |

LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | |
|-----------|-----------------|--------------------|----------------|------------|--------------|
| AP | B BRAUN | 200MG/100ML | N019830 | 002 | Apr 08, 1992 |
| AP | BAXTER HLTHCARE | 200MG/100ML | N018461 | 002 | |

LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | |
|-----------|-----------------|--------------------|----------------|------------|--------------|
| AP | B BRAUN | 400MG/100ML | N019830 | 003 | Apr 08, 1992 |
| AP | BAXTER HLTHCARE | 400MG/100ML | N018461 | 003 | |

LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | |
|-----------|-----------------|--------------------|----------------|------------|--------------|
| AP | B BRAUN | 800MG/100ML | N019830 | 004 | Apr 08, 1992 |
| AP | BAXTER HLTHCARE | 800MG/100ML | N018461 | 004 | Feb 22, 1982 |

LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER

| | | | | | |
|-----------|--------------------|-------------|----------------|------------|--------------|
| AP | FRESENIUS KABI USA | 1% | A088586 | 001 | Jul 24, 1985 |
| AP | HOSPIRA | 0.5% | A088325 | 001 | Jul 31, 1984 |
| AP | | 1% | A088299 | 001 | Jul 31, 1984 |
| AP | | 2% | A088327 | 001 | Jul 31, 1984 |

LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE

| | | | | | |
|-----------|----------------------|-------------|----------------|------------|--------------|
| AP | AUROBINDO PHARMA LTD | 1% | A203040 | 001 | Mar 14, 2013 |
| AP | | 1% | A203082 | 001 | Mar 14, 2013 |
| AP | | 2% | A203040 | 002 | Mar 14, 2013 |
| AP | | 2% | A203082 | 002 | Mar 14, 2013 |
| AP | FRESENIUS KABI USA | 1% | A080404 | 002 | |
| AP | | 2% | A080404 | 003 | |
| AP | | 2% | N017584 | 001 | |
| AP | | 4% | N017584 | 002 | |
| AP | HOSPIRA | 1% | A080408 | 001 | |
| AP | | 1.5% | A080408 | 002 | |

LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER

| | | | | | |
|-----------|---------|-----------|----------------|------------|--------------|
| AP | HOSPIRA | 1% | A040302 | 001 | Sep 28, 1998 |
| AP | | 2% | A040302 | 002 | Sep 28, 1998 |

XYLOCAINE

| | | | | | |
|-----------|-----------------------|-------------|----------------|------------|--|
| AP | +! FRESENIUS KABI USA | 0.5% | N006488 | 008 | |
| AP | +! | 1% | N006488 | 007 | |
| AP | +! | 1.5% | N006488 | 010 | |
| AP | +! | 2% | N006488 | 002 | |

XYLOCAINE PRESERVATIVE FREE

| | | | | | |
|-----------|-----------------------|------------|----------------|------------|--------------|
| AP | +! FRESENIUS KABI USA | 1% | N016801 | 005 | Jan 19, 1988 |
| AP | +! | 2% | N016801 | 001 | |
| AP | +! | 4% | N016801 | 002 | |
| AP | +! | 20% | N016801 | 004 | |

PRESCRIPTION DRUG PRODUCT LIST

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE

! HOSPIRA 4% A088295 001 May 17, 1984

INJECTABLE; SPINAL

LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%

! HOSPIRA 5% A083914 001

JELLY; TOPICAL

GLYDO**AT** SAGENT PHARMS **2%** **A201094 001** Apr 28, 2014LIDOCAINE HYDROCHLORIDE**AT** AKORN **2%** **A040433 001** Feb 12, 2003**AT** INTL MEDICATION **2%** **A086283 001**XYLOCAINE**AT** +! OAK PHARMS **2%** **N008816 001**

SOLUTION; ORAL

LIDOCAINE HYDROCHLORIDE**AT** HI TECH PHARMA **2%** **A040014 001** Jul 10, 1995**AT** WOCKHARDT BIO AG **2%** **A087872 001** Nov 18, 1982LIDOCAINE HYDROCHLORIDE VISCOUS**AT** ! LANNETT CO INC **2%** **A040708 001** Feb 27, 2007LIDOCAINE VISCOUS**AT** WEST-WARD PHARMS **2%** **A088802 001** Apr 26, 1985

INT

SOLUTION; TOPICAL

LARYNG-O-JET KIT**AT** INTL MEDICATION **4%** **A086364 001**LIDOCAINE HYDROCHLORIDE**AT** TELIGENT PHARMA INC **4%** **A204494 001** Mar 12, 2014**AT** ! WEST-WARD PHARMS **4%** **A088803 001** Apr 03, 1985

INT

SYSTEM; INTRADERMAL

ZINGO

POWDER PHARMS 0.5MG N022114 001 Aug 16, 2007

LIDOCAINE; PRILOCAINE

CREAM; TOPICAL

EMLA**AB** +! ACTAVIS LABS UT INC **2.5%; 2.5%** **N019941 001** Dec 30, 1992LIDOCAINE AND PRILOCAINE**AB** FOUGERA PHARMS **2.5%; 2.5%** **A076453 001** Aug 18, 2003**AB** HI TECH PHARMA **2.5%; 2.5%** **A076290 001** Sep 25, 2003**AB** TELIGENT PHARMA INC **2.5%; 2.5%** **A205887 001** Jun 29, 2018**AB** TOLMAR **2.5%; 2.5%** **A076320 001** Aug 27, 2003

GEL; PERIODONTAL

ORAQIX

+! DENTSPLY PHARM 2.5%; 2.5% N021451 001 Dec 19, 2003

LIDOCAINE; TETRACAINE

CREAM; TOPICAL

PLIAGLIS

+! TARO PHARMS 7%; 7% N021717 001 Jun 29, 2006

PATCH; TOPICAL

SYNERA

+! GALEN SPECIALTY 70MG; 70MG N021623 001 Jun 23, 2005

LIFITEGRAST

SOLUTION/DROPS; OPHTHALMIC

XIIDRA

+! SHIRE DEV LLC 5% N208073 001 Jul 11, 2016

LINACLOTIDE

CAPSULE; ORAL

LINZESS

+! ALLERGAN SALES LLC 72MCG N202811 003 Jan 25, 2017

+! 145MCG N202811 001 Aug 30, 2012

+ 290MCG N202811 002 Aug 30, 2012

LINAGLIPTIN

TABLET; ORAL

TRADJENTA

+! BOEHRINGER 5MG N201280 001 May 02, 2011

INGELHEIM

PRESCRIPTION DRUG PRODUCT LIST

LINAGLIPTIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

JENTADUETO

| | | | | |
|---|-------------------------|-------------|-------------|--------------|
| + | BOEHRINGER INGELHEIM | 2.5MG;500MG | N201281 001 | Jan 30, 2012 |
| + | | 2.5MG;850MG | N201281 002 | Jan 30, 2012 |
| + | ! | 2.5MG;1GM | N201281 003 | Jan 30, 2012 |

TABLET, EXTENDED RELEASE; ORAL

JENTADUETO XR

| | | | | |
|---|-------------------------|-----------|-------------|--------------|
| + | BOEHRINGER INGELHEIM | 2.5MG;1GM | N208026 001 | May 27, 2016 |
| + | ! | 5MG;1GM | N208026 002 | May 27, 2016 |

LINCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

LINCOCIN

| | | | | | |
|-----------|---|-------------------------|--------------------------------|---------------------------|--|
| AP | + | PHARMACIA AND UPJOHN | <u>EQ 300MG BASE/ML</u> | <u>N050317 001</u> | |
|-----------|---|-------------------------|--------------------------------|---------------------------|--|

LINCOMYCIN

| | | | | | |
|-----------|--|------------------|--------------------------------|---------------------------|--------------|
| AP | | X-GEN PHARMS INC | <u>EQ 300MG BASE/ML</u> | <u>A201746 001</u> | Jun 04, 2015 |
|-----------|--|------------------|--------------------------------|---------------------------|--------------|

LINDANE

LOTION; TOPICAL

LINDANE

| | | | | |
|---|-------------|----|-------------|--|
| ! | OLTA PHARMS | 1% | A087313 001 | |
|---|-------------|----|-------------|--|

SHAMPOO; TOPICAL

LINDANE

| | | | | | |
|-----------|---|------------------|------------------|---------------------------|--------------|
| AT | | OLTA PHARMS | <u>1%</u> | <u>A087266 001</u> | |
| AT | ! | WOCKHARDT BIO AG | <u>1%</u> | <u>A088191 001</u> | Sep 18, 1984 |

LINEZOLID

FOR SUSPENSION; ORAL

LINEZOLID

| | | | | | |
|-----------|--|-------------------------|-------------------------|---------------------------|--------------|
| AB | | WEST-WARD PHARMS INT | <u>100MG/5ML</u> | <u>A200068 001</u> | Jun 03, 2015 |
|-----------|--|-------------------------|-------------------------|---------------------------|--------------|

ZYVOX

| | | | | | |
|-----------|---|-------------------------|-------------------------|---------------------------|--------------|
| AB | + | PHARMACIA AND UPJOHN | <u>100MG/5ML</u> | <u>N021132 001</u> | Apr 18, 2000 |
|-----------|---|-------------------------|-------------------------|---------------------------|--------------|

SOLUTION; INTRAVENOUS

LINEZOLID

| | | | | | |
|-----------|--|-------------------------|------------------------------------|---------------------------|--------------|
| AP | | AUROBINDO PHARMA LTD | <u>600MG/300ML (2MG/ML)</u> | <u>A206917 001</u> | Aug 04, 2016 |
| AP | | FRESENIUS KABI USA | <u>600MG/300ML (2MG/ML)</u> | <u>A204764 001</u> | Mar 15, 2016 |
| AP | | HOSPIRA INC | <u>600MG/300ML (2MG/ML)</u> | <u>A205442 001</u> | Jul 07, 2015 |
| AP | | HQ SPCLT PHARMA | <u>200MG/100ML (2MG/ML)</u> | <u>A207001 001</u> | Jul 07, 2017 |
| AP | | | <u>600MG/300ML (2MG/ML)</u> | <u>A207001 002</u> | Jul 07, 2017 |
| AP | | MYLAN LABS LTD | <u>200MG/100ML (2MG/ML)</u> | <u>A205154 001</u> | Dec 06, 2017 |
| AP | | | <u>600MG/300ML (2MG/ML)</u> | <u>A205154 002</u> | Dec 06, 2017 |
| AP | | NANG KUANG PHARM CO | <u>200MG/100ML (2MG/ML)</u> | <u>A207354 001</u> | Dec 20, 2016 |
| AP | | | <u>600MG/300ML (2MG/ML)</u> | <u>A207354 002</u> | Dec 20, 2016 |
| AP | | SAGENT PHARMS | <u>200MG/100ML (2MG/ML)</u> | <u>A204696 001</u> | Mar 02, 2017 |
| AP | | | <u>600MG/300ML (2MG/ML)</u> | <u>A204696 002</u> | Mar 02, 2017 |
| AP | | SANDOZ INC | <u>200MG/100ML (2MG/ML)</u> | <u>A200904 001</u> | Jul 16, 2015 |
| AP | | | <u>600MG/300ML (2MG/ML)</u> | <u>A200904 002</u> | Jul 16, 2015 |
| AP | | TEVA PHARMS | <u>600MG/300ML (2MG/ML)</u> | <u>A200222 001</u> | Jun 27, 2012 |

ZYVOX

| | | | | | |
|-----------|---|-------------------------|------------------------------------|---------------------------|--------------|
| AP | + | PHARMACIA AND UPJOHN | <u>200MG/100ML (2MG/ML)</u> | <u>N021131 001</u> | Apr 18, 2000 |
|-----------|---|-------------------------|------------------------------------|---------------------------|--------------|

| | | | | | |
|-----------|---|---|------------------------------------|---------------------------|--------------|
| AP | + | ! | <u>600MG/300ML (2MG/ML)</u> | <u>N021131 003</u> | Apr 18, 2000 |
|-----------|---|---|------------------------------------|---------------------------|--------------|

LINEZOLID IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | |
|---|-------------|----------------------|-------------|--------------|
| + | HOSPIRA INC | 600MG/300ML (2MG/ML) | N206473 001 | Jun 18, 2015 |
|---|-------------|----------------------|-------------|--------------|

TABLET; ORAL

LINEZOLID

| | | | | | |
|-----------|--|-------------------------|---------------------|---------------------------|--------------|
| AB | | ALEMBIC PHARMS LTD | <u>600MG</u> | <u>A205233 001</u> | Dec 21, 2015 |
| AB | | ALKEM LABS LTD | <u>600MG</u> | <u>A205517 001</u> | Dec 21, 2015 |
| AB | | AMNEAL PHARMS | <u>600MG</u> | <u>A204536 001</u> | Dec 21, 2015 |
| AB | | GATE PHARMS | <u>600MG</u> | <u>A091210 001</u> | Feb 05, 2016 |
| AB | | GLENMARK PHARMS | <u>600MG</u> | <u>A078987 001</u> | Dec 21, 2015 |
| AB | | HETERO LABS LTD V | <u>600MG</u> | <u>A204239 001</u> | Dec 21, 2015 |
| AB | | MYLAN PHARMS INC | <u>600MG</u> | <u>A078845 001</u> | Dec 21, 2015 |
| AB | | NOVEL LABS INC | <u>600MG</u> | <u>A207526 001</u> | Aug 22, 2016 |
| AB | | TEVA PHARMS USA | <u>600MG</u> | <u>A078061 001</u> | May 18, 2015 |
| AB | | ZYDUS PHARMS USA INC | <u>600MG</u> | <u>A206097 001</u> | Feb 22, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

LINEZOLID

TABLET; ORAL

ZYVOX

| | | | | | |
|-----------|------------|-------------------------|--------------|--------------------|--------------|
| AB | + ! | PHARMACIA AND UPJOHN | 600MG | N021130 002 | Apr 18, 2000 |
|-----------|------------|-------------------------|--------------|--------------------|--------------|

LIOTHYRONINE SODIUM

INJECTABLE; INJECTION

LIOTHYRONINE SODIUM

| | | | | | |
|-----------|--|--------------|--------------------------|--------------------|--------------|
| AP | | X GEN PHARMS | EQ 0.01MG BASE/ML | A076923 001 | Aug 17, 2005 |
|-----------|--|--------------|--------------------------|--------------------|--------------|

TRIOSTAT

| | | | | | |
|-----------|------------|-------------------------|--------------------------|--------------------|--------------|
| AP | + ! | PAR STERILE PRODUCTS | EQ 0.01MG BASE/ML | N020105 001 | Dec 31, 1991 |
|-----------|------------|-------------------------|--------------------------|--------------------|--------------|

TABLET; ORAL

CYTOMEL

| | | | | | |
|-----------|----------|-------------|------------------------|--------------------|--|
| AB | + | KING PHARMS | EQ 0.005MG BASE | N010379 001 | |
|-----------|----------|-------------|------------------------|--------------------|--|

| | | | | | |
|-----------|----------|--|------------------------|--------------------|--|
| AB | + | | EQ 0.025MG BASE | N010379 002 | |
|-----------|----------|--|------------------------|--------------------|--|

| | | | | | |
|-----------|------------|--|-----------------------|--------------------|--|
| AB | + ! | | EQ 0.05MG BASE | N010379 003 | |
|-----------|------------|--|-----------------------|--------------------|--|

LIOTHYRONINE SODIUM

| | | | | | |
|-----------|--|------------------|------------------------|--------------------|--------------|
| AB | | MAYNE PHARMA INC | EQ 0.005MG BASE | A090097 001 | Mar 20, 2009 |
|-----------|--|------------------|------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|------------------------|--------------------|--------------|
| AB | | | EQ 0.025MG BASE | A090097 002 | Mar 20, 2009 |
|-----------|--|--|------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-----------------------|--------------------|--------------|
| AB | | | EQ 0.05MG BASE | A090097 003 | Mar 20, 2009 |
|-----------|--|--|-----------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------|------------------------|--------------------|--------------|
| AB | | MYLAN | EQ 0.005MG BASE | A090326 001 | Jul 14, 2009 |
|-----------|--|-------|------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|------------------------|--------------------|--------------|
| AB | | | EQ 0.025MG BASE | A090326 002 | Jul 14, 2009 |
|-----------|--|--|------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-----------------------|--------------------|--------------|
| AB | | | EQ 0.05MG BASE | A090326 003 | Jul 14, 2009 |
|-----------|--|--|-----------------------|--------------------|--------------|

| | | | | | |
|-----------|--|---------------------|------------------------|--------------------|--------------|
| AB | | SIGMAPHARM LABS LLC | EQ 0.005MG BASE | A200295 001 | Nov 29, 2012 |
|-----------|--|---------------------|------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|------------------------|--------------------|--------------|
| AB | | | EQ 0.025MG BASE | A200295 002 | Nov 29, 2012 |
|-----------|--|--|------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-----------------------|--------------------|--------------|
| AB | | | EQ 0.05MG BASE | A200295 003 | Nov 29, 2012 |
|-----------|--|--|-----------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------------|------------------------|--------------------|--------------|
| AB | | SUN PHARM INDS LTD | EQ 0.005MG BASE | A091382 001 | Apr 20, 2016 |
|-----------|--|--------------------|------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|------------------------|--------------------|--------------|
| AB | | | EQ 0.025MG BASE | A091382 002 | Apr 20, 2016 |
|-----------|--|--|------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-----------------------|--------------------|--------------|
| AB | | | EQ 0.05MG BASE | A091382 003 | Apr 20, 2016 |
|-----------|--|--|-----------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-----------------|------------------------|--------------------|--------------|
| AB | | TEVA PHARMS USA | EQ 0.005MG BASE | A211510 001 | Oct 26, 2018 |
|-----------|--|-----------------|------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|------------------------|--------------------|--------------|
| AB | | | EQ 0.025MG BASE | A211510 002 | Oct 26, 2018 |
|-----------|--|--|------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-----------------------|--------------------|--------------|
| AB | | | EQ 0.05MG BASE | A211510 003 | Oct 26, 2018 |
|-----------|--|--|-----------------------|--------------------|--------------|

LIRAGLUTIDE RECOMBINANT

SOLUTION; SUBCUTANEOUS

SAXENDA

+! NOVO

18MG/3ML (6MG/ML)

N206321 001 Dec 23, 2014

VICTOZA

+! NOVO NORDISK INC

18MG/3ML (6MG/ML)

N022341 001 Jan 25, 2010

LISDEXAMFETAMINE DIMESYLATE

CAPSULE; ORAL

VYVANSE

+ SHIRE DEVELOPMENT

10MG

N021977 007 Oct 30, 2014

+

20MG

N021977 004 Dec 10, 2007

+

30MG

N021977 001 Feb 23, 2007

+

40MG

N021977 005 Dec 10, 2007

+

50MG

N021977 002 Feb 23, 2007

+

60MG

N021977 006 Dec 10, 2007

+!

70MG

N021977 003 Feb 23, 2007

TABLET, CHEWABLE; ORAL

VYVANSE

+ SHIRE DEV LLC

10MG

N208510 001 Jan 28, 2017

+

20MG

N208510 002 Jan 28, 2017

+

30MG

N208510 003 Jan 28, 2017

+

40MG

N208510 004 Jan 28, 2017

+

50MG

N208510 005 Jan 28, 2017

+!

60MG

N208510 006 Jan 28, 2017

LISINOPRIL

SOLUTION; ORAL

QBRELIS

+! SILVERGATE PHARMS

1MG/ML

N208401 001 Jul 29, 2016

TABLET; ORAL

LISINOPRIL

| | | | | | |
|-----------|--|-----------------|--------------|--------------------|--------------|
| AB | | ACCORD HLTHCARE | 2.5MG | A202554 001 | Jul 30, 2013 |
|-----------|--|-----------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|------------|--------------------|--------------|
| AB | | | 5MG | A202554 002 | Jul 30, 2013 |
|-----------|--|--|------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 10MG | A202554 003 | Jul 30, 2013 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 20MG | A202554 004 | Jul 30, 2013 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 30MG | A202554 005 | Jul 30, 2013 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 40MG | A202554 006 | Jul 30, 2013 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------|--------------|--------------------|--------------|
| AB | | APOTEX INC | 2.5MG | A076102 001 | Sep 30, 2002 |
|-----------|--|------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|------------|--------------------|--------------|
| AB | | | 5MG | A076102 002 | Sep 30, 2002 |
|-----------|--|--|------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

LISINOPRIL

TABLET; ORAL

LISINOPRIL

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| <u>AB</u> | | <u>10MG</u> | <u>A076102 003</u> | Sep 30, 2002 |
| <u>AB</u> | | <u>20MG</u> | <u>A076102 004</u> | Sep 30, 2002 |
| <u>AB</u> | | <u>30MG</u> | <u>A076102 005</u> | Sep 30, 2002 |
| <u>AB</u> | | <u>40MG</u> | <u>A076102 006</u> | Sep 30, 2002 |
| <u>AB</u> | AUROBINDO | <u>2.5MG</u> | <u>A077622 001</u> | Feb 22, 2006 |
| <u>AB</u> | | <u>5MG</u> | <u>A077622 002</u> | Feb 22, 2006 |
| <u>AB</u> | | <u>10MG</u> | <u>A077622 003</u> | Feb 22, 2006 |
| <u>AB</u> | | <u>20MG</u> | <u>A077622 004</u> | Feb 22, 2006 |
| <u>AB</u> | | <u>30MG</u> | <u>A077622 005</u> | Feb 22, 2006 |
| <u>AB</u> | | <u>40MG</u> | <u>A077622 006</u> | Feb 22, 2006 |
| <u>AB</u> | CASI PHARMS INC | <u>2.5MG</u> | <u>A075994 001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>5MG</u> | <u>A075994 002</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>10MG</u> | <u>A075994 003</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>20MG</u> | <u>A075994 004</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>30MG</u> | <u>A075994 005</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>40MG</u> | <u>A075994 006</u> | Jul 01, 2002 |
| <u>AB</u> | HIKMA INTL PHARMS | <u>2.5MG</u> | <u>A076063 001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>5MG</u> | <u>A076063 002</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>10MG</u> | <u>A076063 003</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>20MG</u> | <u>A076063 004</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>30MG</u> | <u>A076063 006</u> | Jun 27, 2003 |
| <u>AB</u> | | <u>40MG</u> | <u>A076063 005</u> | Jul 01, 2002 |
| <u>AB</u> | INVAGEN PHARMS | <u>2.5MG</u> | <u>A203508 001</u> | Oct 29, 2013 |
| <u>AB</u> | | <u>5MG</u> | <u>A203508 002</u> | Oct 29, 2013 |
| <u>AB</u> | | <u>10MG</u> | <u>A203508 003</u> | Oct 29, 2013 |
| <u>AB</u> | | <u>20MG</u> | <u>A203508 004</u> | Oct 29, 2013 |
| <u>AB</u> | | <u>30MG</u> | <u>A203508 005</u> | Oct 29, 2013 |
| <u>AB</u> | | <u>40MG</u> | <u>A203508 006</u> | Oct 29, 2013 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>2.5MG</u> | <u>A075752 001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>5MG</u> | <u>A075752 002</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>10MG</u> | <u>A075752 003</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>20MG</u> | <u>A075752 004</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>30MG</u> | <u>A075752 005</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>40MG</u> | <u>A075752 006</u> | Jul 01, 2002 |
| <u>AB</u> | LUPIN | <u>2.5MG</u> | <u>A077321 001</u> | Sep 09, 2005 |
| <u>AB</u> | | <u>5MG</u> | <u>A077321 002</u> | Sep 09, 2005 |
| <u>AB</u> | | <u>10MG</u> | <u>A077321 003</u> | Sep 09, 2005 |
| <u>AB</u> | | <u>20MG</u> | <u>A077321 004</u> | Sep 09, 2005 |
| <u>AB</u> | | <u>30MG</u> | <u>A077321 005</u> | Sep 09, 2005 |
| <u>AB</u> | | <u>40MG</u> | <u>A077321 006</u> | Sep 09, 2005 |
| <u>AB</u> | MYLAN | <u>2.5MG</u> | <u>A076071 001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>5MG</u> | <u>A076071 002</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>10MG</u> | <u>A076071 003</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>20MG</u> | <u>A076071 004</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>30MG</u> | <u>A076071 005</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>40MG</u> | <u>A076071 006</u> | Jul 01, 2002 |
| <u>AB</u> | PRINSTON INC | <u>2.5MG</u> | <u>A075743 001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>2.5MG</u> | <u>A076180 001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>5MG</u> | <u>A075743 002</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>5MG</u> | <u>A076180 002</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>10MG</u> | <u>A075743 003</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>10MG</u> | <u>A076180 003</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>20MG</u> | <u>A075743 004</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>20MG</u> | <u>A076164 001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>30MG</u> | <u>A075743 005</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>30MG</u> | <u>A076164 002</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>40MG</u> | <u>A075743 006</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>40MG</u> | <u>A076164 003</u> | Jul 01, 2002 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>2.5MG</u> | <u>A075944 001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>5MG</u> | <u>A075944 002</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>10MG</u> | <u>A075944 003</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>20MG</u> | <u>A075944 004</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>30MG</u> | <u>A075944 006</u> | Feb 11, 2003 |
| <u>AB</u> | | <u>40MG</u> | <u>A075944 005</u> | Jul 01, 2002 |
| <u>AB</u> | WATSON LABS | <u>2.5MG</u> | <u>A076059 001</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>5MG</u> | <u>A076059 002</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>10MG</u> | <u>A076059 003</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>20MG</u> | <u>A076059 004</u> | Jul 01, 2002 |
| <u>AB</u> | | <u>30MG</u> | <u>A076059 005</u> | Jul 01, 2002 |

PRESCRIPTION DRUG PRODUCT LIST

LISINOPRIL

TABLET; ORAL

LISINOPRIL

| | | | | |
|-----------|-----------|--------------|--------------------|--------------|
| <u>AB</u> | | <u>40MG</u> | <u>A076059 006</u> | Jul 01, 2002 |
| <u>AB</u> | WOCKHARDT | <u>2.5MG</u> | <u>A078402 001</u> | Apr 19, 2007 |
| <u>AB</u> | | <u>5MG</u> | <u>A078402 002</u> | Apr 19, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A078402 003</u> | Apr 19, 2007 |
| <u>AB</u> | | <u>20MG</u> | <u>A078402 004</u> | Apr 19, 2007 |
| <u>AB</u> | | <u>30MG</u> | <u>A078402 005</u> | Apr 19, 2007 |
| <u>AB</u> | | <u>40MG</u> | <u>A078402 006</u> | Apr 19, 2007 |

PRINIVIL

| | | | | |
|-----------|-------|-------------|--------------------|--------------|
| <u>AB</u> | MERCK | <u>5MG</u> | <u>N019558 001</u> | Dec 29, 1987 |
| <u>AB</u> | | <u>10MG</u> | <u>N019558 002</u> | Dec 29, 1987 |
| <u>AB</u> | | <u>20MG</u> | <u>N019558 003</u> | Dec 29, 1987 |
| <u>AB</u> | | <u>40MG</u> | <u>N019558 004</u> | Oct 25, 1988 |

ZESTRIL

| | | | | | |
|-----------|---|---------------|--------------|--------------------|--------------|
| <u>AB</u> | + | ALVOGEN MALTA | <u>2.5MG</u> | <u>N019777 005</u> | Apr 29, 1993 |
| <u>AB</u> | + | | <u>5MG</u> | <u>N019777 001</u> | May 19, 1988 |
| <u>AB</u> | + | | <u>10MG</u> | <u>N019777 002</u> | May 19, 1988 |
| <u>AB</u> | + | | <u>20MG</u> | <u>N019777 003</u> | May 19, 1988 |
| <u>AB</u> | + | | <u>30MG</u> | <u>N019777 006</u> | Jan 20, 1999 |
| <u>AB</u> | + | | <u>40MG</u> | <u>N019777 004</u> | May 19, 1988 |

LITHIUM CARBONATE

CAPSULE; ORAL

LITHIUM CARBONATE

| | | | | | |
|-----------|---------------------|-------------------------|--------------------|--------------------|--------------|
| <u>AB</u> | ALEMBIC LTD | <u>150MG</u> | <u>A079159 001</u> | Jan 12, 2009 | |
| <u>AB</u> | | <u>300MG</u> | <u>A079159 002</u> | Jan 12, 2009 | |
| <u>AB</u> | | <u>600MG</u> | <u>A079159 003</u> | Jan 12, 2009 | |
| <u>AB</u> | GLENMARK GENERICS | <u>150MG</u> | <u>A079139 001</u> | Feb 03, 2009 | |
| <u>AB</u> | | <u>300MG</u> | <u>A079139 002</u> | Feb 03, 2009 | |
| <u>AB</u> | | <u>600MG</u> | <u>A079139 003</u> | Feb 03, 2009 | |
| <u>AB</u> | HETERO LABS LTD III | <u>150MG</u> | <u>A090702 001</u> | Sep 25, 2009 | |
| <u>AB</u> | | <u>300MG</u> | <u>A090702 002</u> | Sep 25, 2009 | |
| <u>AB</u> | | <u>600MG</u> | <u>A090702 003</u> | Sep 25, 2009 | |
| <u>AB</u> | MYLAN PHARMS INC | <u>150MG</u> | <u>A076243 002</u> | Feb 24, 2003 | |
| <u>AB</u> | | <u>300MG</u> | <u>A076243 001</u> | Jun 27, 2002 | |
| <u>AB</u> | | <u>600MG</u> | <u>A078763 001</u> | Apr 15, 2008 | |
| <u>AB</u> | + | WEST-WARD PHARMS INT | <u>150MG</u> | <u>N017812 002</u> | Jan 28, 1987 |
| <u>AB</u> | + | | <u>300MG</u> | <u>N017812 001</u> | |
| <u>AB</u> | + | | <u>600MG</u> | <u>N017812 003</u> | Jan 28, 1987 |

TABLET; ORAL

LITHIUM CARBONATE

| | | | | | |
|-----------|--------------------|-------------------------|--------------------|--------------------|--------------|
| <u>AB</u> | SUN PHARM INDS INC | <u>300MG</u> | <u>A091027 001</u> | Jun 24, 2010 | |
| <u>AB</u> | + | WEST-WARD PHARMS INT | <u>300MG</u> | <u>N018558 001</u> | Jan 29, 1982 |

TABLET, EXTENDED RELEASE; ORAL

LITHIUM CARBONATE

| | | | | | |
|-----------|-------------------------|----------------|--------------------|--------------------|--|
| <u>AB</u> | ALEMBIC PHARMS LTD | <u>300MG</u> | <u>A204445 001</u> | Jun 10, 2015 | |
| <u>AB</u> | GLENMARK GENERICS | <u>450MG</u> | <u>A091616 001</u> | Feb 14, 2011 | |
| <u>AB</u> | GLENMARK PHARMS INC | <u>300MG</u> | <u>A091544 001</u> | Dec 27, 2010 | |
| <u>AB</u> | HERITAGE PHARMA | <u>300MG</u> | <u>A205532 001</u> | Sep 29, 2016 | |
| <u>AB</u> | MYLAN PHARMS INC | <u>300MG</u> | <u>A202288 001</u> | Jun 29, 2012 | |
| <u>AB</u> | | <u>450MG</u> | <u>A202219 001</u> | Aug 08, 2012 | |
| <u>AB</u> | UNIQUE PHARM LABS | <u>300MG</u> | <u>A204779 001</u> | Jul 27, 2015 | |
| <u>AB</u> | | <u>450MG</u> | <u>A205663 001</u> | Jun 05, 2017 | |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>300MG</u> | <u>A076832 001</u> | Oct 28, 2004 | |
| <u>AB</u> | ! | <u>450MG</u> | <u>A076691 001</u> | Jan 05, 2004 | |
| <u>AB</u> | + | ANI PHARMS INC | <u>300MG</u> | <u>N018027 001</u> | |

LITHIUM CITRATE

SYRUP; ORAL

LITHIUM CITRATE

| | | | | | |
|-----------|---|-------------------------|-------------------------------|--------------------|--------------|
| <u>AA</u> | + | WEST-WARD PHARMS INT | <u>EQ 300MG CARBONATE/5ML</u> | <u>N018421 001</u> | |
| <u>AA</u> | | WOCKHARDT BIO AG | <u>EQ 300MG CARBONATE/5ML</u> | <u>A070755 001</u> | May 21, 1986 |

PRESCRIPTION DRUG PRODUCT LIST

LIXISENATIDE

SOLUTION; SUBCUTANEOUS

ADLYXIN

| | | | | | | |
|---|---|-------------------|------------------------|---------|-----|--------------|
| + | ! | SANOFI-AVENTIS US | 0.15MG/3ML (0.05MG/ML) | N208471 | 001 | Jul 27, 2016 |
| + | ! | | 0.3MG/3ML (0.1MG/ML) | N208471 | 002 | Jul 27, 2016 |

LODOXAMIDE TROMETHAMINE

SOLUTION/DROPS; OPHTHALMIC

ALOMIDE

| | | | | | | |
|---|---|-------------------------|--------------|---------|-----|--------------|
| + | ! | NOVARTIS PHARMS CORP | EQ 0.1% BASE | N020191 | 001 | Sep 23, 1993 |
|---|---|-------------------------|--------------|---------|-----|--------------|

LOFEXIDINE HYDROCHLORIDE

TABLET; ORAL

LUCEMYRA

| | | | | | | |
|---|---|------------------|----------------|---------|-----|--------------|
| + | ! | US WORLDMEDS LLC | EQ 0.18MG BASE | N209229 | 001 | May 16, 2018 |
|---|---|------------------|----------------|---------|-----|--------------|

LOMITAPIDE MESYLATE

CAPSULE; ORAL

JUXTAPID

| | | | | | | |
|---|---|----------|--------------|---------|-----|--------------|
| + | | AEGERION | EQ 5MG BASE | N203858 | 001 | Dec 21, 2012 |
| + | | | EQ 10MG BASE | N203858 | 002 | Dec 21, 2012 |
| + | | | EQ 20MG BASE | N203858 | 003 | Dec 21, 2012 |
| + | | | EQ 30MG BASE | N203858 | 004 | Apr 23, 2015 |
| + | | | EQ 40MG BASE | N203858 | 005 | Apr 23, 2015 |
| + | ! | | EQ 60MG BASE | N203858 | 006 | Apr 23, 2015 |

LOMUSTINE

CAPSULE; ORAL

GLEOSTINE

| | | | | | | |
|---|---|---------------|-------|---------|-----|--|
| + | | CORDEN PHARMA | 10MG | N017588 | 001 | |
| + | | | 40MG | N017588 | 002 | |
| + | ! | | 100MG | N017588 | 003 | |

LOPERAMIDE HYDROCHLORIDE

CAPSULE; ORAL

LOPERAMIDE HYDROCHLORIDE

| | | | | | | |
|-----------|---|-------|------------|----------------|------------|--------------|
| AB | ! | MYLAN | 2MG | A072741 | 001 | Sep 18, 1991 |
| AB | | TEVA | 2MG | A073192 | 001 | Apr 30, 1992 |

LOPINAVIR; RITONAVIR

SOLUTION; ORAL

KALETRA

| | | | | | | | |
|-----------|---|---|--------|-------------------------|----------------|------------|--------------|
| AA | + | ! | ABEVIE | 80MG/ML; 20MG/ML | N021251 | 001 | Sep 15, 2000 |
|-----------|---|---|--------|-------------------------|----------------|------------|--------------|

LOPINAVIR AND RITONAVIR

| | | | | | | | |
|-----------|--|--|----------------|-------------------------|----------------|------------|--------------|
| AA | | | LANNETT CO INC | 80MG/ML; 20MG/ML | A207407 | 001 | Dec 27, 2016 |
|-----------|--|--|----------------|-------------------------|----------------|------------|--------------|

TABLET; ORAL

KALETRA

| | | | | | | |
|---|---|--------|-------------|---------|-----|--------------|
| + | | ABEVIE | 100MG; 25MG | N021906 | 002 | Nov 09, 2007 |
| + | ! | | 200MG; 50MG | N021906 | 001 | Oct 28, 2005 |

LORAZEPAM

CONCENTRATE; ORAL

LORAZEPAM

| | | | | | | | |
|-----------|--|--|-------------------|---------------|----------------|------------|--------------|
| AA | | | AMNEAL PHARMS | 2MG/ML | A091383 | 001 | Dec 23, 2009 |
| AA | | | ANDA REPOSITORY | 2MG/ML | A079244 | 001 | Apr 28, 2009 |
| AA | | | HI-TECH PHARMA CO | 2MG/ML | A200169 | 001 | Jan 30, 2012 |
| AA | | | LUPIN LTD | 2MG/ML | A091407 | 001 | Feb 19, 2013 |
| AA | | | PHARM ASSOC | 2MG/ML | A090260 | 001 | Jun 15, 2010 |

LORAZEPAM INTENSOL

| | | | | | | | |
|-----------|---|--|-------------------------|---------------|----------------|------------|--------------|
| AA | ! | | WEST-WARD PHARMS INT | 2MG/ML | A072755 | 001 | Jun 28, 1991 |
|-----------|---|--|-------------------------|---------------|----------------|------------|--------------|

INJECTABLE; INJECTION

ATIVAN

| | | | | | | | |
|-----------|---|---|-------------------------|---------------|----------------|------------|--|
| AP | + | ! | WEST-WARD PHARMS INT | 2MG/ML | N018140 | 001 | |
|-----------|---|---|-------------------------|---------------|----------------|------------|--|

| | | | | | | | |
|-----------|---|---|--|---------------|----------------|------------|--|
| AP | + | ! | | 4MG/ML | N018140 | 002 | |
|-----------|---|---|--|---------------|----------------|------------|--|

LORAZEPAM

| | | | | | | | |
|-----------|--|--|---------------------|---------------|----------------|------------|--------------|
| AP | | | AKORN | 2MG/ML | A075025 | 001 | Jul 23, 1998 |
| AP | | | HOSPIRA | 2MG/ML | A074243 | 001 | Apr 12, 1994 |
| AP | | | | 2MG/ML | A074282 | 001 | May 27, 1994 |
| AP | | | | 4MG/ML | A074243 | 002 | Apr 12, 1994 |
| AP | | | | 4MG/ML | A074282 | 002 | May 27, 1994 |
| AP | | | INTL MEDICATION SYS | 2MG/ML | A076150 | 001 | Nov 15, 2004 |

PRESCRIPTION DRUG PRODUCT LIST

LORAZEPAM

TABLET;ORAL

ATIVAN

| | | | | | | |
|-----------|----------|--------------|---------------------|-----------------------|-------------------|--|
| AB | + | VALEANT INTL | <u>0.5MG</u> | <u>N017794</u> | <u>001</u> | |
| AB | + | | <u>1MG</u> | <u>N017794</u> | <u>002</u> | |
| AB | + | ! | <u>2MG</u> | <u>N017794</u> | <u>003</u> | |

LORAZEPAM

| | | | | | | |
|-----------|--|---------------------|---------------------|-----------------------|-------------------|--------------|
| AB | | AMNEAL PHARMS | <u>0.5MG</u> | <u>A078826</u> | <u>001</u> | Jun 23, 2010 |
| AB | | | <u>1MG</u> | <u>A078826</u> | <u>002</u> | Jun 23, 2010 |
| AB | | | <u>2MG</u> | <u>A078826</u> | <u>003</u> | Jun 23, 2010 |
| AB | | ANI PHARMS INC | <u>0.5MG</u> | <u>A077396</u> | <u>001</u> | Dec 13, 2006 |
| AB | | | <u>1MG</u> | <u>A077396</u> | <u>002</u> | Dec 13, 2006 |
| AB | | | <u>2MG</u> | <u>A077396</u> | <u>003</u> | Dec 13, 2006 |
| AB | | AUROLIFE PHARMA LLC | <u>0.5MG</u> | <u>A203572</u> | <u>001</u> | Dec 22, 2017 |
| AB | | | <u>1MG</u> | <u>A203572</u> | <u>002</u> | Dec 22, 2017 |
| AB | | | <u>2MG</u> | <u>A203572</u> | <u>003</u> | Dec 22, 2017 |
| AB | | LEADING PHARMA LLC | <u>0.5MG</u> | <u>A078203</u> | <u>001</u> | Jul 30, 2007 |
| AB | | | <u>1MG</u> | <u>A078203</u> | <u>002</u> | Jul 30, 2007 |
| AB | | | <u>2MG</u> | <u>A078203</u> | <u>003</u> | Jul 30, 2007 |
| AB | | MYLAN | <u>0.5MG</u> | <u>A077657</u> | <u>001</u> | Mar 16, 2006 |
| AB | | | <u>1MG</u> | <u>A077657</u> | <u>002</u> | Mar 16, 2006 |
| AB | | | <u>2MG</u> | <u>A077657</u> | <u>003</u> | Mar 16, 2006 |
| AB | | OXFORD PHARMS | <u>0.5MG</u> | <u>A077754</u> | <u>001</u> | May 10, 2006 |
| AB | | | <u>1MG</u> | <u>A077754</u> | <u>002</u> | May 10, 2006 |
| AB | | | <u>2MG</u> | <u>A077754</u> | <u>003</u> | May 10, 2006 |
| AB | | SANDOZ | <u>0.5MG</u> | <u>A071141</u> | <u>002</u> | Apr 21, 1987 |
| AB | | | <u>1MG</u> | <u>A071141</u> | <u>003</u> | Apr 21, 1987 |
| AB | | | <u>2MG</u> | <u>A071141</u> | <u>001</u> | Apr 21, 1987 |
| AB | | WATSON LABS | <u>0.5MG</u> | <u>A072926</u> | <u>001</u> | Oct 31, 1991 |
| AB | | | <u>1MG</u> | <u>A072927</u> | <u>001</u> | Oct 31, 1991 |
| AB | | | <u>2MG</u> | <u>A072928</u> | <u>001</u> | Oct 31, 1991 |

LORCASERIN HYDROCHLORIDE

TABLET;ORAL

BELVIQ

| | | | | | | |
|--|----------|-----------|------|---------|-----|--------------|
| | + | EISAI INC | 10MG | N022529 | 001 | Jun 27, 2012 |
|--|----------|-----------|------|---------|-----|--------------|

TABLET, EXTENDED RELEASE;ORAL

| | | | | | | |
|--|----------|-----------|------|---------|-----|--------------|
| | + | EISAI INC | 20MG | N208524 | 001 | Jul 15, 2016 |
|--|----------|-----------|------|---------|-----|--------------|

LORLATINIB

TABLET;ORAL

LORBRENA

| | | | | | | |
|--|----------|------------|-------|---------|-----|--------------|
| | + | PFIZER INC | 25MG | N210868 | 001 | Nov 02, 2018 |
| | + | ! | 100MG | N210868 | 002 | Nov 02, 2018 |

LOSARTAN POTASSIUM

TABLET;ORAL

COZAAR

| | | | | | | |
|-----------|----------|-------------------|---------------------|-----------------------|-------------------|--------------|
| AB | + | MERCK SHARP DOHME | <u>100MG</u> | <u>N020386</u> | <u>003</u> | Oct 13, 1998 |
|-----------|----------|-------------------|---------------------|-----------------------|-------------------|--------------|

LOSARTAN POTASSIUM

| | | | | | | |
|-----------|--|---------------------|---------------------|-----------------------|-------------------|--------------|
| AB | | ALEMBIC PHARMS LTD | <u>25MG</u> | <u>A090428</u> | <u>001</u> | Oct 06, 2010 |
| AB | | | <u>50MG</u> | <u>A090428</u> | <u>002</u> | Oct 06, 2010 |
| AB | | | <u>100MG</u> | <u>A090428</u> | <u>003</u> | Oct 06, 2010 |
| AB | | AUROBINDO PHARMA | <u>25MG</u> | <u>A090083</u> | <u>001</u> | Oct 06, 2010 |
| AB | | | <u>50MG</u> | <u>A090083</u> | <u>002</u> | Oct 06, 2010 |
| AB | | | <u>100MG</u> | <u>A090083</u> | <u>003</u> | Oct 06, 2010 |
| AB | | CADISTA PHARMS | <u>25MG</u> | <u>A201170</u> | <u>001</u> | Sep 18, 2012 |
| AB | | | <u>50MG</u> | <u>A201170</u> | <u>002</u> | Sep 18, 2012 |
| AB | | | <u>100MG</u> | <u>A201170</u> | <u>003</u> | Sep 18, 2012 |
| AB | | HETERO LABS LTD V | <u>25MG</u> | <u>A203835</u> | <u>001</u> | Aug 12, 2015 |
| AB | | | <u>50MG</u> | <u>A203835</u> | <u>002</u> | Aug 12, 2015 |
| AB | | | <u>100MG</u> | <u>A203835</u> | <u>003</u> | Aug 12, 2015 |
| AB | | IPCA LABS LTD | <u>25MG</u> | <u>A200290</u> | <u>001</u> | Aug 30, 2013 |
| AB | | | <u>50MG</u> | <u>A200290</u> | <u>002</u> | Aug 30, 2013 |
| AB | | | <u>100MG</u> | <u>A200290</u> | <u>003</u> | Aug 30, 2013 |
| AB | | LUPIN LTD | <u>25MG</u> | <u>A078232</u> | <u>001</u> | Oct 06, 2010 |
| AB | | | <u>50MG</u> | <u>A078232</u> | <u>002</u> | Oct 06, 2010 |
| AB | | | <u>100MG</u> | <u>A078232</u> | <u>003</u> | Oct 06, 2010 |
| AB | | MACLEODS PHARMS LTD | <u>25MG</u> | <u>A202230</u> | <u>001</u> | May 30, 2012 |
| AB | | | <u>50MG</u> | <u>A202230</u> | <u>002</u> | May 30, 2012 |
| AB | | | <u>100MG</u> | <u>A202230</u> | <u>003</u> | May 30, 2012 |
| AB | | MYLAN | <u>25MG</u> | <u>A091590</u> | <u>001</u> | Oct 06, 2010 |
| AB | | | <u>50MG</u> | <u>A091590</u> | <u>002</u> | Oct 06, 2010 |

PRESCRIPTION DRUG PRODUCT LIST

LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | <u>100MG</u> | <u>A091590</u> | <u>003</u> | Oct 06, 2010 |
| <u>AB</u> | PRINSTON INC | <u>25MG</u> | <u>A091497</u> | <u>001</u> | Jun 06, 2011 |
| <u>AB</u> | | <u>50MG</u> | <u>A091497</u> | <u>002</u> | Jun 06, 2011 |
| <u>AB</u> | | <u>100MG</u> | <u>A091497</u> | <u>003</u> | Jun 06, 2011 |
| <u>AB</u> | SANDOZ | <u>25MG</u> | <u>A077424</u> | <u>001</u> | Oct 06, 2010 |
| <u>AB</u> | | <u>50MG</u> | <u>A077424</u> | <u>002</u> | Oct 06, 2010 |
| <u>AB</u> | | <u>100MG</u> | <u>A077424</u> | <u>003</u> | Oct 06, 2010 |
| <u>AB</u> | STRIDES VIVIMED | <u>25MG</u> | <u>A090382</u> | <u>001</u> | Oct 06, 2010 |
| <u>AB</u> | | <u>50MG</u> | <u>A090382</u> | <u>002</u> | Oct 06, 2010 |
| <u>AB</u> | | <u>100MG</u> | <u>A090382</u> | <u>003</u> | Oct 06, 2010 |
| <u>AB</u> | TEVA | <u>25MG</u> | <u>A076958</u> | <u>001</u> | Apr 06, 2010 |
| <u>AB</u> | | <u>50MG</u> | <u>A076958</u> | <u>002</u> | Apr 06, 2010 |
| <u>AB</u> | | <u>100MG</u> | <u>A076958</u> | <u>003</u> | Apr 06, 2010 |
| <u>AB</u> | TORRENT PHARMS | <u>25MG</u> | <u>A090467</u> | <u>001</u> | Oct 06, 2010 |
| <u>AB</u> | | <u>50MG</u> | <u>A090467</u> | <u>002</u> | Oct 06, 2010 |
| <u>AB</u> | | <u>100MG</u> | <u>A090467</u> | <u>003</u> | Oct 06, 2010 |
| <u>AB</u> | UNICHEM LABS LTD | <u>25MG</u> | <u>A203030</u> | <u>001</u> | Oct 14, 2015 |
| <u>AB</u> | | <u>50MG</u> | <u>A203030</u> | <u>002</u> | Oct 14, 2015 |
| <u>AB</u> | | <u>100MG</u> | <u>A203030</u> | <u>003</u> | Oct 14, 2015 |
| <u>AB</u> | UPSHER SMITH LABS | <u>25MG</u> | <u>A090544</u> | <u>001</u> | Oct 06, 2010 |
| <u>AB</u> | | <u>50MG</u> | <u>A090544</u> | <u>002</u> | Oct 06, 2010 |
| <u>AB</u> | | <u>100MG</u> | <u>A090544</u> | <u>003</u> | Oct 06, 2010 |
| <u>AB</u> | VIVA HLTHCARE | <u>25MG</u> | <u>A091541</u> | <u>001</u> | Sep 24, 2012 |
| <u>AB</u> | | <u>50MG</u> | <u>A091541</u> | <u>002</u> | Sep 24, 2012 |
| <u>AB</u> | | <u>100MG</u> | <u>A091541</u> | <u>003</u> | Sep 24, 2012 |
| <u>AB</u> | WATSON LABS | <u>25MG</u> | <u>A091129</u> | <u>001</u> | Oct 06, 2010 |
| <u>AB</u> | | <u>50MG</u> | <u>A091129</u> | <u>002</u> | Oct 06, 2010 |
| <u>AB</u> | | <u>100MG</u> | <u>A091129</u> | <u>003</u> | Oct 06, 2010 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>25MG</u> | <u>A077459</u> | <u>001</u> | Oct 06, 2010 |
| <u>AB</u> | | <u>50MG</u> | <u>A077459</u> | <u>002</u> | Oct 06, 2010 |
| <u>AB</u> | | <u>100MG</u> | <u>A077459</u> | <u>003</u> | Oct 06, 2010 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>25MG</u> | <u>A078243</u> | <u>001</u> | Oct 06, 2010 |
| <u>AB</u> | | <u>50MG</u> | <u>A078243</u> | <u>002</u> | Oct 06, 2010 |
| <u>AB</u> | | <u>100MG</u> | <u>A078243</u> | <u>003</u> | Oct 06, 2010 |

LOTEPREDNOL ETABONATE

GEL; OPHTHALMIC

LOTEMAX

+! BAUSCH AND LOMB INC 0.5% N202872 001 Sep 28, 2012

OINTMENT; OPHTHALMIC

LOTEMAX

+! BAUSCH AND LOMB 0.5% N200738 001 Apr 15, 2011

SUSPENSION/DROPS; OPHTHALMIC

ALREX

+! BAUSCH AND LOMB 0.2% N020803 001 Mar 09, 1998

INVELTYS

+! KALA PHARMS INC 1% N210565 001 Aug 22, 2018

LOTEMAX

+! BAUSCH AND LOMB 0.5% N020583 001 Mar 09, 1998

LOTEPREDNOL ETABONATE; TOBRAMYCIN

SUSPENSION/DROPS; OPHTHALMIC

ZYLET

+! BAUSCH AND LOMB 0.5%; 0.3% N050804 001 Dec 14, 2004

LOVASTATIN

TABLET; ORAL

LOVASTATIN

| | | | | | |
|-----------|-------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | ACTAVIS ELIZABETH | <u>10MG</u> | <u>A075828</u> | <u>001</u> | Dec 17, 2001 |
| <u>AB</u> | | <u>20MG</u> | <u>A075828</u> | <u>002</u> | Dec 17, 2001 |
| <u>AB</u> | | <u>40MG</u> | <u>A075828</u> | <u>003</u> | Dec 17, 2001 |
| <u>AB</u> | APOTEX INC | <u>10MG</u> | <u>A077748</u> | <u>001</u> | Feb 28, 2007 |
| <u>AB</u> | | <u>20MG</u> | <u>A077748</u> | <u>002</u> | Feb 28, 2007 |
| <u>AB</u> | | <u>40MG</u> | <u>A077748</u> | <u>003</u> | Feb 28, 2007 |
| <u>AB</u> | CARLSBAD | <u>10MG</u> | <u>A075991</u> | <u>001</u> | Jun 05, 2002 |
| <u>AB</u> | | <u>20MG</u> | <u>A075991</u> | <u>002</u> | Jun 05, 2002 |
| <u>AB</u> | ! | <u>40MG</u> | <u>A075991</u> | <u>003</u> | Jun 05, 2002 |
| <u>AB</u> | LUPIN | <u>10MG</u> | <u>A078296</u> | <u>001</u> | Mar 14, 2008 |
| <u>AB</u> | | <u>20MG</u> | <u>A078296</u> | <u>002</u> | Nov 01, 2007 |

PRESCRIPTION DRUG PRODUCT LIST

LOVASTATIN

TABLET; ORAL

LOVASTATIN

| | | | | |
|-----------|-------------------------|-------------|--------------------|--------------|
| <u>AB</u> | | <u>40MG</u> | <u>A078296 003</u> | Nov 01, 2007 |
| <u>AB</u> | MYLAN | <u>10MG</u> | <u>A075451 001</u> | Dec 17, 2001 |
| <u>AB</u> | | <u>20MG</u> | <u>A075451 002</u> | Dec 17, 2001 |
| <u>AB</u> | | <u>40MG</u> | <u>A075451 003</u> | Dec 17, 2001 |
| <u>AB</u> | SANDOZ | <u>10MG</u> | <u>A075300 001</u> | Dec 17, 2001 |
| <u>AB</u> | | <u>10MG</u> | <u>A075636 001</u> | Dec 17, 2001 |
| <u>AB</u> | | <u>20MG</u> | <u>A075300 002</u> | Dec 17, 2001 |
| <u>AB</u> | | <u>20MG</u> | <u>A075636 002</u> | Dec 17, 2001 |
| <u>AB</u> | | <u>40MG</u> | <u>A075300 003</u> | Dec 17, 2001 |
| <u>AB</u> | | <u>40MG</u> | <u>A075636 003</u> | Dec 17, 2001 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>10MG</u> | <u>A077520 001</u> | Apr 14, 2006 |
| <u>AB</u> | | <u>20MG</u> | <u>A077520 002</u> | Apr 14, 2006 |
| <u>AB</u> | | <u>40MG</u> | <u>A077520 003</u> | Apr 14, 2006 |
| <u>AB</u> | TEVA | <u>10MG</u> | <u>A075551 003</u> | Dec 17, 2001 |
| <u>AB</u> | | <u>20MG</u> | <u>A075551 002</u> | Dec 17, 2001 |
| <u>AB</u> | | <u>40MG</u> | <u>A075551 001</u> | Dec 17, 2001 |

TABLET, EXTENDED RELEASE; ORAL

ALTOPREV

| | | | | |
|---|-----------------|------|-------------|--------------|
| + | COVIS PHARMA BV | 20MG | N021316 002 | Jun 26, 2002 |
| + | | 40MG | N021316 003 | Jun 26, 2002 |
| + | ! | 60MG | N021316 004 | Jun 26, 2002 |

LOXAPINE

POWDER; INHALATION

ADASUVE

| | | | | | |
|---|---|----------|------|-------------|--------------|
| + | ! | GALEN UK | 10MG | N022549 001 | Dec 21, 2012 |
|---|---|----------|------|-------------|--------------|

LOXAPINE SUCCINATE

CAPSULE; ORAL

LOXAPINE SUCCINATE

| | | | | |
|-----------|----------------|---------------------|--------------------|--------------|
| <u>AB</u> | ELITE LABS INC | <u>EQ 5MG BASE</u> | <u>A076868 001</u> | Aug 04, 2005 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A076868 002</u> | Aug 04, 2005 |
| <u>AB</u> | | <u>EQ 25MG BASE</u> | <u>A076868 003</u> | Aug 04, 2005 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A076868 004</u> | Aug 04, 2005 |
| <u>AB</u> | LANNETT CO INC | <u>EQ 5MG BASE</u> | <u>A090695 001</u> | Sep 26, 2011 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A090695 002</u> | Sep 26, 2011 |
| <u>AB</u> | | <u>EQ 25MG BASE</u> | <u>A090695 003</u> | Sep 26, 2011 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A090695 004</u> | Sep 26, 2011 |
| <u>AB</u> | MYLAN | <u>EQ 5MG BASE</u> | <u>A076762 001</u> | Nov 01, 2004 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A076762 002</u> | Nov 01, 2004 |
| <u>AB</u> | | <u>EQ 25MG BASE</u> | <u>A076762 003</u> | Nov 01, 2004 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A076762 004</u> | Nov 01, 2004 |
| <u>AB</u> | WATSON LABS | <u>EQ 5MG BASE</u> | <u>A072204 001</u> | Jun 15, 1988 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A072205 001</u> | Jun 15, 1988 |
| <u>AB</u> | ! | <u>EQ 25MG BASE</u> | <u>A072206 001</u> | Jun 15, 1988 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A072062 001</u> | Jun 15, 1988 |

LUBIPROSTONE

CAPSULE; ORAL

AMITIZA

| | | | | |
|---|--------------------|-------|-------------|--------------|
| + | SUCAMPO PHARMA LLC | 8MCG | N021908 002 | Apr 29, 2008 |
| + | ! | 24MCG | N021908 001 | Jan 31, 2006 |

LULICONAZOLE

CREAM; TOPICAL

LUZU

| | | | | | |
|---|---|---------|----|-------------|--------------|
| + | ! | MEDICIS | 1% | N204153 001 | Nov 14, 2013 |
|---|---|---------|----|-------------|--------------|

LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LATUDA

| | | | | | |
|-----------|---|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | + | SUNOVION PHARMS INC | <u>20MG</u> | <u>N200603 003</u> | Dec 07, 2011 |
| <u>AB</u> | + | ! | <u>40MG</u> | <u>N200603 001</u> | Oct 28, 2010 |
| <u>AB</u> | + | | <u>60MG</u> | <u>N200603 005</u> | Jul 12, 2013 |
| <u>AB</u> | + | | <u>80MG</u> | <u>N200603 002</u> | Oct 28, 2010 |
| <u>AB</u> | + | | <u>120MG</u> | <u>N200603 004</u> | Apr 26, 2012 |

LURASIDONE HYDROCHLORIDE

| | | | | | |
|-----------|--|-----------------|-------------|--------------------|--------------|
| <u>AB</u> | | ACCORD HLTHCARE | <u>20MG</u> | <u>A208049 001</u> | Jan 03, 2019 |
| <u>AB</u> | | | <u>40MG</u> | <u>A208049 002</u> | Jan 03, 2019 |
| <u>AB</u> | | | <u>60MG</u> | <u>A208049 003</u> | Jan 03, 2019 |
| <u>AB</u> | | | <u>80MG</u> | <u>A208049 004</u> | Jan 03, 2019 |

PRESCRIPTION DRUG PRODUCT LIST

LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LURASIDONE HYDROCHLORIDE

| | | | | |
|-----------|--------------------|--------------|--------------------|--------------|
| AB | | 120MG | A208049 005 | Jan 03, 2019 |
| AB | AMNEAL PHARMS CO | 20MG | A208002 001 | Jan 03, 2019 |
| AB | | 40MG | A208002 002 | Jan 03, 2019 |
| AB | | 60MG | A208002 003 | Jan 03, 2019 |
| AB | | 80MG | A208002 004 | Jan 03, 2019 |
| AB | | 120MG | A208002 005 | Jan 03, 2019 |
| AB | INVAGEN PHARMS | 20MG | A208028 001 | Jan 03, 2019 |
| AB | | 40MG | A208028 002 | Jan 03, 2019 |
| AB | | 60MG | A208028 003 | Jan 03, 2019 |
| AB | | 80MG | A208028 004 | Jan 03, 2019 |
| AB | | 120MG | A208028 005 | Jan 03, 2019 |
| AB | LUPIN LTD | 20MG | A208031 001 | Jan 03, 2019 |
| AB | | 40MG | A208031 002 | Jan 03, 2019 |
| AB | | 60MG | A208031 003 | Jan 03, 2019 |
| AB | | 80MG | A208031 004 | Jan 03, 2019 |
| AB | | 120MG | A208031 005 | Jan 03, 2019 |
| AB | SUN PHARMA GLOBAL | 20MG | A208066 001 | Jan 04, 2019 |
| AB | | 40MG | A208066 002 | Jan 04, 2019 |
| AB | | 60MG | A208066 003 | Jan 04, 2019 |
| AB | | 80MG | A208066 004 | Jan 04, 2019 |
| AB | | 120MG | A208066 005 | Jan 04, 2019 |
| AB | TORRENT PHARMS LTD | 20MG | A208055 001 | Jan 03, 2019 |
| AB | | 40MG | A208055 002 | Jan 03, 2019 |
| AB | | 80MG | A208055 003 | Jan 03, 2019 |
| AB | | 120MG | A208055 004 | Jan 03, 2019 |

LUSUTROMBOPAG

TABLET; ORAL

MULPLETA

+! SHIONOGI INC

3MG

N210923 001 Jul 31, 2018

LUTETIUM DOTATATE LU-177

SOLUTION; INTRAVENOUS

LUTATHERA

+! AAA USA INC

10mCi/ML

N208700 001 Jan 26, 2018

MACIMORELIN ACETATE

FOR SOLUTION; ORAL

MACRILEN

+! NOVO

EQ 60MG BASE/POUCH

N205598 001 Dec 20, 2017

MACITENTAN

TABLET; ORAL

OPSUMIT

+! ACTELION PHARMS LTD

10MG

N204410 001 Oct 18, 2013

MAFENIDE ACETATE

CREAM; TOPICAL

SULFAMYLON

+! MYLAN INSTITUTIONAL

EQ 85MG BASE/GM

N016763 001

FOR SOLUTION; TOPICAL

MAFENIDE ACETATE

| | | | | |
|-----------|-------------|-----------|--------------------|--------------|
| AT | NOVAST LABS | 5% | A206716 001 | Jul 31, 2017 |
| AT | PAR FORM | 5% | A201511 001 | Feb 12, 2013 |

SULFAMYLON

| | | | | |
|-----------|------------------------|-----------|--------------------|--------------|
| AT | +! MYLAN INSTITUTIONAL | 5% | N019832 003 | Jun 05, 1998 |
|-----------|------------------------|-----------|--------------------|--------------|

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

ISOLYTE S PH 7.4 IN PLASTIC CONTAINER

+! B BRAUN

30MG/100ML; 37MG/100ML; 0.82MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML; 12MG/100ML

N019696 001 Sep 29, 1989

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

ISOLYTE S IN PLASTIC CONTAINER

B BRAUN

30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML

N019711 001 Sep 29, 1989

NORMOSOL-R IN PLASTIC CONTAINER

ICU MEDICAL INC

30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML

N017586 001

PRESCRIPTION DRUG PRODUCT LIST

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER

| | | | | | | |
|---|---|-----------------|---------------------------------------------------------------|---------|-----|--|
| + | ! | BAXTER HLTHCARE | 30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML | N017378 | 001 | |
|---|---|-----------------|---------------------------------------------------------------|---------|-----|--|

PLASMA-LYTE A IN PLASTIC CONTAINER

| | | | | | | |
|---|---|-----------------|---------------------------------------------------------------|---------|-----|--------------|
| + | ! | BAXTER HLTHCARE | 30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML | N017378 | 002 | Nov 22, 1982 |
|---|---|-----------------|---------------------------------------------------------------|---------|-----|--------------|

SOLUTION; IRRIGATION

PHYSIOLYTE IN PLASTIC CONTAINER

| | | | | | | |
|--|--|---------|---------------------------------------------------------------|---------|-----|--------------|
| | | B BRAUN | 30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML | N019024 | 001 | Jun 08, 1984 |
|--|--|---------|---------------------------------------------------------------|---------|-----|--------------|

PHYSIOSOL IN PLASTIC CONTAINER

| | | | | | | |
|--|--|-----------------|---------------------------------------------------------------|---------|-----|--------------|
| | | ICU MEDICAL INC | 30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG/100ML; 502MG/100ML | N017637 | 002 | Jul 08, 1982 |
|--|--|-----------------|---------------------------------------------------------------|---------|-----|--------------|

MAGNESIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

SOLUTION; INJECTION

NORMOCARB HF 25

| | | | | | | |
|---|---|---------------|-----------------------------------------|---------|-----|--------------|
| + | ! | DIALYSIS SUPS | 0.21GM/100ML; 2.8GM/100ML; 9.07GM/100ML | N021910 | 001 | Jul 26, 2006 |
|---|---|---------------|-----------------------------------------|---------|-----|--------------|

NORMOCARB HF 35

| | | | | | | |
|---|---|---------------|-----------------------------------------|---------|-----|--------------|
| + | ! | DIALYSIS SUPS | 0.21GM/100ML; 3.97GM/100ML; 8.3GM/100ML | N021910 | 002 | Jul 26, 2006 |
|---|---|---------------|-----------------------------------------|---------|-----|--------------|

MAGNESIUM SULFATE

INJECTABLE; INJECTION

MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | | |
|-----------|---|--------------------|------------------|----------------|------------|--------------|
| <u>AP</u> | | FRESENIUS KABI USA | <u>1GM/100ML</u> | <u>A206486</u> | <u>001</u> | Mar 07, 2016 |
| <u>AP</u> | + | HOSPIRA | <u>1GM/100ML</u> | <u>N020488</u> | <u>001</u> | Jul 11, 1995 |
| <u>AP</u> | | HQ SPCLT PHARMA | <u>1GM/100ML</u> | <u>A207349</u> | <u>001</u> | Mar 02, 2016 |
| <u>AP</u> | | MYLAN LABS LTD | <u>1GM/100ML</u> | <u>A209932</u> | <u>001</u> | Sep 10, 2018 |

MAGNESIUM SULFATE IN PLASTIC CONTAINER

| | | | | | | |
|-----------|---|--------------------|------------------------------|----------------|------------|--------------|
| <u>AP</u> | | FRESENIUS KABI USA | <u>4GM/100ML (40MG/ML)</u> | <u>A206485</u> | <u>001</u> | Mar 15, 2016 |
| <u>AP</u> | | | <u>4GM/50ML (80MG/ML)</u> | <u>A206485</u> | <u>002</u> | Mar 15, 2016 |
| <u>AP</u> | | | <u>2GM/50ML (40MG/ML)</u> | <u>A206485</u> | <u>003</u> | Mar 15, 2016 |
| <u>AP</u> | | | <u>20GM/500ML (40MG/ML)</u> | <u>A206485</u> | <u>004</u> | Mar 15, 2016 |
| <u>AP</u> | | | <u>40GM/1000ML (40MG/ML)</u> | <u>A206485</u> | <u>005</u> | Mar 15, 2016 |
| <u>AP</u> | + | HOSPIRA | <u>2GM/50ML (40MG/ML)</u> | <u>N020309</u> | <u>003</u> | Jan 26, 2007 |
| <u>AP</u> | + | ! | <u>4GM/100ML (40MG/ML)</u> | <u>N020309</u> | <u>001</u> | Jun 24, 1994 |
| <u>AP</u> | + | ! | <u>4GM/50ML (80MG/ML)</u> | <u>N020309</u> | <u>002</u> | Jun 24, 1994 |
| <u>AP</u> | + | | <u>20GM/500ML (40MG/ML)</u> | <u>N020309</u> | <u>004</u> | Jan 18, 1995 |
| <u>AP</u> | + | | <u>40GM/1000ML (40MG/ML)</u> | <u>N020309</u> | <u>005</u> | Jan 18, 1995 |
| <u>AP</u> | | HQ SPCLT PHARMA | <u>2GM/50ML (40MG/ML)</u> | <u>A207350</u> | <u>001</u> | Dec 06, 2017 |
| <u>AP</u> | | | <u>4GM/100ML (40MG/ML)</u> | <u>A207350</u> | <u>002</u> | Dec 06, 2017 |
| <u>AP</u> | | | <u>4GM/50ML (80MG/ML)</u> | <u>A207350</u> | <u>003</u> | Dec 06, 2017 |
| <u>AP</u> | | | <u>20GM/500ML (40MG/ML)</u> | <u>A207350</u> | <u>004</u> | Dec 06, 2017 |
| <u>AP</u> | | | <u>40GM/1000ML (40MG/ML)</u> | <u>A207350</u> | <u>005</u> | Dec 06, 2017 |
| <u>AP</u> | | MYLAN LABS LTD | <u>2GM/50ML (40MG/ML)</u> | <u>A209911</u> | <u>001</u> | Sep 14, 2018 |
| <u>AP</u> | | | <u>4GM/100ML (40MG/ML)</u> | <u>A209911</u> | <u>002</u> | Sep 14, 2018 |

MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | | |
|---|---|---------|-----------|---------|-----|--------------|
| + | ! | HOSPIRA | 2GM/100ML | N020488 | 002 | Jul 11, 1995 |
|---|---|---------|-----------|---------|-----|--------------|

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

MAGNESIUM SULFATE

| | | | | | | |
|-----------|---|----------------------|----------------------------|----------------|------------|--------------|
| <u>AP</u> | | EXELA PHARMA SCS LLC | <u>5GM/10ML (500MG/ML)</u> | <u>A206039</u> | <u>001</u> | Dec 18, 2014 |
| <u>AP</u> | + | FRESENIUS KABI USA | <u>5GM/10ML (500MG/ML)</u> | <u>N019316</u> | <u>001</u> | Sep 08, 1986 |
| <u>AP</u> | ! | HOSPIRA | <u>5GM/10ML (500MG/ML)</u> | <u>A075151</u> | <u>001</u> | Apr 25, 2000 |
| | + | FRESENIUS KABI USA | 10GM/20ML (500MG/ML) | N019316 | 003 | Jan 29, 2016 |
| | + | | 25GM/50ML (500MG/ML) | N019316 | 004 | Jan 29, 2016 |
| | | HOSPIRA INC | 10GM/20ML (500MG/ML) | A202411 | 001 | May 14, 2015 |

MAGNESIUM SULFATE

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

MAGNESIUM SULFATE

| | | | | | | |
|---|---|--------------------|--------------------|---------|-----|--------------|
| + | ! | FRESENIUS KABI USA | 1GM/2ML (500MG/ML) | N019316 | 002 | Sep 08, 1986 |
|---|---|--------------------|--------------------|---------|-----|--------------|

MAGNESIUM SULFATE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

TIS-U-SOL

| | | | | | | |
|-----------|--|-----------------|------------------------------------------------------------------------|----------------|------------|--------------|
| <u>AT</u> | | BAXTER HLTHCARE | <u>20MG/100ML; 40MG/100ML; 6.25MG/100ML; 800MG/100ML; 8.75MG/100ML</u> | <u>N018508</u> | <u>001</u> | Feb 19, 1982 |
|-----------|--|-----------------|------------------------------------------------------------------------|----------------|------------|--------------|

TIS-U-SOL IN PLASTIC CONTAINER

| | | | | | | |
|-----------|--|-----------------|------------------------------------------------------------------------|----------------|------------|--|
| <u>AT</u> | | BAXTER HLTHCARE | <u>20MG/100ML; 40MG/100ML; 6.25MG/100ML; 800MG/100ML; 8.75MG/100ML</u> | <u>N018336</u> | <u>001</u> | |
|-----------|--|-----------------|------------------------------------------------------------------------|----------------|------------|--|

PRESCRIPTION DRUG PRODUCT LIST

MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE

POWDER; ORAL

COLPREP KIT

+! GATOR PHARMS 1.6GM/BOT;3.13GM/BOT;17.5GM/BOT N204553 001 Dec 27, 2016

SOLUTION; ORAL

SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE**AA** NOVEL LABS INC 1.6GM/BOT;3.13GM/BOT;17.5GM/BOT **A202511 001** Feb 23, 2017SUPREP BOWEL PREP KIT**AA** +! BRAINTREE LABS 1.6GM/BOT;3.13GM/BOT;17.5GM/BOT **N022372 001** Aug 05, 2010MALATHION

LOTION; TOPICAL

MALATHION**AT** SUVEN LIFE 0.5% **A091559 001** May 23, 2012OVIDE**AT** +! TARO PHARM INDS LTD 0.5% **N018613 001** Aug 02, 1982MANGANESE CHLORIDE

INJECTABLE; INJECTION

MANGANESE CHLORIDE IN PLASTIC CONTAINER

+! HOSPIRA EQ 0.1MG MANGANESE/ML N018962 001 Jun 26, 1986

MANNITOL

INJECTABLE; INJECTION

MANNITOL 10% IN PLASTIC CONTAINER**AP** B BRAUN 10GM/100ML **N020006 002** Jul 26, 1993MANNITOL 15% IN PLASTIC CONTAINER**AP** B BRAUN 15GM/100ML **N020006 003** Jul 26, 1993MANNITOL 20% IN PLASTIC CONTAINER**AP** B BRAUN 20GM/100ML **N020006 004** Jul 26, 1993**AP** ICU MEDICAL INC 20GM/100ML **N019603 004** Jan 08, 1990MANNITOL 25%**AP** FRESENIUS KABI USA 12.5GM/50ML **A080677 001****AP** HOSPIRA 12.5GM/50ML **N016269 006** Aug 25, 1994**AP** INTL MEDICATION 12.5GM/50ML **A083051 001****AP** LUITPOLD 12.5GM/50ML **A087409 001** Jan 21, 1982MANNITOL 5% IN PLASTIC CONTAINER**AP** B BRAUN 5GM/100ML **N020006 001** Jul 26, 1993OSMITROL 10% IN WATER**AP** BAXTER HLTHCARE 10GM/100ML **N013684 002**OSMITROL 10% IN WATER IN PLASTIC CONTAINER**AP** BAXTER HLTHCARE 10GM/100ML **N013684 006**OSMITROL 15% IN WATER**AP** BAXTER HLTHCARE 15GM/100ML **N013684 004**OSMITROL 15% IN WATER IN PLASTIC CONTAINER**AP** BAXTER HLTHCARE 15GM/100ML **N013684 008**OSMITROL 20% IN WATER**AP** BAXTER HLTHCARE 20GM/100ML **N013684 003**OSMITROL 20% IN WATER IN PLASTIC CONTAINER**AP** BAXTER HLTHCARE 20GM/100ML **N013684 007**OSMITROL 5% IN WATER**AP** BAXTER HLTHCARE 5GM/100ML **N013684 001**OSMITROL 5% IN WATER IN PLASTIC CONTAINER**AP** BAXTER HLTHCARE 5GM/100ML **N013684 005**

POWDER; INHALATION

ARIDOL KIT

+! PHARMAXIS LTD N/A, 5MG, 10MG, 20MG, 40MG N022368 001 Oct 05, 2010

SOLUTION; IRRIGATION

RESECTISOL IN PLASTIC CONTAINER

B BRAUN 5GM/100ML N016772 002

MANNITOL; SORBITOL

SOLUTION; IRRIGATION

SORBITOL-MANNITOL IN PLASTIC CONTAINER

+! ICU MEDICAL INC 540MG/100ML;2.7GM/100ML N018316 001

MAPROTIline HYDROCHLORIDE

TABLET; ORAL

MAPROTIline HYDROCHLORIDE

MYLAN 25MG A072285 002 Oct 03, 1988

! 50MG A072285 001 Oct 03, 1988

75MG A072285 003 Oct 03, 1988

PRESCRIPTION DRUG PRODUCT LIST

MARAVIROC

SOLUTION;ORAL

SELZENTRY

+! VIIV HLTHCARE 20MG/ML N208984 001 Nov 04, 2016

TABLET;ORAL

SELZENTRY

+ VIIV HLTHCARE 25MG N022128 003 Nov 04, 2016

+ 75MG N022128 004 Nov 04, 2016

+ 150MG N022128 001 Aug 06, 2007

+! 300MG N022128 002 Aug 06, 2007

MEBENDAZOLE

TABLET, CHEWABLE;ORAL

EMVERM

! IMPAX LABS INC 100MG A073580 001 Jan 04, 1995

MECAMYLAMINE HYDROCHLORIDE

TABLET;ORAL

MECAMYLAMINE HYDROCHLORIDE

! NEXGEN PHARMA 2.5MG A204054 001 Mar 19, 2013

MECASERMIN RECOMBINANT

INJECTABLE;SUBCUTANEOUS

INCRELEX

+! IPSEN INC 40MG/4ML (10MG/ML) N021839 001 Aug 30, 2005

MECHLORETHAMINE HYDROCHLORIDE

GEL;TOPICAL

VALCHLOR

+! HELSINN EQ 0.016% BASE N202317 001 Aug 23, 2013

MECLIZINE HYDROCHLORIDE

TABLET;ORAL

MECLIZINE HYDROCHLORIDE**AA** AMNEAL PHARMS **12.5MG** **A201451 001** Feb 23, 2011**AA** **25MG** **A201451 002** Feb 23, 2011**AA** EPIC PHARMA LLC **12.5MG** **A200294 001** Apr 13, 2012**AA** **25MG** **A200294 002** Apr 13, 2012**AA** JUBILANT CADISTA **12.5MG** **A040659 001** Jun 04, 2010**AA** **25MG** **A040659 002** Jun 04, 2010**AA** MYLAN PHARMS INC **12.5MG** **A202640 001** Sep 17, 2012**AA** **25MG** **A202640 002** Sep 17, 2012**AA** PAR PHARM **12.5MG** **A087127 001****AA** **25MG** **A087128 001****AA** SANDOZ **12.5MG** **A084843 002** May 22, 1989**AA** **25MG** **A084092 003** May 22, 1989MECLOFENAMATE SODIUM

CAPSULE;ORAL

MECLOFENAMATE SODIUM

MYLAN EQ 50MG BASE A071081 002 Sep 03, 1986

! EQ 100MG BASE A071081 001 Sep 03, 1986

MEDROXYPROGESTERONE ACETATE

INJECTABLE;INJECTION

DEPO-PROVERA**AB** +! PHARMACIA AND **150MG/ML** **N020246 001** Oct 29, 1992
UPJOHN**MEDROXYPROGESTERONE ACETATE****AB** AMPHASTAR PHARMS **150MG/ML** **A077235 001** Nov 28, 2017
INC**AB** **150MG/ML** **A077334 001** Nov 28, 2017**AB** MYLAN LABS LTD **150MG/ML** **A210227 001** Oct 12, 2018**AB** TEVA PHARMS USA **150MG/ML** **A076553 001** Jul 28, 2004

DEPO-PROVERA

+! PHARMACIA AND 400MG/ML N012541 003
UPJOHN

INJECTABLE;SUBCUTANEOUS

DEPO-SUBQ PROVERA 104

+! PHARMACIA AND 104MG/0.65ML N021583 001 Dec 17, 2004
UPJOHN

TABLET;ORAL

MEDROXYPROGESTERONE ACETATE**AB** BARR **2.5MG** **A040159 001** Aug 09, 1996**AB** **5MG** **A040159 002** Aug 09, 1996**AB** **10MG** **A040159 003** Aug 09, 1996

PRESCRIPTION DRUG PRODUCT LIST

MEDROXYPROGESTERONE ACETATE

TABLET; ORAL

PROVERA

| | | | | | | |
|-----------|---|-------------------------|--------------|----------------|------------|--|
| <u>AB</u> | + | PHARMACIA AND UPJOHN | <u>2.5MG</u> | <u>N011839</u> | <u>001</u> | |
| <u>AB</u> | + | | <u>5MG</u> | <u>N011839</u> | <u>003</u> | |
| <u>AB</u> | + | ! | <u>10MG</u> | <u>N011839</u> | <u>004</u> | |

MEFENAMIC ACID

CAPSULE; ORAL

MEFENAMIC ACID

| | | | | | | |
|-----------|--|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | BELCHER PHARMS LLC | <u>250MG</u> | <u>A091608</u> | <u>001</u> | Jun 02, 2014 |
| <u>AB</u> | | BRECKENRIDGE PHARM | <u>250MG</u> | <u>A090359</u> | <u>001</u> | Feb 05, 2013 |
| <u>AB</u> | | LUPIN LTD | <u>250MG</u> | <u>A091322</u> | <u>001</u> | Jul 22, 2011 |
| <u>AB</u> | | MICRO LABS | <u>250MG</u> | <u>A090562</u> | <u>001</u> | Nov 19, 2010 |

PONSTEL

| | | | | | | |
|-----------|---|---|--------------|--------------|----------------|------------|
| <u>AB</u> | + | ! | SHIONOGI INC | <u>250MG</u> | <u>N015034</u> | <u>003</u> |
|-----------|---|---|--------------|--------------|----------------|------------|

MEFLOQUINE HYDROCHLORIDE

TABLET; ORAL

MEFLOQUINE HYDROCHLORIDE

| | | | | | | |
|-----------|---|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ! | BARR | <u>250MG</u> | <u>A076392</u> | <u>001</u> | Dec 29, 2003 |
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>250MG</u> | <u>A076523</u> | <u>001</u> | Oct 01, 2004 |

MEGESTROL ACETATE

SUSPENSION; ORAL

MEGACE ES

| | | | | | | | |
|-----------|---|---|-----------------|-----------------|----------------|------------|--------------|
| <u>AB</u> | + | ! | ENDO PHARMS INC | <u>125MG/ML</u> | <u>N021778</u> | <u>001</u> | Jul 05, 2005 |
|-----------|---|---|-----------------|-----------------|----------------|------------|--------------|

MEGESTROL ACETATE

| | | | | | | |
|-----------|---|-------------------------|-----------------|----------------|------------|--------------|
| <u>AB</u> | | BRECKENRIDGE PHARM | <u>125MG/ML</u> | <u>A204688</u> | <u>001</u> | Dec 01, 2017 |
| <u>AB</u> | | HI-TECH PHARMACAL | <u>40MG/ML</u> | <u>A203960</u> | <u>001</u> | Jun 09, 2017 |
| <u>AB</u> | | PAR PHARM | <u>40MG/ML</u> | <u>A075671</u> | <u>001</u> | Jul 25, 2001 |
| <u>AB</u> | | TEVA PHARMS | <u>40MG/ML</u> | <u>A075681</u> | <u>001</u> | May 05, 2003 |
| <u>AB</u> | | TWI PHARMS | <u>125MG/ML</u> | <u>A203139</u> | <u>001</u> | Aug 27, 2014 |
| <u>AB</u> | ! | WEST-WARD PHARMS INT | <u>40MG/ML</u> | <u>A075997</u> | <u>001</u> | Feb 15, 2002 |
| <u>AB</u> | | WOCKHARDT BIO AG | <u>40MG/ML</u> | <u>A076721</u> | <u>001</u> | Nov 01, 2004 |

TABLET; ORAL

MEGESTROL ACETATE

| | | | | | | |
|-----------|---|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | BARR | <u>20MG</u> | <u>A074621</u> | <u>002</u> | Aug 16, 1996 |
| <u>AB</u> | | | <u>40MG</u> | <u>A074621</u> | <u>001</u> | Nov 30, 1995 |
| <u>AB</u> | | PAR PHARM | <u>20MG</u> | <u>A072422</u> | <u>001</u> | Aug 08, 1988 |
| <u>AB</u> | ! | | <u>40MG</u> | <u>A072423</u> | <u>001</u> | Aug 08, 1988 |
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>20MG</u> | <u>A074458</u> | <u>001</u> | Sep 29, 1995 |
| <u>AB</u> | | | <u>40MG</u> | <u>A074458</u> | <u>002</u> | Sep 29, 1995 |

MELOXICAM

CAPSULE; ORAL

VIVLODEX

| | | | | | |
|---|------------------|------|---------|-----|--------------|
| + | IROKO PHARMS LLC | 5MG | N207233 | 001 | Oct 22, 2015 |
| + | ! | 10MG | N207233 | 002 | Oct 22, 2015 |

TABLET; ORAL

MELOXICAM

| | | | | | | |
|-----------|--|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | APOTEX INC | <u>7.5MG</u> | <u>A077882</u> | <u>001</u> | Jul 20, 2006 |
| <u>AB</u> | | | <u>15MG</u> | <u>A077882</u> | <u>002</u> | Jul 20, 2006 |
| <u>AB</u> | | AUROBINDO PHARMA | <u>7.5MG</u> | <u>A078008</u> | <u>001</u> | Oct 02, 2006 |
| <u>AB</u> | | | <u>15MG</u> | <u>A078008</u> | <u>002</u> | Oct 02, 2006 |
| <u>AB</u> | | BRECKENRIDGE PHARM | <u>7.5MG</u> | <u>A077920</u> | <u>001</u> | Jul 19, 2006 |
| <u>AB</u> | | | <u>15MG</u> | <u>A077920</u> | <u>002</u> | Jul 19, 2006 |
| <u>AB</u> | | CIPLA | <u>7.5MG</u> | <u>A077929</u> | <u>001</u> | Jul 19, 2006 |
| <u>AB</u> | | | <u>15MG</u> | <u>A077929</u> | <u>002</u> | Jul 19, 2006 |
| <u>AB</u> | | DR REDDYS LABS INC | <u>7.5MG</u> | <u>A077931</u> | <u>001</u> | Jul 25, 2006 |
| <u>AB</u> | | | <u>15MG</u> | <u>A077931</u> | <u>002</u> | Jul 25, 2006 |
| <u>AB</u> | | GLENMARK GENERICS | <u>7.5MG</u> | <u>A077932</u> | <u>001</u> | Jul 19, 2006 |
| <u>AB</u> | | | <u>15MG</u> | <u>A077932</u> | <u>002</u> | Jul 19, 2006 |
| <u>AB</u> | | LUPIN PHARMS | <u>7.5MG</u> | <u>A077944</u> | <u>001</u> | Jul 19, 2006 |
| <u>AB</u> | | | <u>15MG</u> | <u>A077944</u> | <u>002</u> | Jul 19, 2006 |
| <u>AB</u> | | PURACAP PHARM | <u>7.5MG</u> | <u>A077938</u> | <u>001</u> | Jul 19, 2006 |
| <u>AB</u> | | | <u>15MG</u> | <u>A077938</u> | <u>002</u> | Jul 19, 2006 |
| <u>AB</u> | | STRIDES PHARMA | <u>7.5MG</u> | <u>A077928</u> | <u>001</u> | May 13, 2009 |
| <u>AB</u> | | | <u>15MG</u> | <u>A077928</u> | <u>002</u> | May 13, 2009 |
| <u>AB</u> | | TARO | <u>7.5MG</u> | <u>A078102</u> | <u>001</u> | Nov 07, 2006 |
| <u>AB</u> | | | <u>15MG</u> | <u>A078102</u> | <u>002</u> | Nov 07, 2006 |

PRESCRIPTION DRUG PRODUCT LIST

MELOXICAM

TABLET; ORAL

MELOXICAM

| | | | | | |
|-----------|------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | TEVA PHARMS | <u>7.5MG</u> | <u>A077936</u> | <u>001</u> | Jul 19, 2006 |
| <u>AB</u> | | <u>15MG</u> | <u>A077936</u> | <u>002</u> | Jul 19, 2006 |
| <u>AB</u> | UNICHEM | <u>7.5MG</u> | <u>A077927</u> | <u>001</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>15MG</u> | <u>A077927</u> | <u>002</u> | Dec 20, 2006 |
| <u>AB</u> | YUNG SHIN PHARM | <u>7.5MG</u> | <u>A077918</u> | <u>001</u> | Dec 07, 2006 |
| <u>AB</u> | | <u>15MG</u> | <u>A077918</u> | <u>002</u> | Dec 07, 2006 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>7.5MG</u> | <u>A077921</u> | <u>001</u> | Jul 19, 2006 |
| <u>AB</u> | | <u>15MG</u> | <u>A077921</u> | <u>002</u> | Jul 19, 2006 |

MOBIC

| | | | | | |
|-----------|--------------------------------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + BOEHRINGER INGELHEIM | <u>7.5MG</u> | <u>N020938</u> | <u>001</u> | Apr 13, 2000 |
| <u>AB</u> | +! | <u>15MG</u> | <u>N020938</u> | <u>002</u> | Aug 23, 2000 |
| | TABLET, ORALLY DISINTEGRATING; ORAL QMIIZ ODT | | | | |
| | + TERSERA THERAPS LLC | 7.5MG | N211210 | 001 | Oct 19, 2018 |
| | +! | 15MG | N211210 | 002 | Oct 19, 2018 |

MELPHALAN

TABLET; ORAL

ALKERAN

| | | | | | |
|-----------|-----------------------------------|------------|----------------|------------|--------------|
| <u>AB</u> | +! APOTEX INC | <u>2MG</u> | <u>N014691</u> | <u>002</u> | |
| <u>AB</u> | <u>MELPHALAN</u> ALVOGEN MALTA | <u>2MG</u> | <u>A207809</u> | <u>001</u> | Mar 22, 2017 |

MELPHALAN HYDROCHLORIDE

INJECTABLE; INJECTION

MELPHALAN HYDROCHLORIDE

| | | | | | |
|-----------|--------------------------------|--------------------------|----------------|------------|--------------|
| <u>AP</u> | ACTAVIS LLC | <u>EQ 50MG BASE/VIAL</u> | <u>A206018</u> | <u>001</u> | Dec 19, 2016 |
| <u>AP</u> | DR REDDYS LABS LTD | <u>EQ 50MG BASE/VIAL</u> | <u>A203655</u> | <u>001</u> | Dec 08, 2017 |
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 50MG BASE/VIAL</u> | <u>A203393</u> | <u>001</u> | Dec 22, 2017 |
| <u>AP</u> | ! MYLAN INSTITUTIONAL | <u>EQ 50MG BASE/VIAL</u> | <u>A090270</u> | <u>001</u> | Jun 09, 2009 |
| <u>AP</u> | PAR STERILE PRODUCTS | <u>EQ 50MG BASE/VIAL</u> | <u>A204773</u> | <u>001</u> | Aug 22, 2016 |
| <u>AP</u> | SAGENT PHARMS | <u>EQ 50MG BASE/VIAL</u> | <u>A201379</u> | <u>001</u> | Feb 28, 2017 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>EQ 50MG BASE/VIAL</u> | <u>A090303</u> | <u>001</u> | Oct 28, 2010 |
| | POWDER; INTRAVENOUS EVOMELA | | | | |
| | +! SPECTRUM PHARMS | EQ 50MG BASE/VIAL | N207155 | 001 | Mar 10, 2016 |

MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MEMANTINE HYDROCHLORIDE

| | | | | | |
|-----------|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | AMNEAL PHARMS | <u>7MG</u> | <u>A205825</u> | <u>001</u> | Oct 12, 2016 |
| <u>AB</u> | | <u>14MG</u> | <u>A205825</u> | <u>002</u> | Oct 12, 2016 |
| <u>AB</u> | | <u>21MG</u> | <u>A205825</u> | <u>003</u> | Oct 12, 2016 |
| <u>AB</u> | | <u>28MG</u> | <u>A205825</u> | <u>004</u> | Oct 12, 2016 |
| <u>AB</u> | ANCHEN PHARMS | <u>7MG</u> | <u>A205784</u> | <u>001</u> | Jun 09, 2017 |
| <u>AB</u> | | <u>14MG</u> | <u>A205784</u> | <u>002</u> | Jun 09, 2017 |
| <u>AB</u> | | <u>21MG</u> | <u>A205784</u> | <u>003</u> | Jun 09, 2017 |
| <u>AB</u> | | <u>28MG</u> | <u>A205784</u> | <u>004</u> | Jun 09, 2017 |
| <u>AB</u> | APOTEX INC | <u>7MG</u> | <u>A206135</u> | <u>001</u> | Nov 22, 2016 |
| <u>AB</u> | | <u>14MG</u> | <u>A206135</u> | <u>002</u> | Nov 22, 2016 |
| <u>AB</u> | | <u>21MG</u> | <u>A206135</u> | <u>003</u> | Nov 22, 2016 |
| <u>AB</u> | | <u>28MG</u> | <u>A206135</u> | <u>004</u> | Nov 22, 2016 |
| <u>AB</u> | LUPIN LTD | <u>7MG</u> | <u>A206028</u> | <u>001</u> | Sep 28, 2016 |
| <u>AB</u> | | <u>14MG</u> | <u>A206028</u> | <u>002</u> | Sep 28, 2016 |
| <u>AB</u> | | <u>21MG</u> | <u>A206028</u> | <u>003</u> | Sep 28, 2016 |
| <u>AB</u> | | <u>28MG</u> | <u>A206028</u> | <u>004</u> | Sep 28, 2016 |
| <u>AB</u> | MYLAN PHARMS INC | <u>7MG</u> | <u>A206032</u> | <u>001</u> | Sep 28, 2016 |
| <u>AB</u> | | <u>14MG</u> | <u>A206032</u> | <u>002</u> | Sep 28, 2016 |
| <u>AB</u> | | <u>21MG</u> | <u>A206032</u> | <u>003</u> | Sep 28, 2016 |
| <u>AB</u> | | <u>28MG</u> | <u>A206032</u> | <u>004</u> | Sep 28, 2016 |
| <u>AB</u> | SUN PHARMA GLOBAL | <u>7MG</u> | <u>A205905</u> | <u>001</u> | Sep 28, 2016 |
| <u>AB</u> | | <u>14MG</u> | <u>A205905</u> | <u>002</u> | Sep 28, 2016 |
| <u>AB</u> | | <u>21MG</u> | <u>A205905</u> | <u>003</u> | Sep 28, 2016 |
| <u>AB</u> | | <u>28MG</u> | <u>A205905</u> | <u>004</u> | Sep 28, 2016 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>7MG</u> | <u>A203293</u> | <u>001</u> | Aug 03, 2017 |
| <u>AB</u> | | <u>14MG</u> | <u>A203293</u> | <u>002</u> | Aug 03, 2017 |
| <u>AB</u> | | <u>21MG</u> | <u>A203293</u> | <u>003</u> | Aug 03, 2017 |
| <u>AB</u> | | <u>28MG</u> | <u>A203293</u> | <u>004</u> | Aug 03, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

NAMENDA XR

| | | | | | | |
|-----------|----|-----------------|-------------|----------------|------------|--------------|
| <u>AB</u> | + | FOREST LABS LLC | <u>7MG</u> | <u>N022525</u> | <u>001</u> | Jun 21, 2010 |
| <u>AB</u> | + | | <u>14MG</u> | <u>N022525</u> | <u>002</u> | Jun 21, 2010 |
| <u>AB</u> | + | | <u>21MG</u> | <u>N022525</u> | <u>003</u> | Jun 21, 2010 |
| <u>AB</u> | +! | | <u>28MG</u> | <u>N022525</u> | <u>004</u> | Jun 21, 2010 |

SOLUTION;ORAL

MEMANTINE HYDROCHLORIDE

| | | | | | | |
|-----------|---|---------------------|---------------|----------------|------------|--------------|
| <u>AA</u> | | APOTEX INC | <u>2MG/ML</u> | <u>A209955</u> | <u>001</u> | Feb 09, 2018 |
| <u>AA</u> | ! | BIO-PHARM INC | <u>2MG/ML</u> | <u>A205446</u> | <u>001</u> | Dec 07, 2015 |
| <u>AA</u> | | LANNETT CO INC | <u>2MG/ML</u> | <u>A204033</u> | <u>001</u> | Oct 13, 2015 |
| <u>AA</u> | | MACLEODS PHARMS LTD | <u>2MG/ML</u> | <u>A202790</u> | <u>001</u> | Oct 13, 2015 |

TABLET;ORAL

MEMANTINE HYDROCHLORIDE

| | | | | | | |
|----------------|----|----------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | AJANTA PHARMA LTD | <u>5MG</u> | <u>A206528</u> | <u>001</u> | Nov 30, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A206528</u> | <u>002</u> | Nov 30, 2015 |
| <u>AB</u> | | ALEMBIC PHARMS LTD | <u>5MG</u> | <u>A200891</u> | <u>001</u> | Oct 13, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A200891</u> | <u>002</u> | Oct 13, 2015 |
| <u>AB</u> | | AMNEAL PHARMS | <u>5MG</u> | <u>A090041</u> | <u>001</u> | Apr 10, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090041</u> | <u>002</u> | Apr 10, 2015 |
| <u>AB</u> | | APOTEX INC | <u>5MG</u> | <u>A090244</u> | <u>001</u> | Jul 11, 2018 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090244</u> | <u>002</u> | Jul 11, 2018 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>5MG</u> | <u>A203175</u> | <u>001</u> | Oct 13, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A203175</u> | <u>002</u> | Oct 13, 2015 |
| <u>AB</u> | | CSPC OUYI PHARM CO | <u>5MG</u> | <u>A209527</u> | <u>001</u> | May 07, 2018 |
| <u>AB</u> | | | <u>10MG</u> | <u>A209527</u> | <u>002</u> | May 07, 2018 |
| <u>AB</u> | | DR REDDYS LABS LTD | <u>5MG</u> | <u>A090048</u> | <u>001</u> | Apr 14, 2010 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090048</u> | <u>002</u> | Apr 14, 2010 |
| <u>AB</u> | | JUBILANT GENERICS | <u>5MG</u> | <u>A091585</u> | <u>001</u> | Oct 13, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A091585</u> | <u>002</u> | Oct 13, 2015 |
| <u>AB</u> | | LANNETT CO INC | <u>5MG</u> | <u>A207236</u> | <u>001</u> | Nov 10, 2016 |
| <u>AB</u> | | | <u>10MG</u> | <u>A207236</u> | <u>002</u> | Nov 10, 2016 |
| <u>AB</u> | | LUPIN LTD | <u>5MG</u> | <u>A090051</u> | <u>001</u> | Apr 10, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090051</u> | <u>002</u> | Apr 10, 2015 |
| <u>AB</u> | | MACLEODS PHARMS LTD | <u>5MG</u> | <u>A202840</u> | <u>001</u> | Oct 13, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A202840</u> | <u>002</u> | Oct 13, 2015 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>5MG</u> | <u>A079225</u> | <u>001</u> | Jan 30, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A079225</u> | <u>002</u> | Jan 30, 2015 |
| <u>AB</u> | | PURACAP PHARM LLC | <u>5MG</u> | <u>A206855</u> | <u>001</u> | Nov 17, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A206855</u> | <u>002</u> | Nov 17, 2015 |
| <u>AB</u> | | STRIDES PHARMA | <u>5MG</u> | <u>A202350</u> | <u>001</u> | May 23, 2017 |
| <u>AB</u> | | | <u>10MG</u> | <u>A202350</u> | <u>002</u> | May 23, 2017 |
| <u>AB</u> | | SUN PHARMA GLOBAL | <u>5MG</u> | <u>A090058</u> | <u>001</u> | May 05, 2010 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090058</u> | <u>002</u> | May 05, 2010 |
| <u>AB</u> | | TEVA PHARMS | <u>5MG</u> | <u>A090052</u> | <u>001</u> | Oct 25, 2011 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090052</u> | <u>002</u> | Oct 25, 2011 |
| <u>AB</u> | | TORRENT PHARMS LTD | <u>5MG</u> | <u>A200155</u> | <u>001</u> | Oct 13, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A200155</u> | <u>002</u> | Oct 13, 2015 |
| <u>AB</u> | | UNICHEM LABS LTD | <u>5MG</u> | <u>A200022</u> | <u>001</u> | Oct 13, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A200022</u> | <u>002</u> | Oct 13, 2015 |
| <u>AB</u> | | UPSHER SMITH LABS | <u>5MG</u> | <u>A090043</u> | <u>001</u> | Jul 31, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090043</u> | <u>002</u> | Jul 31, 2015 |
| <u>AB</u> | | WOCKHARDT LTD | <u>5MG</u> | <u>A090073</u> | <u>001</u> | Sep 04, 2015 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090073</u> | <u>002</u> | Sep 04, 2015 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>5MG</u> | <u>A090961</u> | <u>001</u> | Jul 10, 2017 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090961</u> | <u>002</u> | Jul 10, 2017 |
| <u>NAMENDA</u> | | | | | | |
| <u>AB</u> | + | ALLERGAN SALES LLC | <u>5MG</u> | <u>N021487</u> | <u>001</u> | Oct 16, 2003 |
| <u>AB</u> | +! | | <u>10MG</u> | <u>N021487</u> | <u>002</u> | Oct 16, 2003 |

MENOTROPINS (FSH;LH)

INJECTABLE;SUBCUTANEOUS

MENOPUR

| | | | | | | |
|---|---|---------|-----------------------|---------|-----|--------------|
| + | ! | FERRING | 75 IU/VIAL;75 IU/VIAL | N021663 | 001 | Oct 29, 2004 |
|---|---|---------|-----------------------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DEMEROL

| | | | | | | |
|-----------|---|---|---------|-----------------|----------------|------------|
| <u>AP</u> | + | ! | HOSPIRA | <u>25MG/ML</u> | <u>N021171</u> | <u>001</u> |
| <u>AP</u> | + | ! | | <u>50MG/ML</u> | <u>N021171</u> | <u>002</u> |
| <u>AP</u> | + | ! | | <u>75MG/ML</u> | <u>N021171</u> | <u>003</u> |
| <u>AP</u> | + | ! | | <u>100MG/ML</u> | <u>N021171</u> | <u>004</u> |

MEPERIDINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|--|-------------------------|-----------------|----------------|------------|
| <u>AP</u> | | | WEST-WARD PHARMS INT | <u>25MG/ML</u> | <u>A080445</u> | <u>001</u> |
| <u>AP</u> | | | | <u>25MG/ML</u> | <u>A080455</u> | <u>007</u> |
| <u>AP</u> | | | | <u>50MG/ML</u> | <u>A080445</u> | <u>002</u> |
| <u>AP</u> | | | | <u>50MG/ML</u> | <u>A080455</u> | <u>008</u> |
| <u>AP</u> | | | | <u>75MG/ML</u> | <u>A080445</u> | <u>003</u> |
| <u>AP</u> | | | | <u>75MG/ML</u> | <u>A080455</u> | <u>009</u> |
| <u>AP</u> | | | | <u>100MG/ML</u> | <u>A080445</u> | <u>004</u> |
| <u>AP</u> | | | | <u>100MG/ML</u> | <u>A080455</u> | <u>010</u> |

MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE

| | | | | | | | |
|---|--|--|-------------------------|---------|---------|-----|--------------|
| ! | | | WEST-WARD PHARMS INT | 10MG/ML | A081002 | 001 | Jul 30, 1993 |
|---|--|--|-------------------------|---------|---------|-----|--------------|

SYRUP; ORAL

MEPERIDINE HYDROCHLORIDE

| | | | | | | | |
|---|--|--|-------------------------|----------|---------|-----|--------------|
| ! | | | WEST-WARD PHARMS INT | 50MG/5ML | A088744 | 001 | Jan 30, 1985 |
|---|--|--|-------------------------|----------|---------|-----|--------------|

TABLET; ORAL

DEMEROL

| | | | | | | |
|-----------|---|---|-------------------|--------------|----------------|------------|
| <u>AA</u> | + | ! | US PHARM HOLDINGS | <u>50MG</u> | <u>N005010</u> | <u>001</u> |
| <u>AA</u> | + | ! | | <u>100MG</u> | <u>N005010</u> | <u>004</u> |

MEPERIDINE HYDROCHLORIDE

| | | | | | | | |
|-----------|--|--|-------------------------|--------------|----------------|------------|--------------|
| <u>AA</u> | | | EPIC PHARMA | <u>50MG</u> | <u>A040331</u> | <u>001</u> | May 28, 1999 |
| <u>AA</u> | | | | <u>100MG</u> | <u>A040331</u> | <u>002</u> | May 28, 1999 |
| <u>AA</u> | | | MIKART | <u>50MG</u> | <u>A040893</u> | <u>001</u> | Jun 24, 2009 |
| <u>AA</u> | | | | <u>100MG</u> | <u>A040893</u> | <u>003</u> | Jun 24, 2009 |
| <u>AA</u> | | | SPECGX LLC | <u>50MG</u> | <u>A040352</u> | <u>001</u> | Jun 13, 2000 |
| <u>AA</u> | | | | <u>100MG</u> | <u>A040352</u> | <u>002</u> | Jun 13, 2000 |
| <u>AA</u> | | | SUN PHARM INDS INC | <u>50MG</u> | <u>A040446</u> | <u>001</u> | Aug 08, 2002 |
| <u>AA</u> | | | | <u>100MG</u> | <u>A040446</u> | <u>002</u> | Aug 08, 2002 |
| <u>AA</u> | | | VINTAGE PHARMS | <u>50MG</u> | <u>A040191</u> | <u>001</u> | Dec 17, 1998 |
| <u>AA</u> | | | | <u>100MG</u> | <u>A040191</u> | <u>002</u> | Dec 17, 1998 |
| <u>AA</u> | | | WEST-WARD PHARMS INT | <u>50MG</u> | <u>A040110</u> | <u>001</u> | Mar 12, 1997 |
| <u>AA</u> | | | | <u>100MG</u> | <u>A040110</u> | <u>002</u> | Mar 12, 1997 |
| | | | MIKART | 75MG | A040893 | 002 | Jun 24, 2009 |
| | | | | 150MG | A040893 | 004 | Jun 24, 2009 |

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CARBOCAINE

| | | | | | | |
|-----------|---|---|---------|-------------|----------------|------------|
| <u>AP</u> | + | ! | HOSPIRA | <u>1%</u> | <u>N012250</u> | <u>001</u> |
| <u>AP</u> | + | ! | | <u>1.5%</u> | <u>N012250</u> | <u>005</u> |
| <u>AP</u> | + | ! | | <u>2%</u> | <u>N012250</u> | <u>002</u> |

POLOCAINE

| | | | | | | | |
|-----------|--|--|--------------------|-----------|----------------|------------|--------------|
| <u>AP</u> | | | FRESENIUS KABI USA | <u>1%</u> | <u>A089407</u> | <u>001</u> | Dec 01, 1986 |
| <u>AP</u> | | | | <u>2%</u> | <u>A089410</u> | <u>001</u> | Dec 01, 1986 |

POLOCAINE-MPF

| | | | | | | | |
|-----------|--|--|--------------------|-------------|----------------|------------|--------------|
| <u>AP</u> | | | FRESENIUS KABI USA | <u>1%</u> | <u>A089406</u> | <u>001</u> | Dec 01, 1986 |
| <u>AP</u> | | | | <u>1.5%</u> | <u>A089408</u> | <u>001</u> | Dec 01, 1986 |
| <u>AP</u> | | | | <u>2%</u> | <u>A089409</u> | <u>001</u> | Dec 01, 1986 |

SCANDONEST PLAIN

| | | | | | | | |
|-----------|---|--|---------|-----------|----------------|------------|--------------|
| <u>AP</u> | ! | | DEPROCO | <u>3%</u> | <u>A088387</u> | <u>001</u> | Oct 10, 1984 |
|-----------|---|--|---------|-----------|----------------|------------|--------------|

MEPROBAMATE

TABLET; ORAL

MEPROBAMATE

| | | | | | | | |
|-----------|---|--|--------------------|--------------|----------------|------------|--------------|
| <u>AA</u> | | | ALEMBIC PHARMS LTD | <u>200MG</u> | <u>A090122</u> | <u>001</u> | Feb 18, 2009 |
| <u>AA</u> | | | | <u>400MG</u> | <u>A090122</u> | <u>002</u> | Feb 18, 2009 |
| <u>AA</u> | | | INVAGEN PHARMS | <u>200MG</u> | <u>A040797</u> | <u>001</u> | Feb 27, 2008 |
| <u>AA</u> | | | | <u>400MG</u> | <u>A040797</u> | <u>002</u> | Feb 27, 2008 |
| <u>AA</u> | ! | | WATSON LABS | <u>200MG</u> | <u>A083304</u> | <u>001</u> | |
| <u>AA</u> | ! | | | <u>400MG</u> | <u>A083308</u> | <u>001</u> | |

PRESCRIPTION DRUG PRODUCT LIST

MERCAPTOPURINE

SUSPENSION; ORAL

PURIXAN

+! NOVA LABS LTD 20MG/ML

N205919 001 Apr 28, 2014

TABLET; ORAL

MERCAPTOPURINE**AB** DR REDDYS LABS SA **50MG****A040461 001** Feb 11, 2004**AB** MYLAN **50MG****A040594 001** Jul 01, 2005**AB** ! WEST-WARD PHARMS **50MG****A040528 001** Feb 13, 2004

INT

PURINETHOL**AB** + STASON PHARMS **50MG****N009053 002**MEROPENEM

INJECTABLE; INJECTION

MEROPENEM**AP** ACS DOBFAR **500MG/VIAL****A091404 001** Oct 26, 2011**AP** **1GM/VIAL****A091404 002** Oct 26, 2011**AP** ACS DOBFAR SPA **500MG/VIAL****A204139 001** Jun 09, 2016**AP** **1GM/VIAL****A204139 002** Jun 09, 2016**AP** AMNEAL PHARMS **500MG/VIAL****A205883 001** Apr 12, 2016**AP** **1GM/VIAL****A205883 002** Apr 12, 2016**AP** AUROBINDO PHARMA **500MG/VIAL****A205835 001** Mar 27, 2017

LTD

AP **1GM/VIAL****A205835 002** Mar 27, 2017**AP** DAEWOONG PHARM CO **500MG/VIAL****A204854 001** Dec 18, 2015**AP** **1GM/VIAL****A204854 002** Dec 18, 2015**AP** GLAND PHARMA LTD **500MG/VIAL****A206141 001** Jun 08, 2016**AP** **1GM/VIAL****A206141 002** Jun 08, 2016**AP** HOSPIRA INC **500MG/VIAL****A090940 001** Jun 22, 2010**AP** **1GM/VIAL****A090940 002** Jun 22, 2010**AP** SAVIOR LIFETEC CORP **500MG/VIAL****A206086 001** Apr 19, 2016**AP** **1GM/VIAL****A206086 002** Apr 19, 2016MERREM**AP** +! PFIZER **500MG/VIAL****N050706 003** Jun 21, 1996**AP** +! **1GM/VIAL****N050706 001** Jun 21, 1996

POWDER; INTRAVENOUS

MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER

B BRAUN MEDICAL INC 500MG/VIAL

N202106 001 Apr 30, 2015

1GM/VIAL

N202106 002 Apr 30, 2015

MEROPENEM; VABORBACTAM

POWDER; INTRAVENOUS

VABOMERE

+! REMPEX PHARMS 1GM/VIAL; 1GM/VIAL

N209776 001 Aug 29, 2017

MESALAMINE

CAPSULE, DELAYED RELEASE; ORAL

DELZICOL

+! APIL 400MG

N204412 001 Feb 01, 2013

CAPSULE, EXTENDED RELEASE; ORAL

APRISO

+! VALEANT PHARMS INTL 375MG

N022301 001 Oct 31, 2008

PENTASA

+ SHIRE 250MG

N020049 001 May 10, 1993

+! 500MG

N020049 002 Jul 08, 2004

ENEMA; RECTAL

MESALAMINE**AB** PERRIGO ISRAEL **4GM/60ML****A076751 001** Sep 17, 2004ROWASA**AB** +! MYLAN SPECIALITY LP **4GM/60ML****N019618 001** Dec 24, 1987SFROWASA**AB** + MYLAN SPECIALITY LP **4GM/60ML****N019618 002** Jun 20, 2008

SUPPOSITORY; RECTAL

CANASA**AB** +! ALLERGAN SALES LLC **1GM****N021252 002** Nov 05, 2004MESALAMINE**AB** MYLAN PHARMS INC **1GM****A204354 001** Nov 24, 2015

TABLET, DELAYED RELEASE; ORAL

ASACOL HD**AB** +! APIL **800MG****N021830 001** May 29, 2008LIALDA**AB** +! SHIRE **1.2GM****N022000 001** Jan 16, 2007

PRESCRIPTION DRUG PRODUCT LIST

MESALAMINE

TABLET, DELAYED RELEASE;ORAL

MESALAMINE

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ACTAVIS LABS FL | <u>1.2GM</u> | <u>A203817</u> | <u>001</u> | Mar 23, 2018 |
| <u>AB</u> | MYLAN PHARMS INC | <u>1.2GM</u> | <u>A203574</u> | <u>001</u> | Nov 20, 2018 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>800MG</u> | <u>A203286</u> | <u>001</u> | Jul 21, 2017 |
| <u>AB</u> | | <u>1.2GM</u> | <u>A091640</u> | <u>001</u> | Jun 05, 2017 |

MESNA

INJECTABLE; INTRAVENOUS

MESNA

| | | | | | |
|-----------|-------------------------|-----------------|----------------|------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>100MG/ML</u> | <u>A075811</u> | <u>001</u> | Apr 26, 2001 |
| <u>AP</u> | GLAND PHARMA LTD | <u>100MG/ML</u> | <u>A206992</u> | <u>001</u> | Dec 18, 2017 |
| <u>AP</u> | SAGENT PHARMS | <u>100MG/ML</u> | <u>A090913</u> | <u>001</u> | Apr 13, 2010 |
| <u>AP</u> | TEVA PHARMS USA | <u>100MG/ML</u> | <u>A075764</u> | <u>001</u> | Apr 27, 2001 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>100MG/ML</u> | <u>A075739</u> | <u>001</u> | Jan 09, 2004 |

MESNEX

| | | | | | | |
|-----------|------------|-----------------|-----------------|----------------|------------|--------------|
| <u>AP</u> | <u>+</u> ! | BAXTER HLTHCARE | <u>100MG/ML</u> | <u>N019884</u> | <u>001</u> | Dec 30, 1988 |
| | | TABLET;ORAL | | | | |
| | | MESNEX | | | | |
| | <u>+</u> ! | BAXTER HLTHCARE | 400MG | N020855 | 001 | Mar 21, 2002 |

MESTRANOL; NORETHINDRONE

TABLET;ORAL-28

NORINYL 1+50 28-DAY

+! ACTAVIS LABS UT INC 0.05MG;1MG N016659 001METAPROTERENOL SULFATE

SYRUP;ORAL

METAPROTERENOL SULFATE

! LANNETT CO INC 10MG/5ML A073632 001 Jul 22, 1992

TABLET;ORAL

METAPROTERENOL SULFATE

PAR PHARM 10MG A072024 001 Jun 28, 1988

! 20MG A072025 001 Jun 28, 1988

METAXALONE

TABLET;ORAL

METAXALONE

| | | | | | |
|-----------|---------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ACTAVIS LABS FL INC | <u>800MG</u> | <u>A203695</u> | <u>001</u> | Jun 15, 2017 |
| <u>AB</u> | AMNEAL PHARMS | <u>800MG</u> | <u>A203399</u> | <u>001</u> | Jun 21, 2013 |
| <u>AB</u> | LANNETT CO INC | <u>800MG</u> | <u>A204770</u> | <u>001</u> | Nov 22, 2016 |
| <u>AB</u> | RISING PHARMS | <u>800MG</u> | <u>A208774</u> | <u>001</u> | Sep 24, 2018 |
| <u>AB</u> | SANDOZ | <u>800MG</u> | <u>A040445</u> | <u>001</u> | Mar 31, 2010 |
| <u>AB</u> | SCIEGEN PHARMS INC | <u>800MG</u> | <u>A207466</u> | <u>001</u> | Aug 31, 2017 |

SKELAXIN

| | | | | | | |
|-----------|------------|-------------|--------------|----------------|------------|--------------|
| <u>AB</u> | <u>+</u> ! | KING PHARMS | <u>800MG</u> | <u>N013217</u> | <u>003</u> | Aug 30, 2002 |
| | | METAXALONE | | | | |
| | | MOUNTAIN | 400MG | A040486 | 001 | Feb 27, 2015 |

METFORMIN HYDROCHLORIDE

SOLUTION;ORAL

RIOMET

+! SUN PHARM INDS LTD 500MG/5ML N021591 001 Sep 11, 2003

TABLET;ORAL

GLUCOPHAGE

| | | | | | | |
|-----------|------------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | <u>+</u> | BRISTOL MYERS SQUIBB | <u>500MG</u> | <u>N020357</u> | <u>001</u> | Mar 03, 1995 |
| <u>AB</u> | <u>+</u> | | <u>850MG</u> | <u>N020357</u> | <u>002</u> | Mar 03, 1995 |
| <u>AB</u> | <u>+</u> ! | | <u>1GM</u> | <u>N020357</u> | <u>005</u> | Nov 05, 1998 |

METFORMIN HYDROCHLORIDE

| | | | | | |
|-----------|------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ALKEM | <u>500MG</u> | <u>A091184</u> | <u>001</u> | Nov 01, 2010 |
| <u>AB</u> | | <u>850MG</u> | <u>A091184</u> | <u>002</u> | Nov 01, 2010 |
| <u>AB</u> | | <u>1GM</u> | <u>A091184</u> | <u>003</u> | Nov 01, 2010 |
| <u>AB</u> | AMNEAL PHARMS NY | <u>500MG</u> | <u>A077880</u> | <u>001</u> | Jun 05, 2006 |
| <u>AB</u> | | <u>850MG</u> | <u>A077880</u> | <u>002</u> | Jun 05, 2006 |
| <u>AB</u> | | <u>1GM</u> | <u>A077880</u> | <u>003</u> | Jun 05, 2006 |
| <u>AB</u> | APOTEX | <u>500MG</u> | <u>A075984</u> | <u>001</u> | Apr 23, 2002 |
| <u>AB</u> | | <u>500MG</u> | <u>A090666</u> | <u>001</u> | Dec 07, 2011 |
| <u>AB</u> | | <u>850MG</u> | <u>A075984</u> | <u>002</u> | Apr 23, 2002 |
| <u>AB</u> | | <u>850MG</u> | <u>A090666</u> | <u>002</u> | Dec 07, 2011 |
| <u>AB</u> | | <u>1GM</u> | <u>A075984</u> | <u>003</u> | Apr 23, 2002 |
| <u>AB</u> | | <u>1GM</u> | <u>A090666</u> | <u>003</u> | Dec 07, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

METFORMIN HYDROCHLORIDE

TABLET; ORAL

METFORMIN HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | ATLAS PHARMS LLC | <u>500MG</u> | <u>A076033 001</u> | Jan 24, 2002 |
| <u>AB</u> | | <u>850MG</u> | <u>A076033 002</u> | Jan 24, 2002 |
| <u>AB</u> | | <u>1GM</u> | <u>A076033 003</u> | Jan 24, 2002 |
| <u>AB</u> | AUROBINDO | <u>500MG</u> | <u>A077095 001</u> | Jan 14, 2005 |
| <u>AB</u> | | <u>850MG</u> | <u>A077095 002</u> | Jan 14, 2005 |
| <u>AB</u> | | <u>1GM</u> | <u>A077095 003</u> | Jan 14, 2005 |
| <u>AB</u> | CHARTWELL LIFE SCI | <u>500MG</u> | <u>A075972 001</u> | Jan 24, 2002 |
| <u>AB</u> | | <u>850MG</u> | <u>A075972 002</u> | Jan 24, 2002 |
| <u>AB</u> | | <u>1GM</u> | <u>A075972 003</u> | Jan 24, 2002 |
| <u>AB</u> | CSPC OUYI PHARM CO | <u>500MG</u> | <u>A205096 001</u> | Jul 11, 2016 |
| <u>AB</u> | | <u>850MG</u> | <u>A205096 002</u> | Jul 11, 2016 |
| <u>AB</u> | | <u>1GM</u> | <u>A205096 003</u> | Jul 11, 2016 |
| <u>AB</u> | DR REDDYS LABS INC | <u>500MG</u> | <u>A077787 001</u> | Aug 23, 2006 |
| <u>AB</u> | | <u>850MG</u> | <u>A077787 002</u> | Aug 23, 2006 |
| <u>AB</u> | | <u>1GM</u> | <u>A077787 003</u> | Aug 23, 2006 |
| <u>AB</u> | GLENMARK GENERICS | <u>500MG</u> | <u>A078170 001</u> | May 23, 2008 |
| <u>AB</u> | | <u>850MG</u> | <u>A078170 002</u> | May 23, 2008 |
| <u>AB</u> | | <u>1GM</u> | <u>A078170 003</u> | May 23, 2008 |
| <u>AB</u> | GRANULES INDIA | <u>500MG</u> | <u>A090564 001</u> | Apr 22, 2010 |
| <u>AB</u> | | <u>850MG</u> | <u>A090564 002</u> | Apr 22, 2010 |
| <u>AB</u> | | <u>1GM</u> | <u>A090564 003</u> | Apr 22, 2010 |
| <u>AB</u> | INDICUS PHARMA | <u>500MG</u> | <u>A079148 001</u> | Nov 25, 2008 |
| <u>AB</u> | | <u>850MG</u> | <u>A079148 002</u> | Nov 25, 2008 |
| <u>AB</u> | | <u>1GM</u> | <u>A079148 003</u> | Nov 25, 2008 |
| <u>AB</u> | LAURUS LABS LTD | <u>500MG</u> | <u>A209882 001</u> | Aug 27, 2018 |
| <u>AB</u> | | <u>850MG</u> | <u>A209882 002</u> | Aug 27, 2018 |
| <u>AB</u> | | <u>1GM</u> | <u>A209882 003</u> | Aug 27, 2018 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>500MG</u> | <u>A205330 001</u> | Oct 31, 2017 |
| <u>AB</u> | | <u>850MG</u> | <u>A205330 002</u> | Oct 31, 2017 |
| <u>AB</u> | | <u>1GM</u> | <u>A205330 003</u> | Oct 31, 2017 |
| <u>AB</u> | MARKSANS PHARMA | <u>500MG</u> | <u>A090888 001</u> | Mar 12, 2012 |
| <u>AB</u> | | <u>850MG</u> | <u>A090888 002</u> | Mar 12, 2012 |
| <u>AB</u> | | <u>1GM</u> | <u>A090888 003</u> | Mar 12, 2012 |
| <u>AB</u> | MYLAN | <u>500MG</u> | <u>A075973 001</u> | Jan 25, 2002 |
| <u>AB</u> | | <u>500MG</u> | <u>A075976 001</u> | Jan 24, 2002 |
| <u>AB</u> | | <u>850MG</u> | <u>A075973 002</u> | Jan 25, 2002 |
| <u>AB</u> | | <u>850MG</u> | <u>A075976 002</u> | Jan 24, 2002 |
| <u>AB</u> | | <u>1GM</u> | <u>A075973 003</u> | Jan 25, 2002 |
| <u>AB</u> | | <u>1GM</u> | <u>A075976 003</u> | Jan 24, 2002 |
| <u>AB</u> | SANDOZ | <u>500MG</u> | <u>A075965 001</u> | Jan 25, 2002 |
| <u>AB</u> | | <u>850MG</u> | <u>A075965 002</u> | Jan 25, 2002 |
| <u>AB</u> | | <u>1GM</u> | <u>A075965 003</u> | Jan 25, 2002 |
| <u>AB</u> | SCIEGEN PHARMS INC | <u>500MG</u> | <u>A203769 001</u> | Sep 11, 2013 |
| <u>AB</u> | | <u>850MG</u> | <u>A203769 002</u> | Sep 11, 2013 |
| <u>AB</u> | | <u>1GM</u> | <u>A203769 003</u> | Sep 11, 2013 |
| <u>AB</u> | SUN PHARM INDS INC | <u>500MG</u> | <u>A075967 001</u> | Jan 29, 2002 |
| <u>AB</u> | | <u>850MG</u> | <u>A075967 002</u> | Jan 29, 2002 |
| <u>AB</u> | | <u>1GM</u> | <u>A075967 003</u> | Jan 29, 2002 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>500MG</u> | <u>A076038 001</u> | Feb 21, 2002 |
| <u>AB</u> | | <u>850MG</u> | <u>A076038 002</u> | Feb 21, 2002 |
| <u>AB</u> | | <u>1GM</u> | <u>A076038 003</u> | Feb 21, 2002 |
| <u>AB</u> | SUNSHINE LAKE | <u>500MG</u> | <u>A208999 001</u> | Oct 12, 2018 |
| <u>AB</u> | | <u>850MG</u> | <u>A208999 002</u> | Oct 12, 2018 |
| <u>AB</u> | | <u>1GM</u> | <u>A208999 003</u> | Oct 12, 2018 |
| <u>AB</u> | TEVA | <u>500MG</u> | <u>A075978 001</u> | Jan 25, 2002 |
| <u>AB</u> | | <u>850MG</u> | <u>A075978 002</u> | Jan 25, 2002 |
| <u>AB</u> | | <u>1GM</u> | <u>A075978 003</u> | Nov 05, 2002 |
| <u>AB</u> | TORRENT PHARMS | <u>500MG</u> | <u>A077711 001</u> | Jan 24, 2007 |
| <u>AB</u> | | <u>850MG</u> | <u>A077711 002</u> | Jan 24, 2007 |
| <u>AB</u> | | <u>1GM</u> | <u>A077711 003</u> | Jan 24, 2007 |
| <u>AB</u> | ZYDUS HLTHCARE | <u>500MG</u> | <u>A203686 001</u> | Aug 28, 2014 |
| <u>AB</u> | | <u>850MG</u> | <u>A203686 002</u> | Aug 28, 2014 |
| <u>AB</u> | | <u>1GM</u> | <u>A203686 003</u> | Aug 28, 2014 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>500MG</u> | <u>A077064 001</u> | Apr 18, 2005 |
| <u>AB</u> | | <u>850MG</u> | <u>A077064 002</u> | Apr 18, 2005 |
| <u>AB</u> | | <u>1GM</u> | <u>A077064 003</u> | Apr 18, 2005 |
| <u>AB</u> | CHARTWELL LIFE SCI | 625MG | A075972 005 | Jan 24, 2002 |
| | | 750MG | A075972 004 | Jan 24, 2002 |

PRESCRIPTION DRUG PRODUCT LIST

METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

GLUCOPHAGE XR

| | | | | | |
|-----------|------------|-------------------------|--------------|--------------------|--------------|
| AB | + ! | BRISTOL MYERS SQUIBB | 750MG | N021202 004 | Apr 11, 2003 |
|-----------|------------|-------------------------|--------------|--------------------|--------------|

METFORMIN HYDROCHLORIDE

| | | | | | |
|-----------|--|--------------------------|--------------|--------------------|--------------|
| AB | | ACTAVIS LABS FL INC | 750MG | A076869 001 | Apr 12, 2005 |
| AB | | ALKEM LABS LTD | 750MG | A206145 002 | Oct 22, 2018 |
| AB | | AMNEAL PHARMS NY | 750MG | A078596 002 | Jan 03, 2008 |
| AB | | APOTEX | 750MG | A076706 002 | Dec 29, 2005 |
| AB | | AUROBINDO PHARMA LTD | 750MG | A079118 002 | Jul 20, 2012 |
| AB | | BARR | 750MG | A076863 001 | Oct 14, 2004 |
| AB | | BEXIMCO PHARMS USA | 750MG | A207427 002 | Dec 13, 2016 |
| AB | | CSPC OUYI PHARM CO | 750MG | A078321 002 | Apr 17, 2008 |
| AB | | GRANULES INDIA LTD | 750MG | A209313 002 | Mar 16, 2018 |
| AB | | INTELLIPHARMACEUTIC S | 750MG | A202306 002 | Feb 23, 2017 |
| AB | | MACLEODS PHARMS LTD | 750MG | A206955 002 | Dec 07, 2016 |
| AB | | MARKSANS PHARMA | 750MG | A090295 002 | Apr 29, 2016 |
| AB | | NOSTRUM PHARMS LLC | 750MG | A076756 002 | Dec 12, 2011 |
| AB | | PRINSTON INC | 750MG | A208880 002 | Sep 10, 2018 |
| AB | | SUN PHARM INDS (IN) | 750MG | A077336 002 | Feb 09, 2006 |
| AB | | TEVA | 750MG | A076864 001 | Apr 12, 2005 |
| AB | | YICHANG HUMANWELL | 750MG | A211052 002 | Sep 24, 2018 |
| AB | | ZYDUS PHARMS USA | 750MG | A077078 001 | Apr 21, 2005 |

GLUCOPHAGE XR

| | | | | | |
|------------|----------|-------------------------|--------------|--------------------|--------------|
| AB1 | + | BRISTOL MYERS SQUIBB | 500MG | N021202 001 | Oct 13, 2000 |
|------------|----------|-------------------------|--------------|--------------------|--------------|

METFORMIN HYDROCHLORIDE

| | | | | | |
|------------|--|--------------------------|--------------|--------------------|--------------|
| AB1 | | ACTAVIS LABS FL INC | 500MG | A076172 001 | Jun 16, 2004 |
| AB1 | | ALIGNSCIENCE PHARMA | 500MG | A209303 001 | Mar 19, 2018 |
| AB1 | | ALKEM LABS LTD | 500MG | A206145 001 | Oct 22, 2018 |
| AB1 | | AMNEAL PHARMS NY | 500MG | A078596 001 | Jan 03, 2008 |
| AB1 | | APOTEX | 500MG | A076706 001 | Dec 14, 2004 |
| AB1 | | AUROBINDO PHARMA LTD | 500MG | A079118 001 | Jul 20, 2012 |
| AB1 | | BEXIMCO PHARMS USA | 500MG | A207427 001 | Dec 13, 2016 |
| AB1 | | CSPC OUYI PHARM CO | 500MG | A078321 001 | Apr 17, 2008 |
| AB1 | | GRANULES INDIA LTD | 500MG | A209313 001 | Mar 16, 2018 |
| AB1 | | INTELLIPHARMACEUTIC S | 500MG | A202306 001 | Feb 23, 2017 |
| AB1 | | INVENTIA HLTHCARE | 500MG | A201991 001 | Jan 18, 2012 |
| AB1 | | MACLEODS PHARMS LTD | 500MG | A206955 001 | Dec 07, 2016 |
| AB1 | | MARKSANS PHARMA | 500MG | A090295 001 | Apr 29, 2016 |
| AB1 | | NOSTRUM PHARMS LLC | 500MG | A076756 001 | Jul 26, 2006 |
| AB1 | | PRINSTON INC | 500MG | A208880 001 | Sep 10, 2018 |
| AB1 | | SANDOZ | 500MG | A076873 001 | Dec 14, 2004 |
| AB1 | | SUN PHARM INDS (IN) | 500MG | A077336 001 | Feb 09, 2006 |
| AB1 | | TEVA | 500MG | A076269 001 | Jun 18, 2004 |
| AB1 | | TORRENT PHARMS LTD | 500MG | A090014 001 | Dec 30, 2009 |
| AB1 | | YICHANG HUMANWELL | 500MG | A211052 001 | Sep 24, 2018 |
| AB1 | | ZYDUS PHARMS USA | 500MG | A077060 001 | Apr 20, 2005 |

FORTAMET

| | | | | | |
|------------|------------|----------------|--------------|--------------------|--------------|
| AB2 | + | ANDRX LABS LLC | 500MG | N021574 001 | Apr 27, 2004 |
| AB2 | + ! | | 1GM | N021574 002 | Apr 27, 2004 |

METFORMIN HYDROCHLORIDE

| | | | | | |
|------------|--|-------------------------|--------------|--------------------|--------------|
| AB2 | | LUPIN LTD | 500MG | A090692 001 | Jun 29, 2011 |
| AB2 | | | 1GM | A090692 002 | Jun 29, 2011 |
| AB2 | | MYLAN PHARMS INC | 500MG | A200690 001 | Aug 01, 2012 |
| AB2 | | | 1GM | A200690 002 | Aug 01, 2012 |
| AB2 | | NOSTRUM LABS INC | 500MG | A203832 001 | Dec 26, 2017 |
| AB2 | | | 1GM | A203832 002 | Dec 26, 2017 |
| AB2 | | NOVAST LABS | 500MG | A209674 001 | Nov 02, 2018 |
| AB2 | | | 1GM | A209674 002 | Nov 02, 2018 |
| AB2 | | QINGDAO BAHEAL PHARM | 500MG | A209993 001 | Dec 27, 2018 |
| AB2 | | | 1GM | A209993 002 | Dec 27, 2018 |

GLUMETZA

| | | | | | |
|------------|------------|--------------|--------------|--------------------|--------------|
| AB3 | + | SANTARUS INC | 500MG | N021748 001 | Jun 03, 2005 |
| AB3 | + ! | | 1GM | N021748 002 | Jun 03, 2005 |

METFORMIN HYDROCHLORIDE

| | | | | | |
|------------|--|---------------------|--------------|--------------------|--------------|
| AB3 | | ACTAVIS LABS FL INC | 500MG | A203755 001 | Aug 01, 2016 |
| AB3 | | | 1GM | A203755 002 | Aug 01, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METFORMIN HYDROCHLORIDE

| | | | | |
|------------|-------------------|--------------|--------------------|--------------|
| <u>AB3</u> | LUPIN LTD | <u>500MG</u> | <u>A091664 001</u> | Jul 19, 2013 |
| <u>AB3</u> | | <u>1GM</u> | <u>A091664 002</u> | Jul 19, 2013 |
| <u>AB3</u> | SUN PHARMA GLOBAL | <u>500MG</u> | <u>A202917 001</u> | Aug 01, 2016 |
| <u>AB3</u> | | <u>1GM</u> | <u>A202917 002</u> | Aug 01, 2016 |

METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

ACTOPLUS MET

| | | | | | |
|-----------|---|-------------------|---------------------------|--------------------|--------------|
| <u>AB</u> | + | TAKEDA PHARMS USA | <u>500MG;EQ 15MG BASE</u> | <u>N021842 001</u> | Aug 29, 2005 |
| <u>AB</u> | + | ! | <u>850MG;EQ 15MG BASE</u> | <u>N021842 002</u> | Aug 29, 2005 |

PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE

| | | | | |
|-----------|----------------------|---------------------------|--------------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>500MG;EQ 15MG BASE</u> | <u>A200823 001</u> | Feb 13, 2013 |
| <u>AB</u> | | <u>850MG;EQ 15MG BASE</u> | <u>A200823 002</u> | Feb 13, 2013 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>500MG;EQ 15MG BASE</u> | <u>A204802 001</u> | Nov 05, 2015 |
| <u>AB</u> | | <u>850MG;EQ 15MG BASE</u> | <u>A204802 002</u> | Nov 05, 2015 |
| <u>AB</u> | MYLAN | <u>500MG;EQ 15MG BASE</u> | <u>A090406 001</u> | Feb 25, 2011 |
| <u>AB</u> | | <u>850MG;EQ 15MG BASE</u> | <u>A090406 002</u> | Feb 25, 2011 |
| <u>AB</u> | SANDOZ | <u>500MG;EQ 15MG BASE</u> | <u>A091273 001</u> | Apr 16, 2013 |
| <u>AB</u> | | <u>850MG;EQ 15MG BASE</u> | <u>A091273 002</u> | Apr 16, 2013 |
| <u>AB</u> | TEVA PHARMS USA | <u>500MG;EQ 15MG BASE</u> | <u>A091155 001</u> | Mar 10, 2014 |
| <u>AB</u> | | <u>850MG;EQ 15MG BASE</u> | <u>A091155 002</u> | Mar 10, 2014 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>500MG;EQ 15MG BASE</u> | <u>A202001 001</u> | Feb 13, 2013 |
| <u>AB</u> | | <u>850MG;EQ 15MG BASE</u> | <u>A202001 002</u> | Feb 13, 2013 |

TABLET, EXTENDED RELEASE;ORAL

ACTOPLUS MET XR

| | | | | |
|---|-------------------|------------------|-------------|--------------|
| + | TAKEDA PHARMS USA | 1GM;EQ 15MG BASE | N022024 001 | May 12, 2009 |
| + | ! | 1GM;EQ 30MG BASE | N022024 002 | May 12, 2009 |

METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

KOMBIGLYZE XR

| | | | | |
|---|----------------|-------------------|-------------|--------------|
| + | ASTRAZENECA AB | 500MG;EQ 5MG BASE | N200678 001 | Nov 05, 2010 |
| + | | 1GM;EQ 2.5MG BASE | N200678 003 | Nov 05, 2010 |
| + | ! | 1GM;EQ 5MG BASE | N200678 002 | Nov 05, 2010 |

METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE

TABLET;ORAL

JANUMET

| | | | | |
|---|-------------------|--------------------|-------------|--------------|
| + | MERCK SHARP DOHME | 500MG;EQ 50MG BASE | N022044 001 | Mar 30, 2007 |
| + | ! | 1GM;EQ 50MG BASE | N022044 002 | Mar 30, 2007 |

TABLET, EXTENDED RELEASE;ORAL

JANUMET XR

| | | | | |
|---|-------------------|--------------------|-------------|--------------|
| + | MERCK SHARP DOHME | 500MG;EQ 50MG BASE | N202270 001 | Feb 02, 2012 |
| + | | 1GM;EQ 50MG BASE | N202270 002 | Feb 02, 2012 |
| + | | 1GM;EQ 100MG BASE | N202270 003 | Feb 02, 2012 |

METHACHOLINE CHLORIDE

FOR SOLUTION;INHALATION

PROVOCHOLINE

| | | | | |
|---|------------|------------|-------------|--------------|
| + | METHAPHARM | 100MG/VIAL | N019193 001 | Oct 31, 1986 |
|---|------------|------------|-------------|--------------|

METHADONE HYDROCHLORIDE

CONCENTRATE;ORAL

METHADONE HYDROCHLORIDE

| | | | | |
|-----------|----------------------|----------------|--------------------|--------------|
| <u>AA</u> | VISTAPHARM | <u>10MG/ML</u> | <u>A040088 001</u> | Nov 30, 1994 |
| <u>AA</u> | WEST-WARD PHARMS INT | <u>10MG/ML</u> | <u>A040180 001</u> | Apr 30, 1998 |

METHADONE HYDROCHLORIDE INTENSOL

| | | | | |
|-----------|----------------------|----------------|--------------------|--------------|
| <u>AA</u> | WEST-WARD PHARMS INT | <u>10MG/ML</u> | <u>A089897 001</u> | Sep 06, 1988 |
|-----------|----------------------|----------------|--------------------|--------------|

METHADOSE

| | | | | |
|-----------|---|------------|----------------|--------------------|
| <u>AA</u> | + | SPECGX LLC | <u>10MG/ML</u> | <u>N017116 002</u> |
|-----------|---|------------|----------------|--------------------|

INJECTABLE;INJECTION

METHADONE HYDROCHLORIDE

| | | | | |
|-----------|-------|---------------------|--------------------|--------------------|
| <u>AP</u> | AKORN | <u>10MG/ML</u> | <u>A208306 001</u> | Oct 27, 2017 |
| <u>AP</u> | + | MYLAN INSTITUTIONAL | <u>10MG/ML</u> | <u>N021624 001</u> |

SOLUTION;ORAL

METHADONE HYDROCHLORIDE

| | | | | |
|-----------|------------|----------------------|--------------------|--------------------|
| <u>AA</u> | VISTAPHARM | <u>5MG/5ML</u> | <u>A090707 001</u> | Jun 30, 2010 |
| <u>AA</u> | | <u>10MG/5ML</u> | <u>A090707 002</u> | Jun 30, 2010 |
| <u>AA</u> | ! | WEST-WARD PHARMS INT | <u>5MG/5ML</u> | <u>A087393 001</u> |

PRESCRIPTION DRUG PRODUCT LIST

METHADONE HYDROCHLORIDE

SOLUTION;ORAL

METHADONE HYDROCHLORIDEAA ! 10MG/5ML A087997 001 Aug 30, 1982

TABLET;ORAL

DOLOPHINE HYDROCHLORIDEAA +! WEST-WARD PHARMS 5MG N006134 002
INTAA +! 10MG N006134 010METHADONE HYDROCHLORIDEAA ASCENT PHARMS INC 5MG A211228 001 Jan 03, 2019AA 10MG A211228 002 Jan 03, 2019AA AUROLIFE PHARMA LLC 5MG A203502 001 Aug 31, 2015AA 10MG A203502 002 Aug 31, 2015AA ELITE LABS INC 5MG A210484 001 Aug 02, 2018AA 10MG A210484 002 Aug 02, 2018AA EPIC PHARMA LLC 5MG A090065 001 Aug 18, 2015AA 10MG A090065 002 Aug 18, 2015AA SPECGX LLC 5MG A040517 001 Apr 27, 2004AA 10MG A040517 002 Apr 27, 2004AA SUN PHARM 5MG A208305 001 Mar 30, 2018

INDUSTRIES

AA 10MG A208305 002 Mar 30, 2018AA THEPHARMANETWORK 10MG A090635 001 Nov 25, 2009

LLC

AA VISTAPHARM 10MG A040241 002 May 29, 1998METHADOSEAA SPECGX LLC 5MG A040050 001 Apr 15, 1993AA 10MG A040050 002 Apr 15, 1993

TABLET, FOR SUSPENSION;ORAL

METHADONE HYDROCHLORIDEAA SPECGX LLC 40MG A077142 001 Jul 12, 2005AA VISTAPHARM 40MG A075082 001 Mar 25, 1998AA +! WEST-WARD PHARMS 40MG N017058 001
INTMETHADOSEAA SPECGX LLC 40MG A074184 001 Apr 29, 1993METHAMPHETAMINE HYDROCHLORIDE

TABLET;ORAL

DESOXYNAA +! RECORDATI RARE 5MG N005378 002METHAMPHETAMINE HYDROCHLORIDEAA MAYNE PHARMA INC 5MG A091189 001 Apr 21, 2010AA WEST-WARD PHARMS 5MG A203846 001 Nov 17, 2015
INT

PRESCRIPTION DRUG PRODUCT LIST

METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

| | | | | | |
|-----------|--------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | CASI PHARMS INC | <u>5MG</u> | <u>A040411</u> | <u>001</u> | Mar 27, 2001 |
| <u>AB</u> | | <u>10MG</u> | <u>A040411</u> | <u>002</u> | Mar 27, 2001 |
| <u>AB</u> | ECI PHARMS LLC | <u>5MG</u> | <u>A040547</u> | <u>001</u> | Feb 18, 2005 |
| <u>AB</u> | | <u>10MG</u> | <u>A040547</u> | <u>002</u> | Feb 18, 2005 |
| <u>AB</u> | HERITAGE PHARMA | <u>5MG</u> | <u>A040734</u> | <u>001</u> | Dec 14, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A040734</u> | <u>002</u> | Dec 14, 2007 |
| <u>AB</u> | MYLAN | <u>5MG</u> | <u>A040350</u> | <u>001</u> | Mar 29, 2000 |
| <u>AB</u> | ! | <u>10MG</u> | <u>A040350</u> | <u>002</u> | Mar 29, 2000 |
| <u>AB</u> | RISING PHARMS | <u>5MG</u> | <u>A202068</u> | <u>001</u> | Mar 07, 2012 |
| <u>AB</u> | | <u>10MG</u> | <u>A202068</u> | <u>002</u> | Mar 07, 2012 |
| <u>AB</u> | SUN PHARM INDS INC | <u>5MG</u> | <u>A040870</u> | <u>001</u> | Sep 25, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A040870</u> | <u>002</u> | Sep 25, 2007 |

TAPAZOLE

| | | | | | |
|-----------|-----------------|-------------|----------------|------------|--------------|
| <u>AB</u> | KING PHARMS LLC | <u>5MG</u> | <u>A040320</u> | <u>001</u> | Mar 31, 2000 |
| <u>AB</u> | | <u>10MG</u> | <u>A040320</u> | <u>002</u> | Mar 31, 2000 |

METHOCARBAMOL

SOLUTION; IM-IV

METHOCARBAMOL

| | | | | | |
|-----------|----------------------|----------------------------|----------------|------------|--------------|
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>1GM/10ML (100MG/ML)</u> | <u>A206128</u> | <u>001</u> | May 27, 2016 |
| <u>AP</u> | FRESENIUS KABI USA | <u>1GM/10ML (100MG/ML)</u> | <u>A209331</u> | <u>001</u> | Apr 17, 2018 |
| <u>AP</u> | GLAND PHARMA LTD | <u>1GM/10ML (100MG/ML)</u> | <u>A211504</u> | <u>001</u> | Oct 26, 2018 |
| <u>AP</u> | LUITPOLD | <u>1GM/10ML (100MG/ML)</u> | <u>A207496</u> | <u>001</u> | Jun 22, 2017 |
| <u>AP</u> | MONTEREY PHARMS LLC | <u>1GM/10ML (100MG/ML)</u> | <u>A205354</u> | <u>001</u> | Oct 27, 2016 |
| <u>AP</u> | MYLAN INSTITUTIONAL | <u>1GM/10ML (100MG/ML)</u> | <u>A204404</u> | <u>001</u> | Dec 05, 2014 |
| <u>AP</u> | NAVINTA LLC | <u>1GM/10ML (100MG/ML)</u> | <u>A206071</u> | <u>001</u> | Nov 24, 2017 |
| <u>AP</u> | RENAISSANCE SSA LLC | <u>1GM/10ML (100MG/ML)</u> | <u>A208116</u> | <u>001</u> | Jan 19, 2017 |
| <u>AP</u> | SAGENT PHARMS | <u>1GM/10ML (100MG/ML)</u> | <u>A205404</u> | <u>001</u> | Jul 18, 2017 |
| <u>AP</u> | SOMERSET THERAPS LLC | <u>1GM/10ML (100MG/ML)</u> | <u>A207522</u> | <u>001</u> | Jul 31, 2017 |

ROBAXIN

| | | | | | |
|-----------|------------------------|----------------------------|----------------|------------|--|
| <u>AP</u> | ! WEST-WARD PHARMS INT | <u>1GM/10ML (100MG/ML)</u> | <u>N011790</u> | <u>001</u> | |
|-----------|------------------------|----------------------------|----------------|------------|--|

TABLET; ORAL

METHOCARBAMOL

| | | | | | |
|-----------|---------------------|--------------|----------------|------------|--------------|
| <u>AA</u> | AUSTARPHARMA LLC | <u>500MG</u> | <u>A200958</u> | <u>001</u> | Oct 21, 2011 |
| <u>AA</u> | | <u>750MG</u> | <u>A200958</u> | <u>002</u> | Oct 21, 2011 |
| <u>AA</u> | BEXIMCO PHARMS USA | <u>500MG</u> | <u>A208507</u> | <u>001</u> | Jul 21, 2017 |
| <u>AA</u> | | <u>750MG</u> | <u>A208507</u> | <u>002</u> | Jul 21, 2017 |
| <u>AA</u> | DBL PHARMS | <u>500MG</u> | <u>A203550</u> | <u>001</u> | Feb 08, 2017 |
| <u>AA</u> | | <u>750MG</u> | <u>A203550</u> | <u>002</u> | Feb 08, 2017 |
| <u>AA</u> | GRANULES INDIA LTD | <u>500MG</u> | <u>A209312</u> | <u>001</u> | May 07, 2018 |
| <u>AA</u> | | <u>750MG</u> | <u>A209312</u> | <u>002</u> | May 07, 2018 |
| <u>AA</u> | HETERO LABS LTD III | <u>500MG</u> | <u>A090200</u> | <u>001</u> | Nov 06, 2009 |
| <u>AA</u> | | <u>750MG</u> | <u>A090200</u> | <u>002</u> | Nov 06, 2009 |
| <u>AA</u> | HIKMA INTL PHARMS | <u>500MG</u> | <u>A085159</u> | <u>001</u> | |
| <u>AA</u> | | <u>750MG</u> | <u>A085123</u> | <u>001</u> | |
| <u>AA</u> | OXFORD PHARMS | <u>500MG</u> | <u>A040489</u> | <u>001</u> | Jan 29, 2003 |
| <u>AA</u> | | <u>750MG</u> | <u>A040489</u> | <u>002</u> | Jan 29, 2003 |
| <u>AA</u> | PRINSTON INC | <u>500MG</u> | <u>A086989</u> | <u>001</u> | |
| <u>AA</u> | | <u>750MG</u> | <u>A086988</u> | <u>001</u> | |
| <u>AA</u> | WATSON LABS | <u>500MG</u> | <u>A084277</u> | <u>001</u> | |
| <u>AA</u> | | <u>750MG</u> | <u>A084276</u> | <u>002</u> | |

ROBAXIN

| | | | | | |
|-----------|-----------------------|--------------|----------------|------------|--|
| <u>AA</u> | ! AUXILIUM PHARMS LLC | <u>500MG</u> | <u>N011011</u> | <u>004</u> | |
|-----------|-----------------------|--------------|----------------|------------|--|

ROBAXIN-750

| | | | | | |
|-----------|-----------------------|--------------|----------------|------------|--|
| <u>AA</u> | ! AUXILIUM PHARMS LLC | <u>750MG</u> | <u>N011011</u> | <u>006</u> | |
|-----------|-----------------------|--------------|----------------|------------|--|

METHOHEXITAL SODIUM

INJECTABLE; INJECTION

BREVITAL SODIUM

| | | | | | |
|---|---|----------------------|------------|---------|-----|
| + | ! | PAR STERILE PRODUCTS | 500MG/VIAL | N011559 | 001 |
| + | ! | | 2.5GM/VIAL | N011559 | 002 |

PRESCRIPTION DRUG PRODUCT LIST

METHOTREXATE

SOLUTION;SUBCUTANEOUS

OTREXUP

| | | | | | | |
|---|---|--------------------|-----------------------------|---------|-----|--------------|
| + | ! | ANTARES PHARMA INC | 10MG/0.4ML (10MG/0.4ML) | N204824 | 001 | Oct 11, 2013 |
| + | ! | | 12.5MG/0.4ML (12.5MG/0.4ML) | N204824 | 006 | Mar 24, 2016 |
| + | ! | | 15MG/0.4ML (15MG/0.4ML) | N204824 | 002 | Oct 11, 2013 |
| + | ! | | 17.5MG/0.4ML (17.5MG/0.4ML) | N204824 | 007 | Mar 24, 2016 |
| + | ! | | 20MG/0.4ML (20MG/0.4ML) | N204824 | 003 | Oct 11, 2013 |
| + | ! | | 22.5MG/0.4ML (22.5MG/0.4ML) | N204824 | 008 | Mar 24, 2016 |
| + | ! | | 25MG/0.4ML (25MG/0.4ML) | N204824 | 004 | Oct 11, 2013 |

RASUVO

| | | | | | | |
|---|---|------------------|-------------------------------|---------|-----|--------------|
| + | ! | MEDAC PHARMA INC | 7.5MG/0.15ML (7.5MG/0.15ML) | N205776 | 001 | Jul 10, 2014 |
| + | ! | | 10MG/0.20ML (10MG/0.20ML) | N205776 | 002 | Jul 10, 2014 |
| + | ! | | 12.5MG/0.25ML (12.5MG/0.25ML) | N205776 | 003 | Jul 10, 2014 |
| + | ! | | 15MG/0.30ML (15MG/0.30ML) | N205776 | 004 | Jul 10, 2014 |
| + | ! | | 17.5MG/0.35ML (17.5MG/0.35ML) | N205776 | 005 | Jul 10, 2014 |
| + | ! | | 20MG/0.4ML (20MG/0.4ML) | N205776 | 006 | Jul 10, 2014 |
| + | ! | | 22.5MG/0.45ML (22.5MG/0.45ML) | N205776 | 007 | Jul 10, 2014 |
| + | ! | | 25MG/0.5ML (25MG/0.5ML) | N205776 | 008 | Jul 10, 2014 |
| + | ! | | 30MG/0.6ML (30MG/0.6ML) | N205776 | 010 | Jul 10, 2014 |

METHOTREXATE SODIUM

INJECTABLE;INJECTION

METHOTREXATE PRESERVATIVE FREE

| | | | | | | |
|-----------|--|--------------------|--------------------------------|-----------------------|-------------------|--------------|
| AP | | FRESENIUS KABI USA | <u>EQ 25MG BASE/ML</u> | <u>A040265</u> | <u>001</u> | Feb 26, 1999 |
| AP | | | <u>EQ 1GM BASE/VIAL</u> | <u>A040266</u> | <u>001</u> | Feb 26, 1999 |

METHOTREXATE SODIUM

| | | | | | | |
|-----------|---|-------------------------|---------------------------------------------------|-----------------------|-------------------|--------------|
| AP | ! | FRESENIUS KABI USA | <u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u> | <u>A040263</u> | <u>001</u> | Feb 26, 1999 |
| AP | + | HOSPIRA | <u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u> | <u>N011719</u> | <u>010</u> | Dec 15, 2004 |
| AP | ! | WEST-WARD PHARMS INT | <u>EQ 100MG BASE/4ML (EQ 25MG BASE/ML)</u> | <u>A089341</u> | <u>001</u> | Sep 16, 1986 |

METHOTREXATE SODIUM PRESERVATIVE FREE

| | | | | | | |
|-----------|---|-------------------------|----------------------------------------------------|-----------------------|-------------------|--------------|
| AP | ! | ACCORD HLTHCARE | <u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u> | <u>A040767</u> | <u>001</u> | Apr 30, 2007 |
| AP | ! | | <u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u> | <u>A040768</u> | <u>001</u> | Apr 30, 2007 |
| AP | ! | | <u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u> | <u>A040716</u> | <u>001</u> | Apr 30, 2007 |
| AP | + | HOSPIRA | <u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u> | <u>N011719</u> | <u>012</u> | Apr 13, 2005 |
| AP | | MYLAN LABS LTD | <u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u> | <u>A201529</u> | <u>001</u> | Mar 29, 2012 |
| AP | | | <u>EQ 100MG BASE/4ML (EQ 25MG BASE/ML)</u> | <u>A201529</u> | <u>002</u> | Mar 29, 2012 |
| AP | | | <u>EQ 200MG BASE/8ML (EQ 25MG BASE/ML)</u> | <u>A201529</u> | <u>003</u> | Mar 29, 2012 |
| AP | | | <u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u> | <u>A201529</u> | <u>004</u> | Mar 29, 2012 |
| AP | | | <u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u> | <u>A201530</u> | <u>001</u> | Mar 29, 2012 |
| AP | | PHARMACHEMIE BV | <u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u> | <u>A040843</u> | <u>002</u> | Jan 11, 2010 |
| AP | | | <u>EQ 100MG BASE/4ML (EQ 25MG BASE/ML)</u> | <u>A040843</u> | <u>003</u> | Feb 27, 2012 |
| AP | | | <u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u> | <u>A040843</u> | <u>004</u> | Jan 11, 2010 |
| AP | | | <u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u> | <u>A040843</u> | <u>001</u> | Jan 11, 2010 |
| AP | | SAGENT PHARMS | <u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u> | <u>A203407</u> | <u>001</u> | Aug 09, 2018 |
| AP | | | <u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u> | <u>A203407</u> | <u>002</u> | Aug 09, 2018 |
| AP | | | <u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u> | <u>A203407</u> | <u>003</u> | Aug 09, 2018 |
| AP | | SANDOZ INC | <u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u> | <u>A090039</u> | <u>001</u> | Mar 31, 2009 |
| AP | | | <u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u> | <u>A090039</u> | <u>002</u> | Mar 31, 2009 |
| AP | | | <u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u> | <u>A090029</u> | <u>001</u> | Mar 31, 2009 |
| AP | ! | WEST-WARD PHARMS INT | <u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u> | <u>A089340</u> | <u>001</u> | Sep 16, 1986 |
| AP | ! | | <u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u> | <u>A089343</u> | <u>001</u> | Sep 16, 1986 |

METHOTREXATE SODIUM

| | | | | | | |
|---|--|-------------------------|-------------------------------------|---------|-----|--------------|
| ! | | WEST-WARD PHARMS INT | EQ 200MG BASE/8ML (EQ 25MG BASE/ML) | A089342 | 001 | Sep 16, 1986 |
|---|--|-------------------------|-------------------------------------|---------|-----|--------------|

METHOTREXATE SODIUM PRESERVATIVE FREE

| | | | | | | |
|---|--|-------------------------|------------------|---------|-----|--------------|
| ! | | WEST-WARD PHARMS INT | EQ 1GM BASE/VIAL | A040632 | 001 | Aug 12, 2005 |
|---|--|-------------------------|------------------|---------|-----|--------------|

SOLUTION;ORAL

XATMEP

| | | | | | | |
|---|---|-------------------|----------------|---------|-----|--------------|
| + | ! | SILVERGATE PHARMS | EQ 2MG BASE/ML | N208400 | 001 | Apr 25, 2017 |
|---|---|-------------------|----------------|---------|-----|--------------|

TABLET;ORAL

METHOTREXATE SODIUM

| | | | | | | |
|-----------|---|-------------------------|-----------------------------|-----------------------|-------------------|--------------|
| AB | | AMNEAL PHARMS | <u>EQ 2.5MG BASE</u> | <u>A210040</u> | <u>001</u> | Dec 22, 2017 |
| AB | | BARR | <u>EQ 2.5MG BASE</u> | <u>A081099</u> | <u>001</u> | Oct 15, 1990 |
| AB | + | DAVA PHARMS INC | <u>EQ 2.5MG BASE</u> | <u>N008085</u> | <u>002</u> | |
| AB | | MYLAN | <u>EQ 2.5MG BASE</u> | <u>A081235</u> | <u>001</u> | May 15, 1992 |
| AB | | SUN PHARMA GLOBAL | <u>EQ 2.5MG BASE</u> | <u>A201749</u> | <u>001</u> | May 21, 2015 |
| AB | | WEST-WARD PHARMS INT | <u>EQ 2.5MG BASE</u> | <u>A040054</u> | <u>001</u> | Aug 01, 1994 |
| AB | | ZYDUS PHARMS USA INC | <u>EQ 2.5MG BASE</u> | <u>A207812</u> | <u>001</u> | Jan 13, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

METHOTREXATE SODIUM

TABLET; ORAL

Trexall

BARR

EQ 5MG BASE
EQ 7.5MG BASE
EQ 10MG BASE
EQ 15MG BASEA040385 001 Mar 21, 2001
A040385 002 Mar 21, 2001
A040385 003 Mar 21, 2001
A040385 004 Mar 21, 2001

!

METHOXSALEN

CAPSULE; ORAL

METHOXSALENAB ACTAVIS INC10MGA202603 001 Jun 09, 2015AB STRIDES PHARMA10MGA202687 001 Jun 05, 2014OXSORALEN-ULTRAAB +! DOW PHARM10MGN019600 001 Oct 30, 1986

INJECTABLE; INJECTION

UVADEX

+! MALLINCKRODT HOSP

0.02MG/ML

N020969 001 Feb 25, 1999

METHSCOPOLAMINE BROMIDE

TABLET; ORAL

METHSCOPOLAMINE BROMIDEAA BAYSHORE PHARMS LLC2.5MGA200602 001 Sep 24, 2012AA5MGA200602 002 Sep 24, 2012AA BRECKENRIDGE PHARM2.5MGA040642 001 Dec 06, 2011AA5MGA040642 002 Dec 06, 2011AA ! VINTAGE PHARMS2.5MGA040624 001 Dec 28, 2006AA !5MGA040624 002 Dec 28, 2006METHSUXIMIDE

CAPSULE; ORAL

CELONTIN

+! PARKE DAVIS

300MG

N010596 008

METHYCLOTHIAZIDE

TABLET; ORAL

METHYCLOTHIAZIDE

! MYLAN PHARMS INC

5MG

A087672 001 Aug 17, 1982

METHYLDOPA

TABLET; ORAL

METHYLDOPAAB ACCORD HLTHCARE250MGA070084 001 Oct 15, 1985AB500MGA070085 001 Oct 15, 1985AB IVAX SUB TEVA
PHARMS250MGA070098 001 Feb 20, 1986AB500MGA070343 001 Feb 20, 1986AB MYLAN250MGA070076 002 Apr 18, 1985AB !500MGA070076 001 Apr 18, 1985AB WATSON LABS500MGA070625 001 Jun 06, 1986METHYLDOPATE HYDROCHLORIDE

INJECTABLE; INJECTION

METHYLDOPATE HYDROCHLORIDE

! LUITPOLD

50MG/ML

A071279 001 Oct 02, 1987

METHYLENE BLUE

SOLUTION; INTRAVENOUS

PROVAYBLUE

+! PROVEPHARM SAS

50MG/10ML (5MG/ML)

N204630 001 Apr 08, 2016

METHYLERGONOVINE MALEATE

INJECTABLE; INJECTION

METHERGINEAP +! EDISON THERAPS LLC0.2MG/MLN006035 004METHYLERGONOVINE MALEATEAP BRECKENRIDGE PHARM0.2MG/MLA040889 001 Sep 13, 2010AP LUITPOLD0.2MG/MLA090193 001 Nov 24, 2008

TABLET; ORAL

METHYLERGONOVINE MALEATEAB AMNEAL PHARMS0.2MGA211483 001 Sep 10, 2018AB GRANULES PHARMS0.2MGA210424 001 May 15, 2018AB ! NOVEL LABS INC0.2MGA091577 001 May 02, 2011

PRESCRIPTION DRUG PRODUCT LIST

METHYLNALTREXONE BROMIDE

SOLUTION;SUBCUTANEOUS

RELISTOR

| | | | | | | |
|---|---|--------------|-------------------------|---------|-----|--------------|
| + | ! | SALIX PHARMS | 8MG/0.4ML (8MG/0.4ML) | N021964 | 002 | Sep 27, 2010 |
| + | ! | | 12MG/0.6ML (12MG/0.6ML) | N021964 | 001 | Apr 24, 2008 |
| + | ! | | 12MG/0.6ML (12MG/0.6ML) | N021964 | 003 | Sep 27, 2010 |

TABLET;ORAL

RELISTOR

| | | | | | | |
|---|---|------------------|-------|---------|-----|--------------|
| + | ! | SALIX PHARMS INC | 150MG | N208271 | 001 | Jul 19, 2016 |
|---|---|------------------|-------|---------|-----|--------------|

METHYLPHENIDATE

FILM, EXTENDED RELEASE;TRANSDERMAL

DAYTRANA

| | | | | | | |
|---|---|------------------|---------------------|---------|-----|--------------|
| + | | NOVEN PHARMS INC | 10MG/9HR (1.1MG/HR) | N021514 | 001 | Apr 06, 2006 |
| + | | | 15MG/9HR (1.6MG/HR) | N021514 | 002 | Apr 06, 2006 |
| + | | | 20MG/9HR (2.2MG/HR) | N021514 | 003 | Apr 06, 2006 |
| + | ! | | 30MG/9HR (3.3MG/HR) | N021514 | 004 | Apr 06, 2006 |

TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE;ORAL

COTEMPLA XR-ODT

| | | | | | | |
|---|--|------------------|--------|---------|-----|--------------|
| + | | NEOS THERAPS INC | 8.6MG | N205489 | 001 | Jun 19, 2017 |
| + | | | 17.3MG | N205489 | 002 | Jun 19, 2017 |
| + | | | 25.9MG | N205489 | 003 | Jun 19, 2017 |

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

| | | | | | | |
|------------|--|---------------|-------------|----------------|------------|--------------|
| <u>AB1</u> | | BARR LABS INC | <u>10MG</u> | <u>A079031</u> | <u>004</u> | Oct 15, 2014 |
| <u>AB1</u> | | | <u>20MG</u> | <u>A079031</u> | <u>001</u> | Jul 13, 2012 |
| <u>AB1</u> | | | <u>30MG</u> | <u>A079031</u> | <u>002</u> | Jul 13, 2012 |
| <u>AB1</u> | | | <u>40MG</u> | <u>A079031</u> | <u>003</u> | Jul 13, 2012 |
| <u>AB1</u> | | MAYNE PHARMA | <u>10MG</u> | <u>A200886</u> | <u>001</u> | Feb 26, 2018 |
| <u>AB1</u> | | | <u>20MG</u> | <u>A078458</u> | <u>001</u> | Dec 01, 2011 |
| <u>AB1</u> | | | <u>30MG</u> | <u>A078458</u> | <u>002</u> | Dec 01, 2011 |
| <u>AB1</u> | | | <u>40MG</u> | <u>A078458</u> | <u>003</u> | Dec 01, 2011 |

RITALIN LA

| | | | | | | |
|------------|---|----------|-------------|----------------|------------|--------------|
| <u>AB1</u> | + | NOVARTIS | <u>10MG</u> | <u>N021284</u> | <u>004</u> | Apr 10, 2004 |
| <u>AB1</u> | + | | <u>20MG</u> | <u>N021284</u> | <u>001</u> | Jun 05, 2002 |
| <u>AB1</u> | + | | <u>30MG</u> | <u>N021284</u> | <u>002</u> | Jun 05, 2002 |
| <u>AB1</u> | + | | <u>40MG</u> | <u>N021284</u> | <u>003</u> | Jun 05, 2002 |

METADATE CD

| | | | | | | |
|------------|---|----------------|-------------|----------------|------------|--------------|
| <u>AB2</u> | + | LANNETT CO INC | <u>10MG</u> | <u>N021259</u> | <u>003</u> | May 27, 2003 |
| <u>AB2</u> | + | | <u>20MG</u> | <u>N021259</u> | <u>001</u> | Apr 03, 2001 |
| <u>AB2</u> | + | | <u>30MG</u> | <u>N021259</u> | <u>002</u> | Jun 19, 2003 |
| <u>AB2</u> | + | | <u>40MG</u> | <u>N021259</u> | <u>004</u> | Feb 19, 2006 |
| <u>AB2</u> | + | | <u>50MG</u> | <u>N021259</u> | <u>005</u> | Feb 19, 2006 |
| <u>AB2</u> | + | ! | <u>60MG</u> | <u>N021259</u> | <u>006</u> | Feb 19, 2006 |

METHYLPHENIDATE HYDROCHLORIDE

| | | | | | | |
|------------|--|----------------|-------------|----------------|------------|--------------|
| <u>AB2</u> | | IMPAX LABS INC | <u>10MG</u> | <u>A205105</u> | <u>001</u> | Jul 28, 2016 |
| <u>AB2</u> | | | <u>20MG</u> | <u>A205105</u> | <u>002</u> | Jul 28, 2016 |
| <u>AB2</u> | | | <u>30MG</u> | <u>A205105</u> | <u>003</u> | Jul 28, 2016 |
| <u>AB2</u> | | | <u>40MG</u> | <u>A205105</u> | <u>004</u> | Jul 28, 2016 |
| <u>AB2</u> | | | <u>50MG</u> | <u>A205105</u> | <u>005</u> | Jul 28, 2016 |
| <u>AB2</u> | | | <u>60MG</u> | <u>A205105</u> | <u>006</u> | Jul 28, 2016 |
| <u>AB2</u> | | SPECGX LLC | <u>10MG</u> | <u>A203583</u> | <u>001</u> | Sep 29, 2015 |
| <u>AB2</u> | | | <u>20MG</u> | <u>A203583</u> | <u>002</u> | Sep 29, 2015 |
| <u>AB2</u> | | | <u>30MG</u> | <u>A203583</u> | <u>003</u> | Sep 29, 2015 |
| <u>AB2</u> | | | <u>40MG</u> | <u>A203583</u> | <u>004</u> | Sep 29, 2015 |
| <u>AB2</u> | | | <u>50MG</u> | <u>A203583</u> | <u>005</u> | Sep 29, 2015 |
| <u>AB2</u> | | | <u>60MG</u> | <u>A203583</u> | <u>006</u> | Sep 29, 2015 |
| <u>AB2</u> | | TEVA PHARMS | <u>10MG</u> | <u>A077707</u> | <u>001</u> | Jul 19, 2012 |
| <u>AB2</u> | | | <u>20MG</u> | <u>A077707</u> | <u>002</u> | Jul 19, 2012 |
| <u>AB2</u> | | | <u>30MG</u> | <u>A077707</u> | <u>003</u> | Jul 19, 2012 |
| <u>AB2</u> | | | <u>40MG</u> | <u>A078873</u> | <u>001</u> | Jul 19, 2012 |
| <u>AB2</u> | | | <u>50MG</u> | <u>A078873</u> | <u>002</u> | Jul 19, 2012 |
| <u>AB2</u> | | | <u>60MG</u> | <u>A078873</u> | <u>003</u> | Jul 19, 2012 |

APTENSIO XR

| | | | | | | |
|------------|---|---------------|-------------|----------------|------------|--------------|
| <u>AB3</u> | + | RHODES PHARMS | <u>10MG</u> | <u>N205831</u> | <u>001</u> | Apr 17, 2015 |
| <u>AB3</u> | + | | <u>15MG</u> | <u>N205831</u> | <u>002</u> | Apr 17, 2015 |
| <u>AB3</u> | + | | <u>20MG</u> | <u>N205831</u> | <u>003</u> | Apr 17, 2015 |
| <u>AB3</u> | + | | <u>30MG</u> | <u>N205831</u> | <u>004</u> | Apr 17, 2015 |
| <u>AB3</u> | + | | <u>40MG</u> | <u>N205831</u> | <u>005</u> | Apr 17, 2015 |
| <u>AB3</u> | + | | <u>50MG</u> | <u>N205831</u> | <u>006</u> | Apr 17, 2015 |
| <u>AB3</u> | + | ! | <u>60MG</u> | <u>N205831</u> | <u>007</u> | Apr 17, 2015 |

PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

| | | | | |
|------------|-------------------|-------------|--------------------|--------------|
| <u>AB3</u> | ACTAVIS ELIZABETH | <u>10MG</u> | <u>A208861 001</u> | Dec 13, 2018 |
| <u>AB3</u> | | <u>15MG</u> | <u>A208861 002</u> | Dec 13, 2018 |
| <u>AB3</u> | | <u>20MG</u> | <u>A208861 003</u> | Dec 13, 2018 |
| <u>AB3</u> | | <u>30MG</u> | <u>A208861 004</u> | Dec 13, 2018 |
| <u>AB3</u> | | <u>40MG</u> | <u>A208861 005</u> | Dec 13, 2018 |
| <u>AB3</u> | | <u>50MG</u> | <u>A208861 006</u> | Dec 13, 2018 |
| <u>AB3</u> | | <u>60MG</u> | <u>A208861 007</u> | Dec 13, 2018 |

JORNAY PM

| | | | | |
|---|------------------|-------|-------------|--------------|
| + | IRONSHORE PHARMS | 20MG | N209311 001 | Aug 08, 2018 |
| + | | 40MG | N209311 002 | Aug 08, 2018 |
| + | | 60MG | N209311 003 | Aug 08, 2018 |
| + | | 80MG | N209311 004 | Aug 08, 2018 |
| + | | 100MG | N209311 005 | Aug 08, 2018 |

METHYLPHENIDATE HYDROCHLORIDE

| | | | | |
|---|--------------|------|-------------|--------------|
| ! | MAYNE PHARMA | 60MG | A078458 004 | Jun 23, 2016 |
|---|--------------|------|-------------|--------------|

FOR SUSPENSION, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

| | | | | |
|-----------|---------------------|---------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS LABS FL INC | <u>5MG/ML</u> | <u>A206049 001</u> | May 17, 2018 |
|-----------|---------------------|---------------|--------------------|--------------|

OUILIVANT XR

| | | | | |
|-----------|-------------------|---------------|--------------------|--------------|
| <u>AB</u> | ! NEXTWAVE PHARMS | <u>5MG/ML</u> | <u>N202100 001</u> | Sep 27, 2012 |
|-----------|-------------------|---------------|--------------------|--------------|

SOLUTION;ORAL

METHYLIN

| | | | | |
|-----------|--------------|----------------|--------------------|--------------|
| <u>AA</u> | ! SPECGX LLC | <u>5MG/5ML</u> | <u>N021419 001</u> | Dec 19, 2002 |
|-----------|--------------|----------------|--------------------|--------------|

| | | | | |
|-----------|---|-----------------|--------------------|--------------|
| <u>AA</u> | ! | <u>10MG/5ML</u> | <u>N021419 002</u> | Dec 19, 2002 |
|-----------|---|-----------------|--------------------|--------------|

METHYLPHENIDATE HYDROCHLORIDE

| | | | | |
|-----------|--------------------|-----------------|--------------------|--------------|
| <u>AA</u> | ABHAI LLC | <u>5MG/5ML</u> | <u>A207485 001</u> | Nov 18, 2016 |
| <u>AA</u> | | <u>10MG/5ML</u> | <u>A207485 002</u> | Nov 18, 2016 |
| <u>AA</u> | BRECKENRIDGE PHARM | <u>5MG/5ML</u> | <u>A201466 001</u> | Nov 12, 2013 |
| <u>AA</u> | | <u>10MG/5ML</u> | <u>A201466 002</u> | Nov 12, 2013 |
| <u>AA</u> | NOVEL LABS INC | <u>5MG/5ML</u> | <u>A204602 001</u> | Aug 14, 2015 |
| <u>AA</u> | | <u>10MG/5ML</u> | <u>A204602 002</u> | Aug 14, 2015 |
| <u>AA</u> | TRIS PHARMA INC | <u>5MG/5ML</u> | <u>A091601 001</u> | Jul 23, 2010 |
| <u>AA</u> | | <u>10MG/5ML</u> | <u>A091601 002</u> | Jul 23, 2010 |
| <u>AA</u> | WES PHARMA INC | <u>5MG/5ML</u> | <u>A210139 001</u> | Oct 03, 2018 |
| <u>AA</u> | | <u>10MG/5ML</u> | <u>A210139 002</u> | Oct 03, 2018 |

TABLET;ORAL

METHYLPHENIDATE HYDROCHLORIDE

| | | | | |
|-----------|---------------------|-------------|--------------------|--------------|
| <u>AB</u> | ABHAI INC | <u>5MG</u> | <u>A206932 001</u> | May 11, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A206932 002</u> | May 11, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A206932 003</u> | May 11, 2017 |
| <u>AB</u> | ACTAVIS LABS FL INC | <u>5MG</u> | <u>A040220 001</u> | Aug 29, 1997 |
| <u>AB</u> | | <u>10MG</u> | <u>A040220 002</u> | Aug 29, 1997 |
| <u>AB</u> | | <u>20MG</u> | <u>A040220 003</u> | Aug 29, 1997 |
| <u>AB</u> | ASCENT PHARMS INC | <u>5MG</u> | <u>A207416 001</u> | Sep 22, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A207416 002</u> | Sep 22, 2015 |
| <u>AB</u> | | <u>20MG</u> | <u>A207416 003</u> | Sep 22, 2015 |
| <u>AB</u> | AUROLIFE PHARMA LLC | <u>5MG</u> | <u>A209276 001</u> | Oct 25, 2018 |
| <u>AB</u> | | <u>10MG</u> | <u>A209276 002</u> | Oct 25, 2018 |
| <u>AB</u> | | <u>20MG</u> | <u>A209276 003</u> | Oct 25, 2018 |
| <u>AB</u> | BIONPHARMA INC | <u>5MG</u> | <u>A209753 001</u> | Mar 02, 2018 |
| <u>AB</u> | | <u>10MG</u> | <u>A209753 002</u> | Mar 02, 2018 |
| <u>AB</u> | | <u>20MG</u> | <u>A209753 003</u> | Mar 02, 2018 |
| <u>AB</u> | BRECKENRIDGE PHARM | <u>5MG</u> | <u>A207587 001</u> | Mar 03, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A207587 002</u> | Mar 03, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A207587 003</u> | Mar 03, 2017 |
| <u>AB</u> | CNTY LINE PHARMS | <u>5MG</u> | <u>A206840 001</u> | Sep 15, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A206840 002</u> | Sep 15, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A206840 003</u> | Sep 15, 2016 |
| <u>AB</u> | LANNETT CO INC | <u>5MG</u> | <u>A086429 001</u> | |
| <u>AB</u> | | <u>10MG</u> | <u>A085799 001</u> | |
| <u>AB</u> | | <u>20MG</u> | <u>A086428 001</u> | |
| <u>AB</u> | MOUNTAIN | <u>5MG</u> | <u>A091159 001</u> | Mar 12, 2014 |
| <u>AB</u> | | <u>10MG</u> | <u>A091159 002</u> | Mar 12, 2014 |
| <u>AB</u> | | <u>20MG</u> | <u>A091159 003</u> | Mar 12, 2014 |
| <u>AB</u> | NOVEL LABS INC | <u>5MG</u> | <u>A207884 001</u> | Nov 13, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A207884 002</u> | Nov 13, 2015 |
| <u>AB</u> | | <u>20MG</u> | <u>A207884 003</u> | Nov 13, 2015 |
| <u>AB</u> | OXFORD PHARMS | <u>5MG</u> | <u>A202892 001</u> | Sep 23, 2014 |
| <u>AB</u> | | <u>10MG</u> | <u>A202892 002</u> | Sep 23, 2014 |

PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

TABLET; ORAL

METHYLPHENIDATE HYDROCHLORIDE

| | | | | | |
|-----------|--------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | <u>20MG</u> | <u>A202892</u> | <u>003</u> | Sep 23, 2014 |
| <u>AB</u> | SPECGX LLC | <u>5MG</u> | <u>A040300</u> | <u>001</u> | Nov 27, 1998 |
| <u>AB</u> | | <u>10MG</u> | <u>A040300</u> | <u>002</u> | Nov 27, 1998 |
| <u>AB</u> | | <u>20MG</u> | <u>A040300</u> | <u>003</u> | Nov 27, 1998 |
| <u>AB</u> | SUN PHARM INDS INC | <u>5MG</u> | <u>A090710</u> | <u>001</u> | Mar 15, 2012 |
| <u>AB</u> | | <u>10MG</u> | <u>A090710</u> | <u>002</u> | Mar 15, 2012 |
| <u>AB</u> | | <u>20MG</u> | <u>A090710</u> | <u>003</u> | Mar 15, 2012 |

RITALIN

| | | | | | |
|-----------|---|----------|-------------|----------------|------------|
| <u>AB</u> | + | NOVARTIS | <u>5MG</u> | <u>N010187</u> | <u>003</u> |
| <u>AB</u> | + | | <u>10MG</u> | <u>N010187</u> | <u>006</u> |
| <u>AB</u> | + | ! | <u>20MG</u> | <u>N010187</u> | <u>010</u> |

TABLET, CHEWABLE; ORAL

METHYLPHENIDATE HYDROCHLORIDE

| | | | | | |
|-----------|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ASCENT PHARMS INC | <u>2.5MG</u> | <u>A210354</u> | <u>001</u> | Dec 29, 2017 |
| <u>AB</u> | | <u>5MG</u> | <u>A210354</u> | <u>002</u> | Dec 29, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A210354</u> | <u>003</u> | Dec 29, 2017 |
| <u>AB</u> | BRECKENRIDGE PHARM | <u>2.5MG</u> | <u>A204954</u> | <u>001</u> | Jan 26, 2017 |
| <u>AB</u> | | <u>5MG</u> | <u>A204954</u> | <u>002</u> | Jan 26, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A204954</u> | <u>003</u> | Jan 26, 2017 |
| <u>AB</u> | NOVEL LABS INC | <u>2.5MG</u> | <u>A204115</u> | <u>001</u> | Feb 25, 2015 |
| <u>AB</u> | | <u>5MG</u> | <u>A204115</u> | <u>002</u> | Feb 25, 2015 |
| <u>AB</u> | ! | <u>10MG</u> | <u>A204115</u> | <u>003</u> | Feb 25, 2015 |
| <u>AB</u> | RISING PHARMS | <u>2.5MG</u> | <u>A205756</u> | <u>001</u> | Nov 07, 2016 |
| <u>AB</u> | | <u>5MG</u> | <u>A205756</u> | <u>002</u> | Nov 07, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A205756</u> | <u>003</u> | Nov 07, 2016 |

TABLET, EXTENDED RELEASE; ORAL

CONCERTA

| | | | | | | |
|-----------|---|----------------|-------------|----------------|------------|--------------|
| <u>AB</u> | + | JANSSEN PHARMS | <u>18MG</u> | <u>N021121</u> | <u>001</u> | Aug 01, 2000 |
| <u>AB</u> | + | | <u>27MG</u> | <u>N021121</u> | <u>004</u> | Apr 01, 2002 |
| <u>AB</u> | + | | <u>36MG</u> | <u>N021121</u> | <u>002</u> | Aug 01, 2000 |
| <u>AB</u> | + | ! | <u>54MG</u> | <u>N021121</u> | <u>003</u> | Dec 08, 2000 |

METADATE ER

| | | | | | | |
|-----------|---|----------------|-------------|----------------|------------|--------------|
| <u>AB</u> | ! | LANNETT CO INC | <u>20MG</u> | <u>A089601</u> | <u>001</u> | Jun 01, 1988 |
|-----------|---|----------------|-------------|----------------|------------|--------------|

METHYLIN ER

| | | | | | | |
|-----------|--|------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | SPECGX LLC | <u>10MG</u> | <u>A075629</u> | <u>001</u> | May 09, 2000 |
| <u>AB</u> | | | <u>20MG</u> | <u>A075629</u> | <u>002</u> | May 09, 2000 |

METHYLPHENIDATE HYDROCHLORIDE

| | | | | | |
|-----------|--------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | ABHAI LLC | <u>10MG</u> | <u>A207488</u> | <u>001</u> | Jun 09, 2015 |
| <u>AB</u> | | <u>20MG</u> | <u>A207488</u> | <u>002</u> | Jun 09, 2015 |
| <u>AB</u> | ACTAVIS LABS FL | <u>18MG</u> | <u>A076772</u> | <u>001</u> | Mar 22, 2018 |
| <u>AB</u> | | <u>27MG</u> | <u>A076772</u> | <u>002</u> | Mar 22, 2018 |
| <u>AB</u> | | <u>36MG</u> | <u>A076772</u> | <u>003</u> | Mar 22, 2018 |
| <u>AB</u> | | <u>54MG</u> | <u>A076655</u> | <u>001</u> | Mar 21, 2018 |
| <u>AB</u> | ALVOGEN PINE BROOK | <u>18MG</u> | <u>A210818</u> | <u>001</u> | Nov 30, 2018 |
| <u>AB</u> | | <u>27MG</u> | <u>A210818</u> | <u>002</u> | Nov 30, 2018 |
| <u>AB</u> | | <u>36MG</u> | <u>A210818</u> | <u>003</u> | Nov 30, 2018 |
| <u>AB</u> | | <u>54MG</u> | <u>A210818</u> | <u>004</u> | Nov 30, 2018 |
| <u>AB</u> | AMNEAL PHARMS | <u>18MG</u> | <u>A207515</u> | <u>001</u> | Feb 01, 2018 |
| <u>AB</u> | | <u>27MG</u> | <u>A207515</u> | <u>002</u> | Feb 01, 2018 |
| <u>AB</u> | | <u>36MG</u> | <u>A207515</u> | <u>003</u> | Feb 01, 2018 |
| <u>AB</u> | | <u>54MG</u> | <u>A207515</u> | <u>004</u> | Feb 01, 2018 |
| <u>AB</u> | ANI PHARMS INC | <u>18MG</u> | <u>A208607</u> | <u>001</u> | Jul 14, 2017 |
| <u>AB</u> | | <u>27MG</u> | <u>A208607</u> | <u>002</u> | Jul 14, 2017 |
| <u>AB</u> | | <u>36MG</u> | <u>A208607</u> | <u>003</u> | Jul 14, 2017 |
| <u>AB</u> | | <u>54MG</u> | <u>A208607</u> | <u>004</u> | Jul 14, 2017 |
| <u>AB</u> | CNTY LINE PHARMS | <u>10MG</u> | <u>A204772</u> | <u>001</u> | Feb 29, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A204772</u> | <u>002</u> | Feb 29, 2016 |
| <u>AB</u> | GRANULES PHARMS | <u>10MG</u> | <u>A210992</u> | <u>001</u> | Nov 21, 2018 |
| <u>AB</u> | | <u>20MG</u> | <u>A210992</u> | <u>002</u> | Nov 21, 2018 |
| <u>AB</u> | MYLAN PHARMS INC | <u>18MG</u> | <u>A206726</u> | <u>001</u> | Oct 21, 2016 |
| <u>AB</u> | | <u>27MG</u> | <u>A206726</u> | <u>002</u> | Oct 21, 2016 |
| <u>AB</u> | | <u>36MG</u> | <u>A206726</u> | <u>003</u> | Oct 21, 2016 |
| <u>AB</u> | | <u>54MG</u> | <u>A206726</u> | <u>004</u> | Oct 21, 2016 |
| <u>AB</u> | OSMOTICA | <u>18MG</u> | <u>A205327</u> | <u>001</u> | Jul 28, 2017 |
| <u>AB</u> | | <u>27MG</u> | <u>A205327</u> | <u>002</u> | Jul 28, 2017 |
| <u>AB</u> | | <u>36MG</u> | <u>A205327</u> | <u>003</u> | Jul 28, 2017 |
| <u>AB</u> | | <u>54MG</u> | <u>A205327</u> | <u>004</u> | Jul 28, 2017 |
| BX | LANNETT CO INC | 18MG | A091695 | 001 | Jul 09, 2013 |
| BX | | 27MG | A091695 | 002 | Jul 09, 2013 |

PRESCRIPTION DRUG PRODUCT LIST

METHYLPHENIDATE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

| | | | | |
|----|------------|------|-------------|--------------|
| BX | | 36MG | A091695 003 | Sep 23, 2013 |
| BX | | 54MG | A091695 004 | Sep 23, 2013 |
| BX | SPECGX LLC | 27MG | A202608 001 | Dec 28, 2012 |
| BX | | 36MG | A202608 002 | Dec 28, 2012 |
| BX | | 54MG | A202608 003 | Dec 28, 2012 |
| ! | OSMOTICA | 72MG | A205327 005 | Jul 28, 2017 |

TABLET, EXTENDED RELEASE, CHEWABLE;ORAL

QUILLICHEW ER

| | | | | |
|---|-----------------|------|-------------|--------------|
| + | NEXTWAVE PHARMS | 20MG | N207960 001 | Dec 04, 2015 |
| + | | 30MG | N207960 002 | Dec 04, 2015 |
| + | ! | 40MG | N207960 003 | Dec 04, 2015 |

METHYLPREDNISOLONE

TABLET;ORAL

MEDROL

| | | | | |
|-----------|---|-------------------------|-------------|--------------------|
| AB | + | PHARMACIA AND UPJOHN | 4MG | N011153 001 |
| AB | + | | 8MG | N011153 004 |
| AB | + | | 16MG | N011153 003 |
| AB | + | ! | 32MG | N011153 006 |

METHYLPREDNISOLONE

| | | | | | |
|-----------|--|-------------------------|-------------|--------------------|--------------|
| AB | | DURAMED PHARMS BARR | 4MG | A088497 001 | Feb 21, 1984 |
| AB | | JUBILANT CADISTA | 4MG | A040189 001 | Oct 31, 1997 |
| AB | | | 8MG | A040189 002 | Oct 31, 1997 |
| AB | | | 16MG | A040189 003 | Jul 20, 2007 |
| AB | | | 32MG | A040189 004 | Jul 20, 2007 |
| AB | | RICONPHARMA LLC | 4MG | A210985 001 | Jan 09, 2019 |
| AB | | SANDOZ | 4MG | A040194 001 | Oct 31, 1997 |
| AB | | VINTAGE PHARMS | 4MG | A040183 001 | Dec 22, 1998 |
| AB | | WATSON LABS | 4MG | A040232 001 | Oct 16, 1997 |
| AB | | ZYDUS PHARMS USA INC | 4MG | A206751 001 | Apr 23, 2018 |
| AB | | | 8MG | A206751 002 | Apr 23, 2018 |
| AB | | | 16MG | A206751 003 | Apr 23, 2018 |
| AB | | | 32MG | A206751 004 | Apr 23, 2018 |

MEDROL

| | | | |
|---|-------------------------|-----|-------------|
| + | PHARMACIA AND UPJOHN | 2MG | N011153 002 |
|---|-------------------------|-----|-------------|

METHYLPREDNISOLONE ACETATE

INJECTABLE; INJECTION

DEPO-MEDROL

| | | | | | |
|-----------|---|---|-------------------------|----------------|--------------------|
| AB | + | ! | PHARMACIA AND UPJOHN | 20MG/ML | N011757 002 |
| AB | + | ! | | 40MG/ML | N011757 001 |
| AB | + | ! | | 80MG/ML | N011757 004 |

METHYLPREDNISOLONE ACETATE

| | | | | | |
|-----------|--|-----------------|----------------|--------------------|--------------|
| AB | | SAGENT PHARMS | 20MG/ML | A201835 001 | Jun 27, 2018 |
| AB | | | 40MG/ML | A201835 002 | Jun 27, 2018 |
| AB | | | 80MG/ML | A201835 003 | Jun 27, 2018 |
| AB | | SANDOZ INC | 40MG/ML | A040719 001 | Jan 29, 2009 |
| AB | | | 40MG/ML | A040794 001 | Mar 05, 2009 |
| AB | | | 80MG/ML | A040719 002 | Jan 29, 2009 |
| AB | | | 80MG/ML | A040794 002 | Mar 05, 2009 |
| AB | | TEVA PHARMS USA | 40MG/ML | A040557 001 | Feb 23, 2005 |
| AB | | | 40MG/ML | A040620 001 | Oct 27, 2006 |
| AB | | | 80MG/ML | A040557 002 | Feb 23, 2005 |
| AB | | | 80MG/ML | A040620 002 | Oct 27, 2006 |

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

A-METHAPRED

| | | | | | |
|-----------|--|---------|---------------------------|--------------------|--------------|
| AP | | HOSPIRA | EQ 40MG BASE/VIAL | A040664 001 | Dec 20, 2005 |
| AP | | | EQ 125MG BASE/VIAL | A040665 001 | Dec 20, 2005 |

METHYLPREDNISOLONE SODIUM SUCCINATE

| | | | | | |
|-----------|--|-------------------------|---------------------------|--------------------|--------------|
| AP | | AMNEAL PHARMS CO | EQ 40MG BASE/VIAL | A207549 001 | Nov 09, 2016 |
| AP | | | EQ 125MG BASE/VIAL | A207549 002 | Nov 09, 2016 |
| AP | | AUROBINDO PHARMA LTD | EQ 40MG BASE/VIAL | A207667 001 | Dec 15, 2015 |
| AP | | | EQ 125MG BASE/VIAL | A207667 002 | Dec 15, 2015 |
| AP | | | EQ 500MG BASE/VIAL | A207667 003 | Dec 15, 2015 |
| AP | | | EQ 2GM BASE/VIAL | A207667 004 | Dec 15, 2015 |

PRESCRIPTION DRUG PRODUCT LIST

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

METHYLPREDNISOLONE SODIUM SUCCINATE

| | | | | |
|-----------|--------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 40MG BASE/VIAL</u> | <u>A040583 001</u> | Jul 30, 2004 |
| <u>AP</u> | | <u>EQ 125MG BASE/VIAL</u> | <u>A040583 002</u> | Jul 30, 2004 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A040612 001</u> | Aug 12, 2004 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>EQ 500MG BASE/VIAL</u> | <u>A202691 001</u> | Feb 16, 2016 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A202691 002</u> | Feb 16, 2016 |
| <u>AP</u> | SAGENT PHARMS | <u>EQ 40MG BASE/VIAL</u> | <u>A040888 001</u> | Jul 18, 2011 |
| <u>AP</u> | | <u>EQ 125MG BASE/VIAL</u> | <u>A040888 002</u> | Jul 18, 2011 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A040888 003</u> | Jul 18, 2011 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A040888 004</u> | Jul 18, 2011 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A040888 005</u> | Jul 18, 2011 |

SOLU-MEDROL

| | | | | |
|-----------|----------------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | +! PHARMACIA AND UPJOHN | <u>EQ 40MG BASE/VIAL</u> | <u>N011856 003</u> | |
| <u>AP</u> | +! | <u>EQ 125MG BASE/VIAL</u> | <u>N011856 004</u> | |
| <u>AP</u> | +! | <u>EQ 500MG BASE/VIAL</u> | <u>N011856 005</u> | |
| <u>AP</u> | +! | <u>EQ 1GM BASE/VIAL</u> | <u>N011856 006</u> | |
| <u>AP</u> | +! | <u>EQ 2GM BASE/VIAL</u> | <u>N011856 007</u> | Feb 27, 1985 |

METHYLTESTOSTERONE

CAPSULE; ORAL

METHYLTESTOSTERONE

| | | | | |
|-----------|----------------------|-------------|--------------------|--------------|
| <u>AB</u> | IMPAX LABS INC | <u>10MG</u> | <u>A204851 001</u> | Sep 21, 2015 |
| | <u>TESTRED</u> | | | |
| <u>AB</u> | ! VALEANT PHARM INTL | <u>10MG</u> | <u>A083976 001</u> | |
| | TABLET; ORAL | | | |
| | ANDROID 25 | | | |
| BP | VALEANT PHARM INTL | 25MG | A087147 001 | |
| | METHYLTESTOSTERONE | | | |
| BP | IMPAX LABS | 10MG | A080767 002 | |

METIPRANOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

METIPRANOLOL

| | | | | |
|-----------|---------------------|-------------|--------------------|--------------|
| <u>AT</u> | SANDOZ INC | <u>0.3%</u> | <u>A075720 001</u> | Aug 06, 2001 |
| | <u>OPTIPRANOLOL</u> | | | |
| <u>AT</u> | +! BAUSCH AND LOMB | <u>0.3%</u> | <u>N019907 001</u> | Dec 29, 1989 |

METOCLOPRAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

METOCLOPRAMIDE

| | | | | |
|-----------|-------------------------------------|-----------------------|--------------------|--------------|
| <u>AP</u> | EMCURE PHARMS LTD | <u>EQ 5MG BASE/ML</u> | <u>A204756 001</u> | Dec 20, 2013 |
| | <u>METOCLOPRAMIDE HYDROCHLORIDE</u> | | | |
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 5MG BASE/ML</u> | <u>A091392 001</u> | Apr 19, 2013 |
| <u>AP</u> | ! HOSPIRA | <u>EQ 5MG BASE/ML</u> | <u>A073118 001</u> | Jan 17, 1991 |
| <u>AP</u> | TEVA PHARMS USA | <u>EQ 5MG BASE/ML</u> | <u>A073135 001</u> | Nov 27, 1991 |

SOLUTION; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

| | | | | |
|-----------|--------------------|------------------------|--------------------|--------------|
| <u>AA</u> | ANI PHARMS | <u>EQ 5MG BASE/5ML</u> | <u>A071402 001</u> | Jun 25, 1993 |
| <u>AA</u> | PHARM ASSOC | <u>EQ 5MG BASE/5ML</u> | <u>A072744 001</u> | May 28, 1991 |
| <u>AA</u> | VISTAPHARM | <u>EQ 5MG BASE/5ML</u> | <u>A075051 001</u> | Jan 26, 2001 |
| <u>AA</u> | ! WOCKHARDT BIO AG | <u>EQ 5MG BASE/5ML</u> | <u>A074703 001</u> | Oct 31, 1997 |

TABLET; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

| | | | | |
|-----------|----------------|---------------------|--------------------|--------------|
| <u>AB</u> | IMPAX LABS INC | <u>EQ 5MG BASE</u> | <u>A071250 002</u> | Dec 28, 1995 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A071250 001</u> | Feb 03, 1988 |
| <u>AB</u> | IPCA LABS LTD | <u>EQ 5MG BASE</u> | <u>A078807 001</u> | Jun 12, 2008 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A078807 002</u> | Jun 12, 2008 |
| <u>AB</u> | PAR PHARM INC | <u>EQ 10MG BASE</u> | <u>A070581 001</u> | Oct 17, 1985 |
| <u>AB</u> | TEVA | <u>EQ 5MG BASE</u> | <u>A072801 001</u> | Jun 15, 1993 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A070184 001</u> | Jul 29, 1985 |
| <u>AB</u> | VINTAGE PHARMS | <u>EQ 5MG BASE</u> | <u>A077878 001</u> | Aug 28, 2006 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A077878 002</u> | Aug 28, 2006 |

REGLAN

| | | | | |
|-----------|--------------|---------------------|--------------------|--------------|
| <u>AB</u> | + ANI PHARMS | <u>EQ 5MG BASE</u> | <u>N017854 002</u> | May 05, 1987 |
| <u>AB</u> | +! | <u>EQ 10MG BASE</u> | <u>N017854 001</u> | |

TABLET, ORALLY DISINTEGRATING; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

| | | | | |
|--|----------------|--------------|-------------|--------------|
| | NOVEL LABS INC | EQ 5MG BASE | A202191 001 | Aug 15, 2014 |
| | ! | EQ 10MG BASE | A202191 002 | Aug 15, 2014 |

PRESCRIPTION DRUG PRODUCT LIST

METOLAZONE

TABLET; ORAL

METOLAZONE

| | | | | |
|-----------|--------|--------------|--------------------|--------------|
| <u>AB</u> | MYLAN | <u>2.5MG</u> | <u>A076698 001</u> | Dec 23, 2003 |
| <u>AB</u> | | <u>5MG</u> | <u>A076698 002</u> | Oct 19, 2004 |
| <u>AB</u> | | <u>10MG</u> | <u>A076698 003</u> | Oct 19, 2004 |
| <u>AB</u> | SANDOZ | <u>2.5MG</u> | <u>A076732 001</u> | Dec 19, 2003 |
| <u>AB</u> | | <u>5MG</u> | <u>A076466 001</u> | Dec 19, 2003 |
| <u>AB</u> | | <u>10MG</u> | <u>A076466 002</u> | Dec 19, 2003 |

ZAROXOLYN

| | | | | |
|-----------|---|----------------|--------------|--------------------|
| <u>AB</u> | + | LANNETT CO INC | <u>2.5MG</u> | <u>N017386 001</u> |
| <u>AB</u> | + | ! | <u>5MG</u> | <u>N017386 002</u> |
| <u>AB</u> | + | ! | <u>10MG</u> | <u>N017386 003</u> |

METOPROLOL SUCCINATE

CAPSULE, EXTENDED RELEASE; ORAL

KAPSPARGO SPRINKLE

| | | | | |
|---|------|-------------------|-------------|--------------|
| + | SPIL | EQ 25MG TARTRATE | N210428 001 | Jan 26, 2018 |
| + | | EQ 50MG TARTRATE | N210428 002 | Jan 26, 2018 |
| + | | EQ 100MG TARTRATE | N210428 003 | Jan 26, 2018 |
| + | ! | EQ 200MG TARTRATE | N210428 004 | Jan 26, 2018 |

TABLET, EXTENDED RELEASE; ORAL

METOPROLOL SUCCINATE

| | | | | |
|-----------|-------------------------|--------------------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS ELIZABETH | <u>EQ 25MG TARTRATE</u> | <u>A204161 001</u> | Nov 25, 2016 |
| <u>AB</u> | | <u>EQ 50MG TARTRATE</u> | <u>A204161 002</u> | Nov 25, 2016 |
| <u>AB</u> | | <u>EQ 100MG TARTRATE</u> | <u>A204161 003</u> | Nov 25, 2016 |
| <u>AB</u> | | <u>EQ 200MG TARTRATE</u> | <u>A204161 004</u> | Nov 25, 2016 |
| <u>AB</u> | ACTAVIS LABS FL INC | <u>EQ 25MG TARTRATE</u> | <u>A076862 002</u> | Aug 03, 2009 |
| <u>AB</u> | | <u>EQ 50MG TARTRATE</u> | <u>A076862 001</u> | Aug 03, 2009 |
| <u>AB</u> | | <u>EQ 100MG TARTRATE</u> | <u>A077298 001</u> | Apr 15, 2010 |
| <u>AB</u> | | <u>EQ 200MG TARTRATE</u> | <u>A077298 002</u> | Apr 15, 2010 |
| <u>AB</u> | CIPLA | <u>EQ 50MG TARTRATE</u> | <u>A207465 001</u> | Oct 26, 2018 |
| <u>AB</u> | | <u>EQ 100MG TARTRATE</u> | <u>A207465 002</u> | Oct 26, 2018 |
| <u>AB</u> | | <u>EQ 200MG TARTRATE</u> | <u>A207465 003</u> | Oct 26, 2018 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 25MG TARTRATE</u> | <u>A090617 001</u> | Aug 01, 2012 |
| <u>AB</u> | | <u>EQ 50MG TARTRATE</u> | <u>A090617 002</u> | Aug 01, 2012 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 25MG TARTRATE</u> | <u>A202033 001</u> | Dec 15, 2011 |
| <u>AB</u> | | <u>EQ 50MG TARTRATE</u> | <u>A202033 002</u> | Dec 15, 2011 |
| <u>AB</u> | | <u>EQ 100MG TARTRATE</u> | <u>A202033 003</u> | Dec 15, 2011 |
| <u>AB</u> | | <u>EQ 200MG TARTRATE</u> | <u>A202033 004</u> | Dec 15, 2011 |
| <u>AB</u> | NOVAST LABS | <u>EQ 25MG TARTRATE</u> | <u>A204106 001</u> | Feb 06, 2018 |
| <u>AB</u> | | <u>EQ 50MG TARTRATE</u> | <u>A204106 002</u> | Feb 06, 2018 |
| <u>AB</u> | | <u>EQ 100MG TARTRATE</u> | <u>A204106 003</u> | Feb 06, 2018 |
| <u>AB</u> | | <u>EQ 200MG TARTRATE</u> | <u>A204106 004</u> | Feb 06, 2018 |
| <u>AB</u> | REDDYS | <u>EQ 100MG TARTRATE</u> | <u>A078889 001</u> | Aug 15, 2012 |
| <u>AB</u> | | <u>EQ 200MG TARTRATE</u> | <u>A078889 002</u> | Aug 15, 2012 |
| <u>AB</u> | TWI PHARMS | <u>EQ 25MG TARTRATE</u> | <u>A207206 001</u> | Dec 19, 2018 |
| <u>AB</u> | | <u>EQ 50MG TARTRATE</u> | <u>A207206 002</u> | Dec 19, 2018 |
| <u>AB</u> | | <u>EQ 100MG TARTRATE</u> | <u>A207206 003</u> | Dec 19, 2018 |
| <u>AB</u> | | <u>EQ 200MG TARTRATE</u> | <u>A207206 004</u> | Dec 19, 2018 |
| <u>AB</u> | WOCKHARDT | <u>EQ 25MG TARTRATE</u> | <u>A090615 001</u> | Jul 22, 2010 |
| <u>AB</u> | | <u>EQ 50MG TARTRATE</u> | <u>A090615 002</u> | Jul 22, 2010 |
| <u>AB</u> | | <u>EQ 100MG TARTRATE</u> | <u>A090615 003</u> | Jul 22, 2010 |
| <u>AB</u> | | <u>EQ 200MG TARTRATE</u> | <u>A090615 004</u> | Jul 22, 2010 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 25MG TARTRATE</u> | <u>A203894 001</u> | Mar 23, 2018 |
| <u>AB</u> | | <u>EQ 50MG TARTRATE</u> | <u>A203894 002</u> | Mar 23, 2018 |
| <u>AB</u> | | <u>EQ 100MG TARTRATE</u> | <u>A203894 003</u> | Mar 23, 2018 |
| <u>AB</u> | | <u>EQ 200MG TARTRATE</u> | <u>A203894 004</u> | Mar 23, 2018 |

TOPROL-XL

| | | | | | |
|-----------|---|---------------|--------------------------|--------------------|--------------|
| <u>AB</u> | + | ARALEZ PHARMS | <u>EQ 25MG TARTRATE</u> | <u>N019962 004</u> | Feb 05, 2001 |
| <u>AB</u> | + | ! | <u>EQ 50MG TARTRATE</u> | <u>N019962 001</u> | Jan 10, 1992 |
| <u>AB</u> | + | | <u>EQ 100MG TARTRATE</u> | <u>N019962 002</u> | Jan 10, 1992 |
| <u>AB</u> | + | ! | <u>EQ 200MG TARTRATE</u> | <u>N019962 003</u> | Jan 10, 1992 |

METOPROLOL TARTRATE

INJECTABLE; INJECTION

LOPRESSOR

| | | | | | |
|-----------|---|----------|---------------|--------------------|--------------|
| <u>AP</u> | + | NOVARTIS | <u>1MG/ML</u> | <u>N018704 001</u> | Mar 30, 1984 |
|-----------|---|----------|---------------|--------------------|--------------|

METOPROLOL TARTRATE

| | | | | |
|-----------|-------------------------|---------------|--------------------|--------------|
| <u>AP</u> | BAXTER HLTHCARE CORP | <u>1MG/ML</u> | <u>A078950 001</u> | Apr 29, 2013 |
| <u>AP</u> | FRESENIUS KABI USA | <u>1MG/ML</u> | <u>A091045 001</u> | Oct 25, 2010 |
| <u>AP</u> | GLAND PHARMA LTD | <u>1MG/ML</u> | <u>A204205 001</u> | Aug 27, 2014 |

PRESCRIPTION DRUG PRODUCT LIST

METOPROLOL TARTRATE

INJECTABLE; INJECTION

METOPROLOL TARTRATE

| | | | | | |
|-----------|-------------------------|---------------|----------------|------------|--------------|
| <u>AP</u> | HIKMA FARMACEUTICA | <u>1MG/ML</u> | <u>A077761</u> | <u>001</u> | May 30, 2007 |
| <u>AP</u> | HOSPIRA | <u>1MG/ML</u> | <u>A074133</u> | <u>001</u> | Dec 21, 1993 |
| <u>AP</u> | | <u>1MG/ML</u> | <u>A075160</u> | <u>001</u> | Jul 06, 1998 |
| <u>AP</u> | | <u>1MG/ML</u> | <u>A078085</u> | <u>001</u> | Apr 29, 2008 |
| <u>AP</u> | LUITPOLD | <u>1MG/ML</u> | <u>A090386</u> | <u>001</u> | Sep 30, 2009 |
| <u>AP</u> | | <u>1MG/ML</u> | <u>A091307</u> | <u>001</u> | Dec 29, 2010 |
| <u>AP</u> | MYLAN ASI | <u>1MG/ML</u> | <u>A090317</u> | <u>001</u> | Apr 19, 2010 |
| <u>AP</u> | SANDOZ INC | <u>1MG/ML</u> | <u>A077360</u> | <u>001</u> | Oct 02, 2007 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>1MG/ML</u> | <u>A076495</u> | <u>001</u> | Jul 07, 2003 |

TABLET; ORAL

LOPRESSOR

| | | | | | |
|-----------|----------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + US PHARMS HOLDINGS I | <u>50MG</u> | <u>N017963</u> | <u>001</u> | |
| <u>AB</u> | + | <u>100MG</u> | <u>N017963</u> | <u>002</u> | |
| | <u>METOPROLOL TARTRATE</u> | | | | |
| <u>AB</u> | ALEMBIC PHARMS LTD | <u>25MG</u> | <u>A202871</u> | <u>001</u> | May 28, 2013 |
| <u>AB</u> | | <u>50MG</u> | <u>A202871</u> | <u>002</u> | May 28, 2013 |
| <u>AB</u> | | <u>100MG</u> | <u>A202871</u> | <u>003</u> | May 28, 2013 |
| <u>AB</u> | AUROBINDO PHARMA | <u>25MG</u> | <u>A077739</u> | <u>001</u> | Sep 11, 2007 |
| <u>AB</u> | | <u>50MG</u> | <u>A077739</u> | <u>002</u> | Sep 11, 2007 |
| <u>AB</u> | | <u>100MG</u> | <u>A077739</u> | <u>003</u> | Sep 11, 2007 |
| <u>AB</u> | IPCA LABS LTD | <u>25MG</u> | <u>A078459</u> | <u>001</u> | Jun 17, 2008 |
| <u>AB</u> | | <u>50MG</u> | <u>A078459</u> | <u>002</u> | Jun 17, 2008 |
| <u>AB</u> | | <u>100MG</u> | <u>A078459</u> | <u>003</u> | Jun 17, 2008 |
| <u>AB</u> | MYLAN | <u>25MG</u> | <u>A076704</u> | <u>001</u> | Jan 16, 2004 |
| <u>AB</u> | | <u>50MG</u> | <u>A076704</u> | <u>002</u> | Jan 16, 2004 |
| <u>AB</u> | ! | <u>100MG</u> | <u>A076704</u> | <u>003</u> | Jan 16, 2004 |
| <u>AB</u> | RUBICON RES PVT LTD | <u>25MG</u> | <u>A200981</u> | <u>001</u> | Oct 28, 2014 |
| <u>AB</u> | | <u>50MG</u> | <u>A200981</u> | <u>002</u> | Oct 28, 2014 |
| <u>AB</u> | | <u>100MG</u> | <u>A200981</u> | <u>003</u> | Oct 28, 2014 |
| <u>AB</u> | SUN PHARM INDS INC | <u>25MG</u> | <u>A076670</u> | <u>001</u> | Jan 15, 2004 |
| <u>AB</u> | | <u>50MG</u> | <u>A074644</u> | <u>001</u> | Dec 10, 1996 |
| <u>AB</u> | | <u>100MG</u> | <u>A074644</u> | <u>002</u> | Dec 10, 1996 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>25MG</u> | <u>A073654</u> | <u>002</u> | Jul 15, 2009 |
| <u>AB</u> | | <u>50MG</u> | <u>A073654</u> | <u>003</u> | Dec 21, 1993 |
| <u>AB</u> | | <u>100MG</u> | <u>A073654</u> | <u>001</u> | Dec 21, 1993 |
| <u>AB</u> | TEVA | <u>50MG</u> | <u>A074141</u> | <u>001</u> | Jan 31, 1995 |
| <u>AB</u> | | <u>100MG</u> | <u>A074141</u> | <u>002</u> | Jan 31, 1995 |
| <u>AB</u> | WATSON LABS | <u>50MG</u> | <u>A074217</u> | <u>001</u> | May 27, 1994 |
| <u>AB</u> | | <u>100MG</u> | <u>A074217</u> | <u>002</u> | May 27, 1994 |
| | MYLAN | 37.5MG | A076704 | 004 | Mar 18, 2015 |
| | | 75MG | A076704 | 005 | Mar 18, 2015 |

METRONIDAZOLE

CAPSULE; ORAL

FLAGYL

| | | | | | |
|-----------|------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | +! GD SEARLE LLC | <u>375MG</u> | <u>N020334</u> | <u>001</u> | May 03, 1995 |
|-----------|------------------|--------------|----------------|------------|--------------|

METRONIDAZOLE

| | | | | | |
|-----------|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ALEMBIC PHARMS LTD | <u>375MG</u> | <u>A079065</u> | <u>001</u> | Jun 23, 2009 |
| <u>AB</u> | PAR PHARM | <u>375MG</u> | <u>A076522</u> | <u>001</u> | Jan 29, 2004 |

CREAM; TOPICAL

METROCREAM

| | | | | | |
|-----------|---------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | +! GALDERMA LABS LP | <u>0.75%</u> | <u>N020531</u> | <u>001</u> | Sep 20, 1995 |
|-----------|---------------------|--------------|----------------|------------|--------------|

METRONIDAZOLE

| | | | | | |
|-----------|----------------|--------------|----------------|------------|--------------|
| <u>AB</u> | FOUGERA PHARMS | <u>0.75%</u> | <u>A076408</u> | <u>001</u> | May 28, 2004 |
| <u>AB</u> | G AND W LABS | <u>0.75%</u> | <u>A077549</u> | <u>001</u> | Dec 19, 2007 |

NORITATE

| | | | | | |
|----|-------------------------|----|---------|-----|--------------|
| +! | VALEANT PHARMS NORTH | 1% | N020743 | 001 | Sep 26, 1997 |
|----|-------------------------|----|---------|-----|--------------|

GEL; TOPICAL

METROGEL

| | | | | | |
|-----------|---------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | +! GALDERMA LABS LP | <u>0.75%</u> | <u>N019737</u> | <u>001</u> | Nov 22, 1988 |
| <u>AB</u> | +! | <u>1%</u> | <u>N021789</u> | <u>001</u> | Jun 30, 2005 |

METRONIDAZOLE

| | | | | | |
|-----------|------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | FOUGERA PHARMS | <u>0.75%</u> | <u>A077018</u> | <u>001</u> | Jun 06, 2006 |
| <u>AB</u> | G AND W LABS INC | <u>0.75%</u> | <u>A078178</u> | <u>001</u> | Jan 19, 2011 |
| <u>AB</u> | TARO | <u>0.75%</u> | <u>A077819</u> | <u>001</u> | Jul 18, 2006 |
| <u>AB</u> | TARO PHARM | <u>1%</u> | <u>A204651</u> | <u>001</u> | Mar 14, 2017 |
| <u>AB</u> | TOLMAR | <u>0.75%</u> | <u>A077547</u> | <u>001</u> | Jul 13, 2006 |

PRESCRIPTION DRUG PRODUCT LIST

METRONIDAZOLE

GEL; TOPICAL

METRONIDAZOLE

| | | | | |
|-----------|--|-----------|--------------------|--------------|
| AB | | 1% | A090903 001 | Jul 22, 2011 |
|-----------|--|-----------|--------------------|--------------|

GEL; VAGINAL

METROGEL-VAGINAL

| | | | | | |
|-----------|----------|----------------|--------------|--------------------|--------------|
| AB | + | MEDICIS | 0.75% | N020208 001 | Aug 17, 1992 |
|-----------|----------|----------------|--------------|--------------------|--------------|

METRONIDAZOLE

| | | | | | |
|-----------|--|---------------|--------------|--------------------|--------------|
| AB | | TOLMAR | 0.75% | A077264 001 | Oct 31, 2006 |
|-----------|--|---------------|--------------|--------------------|--------------|

VANDAZOLE

| | | | | | |
|-----------|--|--------------------|--------------|--------------------|--------------|
| BX | | TEVA PHARMS | 0.75% | N021806 001 | May 20, 2005 |
|-----------|--|--------------------|--------------|--------------------|--------------|

NUVESSA

| | | | | |
|----------|--------------------------|-------------|--------------------|--------------|
| + | CHEMO RESEARCH SL | 1.3% | N205223 001 | Mar 24, 2014 |
|----------|--------------------------|-------------|--------------------|--------------|

INJECTABLE; INJECTION

FLAGYL I.V. RTU IN PLASTIC CONTAINER

| | | | | | |
|-----------|----------|------------------------|--------------------|--------------------|--|
| AP | + | BAXTER HLTHCARE | 500MG/100ML | N018657 001 | |
|-----------|----------|------------------------|--------------------|--------------------|--|

METRO I.V. IN PLASTIC CONTAINER

| | | | | | |
|-----------|----------|----------------|--------------------|--------------------|--------------|
| AP | + | B BRAUN | 500MG/100ML | N018900 001 | Sep 29, 1983 |
|-----------|----------|----------------|--------------------|--------------------|--------------|

METRONIDAZOLE IN PLASTIC CONTAINER

| | | | | | |
|-----------|--|------------------------|--------------------|--------------------|--------------|
| AP | | BAXTER HLTHCARE | 500MG/100ML | A078084 001 | Mar 31, 2008 |
|-----------|--|------------------------|--------------------|--------------------|--------------|

CORP

| | | | | | |
|-----------|----------|----------------|--------------------|--------------------|--------------|
| AP | + | HOSPIRA | 500MG/100ML | N018890 002 | Nov 18, 1983 |
|-----------|----------|----------------|--------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-----------------------|--------------------|--------------------|--------------|
| AP | | MYLAN LABS LTD | 500MG/100ML | A205531 001 | May 09, 2017 |
|-----------|--|-----------------------|--------------------|--------------------|--------------|

LOTION; TOPICAL

METROLOTION

| | | | | | |
|-----------|----------|-------------------------|--------------|--------------------|--------------|
| AB | + | GALDERMA LABS LP | 0.75% | N020901 001 | Nov 24, 1998 |
|-----------|----------|-------------------------|--------------|--------------------|--------------|

METRONIDAZOLE

| | | | | | |
|-----------|--|-----------------------|--------------|--------------------|--------------|
| AB | | FOUGERA PHARMS | 0.75% | A077197 001 | May 24, 2006 |
|-----------|--|-----------------------|--------------|--------------------|--------------|

TABLET; ORAL

FLAGYL

| | | | | | |
|-----------|----------|----------------------|--------------|--------------------|--|
| AB | + | GD SEARLE LLC | 250MG | N012623 001 | |
|-----------|----------|----------------------|--------------|--------------------|--|

| | | | | | |
|-----------|----------|--|--------------|--------------------|--|
| AB | + | | 500MG | N012623 003 | |
|-----------|----------|--|--------------|--------------------|--|

METRONIDAZOLE

| | | | | | |
|-----------|--|---------------------------|--------------|--------------------|--------------|
| AB | | ALEMBIC PHARMS LTD | 250MG | A079067 001 | Mar 13, 2009 |
|-----------|--|---------------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 500MG | A079067 002 | Mar 13, 2009 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------------------------|--------------|--------------------|--------------|
| AB | | AUROBINDO PHARMA | 250MG | A203974 001 | May 29, 2015 |
|-----------|--|-------------------------|--------------|--------------------|--------------|

LTD

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 500MG | A203974 002 | May 29, 2015 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------------------|--------------|--------------------|--------------|
| AB | | CADILA PHARMS LTD | 250MG | A209794 001 | Dec 12, 2017 |
|-----------|--|--------------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 500MG | A209794 002 | Dec 12, 2017 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------------------|--------------|--------------------|--------------|
| AB | | FLAMINGO PHARMS | 250MG | A207309 001 | May 16, 2016 |
|-----------|--|------------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 500MG | A207309 002 | May 16, 2016 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------------------|--------------|--------------------|--------------|
| AB | | HERITAGE PHARMS INC | 250MG | A205245 001 | Sep 23, 2015 |
|-----------|--|----------------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 500MG | A205245 002 | Sep 23, 2015 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------------|--------------|--------------------|--------------|
| AB | | INNOGENIX | 250MG | A070772 001 | Jul 16, 1986 |
|-----------|--|------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 500MG | A070772 002 | Jul 16, 1986 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------------|--------------|--------------------|--------------|
| AB | | LUPIN LTD | 250MG | A209096 001 | Sep 12, 2017 |
|-----------|--|------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 500MG | A209096 002 | Sep 12, 2017 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------------|--------------|--------------------|--------------|
| AB | | ORIT LABS LLC | 250MG | A208681 001 | Jun 20, 2017 |
|-----------|--|----------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 500MG | A208681 002 | Jun 20, 2017 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------|--------------|--------------------|--------------|
| AB | | PLIVA | 500MG | A070033 001 | Dec 06, 1984 |
|-----------|--|--------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|-----------------------|--------------|--------------------|--------------|
| AB | | STRIDES PHARMA | 250MG | A208162 001 | May 25, 2016 |
|-----------|--|-----------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 500MG | A208162 002 | May 25, 2016 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------------------|--------------|--------------------|--------------|
| AB | | STRIDES VIVIMED | 250MG | A070040 001 | Jan 29, 1985 |
|-----------|--|------------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 500MG | A070039 001 | Jan 29, 1985 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------------------|--------------|--------------------|--------------|
| AB | | TEVA PHARMS USA | 250MG | A070027 001 | Nov 06, 1984 |
|-----------|--|------------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------------------------|--------------|--------------------|--------------|
| AB | | UNICHEM LABS LTD | 250MG | A203458 001 | Jan 22, 2014 |
|-----------|--|-------------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 500MG | A203458 002 | Jan 22, 2014 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------------|--------------|--------------------|--------------|
| AB | | WATSON LABS | 250MG | A070035 001 | Dec 20, 1984 |
|-----------|--|--------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------------------|--------------|--------------------|--------------|
| AB | | WATSON LABS INC | 500MG | A070044 001 | Feb 08, 1985 |
|-----------|--|------------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------------------------|--------------|--------------------|--------------|
| AB | | ZYDUS PHARMS USA | 250MG | A206560 001 | Nov 16, 2016 |
|-----------|--|-------------------------|--------------|--------------------|--------------|

INC

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 500MG | A206560 002 | Nov 16, 2016 |
|-----------|--|--|--------------|--------------------|--------------|

METYRAPONE

CAPSULE; ORAL

METOPIRONE

| | | | | |
|----------|------------------------|--------------|--------------------|--------------|
| + | LABORATORIE HRA | 250MG | N012911 002 | Aug 09, 1996 |
|----------|------------------------|--------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

METYROSINE

CAPSULE; ORAL

DEMSEER

+! ATON PHARMA VPNA 250MG N017871 001

MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

TEVA

150MG

A074377 001 May 16, 1995

200MG

A074377 002 May 16, 1995

!

250MG

A074377 003 May 16, 1995

MICAFUNGIN SODIUM

INJECTABLE; INTRAVENOUS

MYCAMINE

+! ASTELLAS

EQ 50MG BASE/VIAL

N021506 002 Mar 16, 2005

+!

EQ 100MG BASE/VIAL

N021506 003 Jun 27, 2006

MICONAZOLE

TABLET; BUCCAL

ORAVIG

+! MIDATECH PHARMA US 50MG

N022404 001 Apr 16, 2010

MICONAZOLE NITRATE

SUPPOSITORY; VAGINAL

MICONAZOLE NITRATEAB ACTAVIS PHARMA200MGA073508 001 Nov 19, 1993MONISTAT 3AB +! MEDTECH PRODUCTS200MGN018888 001 Aug 15, 1984MICONAZOLE NITRATE; PETROLATUM, WHITE; ZINC OXIDE

OINTMENT; TOPICAL

VUSION

+! MYLAN PHARMS INC 0.25%; 81.35%; 15%

N021026 001 Feb 16, 2006

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDEAP AKORN INCEQ 1MG BASE/MLA075494 001 Jun 30, 2000APEQ 5MG BASE/MLA075494 002 Jun 30, 2000AP FRESENIUS KABI USAEQ 1MG BASE/MLA075154 002 Jun 20, 2000APEQ 5MG BASE/MLA075154 001 Jun 20, 2000AP GLAND PHARMA LTDEQ 1MG BASE/MLA090696 001 Feb 29, 2012APEQ 5MG BASE/MLA090850 001 Jan 25, 2012AP ! HOSPIRAEQ 1MG BASE/MLA075293 001 Jun 20, 2000APEQ 1MG BASE/MLA075856 001 Jun 13, 2002AP !EQ 5MG BASE/MLA075293 002 Jun 20, 2000APEQ 5MG BASE/MLA075856 002 Jun 13, 2002AP WEST-WARD PHARMSEQ 1MG BASE/MLA075243 001 Jun 20, 2000

INT

APEQ 1MG BASE/MLA075247 002 Jun 23, 2000APEQ 1MG BASE/MLA075324 001 Jun 20, 2000APEQ 1MG BASE/MLA075421 002 Jun 20, 2000APEQ 5MG BASE/MLA075243 002 Jun 20, 2000APEQ 5MG BASE/MLA075247 001 Jun 23, 2000APEQ 5MG BASE/MLA075324 002 Jun 20, 2000APEQ 5MG BASE/MLA075421 001 Jun 20, 2000MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREEAP FRESENIUS KABI USAEQ 1MG BASE/MLA203460 001 Aug 22, 2014APEQ 5MG BASE/MLA203460 002 Aug 22, 2014AP ! HOSPIRAEQ 1MG BASE/MLA075857 001 Jul 22, 2002AP !EQ 5MG BASE/MLA075857 002 Jul 22, 2002AP MYLAN ASIEQ 1MG BASE/MLA090315 001 Nov 29, 2010APEQ 5MG BASE/MLA090315 002 Nov 29, 2010MIDOZALAM HYDROCHLORIDEAP SAGENT STRIDESEQ 1MG BASE/MLA090316 001 May 04, 2011APEQ 5MG BASE/MLA090316 002 May 04, 2011

MIDAZOLAM HYDROCHLORIDE

FRESENIUS KABI USA

EQ 5MG BASE/ML

A208878 001 Mar 28, 2017

SOLUTION; INTRAMUSCULAR

SEIZALAM

+! MERIDIAN MEDCL

EQ 50MG BASE/10ML (EQ 5MG BASE/ML)

N209566 001 Sep 14, 2018

TECHN

PRESCRIPTION DRUG PRODUCT LIST

MIDAZOLAM HYDROCHLORIDE

SYRUP; ORAL

MIDAZOLAM HYDROCHLORIDE

| | | | | |
|-----------|---------------------------|------------------------------|---------------------------|--------------|
| AA | HI TECH PHARMA | <u>EQ 2MG BASE/ML</u> | <u>A075958 001</u> | Sep 04, 2003 |
| AA | PADDOCK LLC | <u>EQ 2MG BASE/ML</u> | <u>A076379 001</u> | May 02, 2005 |
| AA | ! WEST-WARD PHARMS INT | <u>EQ 2MG BASE/ML</u> | <u>A075873 001</u> | Apr 30, 2002 |

MIDODRINE HYDROCHLORIDE

TABLET; ORAL

MIDODRINE HYDROCHLORIDE

| | | | | |
|-----------------------|-------------------|---------------------|---------------------------|--------------|
| AB | APOTEX INC | <u>2.5MG</u> | <u>A077746 001</u> | Sep 12, 2006 |
| AB | | <u>5MG</u> | <u>A077746 002</u> | Sep 12, 2006 |
| AB | | <u>10MG</u> | <u>A077746 003</u> | Sep 12, 2006 |
| AB | CASI PHARMS INC | <u>2.5MG</u> | <u>A076514 001</u> | Sep 11, 2003 |
| AB | | <u>5MG</u> | <u>A076514 002</u> | Sep 11, 2003 |
| AB | | <u>10MG</u> | <u>A076514 003</u> | Jul 02, 2004 |
| AB | IMPAX PHARMS | <u>2.5MG</u> | <u>A076449 001</u> | May 27, 2004 |
| AB | | <u>5MG</u> | <u>A076449 002</u> | May 27, 2004 |
| AB | | <u>10MG</u> | <u>A076449 003</u> | Dec 16, 2005 |
| AB | MYLAN PHARMS INC | <u>2.5MG</u> | <u>A076577 001</u> | Sep 10, 2003 |
| AB | ! | <u>5MG</u> | <u>A076577 002</u> | Sep 10, 2003 |
| AB | | <u>10MG</u> | <u>A076577 003</u> | Sep 10, 2003 |
| AB | PAR PHARM INC | <u>2.5MG</u> | <u>A207169 001</u> | Oct 29, 2018 |
| AB | | <u>5MG</u> | <u>A207169 002</u> | Oct 29, 2018 |
| AB | | <u>10MG</u> | <u>A207169 003</u> | Oct 29, 2018 |
| AB | UNIQUE PHARM LABS | <u>2.5MG</u> | <u>A207613 001</u> | Nov 02, 2018 |
| AB | | <u>5MG</u> | <u>A207613 002</u> | Nov 02, 2018 |
| AB | | <u>10MG</u> | <u>A207613 003</u> | Nov 02, 2018 |
| <u>ORVATEN</u> | | | | |
| AB | UPSHER SMITH LABS | <u>2.5MG</u> | <u>A076725 001</u> | Nov 03, 2004 |
| AB | | <u>5MG</u> | <u>A076725 002</u> | Nov 03, 2004 |
| AB | | <u>10MG</u> | <u>A076725 003</u> | Nov 03, 2004 |

MIDOSTAURIN

CAPSULE; ORAL

RYDAPT

| | | | | | |
|---|---|-------------------------|------|-------------|--------------|
| + | ! | NOVARTIS PHARMS CORP | 25MG | N207997 001 | Apr 28, 2017 |
|---|---|-------------------------|------|-------------|--------------|

MIFEPRISTONE

TABLET; ORAL

KORLYM

| | | | | | |
|---|---|----------------|-------|-------------|--------------|
| + | ! | CORCEPT THERAP | 300MG | N202107 001 | Feb 17, 2012 |
|---|---|----------------|-------|-------------|--------------|

MIFEPREX

| | | | | | |
|---|---|----------------|-------|-------------|--------------|
| + | ! | DANCO LABS LLC | 200MG | N020687 001 | Sep 28, 2000 |
|---|---|----------------|-------|-------------|--------------|

MIGALASTAT HYDROCHLORIDE

CAPSULE; ORAL

GALAFOLD

| | | | | | |
|---|---|-------------------|---------------|-------------|--------------|
| + | ! | AMICUS THERAPS US | EQ 123MG BASE | N208623 001 | Aug 10, 2018 |
|---|---|-------------------|---------------|-------------|--------------|

MIGLITOL

TABLET; ORAL

GLYSET

| | | | | | | |
|-----------|---|---|-------------------------|---------------------|---------------------------|--------------|
| AB | + | ! | PHARMACIA AND UPJOHN | <u>25MG</u> | <u>N020682 001</u> | Dec 18, 1996 |
| AB | + | | | <u>50MG</u> | <u>N020682 002</u> | Dec 18, 1996 |
| AB | + | | | <u>100MG</u> | <u>N020682 003</u> | Dec 18, 1996 |

MIGLITOL

| | | | | | |
|-----------|--|-------------------------|---------------------|---------------------------|--------------|
| AB | | ORIENT PHARMA CO LTD | <u>25MG</u> | <u>A203965 001</u> | Feb 24, 2015 |
| AB | | | <u>50MG</u> | <u>A203965 002</u> | Feb 24, 2015 |
| AB | | | <u>100MG</u> | <u>A203965 003</u> | Feb 24, 2015 |

MIGLUSTAT

CAPSULE; ORAL

MIGLUSTAT

| | | | | | |
|-----------|--|---------------------|---------------------|---------------------------|--------------|
| AB | | AMERIGEN PHARMS LTD | <u>100MG</u> | <u>A208342 001</u> | Apr 17, 2018 |
|-----------|--|---------------------|---------------------|---------------------------|--------------|

ZAVESCA

| | | | | | | |
|-----------|---|---|---------------------|---------------------|---------------------------|--------------|
| AB | + | ! | ACTELION PHARMS LTD | <u>100MG</u> | <u>N021348 001</u> | Jul 31, 2003 |
|-----------|---|---|---------------------|---------------------|---------------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

MILNACIPRAN HYDROCHLORIDE

TABLET; ORAL

SAVELLA

| | | | | | |
|---|--------------------|--------|---------|-----|--------------|
| + | ALLERGAN SALES LLC | 12.5MG | N022256 | 001 | Jan 14, 2009 |
| + | | 25MG | N022256 | 002 | Jan 14, 2009 |
| + | ! | 50MG | N022256 | 003 | Jan 14, 2009 |
| + | | 100MG | N022256 | 004 | Jan 14, 2009 |

MILRINONE LACTATE

INJECTABLE; INJECTION

MILRINONE LACTATE

| | | | | | |
|-----------|-------------------------|-----------------------|----------------|------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 1MG BASE/ML</u> | <u>A075936</u> | <u>001</u> | May 28, 2002 |
| <u>AP</u> | GLAND PHARMA LTD | <u>EQ 1MG BASE/ML</u> | <u>A077190</u> | <u>001</u> | Oct 31, 2006 |
| <u>AP</u> | ! | <u>EQ 1MG BASE/ML</u> | <u>A077966</u> | <u>001</u> | Dec 03, 2010 |
| <u>AP</u> | HOSPIRA INC | <u>EQ 1MG BASE/ML</u> | <u>A203280</u> | <u>001</u> | Sep 03, 2014 |
| <u>AP</u> | INTL MEDICATED | <u>EQ 1MG BASE/ML</u> | <u>A076013</u> | <u>001</u> | Aug 02, 2002 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>EQ 1MG BASE/ML</u> | <u>A075530</u> | <u>001</u> | May 28, 2002 |

AP EQ 1MG BASE/ML A075660 001 May 28, 2002MILRINONE LACTATE IN DEXTROSE 5%

| | | | | | |
|-----------|---------------------|----------------------------------------------|----------------|------------|--------------|
| <u>AP</u> | RENAISSANCE SSA LLC | <u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u> | <u>A077151</u> | <u>002</u> | Jul 20, 2005 |
|-----------|---------------------|----------------------------------------------|----------------|------------|--------------|

MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | | |
|-----------|---|-------------------------|----------------------------------------------|----------------|------------|--------------|
| <u>AP</u> | ! | <u>BAXTER HLTHCARE</u> | <u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u> | <u>A075834</u> | <u>001</u> | May 28, 2002 |
| <u>AP</u> | ! | | <u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u> | <u>A075834</u> | <u>002</u> | May 28, 2002 |
| <u>AP</u> | | HOSPIRA | <u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u> | <u>A075885</u> | <u>001</u> | May 28, 2002 |
| <u>AP</u> | | | <u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u> | <u>A075885</u> | <u>002</u> | May 28, 2002 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u> | <u>A078113</u> | <u>001</u> | May 21, 2008 |
| <u>AP</u> | | | <u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u> | <u>A078113</u> | <u>002</u> | May 21, 2008 |

MILRINONE LACTATE IN PLASTIC CONTAINER

| | | | | | | |
|-----------|--|--------------------|----------------------------------------------|----------------|------------|--------------|
| <u>AP</u> | | HIKMA FARMACEUTICA | <u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u> | <u>A090038</u> | <u>001</u> | Jan 21, 2010 |
| <u>AP</u> | | | <u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u> | <u>A090038</u> | <u>002</u> | Jan 21, 2010 |

MILTEFOSINE

CAPSULE; ORAL

IMPAVIDO

| | | | | | | |
|---|---|----------------|------|---------|-----|--------------|
| + | ! | KNIGHT THERAPS | 50MG | N204684 | 001 | Mar 19, 2014 |
|---|---|----------------|------|---------|-----|--------------|

MINOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL

DYNACIN

| | | | | | | |
|-----------|--|------------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | | CNTY LINE PHARMS | <u>EQ 50MG BASE</u> | <u>A063067</u> | <u>003</u> | Aug 14, 1990 |
| <u>AB</u> | | | <u>EQ 75MG BASE</u> | <u>A063067</u> | <u>002</u> | Sep 15, 1999 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A063067</u> | <u>001</u> | Jul 31, 1990 |

MINOCIN

| | | | | | | |
|-----------|---|------------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | + | PRECISION DERMAT | <u>EQ 50MG BASE</u> | <u>N050649</u> | <u>001</u> | May 31, 1990 |
| <u>AB</u> | + | | <u>EQ 100MG BASE</u> | <u>N050649</u> | <u>002</u> | May 31, 1990 |

MINOCYCLINE HYDROCHLORIDE

| | | | | | | |
|-----------|---|--------------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | | AUROBINDO PHARMA | <u>EQ 50MG BASE</u> | <u>A065470</u> | <u>001</u> | Mar 11, 2008 |
| <u>AB</u> | | | <u>EQ 75MG BASE</u> | <u>A065470</u> | <u>002</u> | Mar 11, 2008 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A065470</u> | <u>003</u> | Mar 11, 2008 |
| <u>AB</u> | | IMPAX LABS | <u>EQ 50MG BASE</u> | <u>A065005</u> | <u>001</u> | Mar 23, 1999 |
| <u>AB</u> | | | <u>EQ 75MG BASE</u> | <u>A065005</u> | <u>003</u> | Apr 18, 2001 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A065005</u> | <u>002</u> | Mar 23, 1999 |
| <u>AB</u> | | SUN PHARM INDS INC | <u>EQ 50MG BASE</u> | <u>A090867</u> | <u>001</u> | May 13, 2013 |
| <u>AB</u> | | | <u>EQ 75MG BASE</u> | <u>A090867</u> | <u>002</u> | May 13, 2013 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A090867</u> | <u>003</u> | May 13, 2013 |
| <u>AB</u> | | TORRENT PHARMA INC | <u>EQ 50MG BASE</u> | <u>A065062</u> | <u>001</u> | Nov 30, 2000 |
| <u>AB</u> | | | <u>EQ 75MG BASE</u> | <u>A065062</u> | <u>002</u> | Nov 30, 2000 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A065062</u> | <u>003</u> | Nov 30, 2000 |
| <u>AB</u> | | WATSON LABS | <u>EQ 75MG BASE</u> | <u>A063065</u> | <u>002</u> | Jun 10, 1999 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A063065</u> | <u>001</u> | Dec 30, 1991 |
| <u>AB</u> | | WATSON LABS TEVA | <u>EQ 50MG BASE</u> | <u>A063181</u> | <u>001</u> | Dec 30, 1991 |
| <u>AB</u> | | ZYDUS WORLDWIDE | <u>EQ 50MG BASE</u> | <u>A063011</u> | <u>001</u> | Mar 02, 1992 |
| <u>AB</u> | | | <u>EQ 75MG BASE</u> | <u>A063009</u> | <u>002</u> | Aug 12, 2003 |
| <u>AB</u> | ! | | <u>EQ 100MG BASE</u> | <u>A063009</u> | <u>001</u> | Mar 02, 1992 |

CAPSULE, EXTENDED RELEASE; ORAL

XIMINO

| | | | | | | |
|--|--|--------------------|---------------|---------|-----|--------------|
| | | SUN PHARM INDS LTD | EQ 45MG BASE | N201922 | 001 | Jul 11, 2012 |
| | | | EQ 90MG BASE | N201922 | 003 | Jul 11, 2012 |
| | | | EQ 135MG BASE | N201922 | 005 | Jul 11, 2012 |

INJECTABLE; INJECTION

MINOCIN

| | | | | | | |
|---|---|----------------|--------------------|---------|-----|--|
| + | ! | REMPLEX PHARMS | EQ 100MG BASE/VIAL | N050444 | 001 | |
|---|---|----------------|--------------------|---------|-----|--|

PRESCRIPTION DRUG PRODUCT LIST

MINOCYCLINE HYDROCHLORIDE

POWDER, EXTENDED RELEASE;DENTAL

ARESTIN

+! ORAPHARMA

EQ 1MG BASE

N050781 001 Feb 16, 2001

TABLET;ORAL

MINOCYCLINE HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 50MG BASE</u> | <u>A065436 001</u> | Dec 26, 2007 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A065436 002</u> | Dec 26, 2007 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A065436 003</u> | Dec 26, 2007 |
| <u>AB</u> | PAR PHARM | <u>EQ 50MG BASE</u> | <u>A065131 001</u> | Apr 16, 2003 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A065131 002</u> | Apr 16, 2003 |
| <u>AB</u> | ! | <u>EQ 100MG BASE</u> | <u>A065131 003</u> | Apr 16, 2003 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>EQ 50MG BASE</u> | <u>A090217 001</u> | Jan 29, 2016 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A090217 002</u> | Jan 29, 2016 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A090217 003</u> | Jan 29, 2016 |
| <u>AB</u> | TORRENT PHARMA INC | <u>EQ 50MG BASE</u> | <u>A065156 001</u> | Jan 06, 2004 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A065156 002</u> | Jan 06, 2004 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A065156 003</u> | Jan 06, 2004 |

TABLET, EXTENDED RELEASE;ORAL

MINOCYCLINE HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | ALKEM LABS LTD | <u>EQ 45MG BASE</u> | <u>A204453 001</u> | Sep 28, 2016 |
| <u>AB</u> | | <u>EQ 65MG BASE</u> | <u>A204453 006</u> | Mar 16, 2018 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A204453 002</u> | Sep 28, 2016 |
| <u>AB</u> | | <u>EQ 90MG BASE</u> | <u>A204453 003</u> | Sep 28, 2016 |
| <u>AB</u> | | <u>EQ 105MG BASE</u> | <u>A204453 004</u> | Sep 28, 2016 |
| <u>AB</u> | | <u>EQ 115MG BASE</u> | <u>A204453 007</u> | Mar 16, 2018 |
| <u>AB</u> | | <u>EQ 135MG BASE</u> | <u>A204453 005</u> | Sep 28, 2016 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 45MG BASE</u> | <u>A202261 001</u> | Nov 19, 2012 |
| <u>AB</u> | | <u>EQ 65MG BASE</u> | <u>A202261 002</u> | Sep 28, 2018 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A202261 006</u> | Jun 13, 2016 |
| <u>AB</u> | | <u>EQ 90MG BASE</u> | <u>A202261 003</u> | Nov 19, 2012 |
| <u>AB</u> | | <u>EQ 105MG BASE</u> | <u>A202261 007</u> | Jun 13, 2016 |
| <u>AB</u> | | <u>EQ 115MG BASE</u> | <u>A202261 004</u> | Sep 28, 2018 |
| <u>AB</u> | | <u>EQ 135MG BASE</u> | <u>A202261 005</u> | Nov 19, 2012 |
| <u>AB</u> | BARR LABS INC | <u>EQ 65MG BASE</u> | <u>A065485 004</u> | May 18, 2012 |
| <u>AB</u> | | <u>EQ 115MG BASE</u> | <u>A065485 005</u> | May 18, 2012 |
| <u>AB</u> | LUPIN LTD | <u>EQ 45MG BASE</u> | <u>A091424 001</u> | Nov 30, 2011 |
| <u>AB</u> | | <u>EQ 55MG BASE</u> | <u>A091424 002</u> | Nov 30, 2011 |
| <u>AB</u> | | <u>EQ 90MG BASE</u> | <u>A091424 003</u> | Nov 30, 2011 |
| <u>AB</u> | | <u>EQ 135MG BASE</u> | <u>A091424 004</u> | Nov 30, 2011 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 80MG BASE</u> | <u>A203443 002</u> | Aug 22, 2014 |
| <u>AB</u> | | <u>EQ 105MG BASE</u> | <u>A203443 003</u> | Aug 22, 2014 |
| <u>AB</u> | SANDOZ | <u>EQ 45MG BASE</u> | <u>A090422 001</u> | Aug 13, 2009 |
| <u>AB</u> | | <u>EQ 90MG BASE</u> | <u>A090422 002</u> | Aug 13, 2009 |
| <u>AB</u> | ! | <u>EQ 135MG BASE</u> | <u>A090422 003</u> | Aug 13, 2009 |
| <u>AB</u> | SIDMAK LABS INDIA | <u>EQ 45MG BASE</u> | <u>A204394 001</u> | Dec 30, 2015 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A204394 004</u> | Dec 30, 2015 |
| <u>AB</u> | | <u>EQ 105MG BASE</u> | <u>A204394 005</u> | Dec 30, 2015 |
| <u>AB</u> | | <u>EQ 135MG BASE</u> | <u>A204394 007</u> | Dec 30, 2015 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>EQ 45MG BASE</u> | <u>A091118 001</u> | Sep 25, 2014 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A091118 004</u> | Sep 25, 2014 |
| <u>AB</u> | | <u>EQ 90MG BASE</u> | <u>A091118 005</u> | Sep 25, 2014 |
| <u>AB</u> | | <u>EQ 105MG BASE</u> | <u>A091118 006</u> | Sep 25, 2014 |
| <u>AB</u> | | <u>EQ 135MG BASE</u> | <u>A091118 008</u> | Sep 25, 2014 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 45MG BASE</u> | <u>A203553 001</u> | Nov 16, 2017 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A203553 004</u> | Nov 16, 2017 |
| <u>AB</u> | | <u>EQ 90MG BASE</u> | <u>A203553 005</u> | Nov 16, 2017 |
| <u>AB</u> | | <u>EQ 105MG BASE</u> | <u>A203553 006</u> | Nov 16, 2017 |
| <u>AB</u> | | <u>EQ 135MG BASE</u> | <u>A203553 008</u> | Nov 16, 2017 |

SOLODYN

| | | | | |
|-----------|-----------|----------------------|--------------------|--------------|
| <u>AB</u> | + MEDICIS | <u>EQ 55MG BASE</u> | <u>N050808 008</u> | Aug 27, 2010 |
| <u>AB</u> | + | <u>EQ 65MG BASE</u> | <u>N050808 004</u> | Jul 23, 2009 |
| <u>AB</u> | + | <u>EQ 80MG BASE</u> | <u>N050808 007</u> | Aug 27, 2010 |
| <u>AB</u> | + | <u>EQ 105MG BASE</u> | <u>N050808 006</u> | Aug 27, 2010 |
| <u>AB</u> | +! | <u>EQ 115MG BASE</u> | <u>N050808 005</u> | Jul 23, 2009 |

MINOLIRA

EPI HLTH

EQ 105MG BASE

N209269 001 May 08, 2017

EQ 135MG BASE

N209269 002 May 08, 2017

PRESCRIPTION DRUG PRODUCT LIST

MINOXIDIL

TABLET; ORAL

MINOXIDIL

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | PAR PHARM | <u>2.5MG</u> | <u>A071826</u> | <u>001</u> | Nov 14, 1988 |
| <u>AB</u> | | <u>10MG</u> | <u>A071839</u> | <u>001</u> | Nov 14, 1988 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>2.5MG</u> | <u>A072709</u> | <u>002</u> | Dec 14, 1995 |
| <u>AB</u> | | <u>10MG</u> | <u>A072709</u> | <u>001</u> | Dec 14, 1995 |
| <u>AB</u> | WATSON LABS | <u>2.5MG</u> | <u>A071344</u> | <u>001</u> | Mar 03, 1987 |
| <u>AB</u> | ! | <u>10MG</u> | <u>A071345</u> | <u>001</u> | Mar 03, 1987 |

MIRABEGRON

TABLET, EXTENDED RELEASE; ORAL

MYRBETRIQ

| | | | | | | |
|---|---|-------|------|---------|-----|--------------|
| + | ! | APGDI | 25MG | N202611 | 001 | Jun 28, 2012 |
| + | ! | | 50MG | N202611 | 002 | Jun 28, 2012 |

MIRTAZAPINE

TABLET; ORAL

MIRTAZAPINE

| | | | | | |
|-----------|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | APOTEX INC | <u>15MG</u> | <u>A077666</u> | <u>001</u> | Aug 22, 2007 |
| <u>AB</u> | | <u>30MG</u> | <u>A077666</u> | <u>002</u> | Aug 22, 2007 |
| <u>AB</u> | | <u>45MG</u> | <u>A077666</u> | <u>003</u> | Aug 22, 2007 |
| <u>AB</u> | AUROBINDO | <u>7.5MG</u> | <u>A076921</u> | <u>001</u> | Oct 22, 2004 |
| <u>AB</u> | | <u>15MG</u> | <u>A076921</u> | <u>002</u> | Oct 22, 2004 |
| <u>AB</u> | | <u>30MG</u> | <u>A076921</u> | <u>003</u> | Oct 22, 2004 |
| <u>AB</u> | | <u>45MG</u> | <u>A076921</u> | <u>004</u> | Oct 22, 2004 |
| <u>AB</u> | MYLAN | <u>15MG</u> | <u>A076122</u> | <u>001</u> | Jun 19, 2003 |
| <u>AB</u> | | <u>30MG</u> | <u>A076122</u> | <u>002</u> | Jun 19, 2003 |
| <u>AB</u> | | <u>45MG</u> | <u>A076122</u> | <u>003</u> | Jun 19, 2003 |
| <u>AB</u> | SUN PHARM INDS INC | <u>7.5MG</u> | <u>A076541</u> | <u>004</u> | Apr 22, 2004 |
| <u>AB</u> | | <u>15MG</u> | <u>A076541</u> | <u>001</u> | Apr 22, 2004 |
| <u>AB</u> | | <u>30MG</u> | <u>A076541</u> | <u>002</u> | Apr 22, 2004 |
| <u>AB</u> | | <u>45MG</u> | <u>A076541</u> | <u>003</u> | Apr 22, 2004 |
| <u>AB</u> | TEVA | <u>15MG</u> | <u>A076119</u> | <u>001</u> | Jan 24, 2003 |
| <u>AB</u> | | <u>30MG</u> | <u>A076119</u> | <u>002</u> | Jan 24, 2003 |
| <u>AB</u> | | <u>45MG</u> | <u>A076119</u> | <u>003</u> | Jun 19, 2003 |
| <u>AB</u> | UPSHER SMITH LABS | <u>15MG</u> | <u>A076219</u> | <u>001</u> | Jun 19, 2003 |
| <u>AB</u> | | <u>30MG</u> | <u>A076219</u> | <u>002</u> | Jun 19, 2003 |
| <u>AB</u> | | <u>45MG</u> | <u>A076219</u> | <u>003</u> | Jun 19, 2003 |
| <u>AB</u> | WATSON LABS | <u>15MG</u> | <u>A076312</u> | <u>001</u> | Jun 19, 2003 |
| <u>AB</u> | | <u>30MG</u> | <u>A076312</u> | <u>002</u> | Jun 19, 2003 |
| <u>AB</u> | | <u>45MG</u> | <u>A076312</u> | <u>003</u> | Jun 19, 2003 |

REMERON

| | | | | | | | |
|-----------|---|---|-----------------|-------------|----------------|------------|--------------|
| <u>AB</u> | + | ! | ORGANON USA INC | <u>15MG</u> | <u>N020415</u> | <u>001</u> | Jun 14, 1996 |
| <u>AB</u> | + | | | <u>30MG</u> | <u>N020415</u> | <u>002</u> | Jun 14, 1996 |

TABLET, ORALLY DISINTEGRATING; ORAL

MIRTAZAPINE

| | | | | | |
|-----------|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>15MG</u> | <u>A077376</u> | <u>002</u> | Dec 08, 2005 |
| <u>AB</u> | | <u>30MG</u> | <u>A077376</u> | <u>003</u> | Dec 08, 2005 |
| <u>AB</u> | | <u>45MG</u> | <u>A077376</u> | <u>004</u> | Feb 28, 2006 |
| <u>AB</u> | IMPAX LABS INC | <u>15MG</u> | <u>A076901</u> | <u>001</u> | Jun 28, 2005 |
| <u>AB</u> | | <u>30MG</u> | <u>A076901</u> | <u>002</u> | Jun 28, 2005 |
| <u>AB</u> | | <u>45MG</u> | <u>A076901</u> | <u>003</u> | Jun 28, 2005 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>15MG</u> | <u>A205798</u> | <u>001</u> | Jun 01, 2017 |
| <u>AB</u> | | <u>30MG</u> | <u>A205798</u> | <u>002</u> | Jun 01, 2017 |
| <u>AB</u> | | <u>45MG</u> | <u>A205798</u> | <u>003</u> | Jun 01, 2017 |

REMERON SOLTAB

| | | | | | | | |
|-----------|---|---|-----------------|-------------|----------------|------------|--------------|
| <u>AB</u> | + | ! | ORGANON USA INC | <u>15MG</u> | <u>N021208</u> | <u>001</u> | Jan 12, 2001 |
| <u>AB</u> | + | | | <u>30MG</u> | <u>N021208</u> | <u>002</u> | Jan 12, 2001 |
| <u>AB</u> | + | | | <u>45MG</u> | <u>N021208</u> | <u>003</u> | Jan 12, 2001 |

MISOPROSTOL

TABLET; ORAL

CYTOTEC

| | | | | | | | |
|-----------|---|---|---------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | | GD SEARLE LLC | <u>0.1MG</u> | <u>N019268</u> | <u>003</u> | Sep 21, 1990 |
| <u>AB</u> | + | ! | | <u>0.2MG</u> | <u>N019268</u> | <u>001</u> | Dec 27, 1988 |

MISOPROSTOL

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>0.1MG</u> | <u>A076095</u> | <u>001</u> | Jul 10, 2002 |
| <u>AB</u> | | <u>0.2MG</u> | <u>A076095</u> | <u>002</u> | Jul 10, 2002 |
| <u>AB</u> | NOVEL LABS INC | <u>0.1MG</u> | <u>A091667</u> | <u>001</u> | Jul 25, 2012 |
| <u>AB</u> | | <u>0.2MG</u> | <u>A091667</u> | <u>002</u> | Jul 25, 2012 |

PRESCRIPTION DRUG PRODUCT LIST

MITOMYCIN

FOR SOLUTION; TOPICAL

MITOSOL

+! MOBIUS THERAP

0.2MG/VIAL

N022572 001 Feb 07, 2012

INJECTABLE; INJECTION

MITOMYCIN

| | | | | | | |
|-----------|---|-------------------------|------------------|----------------|------------|--------------|
| <u>AP</u> | ! | ACCORD HLTHCARE | <u>5MG/VIAL</u> | <u>A064144</u> | <u>001</u> | Apr 30, 1998 |
| <u>AP</u> | ! | | <u>20MG/VIAL</u> | <u>A064144</u> | <u>002</u> | Apr 30, 1998 |
| <u>AP</u> | ! | | <u>40MG/VIAL</u> | <u>A064144</u> | <u>003</u> | Aug 11, 2009 |
| <u>AP</u> | | MYLAN LABS LTD | <u>5MG/VIAL</u> | <u>A202670</u> | <u>001</u> | Oct 13, 2017 |
| <u>AP</u> | | | <u>20MG/VIAL</u> | <u>A202670</u> | <u>002</u> | Oct 13, 2017 |
| <u>AP</u> | | | <u>40MG/VIAL</u> | <u>A203386</u> | <u>001</u> | Oct 13, 2017 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>5MG/VIAL</u> | <u>A064180</u> | <u>001</u> | Dec 23, 1999 |
| <u>AP</u> | | | <u>20MG/VIAL</u> | <u>A064180</u> | <u>002</u> | Dec 23, 1999 |

MITOTANE

TABLET; ORAL

LYSODREN

+! LABORATOIRE HRA

500MG

N016885 001

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

MITOXANTRONE HYDROCHLORIDE

| | | | | | | |
|-----------|---|-------------------------|---------------------------------------------|----------------|------------|--------------|
| <u>AP</u> | | FRESENIUS KABI USA | <u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u> | <u>A077496</u> | <u>001</u> | Apr 11, 2006 |
| <u>AP</u> | | | <u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u> | <u>A077496</u> | <u>002</u> | Apr 11, 2006 |
| <u>AP</u> | | | <u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u> | <u>A077496</u> | <u>003</u> | Apr 11, 2006 |
| <u>AP</u> | ! | HOSPIRA | <u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u> | <u>A076871</u> | <u>001</u> | Apr 11, 2006 |
| <u>AP</u> | ! | | <u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u> | <u>A076871</u> | <u>002</u> | Apr 11, 2006 |
| <u>AP</u> | ! | | <u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u> | <u>A076871</u> | <u>003</u> | Apr 11, 2006 |
| <u>AP</u> | | MYLAN INSTITUTIONAL | <u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u> | <u>A078980</u> | <u>001</u> | Apr 13, 2009 |
| <u>AP</u> | | | <u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u> | <u>A078980</u> | <u>002</u> | Apr 13, 2009 |
| <u>AP</u> | | MYLAN LABS LTD | <u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u> | <u>A201014</u> | <u>001</u> | Dec 11, 2012 |
| <u>AP</u> | | TEVA PHARMS USA | <u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u> | <u>A077356</u> | <u>001</u> | Apr 11, 2006 |
| <u>AP</u> | | | <u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u> | <u>A077356</u> | <u>002</u> | Apr 11, 2006 |
| <u>AP</u> | | | <u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u> | <u>A077356</u> | <u>003</u> | Apr 11, 2006 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u> | <u>A076611</u> | <u>001</u> | Apr 11, 2006 |
| <u>AP</u> | | | <u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u> | <u>A076611</u> | <u>002</u> | Apr 11, 2006 |
| <u>AP</u> | | | <u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u> | <u>A076611</u> | <u>003</u> | Apr 11, 2006 |

MODAFINIL

TABLET; ORAL

MODAFINIL

| | | | | | | |
|-----------|--|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | ALEMBIC PHARMS LTD | <u>100MG</u> | <u>A202700</u> | <u>001</u> | Oct 18, 2012 |
| <u>AB</u> | | | <u>200MG</u> | <u>A202700</u> | <u>002</u> | Oct 18, 2012 |
| <u>AB</u> | | APOTEX INC | <u>100MG</u> | <u>A077667</u> | <u>001</u> | Feb 03, 2014 |
| <u>AB</u> | | | <u>200MG</u> | <u>A077667</u> | <u>002</u> | Feb 03, 2014 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>100MG</u> | <u>A202566</u> | <u>001</u> | Sep 27, 2012 |
| <u>AB</u> | | | <u>200MG</u> | <u>A202566</u> | <u>002</u> | Sep 27, 2012 |
| <u>AB</u> | | HERITAGE PHARMS INC | <u>100MG</u> | <u>A207196</u> | <u>001</u> | Aug 16, 2017 |
| <u>AB</u> | | | <u>200MG</u> | <u>A207196</u> | <u>002</u> | Aug 16, 2017 |
| <u>AB</u> | | HIKMA PHARMS | <u>100MG</u> | <u>A090543</u> | <u>001</u> | Sep 26, 2012 |
| <u>AB</u> | | | <u>200MG</u> | <u>A090543</u> | <u>002</u> | Sep 26, 2012 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>100MG</u> | <u>A076594</u> | <u>001</u> | Jul 16, 2012 |
| <u>AB</u> | | | <u>200MG</u> | <u>A076594</u> | <u>002</u> | Jul 16, 2012 |
| <u>AB</u> | | ORCHID HLTHCARE | <u>100MG</u> | <u>A078963</u> | <u>001</u> | Sep 26, 2012 |
| <u>AB</u> | | | <u>200MG</u> | <u>A078963</u> | <u>002</u> | Sep 26, 2012 |
| <u>AB</u> | | WATSON LABS INC | <u>100MG</u> | <u>A076715</u> | <u>001</u> | Nov 01, 2012 |
| <u>AB</u> | | | <u>200MG</u> | <u>A076715</u> | <u>002</u> | Nov 01, 2012 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>100MG</u> | <u>A209966</u> | <u>001</u> | Sep 14, 2017 |
| <u>AB</u> | | | <u>200MG</u> | <u>A209966</u> | <u>002</u> | Sep 14, 2017 |

PROVIGIL

| | | | | | | |
|-----------|----|----------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | CEPHALON | <u>100MG</u> | <u>N020717</u> | <u>001</u> | Dec 24, 1998 |
| <u>AB</u> | +! | | <u>200MG</u> | <u>N020717</u> | <u>002</u> | Dec 24, 1998 |

MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE

| | | | | | | |
|-----------|--|--------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | APOTEX INC | <u>7.5MG</u> | <u>A078454</u> | <u>001</u> | Jun 02, 2008 |
| <u>AB</u> | | | <u>15MG</u> | <u>A078454</u> | <u>002</u> | Jun 02, 2008 |
| <u>AB</u> | | CHARTWELL RX | <u>7.5MG</u> | <u>A077536</u> | <u>001</u> | Nov 30, 2006 |
| <u>AB</u> | | | <u>15MG</u> | <u>A077536</u> | <u>002</u> | Nov 30, 2006 |

PRESCRIPTION DRUG PRODUCT LIST

MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE

| | | | | | |
|-----------|-------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | GLENMARK GENERICS | <u>7.5MG</u> | <u>A090416</u> | <u>001</u> | Mar 30, 2010 |
| <u>AB</u> | | <u>15MG</u> | <u>A090416</u> | <u>002</u> | Mar 30, 2010 |
| <u>AB</u> | TEVA | <u>7.5MG</u> | <u>A076204</u> | <u>001</u> | May 08, 2003 |
| <u>AB</u> | ! | <u>15MG</u> | <u>A076204</u> | <u>002</u> | May 08, 2003 |

MOLINDONE HYDROCHLORIDE

TABLET; ORAL

MOLINDONE HYDROCHLORIDE

EPIC PHARMA LLC

5MG

A090453 001 Mar 20, 2015

10MG

A090453 002 Mar 20, 2015

!

25MG

A090453 003 Mar 20, 2015

MOMETASONE FUROATE

AEROSOL, METERED; INHALATION

ASMANEX HFA

+ MERCK SHARP DOHME

0.10MG/INH

N205641 001 Apr 25, 2014

+!

0.20MG/INH

N205641 002 Apr 25, 2014

CREAM; TOPICAL

ELOCON

| | | | | | | |
|-----------|----|-------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | +! | MERCK SHARP DOHME | <u>0.1%</u> | <u>N019625</u> | <u>002</u> | Apr 19, 2013 |
|-----------|----|-------------------|-------------|----------------|------------|--------------|

MOMETASONE FUROATE

| | | | | | |
|-----------|-------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | ANDA REPOSITORY | <u>0.1%</u> | <u>A076591</u> | <u>001</u> | Apr 18, 2007 |
| <u>AB</u> | FOUGERA PHARMS | <u>0.1%</u> | <u>A076171</u> | <u>001</u> | Apr 08, 2005 |
| <u>AB</u> | G AND W LABS | <u>0.1%</u> | <u>A077447</u> | <u>001</u> | May 22, 2006 |
| <u>AB</u> | GLENMARK GENERICS | <u>0.1%</u> | <u>A078541</u> | <u>001</u> | May 28, 2008 |
| <u>AB</u> | TARO | <u>0.1%</u> | <u>A076679</u> | <u>001</u> | Dec 21, 2004 |

IMPLANT; IMPLANTATION

SINUVA

+ INTERSECT ENT INC

1.35MG

N209310 001 Dec 08, 2017

LOTION; TOPICAL

ELOCON

| | | | | | | |
|-----------|----|-------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | +! | MERCK SHARP DOHME | <u>0.1%</u> | <u>N019796</u> | <u>001</u> | Mar 30, 1989 |
|-----------|----|-------------------|-------------|----------------|------------|--------------|

MOMETASONE FUROATE

| | | | | | |
|-----------|-------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | ANDA REPOSITORY | <u>0.1%</u> | <u>A076499</u> | <u>001</u> | Nov 21, 2007 |
| <u>AB</u> | FOUGERA PHARMS | <u>0.1%</u> | <u>A075919</u> | <u>001</u> | Nov 29, 2007 |
| <u>AB</u> | G AND W LABS | <u>0.1%</u> | <u>A077678</u> | <u>001</u> | Nov 21, 2007 |
| <u>AB</u> | GLENMARK GENERICS | <u>0.1%</u> | <u>A090506</u> | <u>001</u> | Aug 09, 2010 |
| <u>AB</u> | PERRIGO ISRAEL | <u>0.1%</u> | <u>A077180</u> | <u>001</u> | Apr 06, 2005 |
| <u>AB</u> | TARO | <u>0.1%</u> | <u>A076788</u> | <u>001</u> | Mar 15, 2006 |

OINTMENT; TOPICAL

ELOCON

| | | | | | | |
|-----------|----|-------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | +! | MERCK SHARP DOHME | <u>0.1%</u> | <u>N019543</u> | <u>001</u> | Apr 30, 1987 |
|-----------|----|-------------------|-------------|----------------|------------|--------------|

MOMETASONE FUROATE

| | | | | | |
|-----------|--------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | ANDA REPOSITORY | <u>0.1%</u> | <u>A076481</u> | <u>001</u> | Nov 14, 2003 |
| <u>AB</u> | FOUGERA PHARMS | <u>0.1%</u> | <u>A077061</u> | <u>001</u> | Mar 28, 2005 |
| <u>AB</u> | G AND W LABS | <u>0.1%</u> | <u>A077401</u> | <u>001</u> | Jun 20, 2006 |
| <u>AB</u> | GLENMARK GENERICS | <u>0.1%</u> | <u>A078571</u> | <u>001</u> | May 28, 2008 |
| <u>AB</u> | PERRIGO NEW YORK | <u>0.1%</u> | <u>A076067</u> | <u>001</u> | Mar 18, 2002 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>0.1%</u> | <u>A207899</u> | <u>001</u> | Jul 13, 2018 |

POWDER; INHALATION

ASMANEX TWISTHALER

+ MERCK SHARP DOHME

0.11MG/INH

N021067 002 Feb 01, 2008

+!

0.22MG/INH

N021067 001 Mar 30, 2005

SPRAY, METERED; NASAL

MOMETASONE FUROATE

| | | | | | |
|-----------|---------------|-----------------------------|----------------|------------|--------------|
| <u>AB</u> | AMNEAL PHARMS | <u>EQ 0.05MG BASE/SPRAY</u> | <u>A207989</u> | <u>001</u> | Apr 03, 2017 |
| <u>AB</u> | APOTEX INC | <u>EQ 0.05MG BASE/SPRAY</u> | <u>A091161</u> | <u>001</u> | Mar 22, 2016 |

NASONEX

| | | | | | | |
|-----------|----|-------------------|-----------------------------|----------------|------------|--------------|
| <u>AB</u> | +! | MERCK SHARP DOHME | <u>EQ 0.05MG BASE/SPRAY</u> | <u>N020762</u> | <u>001</u> | Oct 01, 1997 |
|-----------|----|-------------------|-----------------------------|----------------|------------|--------------|

MONTELUKAST SODIUM

GRANULE; ORAL

MONTELUKAST SODIUM

| | | | | | |
|-----------|--------------------|---------------------------|----------------|------------|--------------|
| <u>AB</u> | AJANTA PHARMA LTD | <u>EQ 4MG BASE/PACKET</u> | <u>A203438</u> | <u>001</u> | Jul 31, 2015 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 4MG BASE/PACKET</u> | <u>A202906</u> | <u>001</u> | Sep 17, 2012 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 4MG BASE/PACKET</u> | <u>A202776</u> | <u>001</u> | Dec 18, 2012 |
| <u>AB</u> | TEVA PHARMS | <u>EQ 4MG BASE/PACKET</u> | <u>A090955</u> | <u>001</u> | Aug 03, 2012 |
| <u>AB</u> | TORRENT PHARMS LLC | <u>EQ 4MG BASE/PACKET</u> | <u>A210431</u> | <u>001</u> | Jul 31, 2018 |

SINGULAIR

| | | | | | | |
|-----------|----|-------|---------------------------|----------------|------------|--------------|
| <u>AB</u> | +! | MERCK | <u>EQ 4MG BASE/PACKET</u> | <u>N021409</u> | <u>001</u> | Jul 26, 2002 |
|-----------|----|-------|---------------------------|----------------|------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

MONTELUKAST SODIUM

TABLET; ORAL

MONTELUKAST SODIUM

| | | | | |
|-----------|----------------------|---------------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>EQ 10MG BASE</u> | <u>A202717 001</u> | Sep 21, 2012 |
| <u>AB</u> | AJANTA PHARMA LTD | <u>EQ 10MG BASE</u> | <u>A203432 001</u> | Jul 31, 2015 |
| <u>AB</u> | AMNEAL PHARMS | <u>EQ 10MG BASE</u> | <u>A204604 001</u> | Sep 04, 2015 |
| <u>AB</u> | ANBISON LAB | <u>EQ 10MG BASE</u> | <u>A205683 001</u> | Jan 12, 2016 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 10MG BASE</u> | <u>A202468 001</u> | Aug 03, 2012 |
| <u>AB</u> | CIPLA | <u>EQ 10MG BASE</u> | <u>A207463 001</u> | Oct 28, 2016 |
| <u>AB</u> | CSPC OUYI PHARM CO | <u>EQ 10MG BASE</u> | <u>A209012 001</u> | Apr 24, 2017 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 10MG BASE</u> | <u>A201582 001</u> | Aug 06, 2012 |
| <u>AB</u> | GLENMARK GENERICS | <u>EQ 10MG BASE</u> | <u>A090926 001</u> | Aug 03, 2012 |
| <u>AB</u> | HETERO LABS LTD V | <u>EQ 10MG BASE</u> | <u>A202843 001</u> | Sep 10, 2014 |
| <u>AB</u> | LANNETT CO INC | <u>EQ 10MG BASE</u> | <u>A201522 001</u> | Aug 03, 2012 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>EQ 10MG BASE</u> | <u>A203366 001</u> | Sep 11, 2014 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 10MG BASE</u> | <u>A079103 001</u> | Aug 03, 2012 |
| <u>AB</u> | PERRIGO R AND D | <u>EQ 10MG BASE</u> | <u>A206112 001</u> | Apr 26, 2017 |
| <u>AB</u> | SANDOZ INC | <u>EQ 10MG BASE</u> | <u>A200889 001</u> | Aug 03, 2012 |
| <u>AB</u> | TEVA PHARMS | <u>EQ 10MG BASE</u> | <u>A078605 001</u> | Aug 03, 2012 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>EQ 10MG BASE</u> | <u>A201515 001</u> | Aug 03, 2012 |
| <u>AB</u> | UNICHEM LABS LTD | <u>EQ 10MG BASE</u> | <u>A204290 001</u> | Oct 08, 2015 |
| <u>AB</u> | UNIMARK REMEDIES LTD | <u>EQ 10MG BASE</u> | <u>A202859 001</u> | Oct 30, 2014 |
| <u>AB</u> | VINTAGE PHARMS LLC | <u>EQ 10MG BASE</u> | <u>A091576 001</u> | Aug 03, 2012 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>EQ 10MG BASE</u> | <u>A090655 001</u> | Aug 03, 2012 |

SINGULAIR

| | | | | |
|-----------|-----------------------|---------------------|--------------------|--------------|
| <u>AB</u> | <u>+</u> MSD MERCK CO | <u>EQ 10MG BASE</u> | <u>N020829 002</u> | Feb 20, 1998 |
|-----------|-----------------------|---------------------|--------------------|--------------|

TABLET, CHEWABLE; ORAL

MONTELUKAST SODIUM

| | | | | |
|-----------|----------------------|--------------------|--------------------|--------------|
| <u>AB</u> | AJANTA PHARMA LTD | <u>EQ 4MG BASE</u> | <u>A203328 001</u> | Jul 31, 2015 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A203328 002</u> | Jul 31, 2015 |
| <u>AB</u> | ANBISON LAB | <u>EQ 4MG BASE</u> | <u>A205695 001</u> | Nov 05, 2015 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A205695 002</u> | Nov 05, 2015 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 4MG BASE</u> | <u>A202096 001</u> | Aug 03, 2012 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A202096 002</u> | Aug 03, 2012 |
| <u>AB</u> | CIPLA | <u>EQ 4MG BASE</u> | <u>A207464 001</u> | Dec 06, 2018 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A207464 002</u> | Dec 06, 2018 |
| <u>AB</u> | CSPC OUYI PHARM CO | <u>EQ 4MG BASE</u> | <u>A209011 001</u> | Apr 18, 2017 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A209011 002</u> | Apr 18, 2017 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 4MG BASE</u> | <u>A201581 001</u> | Aug 06, 2012 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A201581 002</u> | Aug 06, 2012 |
| <u>AB</u> | HETERO LABS LTD V | <u>EQ 4MG BASE</u> | <u>A204093 001</u> | May 22, 2015 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A204093 002</u> | May 22, 2015 |
| <u>AB</u> | JUBILANT GENERICS | <u>EQ 4MG BASE</u> | <u>A203795 001</u> | Feb 27, 2015 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A203795 002</u> | Feb 27, 2015 |
| <u>AB</u> | LANNETT CO INC | <u>EQ 4MG BASE</u> | <u>A200405 001</u> | Aug 03, 2012 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A200405 002</u> | Aug 03, 2012 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>EQ 4MG BASE</u> | <u>A203582 001</u> | Mar 12, 2015 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A203582 002</u> | Mar 12, 2015 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 4MG BASE</u> | <u>A079142 001</u> | Aug 03, 2012 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A079142 002</u> | Aug 03, 2012 |
| <u>AB</u> | SANDOZ INC | <u>EQ 4MG BASE</u> | <u>A091414 001</u> | Aug 03, 2012 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A091414 002</u> | Aug 03, 2012 |
| <u>AB</u> | TEVA PHARMS | <u>EQ 4MG BASE</u> | <u>A078723 001</u> | Aug 03, 2012 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A078723 002</u> | Aug 03, 2012 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>EQ 4MG BASE</u> | <u>A090984 001</u> | Aug 03, 2012 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A090984 002</u> | Aug 03, 2012 |
| <u>AB</u> | UNICHEM LABS LTD | <u>EQ 4MG BASE</u> | <u>A208621 001</u> | Jul 02, 2018 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A208621 002</u> | Jul 02, 2018 |
| <u>AB</u> | UNIMARK REMEDIES LTD | <u>EQ 4MG BASE</u> | <u>A203037 001</u> | Oct 30, 2014 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A203037 002</u> | Oct 30, 2014 |
| <u>AB</u> | VINTAGE PHARMS LLC | <u>EQ 4MG BASE</u> | <u>A091588 001</u> | Aug 03, 2012 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A091588 002</u> | Aug 03, 2012 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>EQ 4MG BASE</u> | <u>A091128 001</u> | Aug 03, 2012 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A091128 002</u> | Aug 03, 2012 |

SINGULAIR

| | | | | |
|-----------|-----------------------|--------------------|--------------------|--------------|
| <u>AB</u> | <u>+</u> MSD MERCK CO | <u>EQ 4MG BASE</u> | <u>N020830 002</u> | Mar 03, 2000 |
| <u>AB</u> | <u>+</u> ! | <u>EQ 5MG BASE</u> | <u>N020830 001</u> | Feb 20, 1998 |

PRESCRIPTION DRUG PRODUCT LIST

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE;ORAL

KADIAN

| | | | | | | |
|-----------|---|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | ALLERGAN SALES LLC | <u>10MG</u> | <u>N020616</u> | <u>008</u> | Apr 20, 2007 |
| <u>AB</u> | + | | <u>20MG</u> | <u>N020616</u> | <u>001</u> | Jul 03, 1996 |
| <u>AB</u> | + | | <u>30MG</u> | <u>N020616</u> | <u>004</u> | Mar 09, 2001 |
| <u>AB</u> | + | | <u>40MG</u> | <u>N020616</u> | <u>009</u> | Jul 09, 2012 |
| <u>AB</u> | + | | <u>50MG</u> | <u>N020616</u> | <u>002</u> | Jul 03, 1996 |
| <u>AB</u> | + | | <u>60MG</u> | <u>N020616</u> | <u>005</u> | Mar 09, 2001 |
| <u>AB</u> | + | | <u>70MG</u> | <u>N020616</u> | <u>010</u> | Jul 09, 2012 |
| <u>AB</u> | + | | <u>80MG</u> | <u>N020616</u> | <u>006</u> | Oct 27, 2006 |
| <u>AB</u> | + | | <u>100MG</u> | <u>N020616</u> | <u>003</u> | Jul 03, 1996 |

MORPHINE SULFATE

| | | | | | | |
|-----------|--|-------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | IMPAX LABS INC | <u>20MG</u> | <u>A200411</u> | <u>001</u> | Apr 12, 2016 |
| <u>AB</u> | | | <u>30MG</u> | <u>A200411</u> | <u>002</u> | Apr 12, 2016 |
| <u>AB</u> | | | <u>50MG</u> | <u>A200411</u> | <u>003</u> | Apr 12, 2016 |
| <u>AB</u> | | | <u>60MG</u> | <u>A200411</u> | <u>004</u> | Apr 12, 2016 |
| <u>AB</u> | | | <u>80MG</u> | <u>A200411</u> | <u>005</u> | Apr 12, 2016 |
| <u>AB</u> | | | <u>100MG</u> | <u>A200411</u> | <u>006</u> | Apr 12, 2016 |
| <u>AB</u> | | PAR PHARM INC | <u>20MG</u> | <u>A200812</u> | <u>001</u> | Nov 10, 2011 |
| <u>AB</u> | | | <u>30MG</u> | <u>A200812</u> | <u>002</u> | Nov 10, 2011 |
| <u>AB</u> | | | <u>50MG</u> | <u>A200812</u> | <u>003</u> | Nov 10, 2011 |
| <u>AB</u> | | | <u>60MG</u> | <u>A200812</u> | <u>004</u> | Nov 10, 2011 |
| <u>AB</u> | | | <u>80MG</u> | <u>A200812</u> | <u>005</u> | Nov 10, 2011 |
| <u>AB</u> | | | <u>100MG</u> | <u>A200812</u> | <u>006</u> | Nov 10, 2011 |
| <u>AB</u> | | TEVA PHARMS USA | <u>20MG</u> | <u>A202718</u> | <u>001</u> | Dec 29, 2014 |
| <u>AB</u> | | | <u>30MG</u> | <u>A202718</u> | <u>002</u> | Dec 29, 2014 |
| <u>AB</u> | | | <u>40MG</u> | <u>A202718</u> | <u>007</u> | Jun 03, 2015 |
| <u>AB</u> | | | <u>50MG</u> | <u>A202718</u> | <u>003</u> | Dec 29, 2014 |
| <u>AB</u> | | | <u>60MG</u> | <u>A202718</u> | <u>004</u> | Dec 29, 2014 |
| <u>AB</u> | | | <u>70MG</u> | <u>A202718</u> | <u>008</u> | Jun 03, 2015 |
| <u>AB</u> | | | <u>80MG</u> | <u>A202718</u> | <u>005</u> | Dec 29, 2014 |
| <u>AB</u> | | | <u>100MG</u> | <u>A202718</u> | <u>006</u> | Dec 29, 2014 |
| <u>AB</u> | | UPSHER SMITH LABS | <u>10MG</u> | <u>A202104</u> | <u>001</u> | Jun 03, 2013 |
| <u>AB</u> | | | <u>20MG</u> | <u>A202104</u> | <u>002</u> | Jun 03, 2013 |
| <u>AB</u> | | | <u>30MG</u> | <u>A202104</u> | <u>003</u> | Jun 03, 2013 |
| <u>AB</u> | | | <u>50MG</u> | <u>A202104</u> | <u>004</u> | Jun 03, 2013 |
| <u>AB</u> | | | <u>60MG</u> | <u>A202104</u> | <u>005</u> | Jun 03, 2013 |
| <u>AB</u> | | | <u>80MG</u> | <u>A202104</u> | <u>006</u> | Jun 03, 2013 |
| <u>AB</u> | | | <u>100MG</u> | <u>A202104</u> | <u>007</u> | Jun 03, 2013 |

KADIAN

| | | | | | | |
|--|---|--------------------|-------|---------|-----|--------------|
| | + | ALLERGAN SALES LLC | 130MG | N020616 | 011 | Jul 09, 2012 |
| | + | | 150MG | N020616 | 012 | Jul 09, 2012 |
| | + | | 200MG | N020616 | 007 | Feb 27, 2007 |

MORPHINE SULFATE

| | | | | | | |
|--|---|-------------------|-------|---------|-----|--------------|
| | | ACTAVIS ELIZABETH | 30MG | A079040 | 001 | Jan 16, 2013 |
| | | | 45MG | A079040 | 002 | Jan 16, 2013 |
| | | | 60MG | A079040 | 003 | Jan 16, 2013 |
| | | | 75MG | A079040 | 004 | Jan 16, 2013 |
| | | | 90MG | A079040 | 005 | Jan 16, 2013 |
| | ! | | 120MG | A079040 | 006 | Jan 16, 2013 |

INJECTABLE; INJECTION

ASTRAMORPH PF

| | | | | | | |
|-----------|--|--------------------|-----------------|----------------|------------|--------------|
| <u>AP</u> | | FRESENIUS KABI USA | <u>0.5MG/ML</u> | <u>A071050</u> | <u>001</u> | Oct 07, 1986 |
| <u>AP</u> | | | <u>0.5MG/ML</u> | <u>A071051</u> | <u>001</u> | Oct 07, 1986 |
| <u>AP</u> | | | <u>1MG/ML</u> | <u>A071052</u> | <u>001</u> | Oct 07, 1986 |
| <u>AP</u> | | | <u>1MG/ML</u> | <u>A071053</u> | <u>001</u> | Oct 07, 1986 |

DURAMORPH PF

| | | | | | | |
|-----------|---|-------------------------|-----------------|----------------|------------|--------------|
| <u>AP</u> | + | WEST-WARD PHARMS INT | <u>0.5MG/ML</u> | <u>N018565</u> | <u>001</u> | Sep 18, 1984 |
| <u>AP</u> | + | | <u>1MG/ML</u> | <u>N018565</u> | <u>002</u> | Sep 18, 1984 |

INFUMORPH

| | | | | | | |
|-----------|---|-------------------------|----------------|----------------|------------|--------------|
| <u>AP</u> | + | WEST-WARD PHARMS INT | <u>10MG/ML</u> | <u>N018565</u> | <u>003</u> | Jul 19, 1991 |
| <u>AP</u> | + | | <u>25MG/ML</u> | <u>N018565</u> | <u>004</u> | Jul 19, 1991 |

MITIGO

| | | | | | | |
|-----------|--|------------------|----------------|----------------|------------|--------------|
| <u>AP</u> | | PIRAMAL CRITICAL | <u>10MG/ML</u> | <u>A204393</u> | <u>001</u> | Jul 16, 2018 |
| <u>AP</u> | | | <u>25MG/ML</u> | <u>A204393</u> | <u>002</u> | Jul 16, 2018 |

MORPHINE SULFATE

| | | | | | | |
|-----------|--|--------------------|-----------------|----------------|------------|--------------|
| <u>AP</u> | | EUROHLTH INTL SARL | <u>4MG/ML</u> | <u>A205758</u> | <u>001</u> | May 21, 2015 |
| <u>AP</u> | | | <u>8MG/ML</u> | <u>A205758</u> | <u>002</u> | May 21, 2015 |
| <u>AP</u> | | | <u>10MG/ML</u> | <u>A205758</u> | <u>003</u> | May 21, 2015 |
| <u>AP</u> | | HOSPIRA | <u>0.5MG/ML</u> | <u>A071849</u> | <u>001</u> | May 11, 1988 |

PRESCRIPTION DRUG PRODUCT LIST

MORPHINE SULFATE

INJECTABLE; INJECTION

MORPHINE SULFATE

| | | | | |
|-----------|----|-----------------------------------------|--------------------|--------------|
| <u>AP</u> | | <u>0.5MG/ML</u> | <u>A073509 001</u> | Sep 30, 1992 |
| <u>AP</u> | | <u>1MG/ML</u> | <u>A071850 001</u> | May 11, 1988 |
| <u>AP</u> | | <u>1MG/ML</u> | <u>A073510 001</u> | Sep 30, 1992 |
| <u>AP</u> | +! | <u>HOSPIRA INC</u> <u>4MG/ML</u> | <u>N202515 002</u> | Nov 14, 2011 |
| <u>AP</u> | +! | <u>8MG/ML</u> | <u>N202515 003</u> | Nov 14, 2011 |
| <u>AP</u> | +! | <u>10MG/ML</u> | <u>N202515 004</u> | Nov 14, 2011 |
| <u>AP</u> | +! | <u>ICU MEDICAL INC</u> <u>1MG/ML</u> | <u>N019916 001</u> | Oct 30, 1992 |
| | +! | <u>HOSPIRA INC</u> <u>2MG/ML</u> | <u>N202515 001</u> | Nov 14, 2011 |
| | +! | <u>ICU MEDICAL INC</u> <u>5MG/ML</u> | <u>N019916 002</u> | Oct 27, 2006 |
| | +! | <u>MERIDIAN MEDCL</u> <u>15MG/ML</u> | <u>N019999 001</u> | Jul 12, 1990 |

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

MORPHINE SULFATE

| | | | | |
|--|----|-----------------------------------------------------|--------------------|--------------|
| | +! | <u>FRESENIUS KABI USA</u> <u>2MG/ML (2MG/ML)</u> | <u>N204223 001</u> | Oct 30, 2013 |
| | +! | <u>4MG/ML (4MG/ML)</u> | <u>N204223 002</u> | Oct 30, 2013 |
| | +! | <u>5MG/ML (5MG/ML)</u> | <u>N204223 003</u> | Oct 30, 2013 |
| | +! | <u>8MG/ML (8MG/ML)</u> | <u>N204223 004</u> | Oct 30, 2013 |
| | +! | <u>10MG/ML (10MG/ML)</u> | <u>N204223 005</u> | Oct 30, 2013 |

SOLUTION; ORAL

MORPHINE SULFATE

| | | | | |
|-----------|----|----------------------------------------------------------|--------------------|--------------|
| <u>AA</u> | | <u>ANI PHARMS INC</u> <u>10MG/5ML</u> | <u>A205509 001</u> | Apr 17, 2018 |
| <u>AA</u> | | <u>20MG/5ML</u> | <u>A205509 002</u> | Apr 17, 2018 |
| <u>AA</u> | | <u>100MG/5ML</u> | <u>A205509 003</u> | Apr 17, 2018 |
| <u>AA</u> | | <u>HI-TECH PHARMACAL</u> <u>100MG/5ML</u> | <u>A208809 001</u> | Jul 06, 2017 |
| <u>AA</u> | | <u>LANNETT CO INC</u> <u>10MG/5ML</u> | <u>A202309 001</u> | Nov 25, 2015 |
| <u>AA</u> | | <u>20MG/5ML</u> | <u>A202310 001</u> | Oct 30, 2015 |
| <u>AA</u> | | <u>NOSTRUM LABS INC</u> <u>10MG/5ML</u> | <u>A201011 001</u> | Feb 05, 2014 |
| <u>AA</u> | | <u>20MG/5ML</u> | <u>A201011 002</u> | Feb 05, 2014 |
| <u>AA</u> | | <u>100MG/5ML</u> | <u>A201011 003</u> | Oct 06, 2016 |
| <u>AA</u> | | <u>PADDOCK LLC</u> <u>100MG/5ML</u> | <u>A201574 001</u> | Aug 06, 2012 |
| <u>AA</u> | | <u>PHARM ASSOC</u> <u>100MG/5ML</u> | <u>A206573 001</u> | Nov 14, 2016 |
| <u>AA</u> | | <u>RHODES PHARMS</u> <u>10MG/5ML</u> | <u>A206308 001</u> | Jun 22, 2017 |
| <u>AA</u> | | <u>20MG/5ML</u> | <u>A206420 001</u> | Jul 12, 2016 |
| <u>AA</u> | | <u>100MG/5ML</u> | <u>A206308 002</u> | Jun 22, 2017 |
| <u>AA</u> | | <u>SPECGX LLC</u> <u>100MG/5ML</u> | <u>A202348 001</u> | Jul 15, 2011 |
| <u>AA</u> | | <u>TRIS PHARMA INC</u> <u>10MG/5ML</u> | <u>A203518 001</u> | May 12, 2015 |
| <u>AA</u> | | <u>20MG/5ML</u> | <u>A203519 001</u> | May 18, 2016 |
| <u>AA</u> | | <u>100MG/5ML</u> | <u>A203518 002</u> | May 12, 2015 |
| <u>AA</u> | | <u>VISTAPHARM</u> <u>10MG/5ML</u> | <u>A201947 001</u> | Jan 05, 2012 |
| <u>AA</u> | | <u>20MG/5ML</u> | <u>A201947 002</u> | Jan 05, 2012 |
| <u>AA</u> | + | <u>WEST-WARD PHARMS</u> <u>INT</u> <u>10MG/5ML</u> | <u>N022195 001</u> | Mar 17, 2008 |
| <u>AA</u> | + | <u>20MG/5ML</u> | <u>N022195 002</u> | Mar 17, 2008 |
| <u>AA</u> | +! | <u>100MG/5ML</u> | <u>N022195 003</u> | Jan 25, 2010 |
| | | <u>LANNETT CO INC</u> <u>100MG/5ML</u> | <u>N201517 001</u> | Jun 23, 2011 |

TABLET; ORAL

MORPHINE SULFATE

| | | | | |
|--|----|------------------------------------------------------|--------------------|--------------|
| | + | <u>WEST-WARD PHARMS</u> <u>INT</u> <u>15MG</u> | <u>N022207 001</u> | Mar 17, 2008 |
| | +! | <u>30MG</u> | <u>N022207 002</u> | Mar 17, 2008 |

TABLET, EXTENDED RELEASE; ORAL

MORPHINE SULFATE

| | | | | |
|-----------|--|-----------------------------------------|--------------------|--------------|
| <u>AB</u> | | <u>ACTAVIS ELIZABETH</u> <u>15MG</u> | <u>A203849 001</u> | Apr 06, 2015 |
| <u>AB</u> | | <u>30MG</u> | <u>A203849 002</u> | Apr 06, 2015 |
| <u>AB</u> | | <u>60MG</u> | <u>A203849 003</u> | Apr 06, 2015 |
| <u>AB</u> | | <u>100MG</u> | <u>A203849 004</u> | Apr 06, 2015 |
| <u>AB</u> | | <u>200MG</u> | <u>A203849 005</u> | Apr 06, 2015 |
| <u>AB</u> | | <u>DAVA PHARMS INC</u> <u>15MG</u> | <u>A075407 001</u> | Jan 28, 2000 |
| <u>AB</u> | | <u>MAYNE PHARMA INC</u> <u>15MG</u> | <u>A205386 001</u> | Oct 28, 2016 |
| <u>AB</u> | | <u>30MG</u> | <u>A205386 002</u> | Oct 28, 2016 |
| <u>AB</u> | | <u>60MG</u> | <u>A205386 003</u> | Oct 28, 2016 |
| <u>AB</u> | | <u>100MG</u> | <u>A205386 004</u> | Oct 28, 2016 |
| <u>AB</u> | | <u>MYLAN PHARMS INC</u> <u>15MG</u> | <u>A200824 001</u> | Oct 18, 2011 |
| <u>AB</u> | | <u>30MG</u> | <u>A200824 002</u> | Oct 18, 2011 |
| <u>AB</u> | | <u>60MG</u> | <u>A200824 003</u> | Oct 18, 2011 |
| <u>AB</u> | | <u>100MG</u> | <u>A200824 004</u> | Oct 18, 2011 |
| <u>AB</u> | | <u>200MG</u> | <u>A200824 005</u> | Oct 18, 2011 |
| <u>AB</u> | | <u>NESHER PHARMS</u> <u>15MG</u> | <u>A076733 001</u> | May 19, 2004 |
| <u>AB</u> | | <u>30MG</u> | <u>A076720 002</u> | Dec 23, 2005 |
| <u>AB</u> | | <u>60MG</u> | <u>A076720 001</u> | May 19, 2004 |

PRESCRIPTION DRUG PRODUCT LIST

MORPHINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

MORPHINE SULFATE

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| <u>AB</u> | | <u>100MG</u> | <u>A077855 001</u> | Sep 27, 2007 |
| <u>AB</u> | | <u>200MG</u> | <u>A077855 002</u> | Sep 27, 2007 |
| <u>AB</u> | NOVEL LABS INC | <u>15MG</u> | <u>A203602 001</u> | Dec 16, 2015 |
| <u>AB</u> | | <u>30MG</u> | <u>A203602 002</u> | Dec 16, 2015 |
| <u>AB</u> | | <u>60MG</u> | <u>A203602 003</u> | Dec 16, 2015 |
| <u>AB</u> | | <u>100MG</u> | <u>A203602 004</u> | Dec 16, 2015 |
| <u>AB</u> | | <u>200MG</u> | <u>A203602 005</u> | Dec 16, 2015 |
| <u>AB</u> | RHODES PHARMS | <u>15MG</u> | <u>A074862 001</u> | Jul 07, 1998 |
| <u>AB</u> | | <u>30MG</u> | <u>A074862 002</u> | Jul 07, 1998 |
| <u>AB</u> | | <u>60MG</u> | <u>A074862 003</u> | Jul 07, 1998 |
| <u>AB</u> | | <u>100MG</u> | <u>A074769 001</u> | Jul 02, 1998 |
| <u>AB</u> | | <u>200MG</u> | <u>A074769 002</u> | Jul 02, 1998 |
| <u>AB</u> | SPECGX LLC | <u>15MG</u> | <u>A076412 001</u> | Jul 31, 2003 |
| <u>AB</u> | | <u>30MG</u> | <u>A076412 002</u> | Jul 31, 2003 |
| <u>AB</u> | | <u>60MG</u> | <u>A076412 003</u> | Jul 31, 2003 |
| <u>AB</u> | | <u>100MG</u> | <u>A076438 001</u> | Jul 03, 2003 |
| <u>AB</u> | | <u>200MG</u> | <u>A076438 002</u> | Jul 03, 2003 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>15MG</u> | <u>A078761 001</u> | May 11, 2012 |
| <u>AB</u> | | <u>30MG</u> | <u>A078761 002</u> | May 11, 2012 |
| <u>AB</u> | | <u>60MG</u> | <u>A078761 003</u> | May 11, 2012 |
| <u>AB</u> | | <u>100MG</u> | <u>A078761 004</u> | May 11, 2012 |
| <u>AB</u> | | <u>200MG</u> | <u>A078761 005</u> | May 11, 2012 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>15MG</u> | <u>A205634 001</u> | Aug 25, 2016 |
| <u>AB</u> | | <u>30MG</u> | <u>A205634 002</u> | Aug 25, 2016 |
| <u>AB</u> | | <u>60MG</u> | <u>A205634 003</u> | Aug 25, 2016 |
| <u>AB</u> | | <u>100MG</u> | <u>A205634 004</u> | Aug 25, 2016 |
| <u>AB</u> | | <u>200MG</u> | <u>A205634 005</u> | Aug 25, 2016 |
| <u>AB</u> | VINTAGE PHARMS LLC | <u>15MG</u> | <u>A075295 001</u> | Oct 28, 1998 |
| <u>AB</u> | | <u>30MG</u> | <u>A075295 002</u> | Oct 28, 1998 |
| <u>AB</u> | | <u>60MG</u> | <u>A075295 003</u> | Oct 28, 1998 |
| <u>AB</u> | | <u>100MG</u> | <u>A075295 004</u> | Sep 15, 2000 |
| <u>AB</u> | | <u>200MG</u> | <u>A075295 005</u> | Sep 15, 2000 |

MS CONTIN

| | | | | | |
|-----------|---|------------------|--------------|--------------------|--------------|
| <u>AB</u> | + | PURDUE PHARMA LP | <u>15MG</u> | <u>N019516 003</u> | Sep 12, 1989 |
| <u>AB</u> | + | | <u>30MG</u> | <u>N019516 001</u> | May 29, 1987 |
| <u>AB</u> | + | | <u>60MG</u> | <u>N019516 002</u> | Apr 08, 1988 |
| <u>AB</u> | + | | <u>100MG</u> | <u>N019516 004</u> | Jan 16, 1990 |
| <u>AB</u> | + | | <u>200MG</u> | <u>N019516 005</u> | Nov 08, 1993 |

MORPHABOND ER

| | | | | |
|---|--------------------|-------|-------------|--------------|
| + | DAIICHI SANKYO INC | 15MG | N206544 001 | Oct 02, 2015 |
| + | | 30MG | N206544 002 | Oct 02, 2015 |
| + | | 60MG | N206544 003 | Oct 02, 2015 |
| + | | 100MG | N206544 004 | Oct 02, 2015 |

MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

EMBEDA

| | | | | |
|---|-----------------|------------|-------------|--------------|
| + | ALPHARMA PHARMS | 20MG;0.8MG | N022321 001 | Aug 13, 2009 |
| + | | 30MG;1.2MG | N022321 002 | Aug 13, 2009 |
| + | | 50MG;2MG | N022321 003 | Aug 13, 2009 |
| + | | 60MG;2.4MG | N022321 004 | Aug 13, 2009 |
| + | | 80MG;3.2MG | N022321 005 | Aug 13, 2009 |
| + | | 100MG;4MG | N022321 006 | Aug 13, 2009 |

MOXIDECTIN

TABLET;ORAL

MOXIDECTIN

| | | | | |
|---|------|-----|-------------|--------------|
| + | MDGH | 2MG | N210867 001 | Jun 13, 2018 |
|---|------|-----|-------------|--------------|

MOXIFLOXACIN HYDROCHLORIDE

SOLUTION;INTRAVENOUS

MOXIFLOXACIN HYDROCHLORIDE

| | | | | |
|---|--------------------|----------------------------------------|-------------|--------------|
| + | FRESENIUS KABI USA | EQ 400MG BASE/250ML (EQ 1.6MG BASE/ML) | N205572 001 | Apr 03, 2015 |
|---|--------------------|----------------------------------------|-------------|--------------|

MOXIFLOXACIN HYDROCHLORIDE IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER

| | | | | |
|---|----------------|------------------------|-------------|--------------|
| ! | MYLAN LABS LTD | 400MG/250ML (1.6MG/ML) | A205833 001 | May 05, 2017 |
|---|----------------|------------------------|-------------|--------------|

SOLUTION/DROPS;OPHTHALMIC

MOXIFLOXACIN HYDROCHLORIDE

| | | | | |
|------------|------------------|---------------------|--------------------|--------------|
| <u>AT1</u> | AKORN | <u>EQ 0.5% BASE</u> | <u>A202916 001</u> | Nov 09, 2017 |
| <u>AT1</u> | APOTEX INC | <u>EQ 0.5% BASE</u> | <u>A090080 001</u> | Jun 30, 2017 |
| <u>AT1</u> | AUROBINDO PHARMA | <u>EQ 0.5% BASE</u> | <u>A206242 001</u> | Oct 04, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

MOXIFLOXACIN HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

MOXIFLOXACIN HYDROCHLORIDE

LTD

AT1 LUPIN LTD **EQ 0.5% BASE** **A202867 001** Sep 04, 2014**AT1** WATSON LABS INC **EQ 0.5% BASE** **A202525 001** Mar 06, 2015VIGAMOX**AT1** +! NOVARTIS PHARMS **EQ 0.5% BASE** **N021598 001** Apr 15, 2003

CORP

MOXEZA**AT2** +! NOVARTIS PHARMS **EQ 0.5% BASE** **N022428 001** Nov 19, 2010

CORP

MOXIFLOXACIN HYDROCHLORIDE**AT2** LUPIN LTD **EQ 0.5% BASE** **A204079 001** May 28, 2015

TABLET;ORAL

AVELOX**AB** +! BAYER HLTHCARE **EQ 400MG BASE** **N021085 001** Dec 10, 1999MOXIFLOXACIN HYDROCHLORIDE**AB** AUROBINDO PHARMA **EQ 400MG BASE** **A202632 001** Mar 04, 2014

LTD

AB CROSSMEDIKA SA **EQ 400MG BASE** **A205348 001** Jan 14, 2016**AB** DR REDDYS LABS LTD **EQ 400MG BASE** **A076938 001** Mar 04, 2014**AB** MSN LABS PVT LTD **EQ 400MG BASE** **A208682 001** Sep 22, 2017**AB** MYLAN PHARMS INC **EQ 400MG BASE** **A204635 001** Aug 31, 2015**AB** NOVEL LABS INC **EQ 400MG BASE** **A207285 001** Feb 13, 2017**AB** SUNSHINE LAKE **EQ 400MG BASE** **A206295 001** Sep 28, 2018**AB** TEVA PHARMS USA **EQ 400MG BASE** **A077437 001** Feb 18, 2014**AB** TORRENT PHARMS LTD **EQ 400MG BASE** **A200160 001** Apr 03, 2014MUPIROCIN

OINTMENT;TOPICAL

MUPIROCIN**AB** FOUGERA PHARMS **2%** **A065192 001** Nov 30, 2005**AB** GLENMARK PHARMS **2%** **A090480 001** Jun 08, 2011**AB** ! PERRIGO NEW YORK **2%** **A065123 001** Nov 07, 2003**AB** TARO **2%** **A065170 001** Sep 23, 2005**AB** TEVA **2%** **A065085 001** Nov 07, 2003

CENTANY

EX PERRIGO NEW YORK **2%** **N050788 001** Dec 04, 2002MUPIROCIN CALCIUM

CREAM;TOPICAL

MUPIROCIN**!** GLENMARK PHARMS INC **EQ 2% BASE** **A201587 001** Jan 24, 2013MYCOPHENOLATE MOFETIL

CAPSULE;ORAL

CELLCEPT**AB** +! ROCHE PALO **250MG** **N050722 001** May 03, 1995MYCOPHENOLATE MOFETIL**AB** ACCORD HLTHCARE **250MG** **A090253 001** May 04, 2009**AB** ALKEM LABS LTD **250MG** **A200197 001** Jun 13, 2013**AB** CONCORD BIOTECH LTD **250MG** **A210181 001** Jan 08, 2019**AB** MYLAN **250MG** **A065520 001** May 04, 2009**AB** SANDOZ **250MG** **A065379 001** Oct 15, 2008**AB** STRIDES PHARMA **250MG** **A090055 001** Jun 10, 2010**AB** TEVA PHARMS **250MG** **A065491 001** May 06, 2009**AB** VINTAGE PHARMS LLC **250MG** **A090111 001** Dec 22, 2009**AB** WEST-WARD PHARMS **250MG** **A065410 001** Jul 29, 2008

INT

AB ZHEJIANG HISUN **250MG** **A204077 001** Nov 13, 2017

PHARM

SUSPENSION;ORAL

CELLCEPT**AB** +! ROCHE PALO **200MG/ML** **N050759 001** Oct 01, 1998MYCOPHENOLATE MOFETIL**AB** ALKEM LABS LTD **200MG/ML** **A203005 001** Nov 14, 2014

TABLET;ORAL

CELLCEPT**AB** +! ROCHE PALO **500MG** **N050723 001** Jun 19, 1997MYCOPHENOLATE MOFETIL**AB** ACCORD HLTHCARE **500MG** **A065416 001** May 04, 2009**AB** ALKEM LABS LTD **500MG** **A091249 001** Nov 04, 2011**AB** MYLAN **500MG** **A065521 001** May 04, 2009**AB** OXFORD PHARMS **500MG** **A090606 001** Jul 16, 2010

PRESCRIPTION DRUG PRODUCT LIST

MYCOPHENOLATE MOFETIL

TABLET; ORAL

MYCOPHENOLATE MOFETIL

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | SANDOZ | <u>500MG</u> | <u>A065451</u> | <u>001</u> | Oct 15, 2008 |
| <u>AB</u> | STRIDES PHARMA | <u>500MG</u> | <u>A090456</u> | <u>001</u> | Jun 10, 2010 |
| <u>AB</u> | TEVA PHARMS | <u>500MG</u> | <u>A065457</u> | <u>001</u> | May 04, 2009 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>500MG</u> | <u>A065413</u> | <u>001</u> | Jul 29, 2008 |
| <u>AB</u> | ZHEJIANG HISUN PHARM | <u>500MG</u> | <u>A204076</u> | <u>001</u> | Nov 16, 2017 |

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

INJECTABLE; INJECTION

CELLCEPT

| | | | | | | |
|-----------|------------|------------|-------------------|----------------|------------|--------------|
| <u>AP</u> | <u>+</u> ! | ROCHE PALO | <u>500MG/VIAL</u> | <u>N050758</u> | <u>001</u> | Aug 12, 1998 |
|-----------|------------|------------|-------------------|----------------|------------|--------------|

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

| | | | | | | |
|-----------|--|-------------------------|-------------------|----------------|------------|--------------|
| <u>AP</u> | | AKORN INC | <u>500MG/VIAL</u> | <u>A204043</u> | <u>001</u> | Feb 28, 2017 |
| <u>AP</u> | | MYLAN LABS LTD | <u>500MG/VIAL</u> | <u>A203859</u> | <u>001</u> | Mar 31, 2017 |
| <u>AP</u> | | PAR STERILE PRODUCTS | <u>500MG/VIAL</u> | <u>A203575</u> | <u>001</u> | Oct 28, 2016 |
| <u>AP</u> | | ZYDUS PHARMS USA INC | <u>500MG/VIAL</u> | <u>A204473</u> | <u>001</u> | Aug 31, 2017 |

MYCOPHENOLIC ACID

TABLET, DELAYED RELEASE; ORAL

MYCOPHENOLIC ACID

| | | | | | | |
|-----------------|------------|------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | ACCORD HLTHCARE | <u>180MG</u> | <u>A202555</u> | <u>001</u> | Aug 23, 2017 |
| <u>AB</u> | | | <u>360MG</u> | <u>A202555</u> | <u>002</u> | Aug 23, 2017 |
| <u>AB</u> | | APOTEX INC | <u>180MG</u> | <u>A091558</u> | <u>001</u> | Aug 21, 2012 |
| <u>AB</u> | | | <u>360MG</u> | <u>A091558</u> | <u>002</u> | Aug 19, 2014 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>180MG</u> | <u>A091248</u> | <u>002</u> | Jan 08, 2014 |
| <u>AB</u> | | | <u>360MG</u> | <u>A091248</u> | <u>001</u> | Jan 08, 2014 |
| <u>AB</u> | | TEVA PHARMS USA | <u>180MG</u> | <u>A202720</u> | <u>001</u> | Oct 30, 2014 |
| <u>AB</u> | | | <u>360MG</u> | <u>A202720</u> | <u>002</u> | Oct 30, 2014 |
| <u>MYFORTIC</u> | | | | | | |
| <u>AB</u> | <u>+</u> | NOVARTIS | <u>180MG</u> | <u>N050791</u> | <u>001</u> | Feb 27, 2004 |
| <u>AB</u> | <u>+</u> ! | | <u>360MG</u> | <u>N050791</u> | <u>002</u> | Feb 27, 2004 |

NABILONE

CAPSULE; ORAL

CESAMET

| | | | | | |
|------------|---------------------|-----|----------------|------------|--------------|
| <u>+</u> ! | MYLAN SPECIALITY LP | 1MG | <u>N018677</u> | <u>001</u> | Dec 26, 1985 |
|------------|---------------------|-----|----------------|------------|--------------|

NABUMETONE

TABLET; ORAL

NABUMETONE

| | | | | | | |
|-----------|----------|---------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | APOTEX INC | <u>500MG</u> | <u>A090427</u> | <u>001</u> | Dec 30, 2011 |
| <u>AB</u> | | | <u>750MG</u> | <u>A090427</u> | <u>002</u> | Dec 30, 2011 |
| <u>AB</u> | | CASI PHARMS INC | <u>500MG</u> | <u>A075280</u> | <u>001</u> | Feb 25, 2002 |
| <u>AB</u> | | | <u>750MG</u> | <u>A075280</u> | <u>002</u> | Feb 25, 2002 |
| <u>AB</u> | | CHARTWELL MOLECULES | <u>500MG</u> | <u>A076009</u> | <u>001</u> | Jan 24, 2003 |
| <u>AB</u> | | | <u>750MG</u> | <u>A076009</u> | <u>002</u> | Jan 24, 2003 |
| <u>AB</u> | | IMPAX LABS INC | <u>500MG</u> | <u>A075189</u> | <u>001</u> | May 26, 2000 |
| <u>AB</u> | <u>!</u> | | <u>750MG</u> | <u>A075189</u> | <u>002</u> | Sep 24, 2001 |
| <u>AB</u> | | INVAGEN PHARMS | <u>500MG</u> | <u>A078671</u> | <u>001</u> | Mar 07, 2008 |
| <u>AB</u> | | | <u>750MG</u> | <u>A078671</u> | <u>002</u> | Mar 07, 2008 |
| <u>AB</u> | | LUPIN LTD | <u>500MG</u> | <u>A090445</u> | <u>001</u> | Jan 12, 2011 |
| <u>AB</u> | | | <u>750MG</u> | <u>A090445</u> | <u>002</u> | Jan 12, 2011 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>500MG</u> | <u>A090516</u> | <u>001</u> | Jul 12, 2010 |
| <u>AB</u> | | | <u>750MG</u> | <u>A090516</u> | <u>002</u> | Jul 12, 2010 |
| <u>AB</u> | | WATSON LABS | <u>500MG</u> | <u>A091083</u> | <u>001</u> | Jun 13, 2011 |
| <u>AB</u> | | | <u>750MG</u> | <u>A091083</u> | <u>002</u> | Jun 13, 2011 |

NADOLOL

TABLET; ORAL

CORGARD

| | | | | | | |
|-----------|------------|------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | <u>+</u> | US WORLDMEDS LLC | <u>20MG</u> | <u>N018063</u> | <u>005</u> | Oct 28, 1986 |
| <u>AB</u> | <u>+</u> | | <u>40MG</u> | <u>N018063</u> | <u>001</u> | |
| <u>AB</u> | <u>+</u> ! | | <u>80MG</u> | <u>N018063</u> | <u>002</u> | |

NADOLOL

| | | | | | | |
|-----------|--|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | AMNEAL PHARMS CO | <u>20MG</u> | <u>A208832</u> | <u>001</u> | Jun 02, 2017 |
| <u>AB</u> | | | <u>40MG</u> | <u>A208832</u> | <u>002</u> | Jun 02, 2017 |
| <u>AB</u> | | | <u>80MG</u> | <u>A208832</u> | <u>003</u> | Jun 02, 2017 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>40MG</u> | <u>A201893</u> | <u>001</u> | Sep 16, 2015 |
| <u>AB</u> | | | <u>80MG</u> | <u>A201893</u> | <u>002</u> | Sep 16, 2015 |

PRESCRIPTION DRUG PRODUCT LIST

NADOLOL

TABLET; ORAL

NADOLOL

| | | | | |
|-----------|-------------------------|-------------|--------------------|--------------|
| <u>AB</u> | BEXIMCO PHARMS USA | <u>20MG</u> | <u>A210955 001</u> | Jul 23, 2018 |
| <u>AB</u> | | <u>40MG</u> | <u>A210955 002</u> | Jul 23, 2018 |
| <u>AB</u> | | <u>80MG</u> | <u>A210955 003</u> | Jul 23, 2018 |
| <u>AB</u> | INVAGEN PHARMS | <u>20MG</u> | <u>A203455 001</u> | Dec 18, 2015 |
| <u>AB</u> | | <u>40MG</u> | <u>A203455 002</u> | Dec 18, 2015 |
| <u>AB</u> | | <u>80MG</u> | <u>A203455 003</u> | Dec 18, 2015 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>20MG</u> | <u>A074229 001</u> | Aug 30, 1996 |
| <u>AB</u> | | <u>40MG</u> | <u>A074229 002</u> | Aug 30, 1996 |
| <u>AB</u> | | <u>80MG</u> | <u>A074255 001</u> | Jan 24, 1996 |
| <u>AB</u> | LUPIN LTD | <u>20MG</u> | <u>A209309 001</u> | Oct 05, 2017 |
| <u>AB</u> | | <u>40MG</u> | <u>A209309 002</u> | Oct 05, 2017 |
| <u>AB</u> | | <u>80MG</u> | <u>A209309 003</u> | Oct 05, 2017 |
| <u>AB</u> | MYLAN | <u>20MG</u> | <u>A074172 001</u> | Oct 31, 1993 |
| <u>AB</u> | | <u>40MG</u> | <u>A074172 002</u> | Oct 31, 1993 |
| <u>AB</u> | | <u>80MG</u> | <u>A074172 003</u> | Oct 31, 1993 |
| <u>AB</u> | NOVAST LABS | <u>20MG</u> | <u>A210786 001</u> | Jun 01, 2018 |
| <u>AB</u> | | <u>40MG</u> | <u>A210786 002</u> | Jun 01, 2018 |
| <u>AB</u> | | <u>80MG</u> | <u>A210786 003</u> | Jun 01, 2018 |
| <u>AB</u> | SANDOZ | <u>20MG</u> | <u>A074501 001</u> | Nov 09, 1995 |
| <u>AB</u> | | <u>40MG</u> | <u>A074501 002</u> | Nov 09, 1995 |
| <u>AB</u> | | <u>80MG</u> | <u>A074501 003</u> | Nov 09, 1995 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>20MG</u> | <u>A207761 001</u> | Jul 28, 2017 |
| <u>AB</u> | | <u>40MG</u> | <u>A207761 002</u> | Jul 28, 2017 |
| <u>AB</u> | | <u>80MG</u> | <u>A207761 003</u> | Jul 28, 2017 |

NAFARELIN ACETATE

SPRAY, METERED; NASAL

SYNAREL

+! GD SEARLE LLC

EQ 0.2MG BASE/SPRAY

N019886 001 Feb 13, 1990

NAFCILLIN SODIUM

INJECTABLE; INJECTION

NAFCILLIN SODIUM

| | | | | |
|------------------------------|-------------------------|--------------------------|--------------------|--------------|
| <u>AP</u> | ANTIBIOTICE | <u>EQ 1GM BASE/VIAL</u> | <u>A090560 001</u> | Oct 03, 2011 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A090560 002</u> | Oct 03, 2011 |
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>EQ 1GM BASE/VIAL</u> | <u>A091613 001</u> | Dec 26, 2012 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A091613 002</u> | Dec 26, 2012 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A091614 001</u> | Dec 26, 2012 |
| <u>AP</u> | ISTITUTO BIO ITA SPA | <u>EQ 1GM BASE/VIAL</u> | <u>A090002 001</u> | Jun 30, 2011 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A090002 002</u> | Jun 30, 2011 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A090005 001</u> | Apr 20, 2011 |
| <u>AP</u> | MYLAN LABS LTD | <u>EQ 1GM BASE/VIAL</u> | <u>A200002 001</u> | Apr 07, 2014 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A200002 002</u> | Apr 07, 2014 |
| <u>AP</u> | SAGENT PHARMS | <u>EQ 1GM BASE/VIAL</u> | <u>A090582 001</u> | Aug 24, 2012 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A090582 002</u> | Aug 24, 2012 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A090580 001</u> | Aug 24, 2012 |
| <u>AP</u> | ! SANDOZ | <u>EQ 1GM BASE/VIAL</u> | <u>A062527 002</u> | Aug 02, 1984 |
| <u>AP</u> | ! | <u>EQ 1GM BASE/VIAL</u> | <u>A062732 001</u> | Dec 23, 1986 |
| <u>AP</u> | ! | <u>EQ 2GM BASE/VIAL</u> | <u>A062527 003</u> | Aug 02, 1984 |
| <u>AP</u> | ! | <u>EQ 2GM BASE/VIAL</u> | <u>A062732 002</u> | Dec 23, 1986 |
| <u>AP</u> | ! | <u>EQ 10GM BASE/VIAL</u> | <u>A062527 004</u> | Aug 02, 1984 |
| NALLPEN IN PLASTIC CONTAINER | | | | |
| | +! BAXTER HLTHCARE | EQ 20MG BASE/ML | N050655 001 | Oct 31, 1989 |
| | +! | EQ 2GM BASE/100ML | N050655 002 | Oct 31, 1989 |

NAFTIFINE HYDROCHLORIDE

CREAM; TOPICAL

NAFTIFINE HYDROCHLORIDE

| | | | | |
|-------------------------|-----------------------|-----------|--------------------|--------------|
| <u>AB</u> | TARO PHARMS | <u>2%</u> | <u>A206901 001</u> | Jan 06, 2016 |
| <u>AB</u> | TOLMAR | <u>2%</u> | <u>A206960 001</u> | Apr 10, 2017 |
| <u>NAFTIN</u> | | | | |
| <u>AB</u> | +! SEBELA IRELAND LTD | <u>2%</u> | <u>N019599 002</u> | Jan 13, 2012 |
| NAFTIFINE HYDROCHLORIDE | | | | |
| | ! TARO PHARMS | 1% | A205975 001 | Sep 08, 2016 |
| GEL; TOPICAL | | | | |
| NAFTIN | | | | |
| | +! SEBELA IRELAND LTD | 1% | N019356 001 | Jun 18, 1990 |
| | +! | 2% | N204286 001 | Jun 27, 2013 |

PRESCRIPTION DRUG PRODUCT LIST

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE HYDROCHLORIDE

| | | | | | |
|-----------|---|----------------|----------------|--------------------|--------------|
| AP | ! | HOSPIRA | 10MG/ML | A070914 001 | Feb 03, 1989 |
| AP | ! | | 10MG/ML | A070915 001 | Feb 03, 1989 |
| AP | ! | | 20MG/ML | A070916 001 | Feb 03, 1989 |
| AP | ! | | 20MG/ML | A070918 001 | Feb 03, 1989 |
| AP | | MYLAN LABS LTD | 10MG/ML | A207595 001 | Jan 11, 2019 |
| AP | | | 20MG/ML | A207595 002 | Jan 11, 2019 |

NALDEMEDINE TOSYLATE

TABLET; ORAL

SYMPROIC

| | | | | | |
|---|---|--------------|---------------|-------------|--------------|
| + | ! | SHIONOGI INC | EQ 0.2MG BASE | N208854 001 | Mar 23, 2017 |
|---|---|--------------|---------------|-------------|--------------|

NALOXEGOL OXALATE

TABLET; ORAL

MOVANTIK

| | | | | | |
|---|---|--------------------|----------------|-------------|--------------|
| + | | ASTRAZENECA PHARMS | EQ 12.5MG BASE | N204760 001 | Sep 16, 2014 |
| + | ! | | EQ 25MG BASE | N204760 002 | Sep 16, 2014 |

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

| | | | | | |
|-----------|--|-------------------------|-----------------|--------------------|--------------|
| AP | | WEST-WARD PHARMS INT | 0.4MG/ML | A070299 001 | Sep 24, 1986 |
|-----------|--|-------------------------|-----------------|--------------------|--------------|

NALOXONE HYDROCHLORIDE

| | | | | | |
|---------------------------------------|---|-------------------------|-----------------------|--------------------|--------------|
| AP | | AKORN | 0.4MG/ML | A208871 001 | Feb 28, 2017 |
| AP | | | 0.4MG/ML | A208872 001 | Mar 14, 2017 |
| AP | ! | HOSPIRA | 0.4MG/ML | A070172 001 | Sep 24, 1986 |
| AP | ! | | 0.4MG/ML | A070254 001 | Jan 07, 1987 |
| AP | ! | | 0.4MG/ML | A070256 001 | Jan 07, 1987 |
| AP | ! | | 0.4MG/ML | A070257 001 | Jan 07, 1987 |
| AP | | INTL MEDICATION | 0.4MG/ML | A070639 001 | Sep 24, 1986 |
| AP | | MYLAN INSTITUTIONAL | 0.4MG/ML | A204997 001 | Mar 06, 2014 |
| AP | | | 0.4MG/ML | A205014 001 | Jun 29, 2016 |
| AP | | RENAISSANCE SSA LLC | 0.4MG/ML | A207846 001 | Dec 17, 2018 |
| AP | | SOMERSET THERAPS LLC | 0.4MG/ML | A207633 001 | Aug 08, 2017 |
| AP | | | 0.4MG/ML | A207634 001 | Jul 26, 2017 |
| ! | | INTL MEDICATION | 1MG/ML | A072076 001 | Mar 24, 1988 |
| SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS | | | | | |
| EVZIO | | | | | |
| + | ! | KALEO INC | 2MG/0.4ML (2MG/0.4ML) | N209862 001 | Oct 19, 2016 |
| SPRAY, METERED; NASAL | | | | | |
| NARCAN | | | | | |
| + | ! | ADAPT | 4MG/SPRAY | N208411 001 | Nov 18, 2015 |

NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE

| | | | | | |
|-----------|---|--------------------|------------------------------------|--------------------|--------------|
| AB | | GAVIS PHARMS | EQ 0.5MG BASE; EQ 50MG BASE | A075735 001 | Jul 11, 2001 |
| AB | | SUN PHARM INDS LTD | EQ 0.5MG BASE; EQ 50MG BASE | A075523 001 | Mar 17, 2000 |
| AB | ! | WATSON LABS | EQ 0.5MG BASE; EQ 50MG BASE | A074736 001 | Jan 21, 1997 |

NALTREXONE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

VIVITROL

| | | | | | |
|---|---|----------|------------|-------------|--------------|
| + | ! | ALKERMES | 380MG/VIAL | N021897 001 | Apr 13, 2006 |
|---|---|----------|------------|-------------|--------------|

NALTREXONE HYDROCHLORIDE

TABLET; ORAL

NALTREXONE HYDROCHLORIDE

| | | | | | |
|-----------|---|-------------------|-------------|--------------------|--------------|
| AB | | ACCORD HLTHCARE | 50MG | A091205 001 | Aug 17, 2011 |
| AB | | APOTEX INC | 50MG | A207905 001 | Jul 21, 2017 |
| AB | | BARR | 50MG | A074918 001 | May 08, 1998 |
| AB | | ELITE LABS | 50MG | A075274 001 | May 26, 1999 |
| AB | ! | SPECGX LLC | 50MG | A076264 002 | Mar 22, 2002 |
| AB | | SUN PHARMA GLOBAL | 50MG | A090356 001 | Feb 24, 2012 |
| | | SPECGX LLC | 25MG | A076264 001 | Mar 22, 2002 |
| | | | 100MG | A076264 003 | Mar 22, 2002 |

PRESCRIPTION DRUG PRODUCT LIST

NANDROLONE DECANOATE

INJECTABLE; INJECTION

NANDROLONE DECANOATE

! LUITPOLD

200MG/ML

A091252 001 Aug 30, 2010

NAPROXEN

SUSPENSION; ORAL

NAPROSYNAB +! ATNAHS PHARMA US25MG/MLN018965 001 Mar 23, 1987NAPROXENAB WEST-WARD PHARMS
INT25MG/MLA074190 001 Mar 30, 1994

TABLET; ORAL

NAPROSYNAB +! ATNAHS PHARMA US500MGN017581 004 Apr 15, 1982NAPROXENAB AMNEAL PHARMS NY250MGA075927 001 Dec 18, 2001AB375MGA075927 002 Dec 18, 2001AB500MGA075927 003 Dec 18, 2001AB AUROBINDO PHARMA
LTD250MGA200429 001 Nov 08, 2011AB375MGA200429 002 Nov 08, 2011AB500MGA200429 003 Nov 08, 2011AB GLENMARK GENERICS250MGA078250 001 Mar 28, 2007AB375MGA078250 002 Mar 28, 2007AB500MGA078250 003 Mar 28, 2007AB INVAGEN PHARMS250MGA091305 001 Aug 24, 2011AB375MGA091305 002 Aug 24, 2011AB500MGA091305 003 Aug 24, 2011AB MARKSANS PHARMA250MGA091416 001 Feb 14, 2011AB375MGA091416 002 Feb 14, 2011AB500MGA091416 003 Feb 14, 2011AB MYLAN250MGA074121 001 Dec 21, 1993AB375MGA074121 002 Dec 21, 1993AB500MGA074121 003 Dec 21, 1993AB PERRIGO R AND D250MGA077339 001 Apr 27, 2005AB375MGA077339 002 Apr 27, 2005AB500MGA077339 003 Apr 27, 2005AB TEVA250MGA074201 001 Dec 21, 1993AB375MGA074201 002 Dec 21, 1993AB500MGA074201 003 Dec 21, 1993AB ZYDUS PHARMS USA250MGA078620 001 Jun 07, 2007AB375MGA078620 002 Jun 07, 2007AB500MGA078620 003 Jun 07, 2007

TABLET, DELAYED RELEASE; ORAL

EC-NAPROSYNAB +! ATNAHS PHARMA US375MGN020067 002 Oct 14, 1994AB +!500MGN020067 003 Oct 14, 1994NAPROXENAB INVAGEN PHARMS375MGA091432 001 Sep 19, 2011AB500MGA091432 002 Sep 19, 2011AB PLIVA375MGA075337 001 May 26, 1999AB500MGA075337 002 May 26, 1999AB TEVA375MGA075227 001 Jun 30, 1998AB500MGA075227 002 Jun 30, 1998NAPROXEN SODIUM

TABLET; ORAL

ANAPROX DSAB +! ATNAHS PHARMA USEQ 500MG BASEN018164 003 Sep 30, 1987NAPROXEN SODIUMAB AMNEAL PHARMS NYEQ 250MG BASEA078432 001 Apr 25, 2007ABEQ 500MG BASEA078432 002 Apr 25, 2007AB AUROBINDO PHARMAEQ 250MG BASEA200629 001 Oct 31, 2011AB LTDEQ 500MG BASEA200629 002 Oct 31, 2011AB DR REDDYS LABS LTDEQ 250MG BASEA078486 001 Jul 26, 2007ABEQ 500MG BASEA078486 002 Jul 26, 2007AB GLENMARK PHARMS LTDEQ 250MG BASEA078314 001 Apr 27, 2007ABEQ 500MG BASEA078314 002 Apr 27, 2007AB TEVAEQ 250MG BASEA074198 001 Dec 21, 1993ABEQ 500MG BASEA074198 002 Dec 21, 1993

PRESCRIPTION DRUG PRODUCT LIST

NAPROXEN SODIUM

TABLET, EXTENDED RELEASE;ORAL

NAPRELAN

| | | | | | |
|-----------|---|---------------|----------------------|--------------------|--------------|
| <u>AB</u> | + | ALVOGEN MALTA | <u>EQ 375MG BASE</u> | <u>N020353 001</u> | Jan 05, 1996 |
| <u>AB</u> | + | | <u>EQ 500MG BASE</u> | <u>N020353 002</u> | Jan 05, 1996 |
| <u>AB</u> | + | ! | <u>EQ 750MG BASE</u> | <u>N020353 003</u> | Jan 05, 1996 |

NAPROXEN SODIUM

| | | | | | |
|-----------|--|---------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS LABS FL INC | <u>EQ 375MG BASE</u> | <u>A075416 002</u> | Apr 23, 2003 |
| <u>AB</u> | | | <u>EQ 500MG BASE</u> | <u>A075416 001</u> | Aug 27, 2002 |
| <u>AB</u> | | | <u>EQ 750MG BASE</u> | <u>A075416 003</u> | Aug 11, 2016 |

NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE

TABLET;ORAL

SUMATRIPTAN AND NAPROXEN SODIUM

| | | | | | |
|-----------|--|----------------------|---------------------------|--------------------|--------------|
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>500MG;EQ 85MG BASE</u> | <u>A207457 001</u> | Feb 15, 2018 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>500MG;EQ 85MG BASE</u> | <u>A090872 001</u> | Sep 04, 2018 |
| <u>AB</u> | | SUN PHARMA GLOBAL | <u>500MG;EQ 85MG BASE</u> | <u>A202803 001</u> | Jul 20, 2018 |

TREXIMET

| | | | | | | |
|-----------|---|---|--------------------|---------------------------|--------------------|--------------|
| <u>AB</u> | + | ! | PERNIX IRELAND LTD | <u>500MG;EQ 85MG BASE</u> | <u>N021926 001</u> | Apr 15, 2008 |
|-----------|---|---|--------------------|---------------------------|--------------------|--------------|

NARATRIPTAN HYDROCHLORIDE

TABLET;ORAL

AMERGE

| | | | | | |
|-----------|---|---------------------|--------------------|--------------------|--------------|
| <u>AB</u> | + | GLAXOSMITHKLINE LLC | <u>EQ 1MG BASE</u> | <u>N020763 002</u> | Feb 10, 1998 |
| <u>AB</u> | + | ! | | <u>N020763 001</u> | Feb 10, 1998 |

NARATRIPTAN

| | | | | | |
|-----------|--|----------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | CASI PHARMS INC | <u>EQ 1MG BASE</u> | <u>A090288 001</u> | Jul 07, 2010 |
| <u>AB</u> | | | <u>EQ 2.5MG BASE</u> | <u>A090288 002</u> | Jul 07, 2010 |
| <u>AB</u> | | HERITAGE PHARMS INC | <u>EQ 1MG BASE</u> | <u>A200502 001</u> | Feb 28, 2011 |
| <u>AB</u> | | | <u>EQ 2.5MG BASE</u> | <u>A200502 002</u> | Feb 28, 2011 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>EQ 1MG BASE</u> | <u>A202431 001</u> | May 31, 2012 |
| <u>AB</u> | | | <u>EQ 2.5MG BASE</u> | <u>A202431 002</u> | May 31, 2012 |
| <u>AB</u> | | ORCHID HLTHCARE | <u>EQ 1MG BASE</u> | <u>A091441 001</u> | Apr 30, 2012 |
| <u>AB</u> | | | <u>EQ 2.5MG BASE</u> | <u>A091441 002</u> | Apr 30, 2012 |
| <u>AB</u> | | PADDOCK LLC | <u>EQ 1MG BASE</u> | <u>A091326 001</u> | Jul 08, 2010 |
| <u>AB</u> | | | <u>EQ 2.5MG BASE</u> | <u>A091326 002</u> | Jul 08, 2010 |
| <u>AB</u> | | SUN PHARM INDS LTD | <u>EQ 2.5MG BASE</u> | <u>A091552 001</u> | Feb 14, 2011 |
| <u>AB</u> | | TEVA PHARMS | <u>EQ 1MG BASE</u> | <u>A078751 001</u> | Jul 07, 2010 |
| <u>AB</u> | | | <u>EQ 2.5MG BASE</u> | <u>A078751 002</u> | Jul 07, 2010 |
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>EQ 1MG BASE</u> | <u>A090381 001</u> | Jul 07, 2010 |
| <u>AB</u> | | | <u>EQ 2.5MG BASE</u> | <u>A090381 002</u> | Jul 07, 2010 |

NATAMYCIN

SUSPENSION;OPHTHALMIC

NATACYN

| | | | | | |
|---|---|----------------------|----|--------------------|--|
| + | ! | NOVARTIS PHARMS CORP | 5% | <u>N050514 001</u> | |
|---|---|----------------------|----|--------------------|--|

NATEGLINIDE

TABLET;ORAL

NATEGLINIDE

| | | | | | |
|-----------|--|----------------------|--------------|--------------------|--------------|
| <u>AB</u> | | ALVOGEN MALTA | <u>60MG</u> | <u>A205055 001</u> | Dec 11, 2015 |
| <u>AB</u> | | | <u>120MG</u> | <u>A205055 002</u> | Dec 11, 2015 |
| <u>AB</u> | | DR REDDYS LABS LTD | <u>60MG</u> | <u>A077461 001</u> | Sep 09, 2009 |
| <u>AB</u> | | | <u>120MG</u> | <u>A077461 002</u> | Sep 09, 2009 |
| <u>AB</u> | | PAR PHARM | <u>60MG</u> | <u>A077463 001</u> | Sep 09, 2009 |
| <u>AB</u> | | | <u>120MG</u> | <u>A077463 002</u> | Sep 09, 2009 |
| <u>AB</u> | | WATSON LABS | <u>60MG</u> | <u>A077462 001</u> | Mar 30, 2011 |
| <u>AB</u> | | | <u>120MG</u> | <u>A077462 002</u> | Mar 30, 2011 |
| <u>AB</u> | | WILSHIRE PHARMS INC | <u>60MG</u> | <u>A205544 001</u> | Jun 18, 2018 |
| <u>AB</u> | | | <u>120MG</u> | <u>A205544 002</u> | Jun 18, 2018 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>60MG</u> | <u>A205248 001</u> | Jul 06, 2016 |
| <u>AB</u> | | | <u>120MG</u> | <u>A205248 002</u> | Jul 06, 2016 |

STARLIX

| | | | | | |
|-----------|---|----------|-------------|--------------------|--------------|
| <u>AB</u> | + | NOVARTIS | <u>60MG</u> | <u>N021204 001</u> | Dec 22, 2000 |
| <u>AB</u> | + | ! | | <u>N021204 002</u> | Dec 22, 2000 |

PRESCRIPTION DRUG PRODUCT LIST

NEBIVOLOL HYDROCHLORIDE

TABLET; ORAL

BYSTOLIC

| | | | | | |
|-----------|---|--------------------|----------------------|--------------------|--------------|
| <u>AB</u> | + | ALLERGAN SALES LLC | <u>EQ 2.5MG BASE</u> | <u>N021742 002</u> | Dec 17, 2007 |
| <u>AB</u> | + | | <u>EQ 5MG BASE</u> | <u>N021742 003</u> | Dec 17, 2007 |
| <u>AB</u> | + | | <u>EQ 10MG BASE</u> | <u>N021742 004</u> | Dec 17, 2007 |
| <u>AB</u> | + | ! | <u>EQ 20MG BASE</u> | <u>N021742 005</u> | Oct 08, 2008 |

NEBIVOLOL HYDROCHLORIDE; VALSARTAN

TABLET; ORAL

BYVALSON

| | | | | | |
|---|---|--------------------|-------------------|-------------|--------------|
| + | ! | ALLERGAN SALES LLC | EQ 5MG BASE; 80MG | N206302 001 | Jun 03, 2016 |
|---|---|--------------------|-------------------|-------------|--------------|

NEDOCROMIL SODIUM

SOLUTION/DROPS; OPHTHALMIC

ALOCRIIL

| | | | | | | |
|-----------|---|---|----------|-----------|--------------------|--------------|
| <u>AT</u> | + | ! | ALLERGAN | <u>2%</u> | <u>N021009 001</u> | Dec 08, 1999 |
| <u>AT</u> | | | | | <u>A090638 001</u> | Aug 22, 2012 |

NEFAZODONE HYDROCHLORIDE

TABLET; ORAL

NEFAZODONE HYDROCHLORIDE

| | | | | | |
|--|--|------|-------|-------------|--------------|
| | | TEVA | 50MG | A076037 001 | Sep 16, 2003 |
| | | | 100MG | A076037 002 | Sep 16, 2003 |
| | | | 150MG | A076037 003 | Sep 16, 2003 |
| | | | 200MG | A076037 004 | Sep 16, 2003 |
| | | ! | 250MG | A076037 005 | Sep 16, 2003 |

NELARABINE

INJECTABLE; INTRAVENOUS

ARRANON

| | | | | | |
|---|---|-----------------|---------------------|-------------|--------------|
| + | ! | NOVARTIS PHARMS | 250MG/50ML (5MG/ML) | N021877 001 | Oct 28, 2005 |
|---|---|-----------------|---------------------|-------------|--------------|

CORP

NELFINAVIR MESYLATE

TABLET; ORAL

VIRACEPT

| | | | | | |
|---|---|----------------|---------------|-------------|--------------|
| + | ! | AGOURON PHARMS | EQ 250MG BASE | N020779 001 | Mar 14, 1997 |
| + | ! | | EQ 625MG BASE | N021503 001 | Apr 30, 2003 |

NEOMYCIN SULFATE

TABLET; ORAL

NEOMYCIN SULFATE

| | | | | | |
|-----------|---|--------------------|--------------|--------------------|--------------|
| <u>AA</u> | | BRECKENRIDGE PHARM | <u>500MG</u> | <u>A065468 001</u> | Mar 29, 2010 |
| <u>AA</u> | | LANNETT CO INC | <u>500MG</u> | <u>A204435 001</u> | Jun 10, 2016 |
| <u>AA</u> | ! | TEVA | <u>500MG</u> | <u>A060304 001</u> | |
| <u>AA</u> | | X GEN PHARMS | <u>500MG</u> | <u>A065220 001</u> | Jul 28, 2006 |

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION; IRRIGATION

NEOMYCIN AND POLYMYXIN B SULFATE

| | | | | | |
|-----------|---|----------------|------------------------------------------|--------------------|--------------|
| <u>AT</u> | | WATSON LABS | <u>EQ 40MG BASE/ML; 200,000 UNITS/ML</u> | <u>A062664 001</u> | Apr 08, 1986 |
| <u>AT</u> | | X GEN PHARMS | <u>EQ 40MG BASE/ML; 200,000 UNITS/ML</u> | <u>A065106 001</u> | Jan 31, 2006 |
| <u>AT</u> | | | <u>EQ 40MG BASE/ML; 200,000 UNITS/ML</u> | <u>A065108 001</u> | Jan 31, 2006 |
| <u>AT</u> | ! | MONARCH PHARMS | <u>EQ 40MG BASE/ML; 200,000 UNITS/ML</u> | <u>A060707 001</u> | |

NEOSTIGMINE METHYLSULFATE

SOLUTION; INTRAVENOUS

BLOXIVERZ

| | | | | | | |
|-----------|---|---|---------------|----------------------------|--------------------|--------------|
| <u>AP</u> | + | ! | AVADEL LEGACY | <u>5MG/10ML (0.5MG/ML)</u> | <u>N204078 001</u> | May 31, 2013 |
| <u>AP</u> | + | ! | | <u>10MG/10ML (1MG/ML)</u> | <u>N204078 002</u> | May 31, 2013 |

NEOSTIGMINE METHYLSULFATE

| | | | | | |
|-----------|--|--------------------|----------------------------|--------------------|--------------|
| <u>AP</u> | | AMNEAL PHARMS CO | <u>5MG/10ML (0.5MG/ML)</u> | <u>A210051 001</u> | Jun 15, 2018 |
| <u>AP</u> | | | <u>10MG/10ML (1MG/ML)</u> | <u>A210051 002</u> | Jun 15, 2018 |
| <u>AP</u> | | AMPHASTAR PHARMS | <u>5MG/10ML (0.5MG/ML)</u> | <u>A209933 001</u> | Sep 25, 2017 |
| <u>AP</u> | | INC | <u>10MG/10ML (1MG/ML)</u> | <u>A209933 002</u> | Sep 25, 2017 |
| <u>AP</u> | | DR REDDYS LABS LTD | <u>5MG/10ML (0.5MG/ML)</u> | <u>A209135 001</u> | Jul 10, 2018 |
| <u>AP</u> | | | <u>10MG/10ML (1MG/ML)</u> | <u>A209135 002</u> | Jul 10, 2018 |
| <u>AP</u> | | EUROHLTH INTL SARL | <u>5MG/10ML (0.5MG/ML)</u> | <u>A207042 001</u> | Dec 28, 2015 |
| <u>AP</u> | | | <u>10MG/10ML (1MG/ML)</u> | <u>A207042 002</u> | Dec 28, 2015 |
| <u>AP</u> | | LUITPOLD | <u>5MG/10ML (0.5MG/ML)</u> | <u>A209182 001</u> | May 04, 2018 |
| <u>AP</u> | | | <u>10MG/10ML (1MG/ML)</u> | <u>A209182 002</u> | May 04, 2018 |
| <u>AP</u> | | PAR STERILE | <u>5MG/10ML (0.5MG/ML)</u> | <u>A208405 001</u> | Apr 26, 2017 |
| | | PRODUCTS | | | |

PRESCRIPTION DRUG PRODUCT LIST

NEOSTIGMINE METHYLSULFATE

SOLUTION; INTRAVENOUS

NEOSTIGMINE METHYLSULFATE

| | | | | |
|-----------|---------------------|----------------------------|--------------------|--------------|
| AP | | 10MG/10ML (1MG/ML) | A208405 002 | Apr 26, 2017 |
| AP | RENAISSANCE SSA LLC | 5MG/10ML (0.5MG/ML) | A210989 001 | Aug 22, 2018 |
| AP | | 10MG/10ML (1MG/ML) | A210989 002 | Aug 22, 2018 |
| | FRESENIUS KABI USA | 3MG/3ML (1MG/ML) | N203629 003 | Sep 18, 2018 |
| | | 5MG/10ML (0.5MG/ML) | N203629 001 | Jan 08, 2015 |
| | | 10MG/10ML (1MG/ML) | N203629 002 | Jan 08, 2015 |

NEPAFENAC

SUSPENSION/DROPS; OPHTHALMIC

ILEVRO

| | | | | |
|---|----------------------|------|-------------|--------------|
| + | NOVARTIS PHARMS CORP | 0.3% | N203491 001 | Oct 16, 2012 |
|---|----------------------|------|-------------|--------------|

NEVANAC

| | | | | |
|---|----------------------|------|-------------|--------------|
| + | NOVARTIS PHARMS CORP | 0.1% | N021862 001 | Aug 19, 2005 |
|---|----------------------|------|-------------|--------------|

NERATINIB MALEATE

TABLET; ORAL

NERLYNX

| | | | | |
|---|--------------|--------------|-------------|--------------|
| + | PUMA BIOTECH | EQ 40MG BASE | N208051 001 | Jul 17, 2017 |
|---|--------------|--------------|-------------|--------------|

NESIRITIDE RECOMBINANT

FOR SOLUTION; INTRAVENOUS

NATRECOR

| | | | | |
|---|-----------|------------|-------------|--------------|
| + | SCIOS LLC | 1.5MG/VIAL | N020920 001 | Aug 10, 2001 |
|---|-----------|------------|-------------|--------------|

NETARSUDIL DIMESYLATE

SOLUTION/DROPS; OPHTHALMIC

RHOPRESSA

| | | | | |
|---|------------------|---------------|-------------|--------------|
| + | AERIE PHARMS INC | EQ 0.02% BASE | N208254 001 | Dec 18, 2017 |
|---|------------------|---------------|-------------|--------------|

NETUPITANT; PALONOSETRON HYDROCHLORIDE

CAPSULE; ORAL

AKYNZEO

| | | | | |
|---|------------------|----------------------|-------------|--------------|
| + | HELSINN HLTHCARE | 300MG; EQ 0.5MG BASE | N205718 001 | Oct 10, 2014 |
|---|------------------|----------------------|-------------|--------------|

NEVIRAPINE

SUSPENSION; ORAL

NEVIRAPINE

| | | | | |
|-----------|-----------|-----------------|--------------------|--------------|
| AA | AUROBINDO | 50MG/5ML | A077702 001 | May 22, 2012 |
| AA | CIPLA | 50MG/5ML | A207684 001 | Aug 03, 2017 |

VIRAMUNE

| | | | | | |
|-----------|---|----------------------|-----------------|--------------------|--------------|
| AA | + | BOEHRINGER INGELHEIM | 50MG/5ML | N020933 001 | Sep 11, 1998 |
|-----------|---|----------------------|-----------------|--------------------|--------------|

TABLET; ORAL

NEVIRAPINE

| | | | | |
|-----------|---------------------|--------------|--------------------|--------------|
| AB | AUROBINDO | 200MG | A077521 001 | May 22, 2012 |
| AB | CIPLA | 200MG | A077956 001 | May 22, 2012 |
| AB | HETERO LABS LTD III | 200MG | A078584 001 | May 22, 2012 |
| AB | MACLEODS PHARMS LTD | 200MG | A090688 001 | Jan 14, 2019 |
| AB | MICRO LABS LTD | 200MG | A203080 001 | May 22, 2012 |
| AB | MYLAN LABS | 200MG | A078864 001 | May 22, 2012 |
| AB | MYLAN PHARMS INC | 200MG | A202523 001 | May 22, 2012 |
| AB | PRINSTON INC | 200MG | A078644 001 | May 22, 2012 |
| AB | STRIDES PHARMA | 200MG | A078195 001 | May 22, 2012 |

VIRAMUNE

| | | | | | |
|-----------|---|----------------------|--------------|--------------------|--------------|
| AB | + | BOEHRINGER INGELHEIM | 200MG | N020636 001 | Jun 21, 1996 |
|-----------|---|----------------------|--------------|--------------------|--------------|

TABLET, EXTENDED RELEASE; ORAL

NEVIRAPINE

| | | | | |
|-----------|----------------------|--------------|--------------------|--------------|
| AB | ALVOGEN MALTA | 100MG | A204621 002 | Nov 09, 2015 |
| AB | | 400MG | A204621 001 | Jul 10, 2015 |
| AB | AUROBINDO PHARMA LTD | 100MG | A208616 001 | Nov 23, 2016 |
| AB | | 400MG | A207698 001 | Feb 28, 2017 |
| AB | CIPLA | 400MG | A206448 001 | Oct 15, 2015 |
| AB | MACLEODS PHARMS LTD | 400MG | A206879 001 | Oct 06, 2017 |
| AB | MYLAN PHARMS INC | 100MG | A206271 001 | Nov 09, 2015 |
| AB | | 400MG | A205651 001 | Oct 27, 2014 |
| AB | SANDOZ INC | 400MG | A203411 001 | Apr 03, 2014 |

VIRAMUNE XR

| | | | | | |
|-----------|---|----------------------|--------------|--------------------|--------------|
| AB | + | BOEHRINGER INGELHEIM | 100MG | N201152 002 | Nov 08, 2012 |
| AB | + | | 400MG | N201152 001 | Mar 25, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

NIACIN

TABLET; ORAL

NIACIN

| | | | | |
|-----------|-----------|--------------|--------------------|--------------|
| AA | WOCKHARDT | 500MG | A081134 001 | Apr 28, 1992 |
|-----------|-----------|--------------|--------------------|--------------|

NIACOR

| | | | | | |
|-----------|---|-----------------|--------------|--------------------|--------------|
| AA | ! | AVONDALE PHARMS | 500MG | A040378 001 | May 03, 2000 |
|-----------|---|-----------------|--------------|--------------------|--------------|

TABLET, EXTENDED RELEASE; ORAL

NIACIN

| | | | | | |
|-----------|--|---------------|--------------|--------------------|--------------|
| AB | | AMNEAL PHARMS | 500MG | A203578 001 | Jul 24, 2015 |
|-----------|--|---------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 750MG | A204178 001 | Dec 11, 2015 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|------------|--------------------|--------------|
| AB | | | 1GM | A203578 002 | Jul 24, 2015 |
|-----------|--|--|------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------------|--------------|--------------------|--------------|
| AB | | AUROBINDO PHARMA LTD | 500MG | A209236 001 | Feb 01, 2018 |
|-----------|--|----------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 750MG | A209236 002 | Feb 01, 2018 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|------------|--------------------|--------------|
| AB | | | 1GM | A209236 003 | Feb 01, 2018 |
|-----------|--|--|------------|--------------------|--------------|

| | | | | | |
|-----------|--|------|--------------|--------------------|--------------|
| AB | | BARR | 500MG | A076378 001 | Apr 26, 2005 |
|-----------|--|------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 750MG | A076378 002 | Apr 26, 2005 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|------------|--------------------|--------------|
| AB | | | 1GM | A076250 001 | Apr 14, 2005 |
|-----------|--|--|------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------------------|--------------|--------------------|--------------|
| AB | | JUBILANT GENERICS | 500MG | A209156 001 | May 14, 2018 |
|-----------|--|-------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 750MG | A209156 002 | May 14, 2018 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|------------|--------------------|--------------|
| AB | | | 1GM | A209156 003 | May 14, 2018 |
|-----------|--|--|------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------|--------------|--------------------|--------------|
| AB | | LANNETT CO INC | 500MG | A203899 001 | Jun 16, 2017 |
|-----------|--|----------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|------------|--------------------|--------------|
| AB | | | 1GM | A203899 002 | Jun 16, 2017 |
|-----------|--|--|------------|--------------------|--------------|

| | | | | | |
|-----------|--|-----------|--------------|--------------------|--------------|
| AB | | LUPIN LTD | 500MG | A090860 001 | Mar 20, 2014 |
|-----------|--|-----------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 750MG | A090892 001 | Mar 20, 2014 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|------------|--------------------|--------------|
| AB | | | 1GM | A090446 001 | Mar 20, 2014 |
|-----------|--|--|------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------------------|--------------|--------------------|--------------|
| AB | | SUN PHARMA GLOBAL | 500MG | A200484 001 | Apr 23, 2014 |
|-----------|--|-------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 750MG | A201273 001 | Apr 23, 2014 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|------------|--------------------|--------------|
| AB | | | 1GM | A200484 002 | Apr 23, 2014 |
|-----------|--|--|------------|--------------------|--------------|

NIASPAN

| | | | | | |
|-----------|---|--------|--------------|--------------------|--------------|
| AB | + | ABEVIE | 500MG | N020381 002 | Jul 28, 1997 |
|-----------|---|--------|--------------|--------------------|--------------|

| | | | | | |
|-----------|---|---|--------------|--------------------|--------------|
| AB | + | ! | 750MG | N020381 003 | Jul 28, 1997 |
|-----------|---|---|--------------|--------------------|--------------|

| | | | | | |
|-----------|---|---|------------|--------------------|--------------|
| AB | + | ! | 1GM | N020381 004 | Jul 28, 1997 |
|-----------|---|---|------------|--------------------|--------------|

NICARDIPINE HYDROCHLORIDE

CAPSULE; ORAL

NICARDIPINE HYDROCHLORIDE

| | | | | | |
|-----------|--|----------------|-------------|--------------------|--------------|
| AB | | ANI PHARMS INC | 20MG | A074439 001 | Dec 10, 1996 |
|-----------|--|----------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 20MG | A074540 001 | Oct 28, 1996 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 30MG | A074439 002 | Dec 10, 1996 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 30MG | A074540 002 | Oct 28, 1996 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------------|-------------|--------------------|--------------|
| AB | | EPIC PHARMA | 20MG | A074928 001 | Mar 19, 1998 |
|-----------|--|-------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 30MG | A074928 002 | Mar 19, 1998 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------|-------------|--------------------|--------------|
| AB | | MYLAN | 20MG | A074642 001 | Jul 18, 1996 |
|-----------|--|-------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|---|-------------|--------------------|--------------|
| AB | | ! | 30MG | A074642 002 | Jul 18, 1996 |
|-----------|--|---|-------------|--------------------|--------------|

INJECTABLE; INJECTION

NICARDIPINE HYDROCHLORIDE

| | | | |
|--------------|----------------------|-------------|--------------|
| EXELA PHARMA | 25MG/10ML (2.5MG/ML) | N022276 001 | Jul 24, 2008 |
| SCIENCE | | | |

INJECTABLE; INTRAVENOUS

CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER

| | | | | | |
|---|---|----------------|-----------------------|-------------|--------------|
| + | ! | CHIESI USA INC | 40MG/200ML (0.2MG/ML) | N019734 004 | Nov 07, 2008 |
|---|---|----------------|-----------------------|-------------|--------------|

CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER

| | | | | | |
|---|---|----------------|-----------------------|-------------|--------------|
| + | ! | CHIESI USA INC | 20MG/200ML (0.1MG/ML) | N019734 003 | Jul 31, 2008 |
|---|---|----------------|-----------------------|-------------|--------------|

CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER

| | | | | | |
|---|---|----------------|-----------------------|-------------|--------------|
| + | ! | CHIESI USA INC | 20MG/200ML (0.1MG/ML) | N019734 002 | Jul 31, 2008 |
|---|---|----------------|-----------------------|-------------|--------------|

NICOTINE

INHALANT; ORAL

NICOTROL

| | | | | | |
|---|---|----------------------|---------------|-------------|--------------|
| + | ! | PHARMACIA AND UPJOHN | 4MG/CARTRIDGE | N020714 001 | May 02, 1997 |
|---|---|----------------------|---------------|-------------|--------------|

SPRAY, METERED; NASAL

NICOTROL

| | | | | | |
|---|---|------------|-------------|-------------|--------------|
| + | ! | PFIZER INC | 0.5MG/SPRAY | N020385 001 | Mar 22, 1996 |
|---|---|------------|-------------|-------------|--------------|

NIFEDIPINE

CAPSULE; ORAL

NIFEDIPINE

| | | | | | |
|-----------|--|-------------------|-------------|--------------------|--------------|
| AB | | ACTAVIS ELIZABETH | 10MG | A072579 001 | Jan 08, 1991 |
|-----------|--|-------------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 20MG | A072556 001 | Sep 20, 1990 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|-----------------|-------------|--------------------|--------------|
| AB | | HERITAGE PHARMA | 10MG | A202644 001 | Apr 25, 2013 |
|-----------|--|-----------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 20MG | A202644 002 | Apr 25, 2013 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------|-------------|--------------------|--------------|
| AB | | INTERGEL PHARM | 10MG | A072781 001 | Jul 30, 1993 |
|-----------|--|----------------|-------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

NIFEDIPINE

CAPSULE;ORAL

NIFEDIPINE

| | | | | |
|-----------|--------------------|-------------|--------------------|--------------|
| AB | LEADING PHARMA LLC | 10MG | A073250 001 | Oct 08, 1991 |
| AB | | 20MG | A074045 001 | Apr 30, 1992 |

PROCARDIA

| | | | | |
|-----------|----|--------|-------------|--------------------|
| AB | +! | PFIZER | 10MG | N018482 001 |
|-----------|----|--------|-------------|--------------------|

TABLET, EXTENDED RELEASE;ORAL

ADALAT CC

| | | | | | |
|------------|---|---------|-------------|--------------------|--------------|
| AB1 | + | ALVOGEN | 30MG | N020198 001 | Apr 21, 1993 |
| AB1 | + | | 60MG | N020198 002 | Apr 21, 1993 |
| AB1 | + | | 90MG | N020198 003 | Apr 21, 1993 |

NIFEDIPINE

| | | | | | |
|------------|--|-------------------------|-------------|--------------------|--------------|
| AB1 | | MYLAN | 30MG | A201071 001 | Dec 03, 2010 |
| AB1 | | | 60MG | A201071 002 | Dec 03, 2010 |
| AB1 | | | 90MG | A201071 003 | Dec 03, 2010 |
| AB1 | | NOVAST LABS | 30MG | A202987 001 | Aug 25, 2016 |
| AB1 | | | 60MG | A202987 002 | Aug 25, 2016 |
| AB1 | | | 90MG | A202987 003 | Aug 25, 2016 |
| AB1 | | PAR PHARM | 30MG | A077899 001 | Dec 13, 2006 |
| AB1 | | | 60MG | A077899 002 | Dec 13, 2006 |
| AB1 | | | 90MG | A077899 003 | May 25, 2012 |
| AB1 | | VALEANT PHARMS NORTH | 30MG | A075269 001 | Dec 04, 2000 |
| AB1 | | | 60MG | A075269 002 | Dec 04, 2000 |
| AB1 | | | 90MG | A076070 001 | Aug 16, 2002 |
| AB1 | | ZYDUS PHARMS USA INC | 30MG | A210184 001 | Jun 29, 2018 |
| AB1 | | | 60MG | A210184 002 | Jun 29, 2018 |
| AB1 | | | 90MG | A210184 003 | Jun 29, 2018 |
| AB2 | | MYLAN | 30MG | A090649 001 | Jun 21, 2010 |
| AB2 | | | 60MG | A090649 002 | Jun 21, 2010 |
| AB2 | | | 90MG | A090649 003 | Jun 21, 2010 |
| AB2 | | OSMOTICA PHARM US | 30MG | A077127 001 | Nov 21, 2005 |
| AB2 | | | 60MG | A077127 002 | Nov 21, 2005 |
| AB2 | | | 90MG | A077410 001 | Oct 03, 2007 |
| AB2 | | TWI PHARMS | 30MG | A203126 001 | Apr 03, 2014 |
| AB2 | | | 60MG | A203126 002 | Apr 03, 2014 |
| AB2 | | | 90MG | A203126 003 | Apr 03, 2014 |
| AB2 | | VALEANT PHARMS NORTH | 30MG | A075289 002 | Feb 06, 2001 |
| AB2 | | | 60MG | A075289 001 | Sep 27, 2000 |
| AB2 | | ZYDUS PHARMS USA INC | 30MG | A210012 001 | Dec 19, 2017 |
| AB2 | | | 60MG | A210012 002 | Dec 19, 2017 |
| AB2 | | | 90MG | A210012 003 | Dec 19, 2017 |

PROCARDIA XL

| | | | | | |
|------------|---|--------|-------------|--------------------|--------------|
| AB2 | + | PFIZER | 30MG | N019684 001 | Sep 06, 1989 |
| AB2 | + | | 60MG | N019684 002 | Sep 06, 1989 |
| AB2 | + | | 90MG | N019684 003 | Sep 06, 1989 |

NILOTINIB HYDROCHLORIDE

CAPSULE;ORAL

TASIGNA

| | | | | |
|---|----------|---------------|-------------|--------------|
| + | NOVARTIS | EQ 50MG BASE | N022068 003 | Mar 22, 2018 |
| + | | EQ 150MG BASE | N022068 002 | Jun 17, 2010 |
| + | | EQ 200MG BASE | N022068 001 | Oct 29, 2007 |

NILUTAMIDE

TABLET;ORAL

NILANDRON

| | | | | | |
|-----------|---|-------------------------|--------------|--------------------|--------------|
| AB | + | CONCORDIA PHARMS INC | 150MG | N020169 002 | Apr 30, 1999 |
|-----------|---|-------------------------|--------------|--------------------|--------------|

NILUTAMIDE

| | | | | | |
|-----------|--|----------------|--------------|--------------------|--------------|
| AB | | ANI PHARMS INC | 150MG | A207631 001 | Jul 15, 2016 |
|-----------|--|----------------|--------------|--------------------|--------------|

NIMODIPINE

CAPSULE;ORAL

NIMODIPINE

| | | | | | |
|-----------|---|-------------------------|-------------|--------------------|--------------|
| AB | ! | BIONPHARMA INC | 30MG | A076740 001 | Jan 17, 2008 |
| AB | | HERITAGE PHARMS INC | 30MG | A077811 001 | May 02, 2007 |
| AB | | SOFGEN PHARMS | 30MG | A201832 001 | Jul 24, 2015 |
| AB | | SUN PHARM INDS INC | 30MG | A077067 001 | Apr 17, 2007 |
| AB | | THEPHARMANETWORK LLC | 30MG | A090103 001 | Apr 07, 2014 |

PRESCRIPTION DRUG PRODUCT LIST

NIMODIPINE

SOLUTION;ORAL

NYMALIZE

+! ARBOR PHARMS LLC 60MG/20ML N203340 001 May 10, 2013

NINTEDANIB ESYLATE

CAPSULE;ORAL

OFEV

+ BOEHRINGER EQ 100MG BASE N205832 001 Oct 15, 2014

INGELHEIM

+! EQ 150MG BASE N205832 002 Oct 15, 2014

NIRAPARIB TOSYLATE

CAPSULE;ORAL

ZEJULA

+! TESARO INC EQ 100MG BASE N208447 001 Mar 27, 2017

NISOLDIPINE

TABLET, EXTENDED RELEASE;ORAL

NISOLDIPINEAB MYLAN 8.5MG A091001 001 Jan 26, 2011AB 17MG A091001 002 Jan 26, 2011AB 34MG A091001 004 Jan 26, 2011SULARAB +! COVIS PHARMA BV 8.5MG N020356 008 Jan 02, 2008AB +! 17MG N020356 007 Jan 02, 2008AB +! 34MG N020356 005 Jan 02, 2008

NISOLDIPINE

MYLAN

20MG A079051 001 Jul 25, 2008

25.5MG A091001 003 Jan 26, 2011

! 30MG A079051 002 Jul 25, 2008

! 40MG A079051 003 Jul 25, 2008

NITAZOXANIDE

FOR SUSPENSION;ORAL

ALINIA

+! ROMARK 100MG/5ML N021498 001 Nov 22, 2002

TABLET;ORAL

ALINIA

+! ROMARK 500MG N021497 001 Jul 21, 2004

NITISINONE

CAPSULE;ORAL

ORFADIN

+ SWEDISH ORPHAN 2MG N021232 001 Jan 18, 2002

+ 5MG N021232 002 Jan 18, 2002

+ 10MG N021232 003 Jan 18, 2002

+! 20MG N021232 004 Jun 13, 2016

SUSPENSION;ORAL

ORFADIN

+! SWEDISH ORPHAN 4MG/ML N206356 001 Apr 22, 2016

TABLET;ORAL

NITYR

+ CYCLE PHARMS LTD 2MG N209449 001 Jul 26, 2017

+ 5MG N209449 002 Jul 26, 2017

+! 10MG N209449 003 Jul 26, 2017

NITRIC OXIDE

GAS;INHALATION

INOMAXAA +! MALLINCKRODT HOSP 800PPM N020845 003 Dec 23, 1999NOXIVENTAA PRAXAIR 800PPM A207141 002 Oct 02, 2018

DISTRIBUTION

100PPM A207141 001 Oct 02, 2018

NITROFURANTOIN

SUSPENSION;ORAL

FURADANTINAB +! CASPER PHARMA LLC 25MG/5ML N009175 001NITROFURANTOINAB ACTAVIS MID 25MG/5ML A205180 001 May 03, 2016

ATLANTIC

AB AMNEAL PHARMS 25MG/5ML A201679 001 May 11, 2011AB NOSTRUM LABS INC 25MG/5ML A201355 001 Aug 14, 2013AB NOVEL LABS INC 25MG/5ML A201693 001 Sep 08, 2014

PRESCRIPTION DRUG PRODUCT LIST

NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACRODANTIN

| | | | | | | |
|-----------|---|---------------|--------------|----------------|------------|--|
| <u>AB</u> | + | ALVOGEN MALTA | <u>25MG</u> | <u>N016620</u> | <u>003</u> | |
| <u>AB</u> | + | | <u>50MG</u> | <u>N016620</u> | <u>001</u> | |
| <u>AB</u> | + | ! | <u>100MG</u> | <u>N016620</u> | <u>002</u> | |

NITROFURANTOIN

| | | | | | | |
|-----------|--|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | ACTAVIS LABS FL INC | <u>25MG</u> | <u>A091095</u> | <u>001</u> | Jun 18, 2015 |
| <u>AB</u> | | | <u>50MG</u> | <u>A091095</u> | <u>002</u> | Jun 18, 2015 |
| <u>AB</u> | | | <u>100MG</u> | <u>A091095</u> | <u>003</u> | Jun 18, 2015 |
| <u>AB</u> | | IMPAX LABS INC | <u>50MG</u> | <u>A073671</u> | <u>001</u> | Jan 28, 1993 |
| <u>AB</u> | | | <u>100MG</u> | <u>A073652</u> | <u>001</u> | Jan 28, 1993 |
| <u>AB</u> | | MYLAN | <u>50MG</u> | <u>A074967</u> | <u>001</u> | Jul 09, 1997 |
| <u>AB</u> | | | <u>100MG</u> | <u>A077025</u> | <u>001</u> | Aug 18, 2004 |
| <u>AB</u> | | NOVEL LABS INC | <u>50MG</u> | <u>A203233</u> | <u>001</u> | Jul 09, 2018 |
| <u>AB</u> | | | <u>100MG</u> | <u>A203233</u> | <u>002</u> | Jul 09, 2018 |
| <u>AB</u> | | SUN PHARM INDUSTRIES | <u>25MG</u> | <u>A201722</u> | <u>001</u> | Feb 16, 2016 |
| <u>AB</u> | | | <u>50MG</u> | <u>A201722</u> | <u>002</u> | Feb 16, 2016 |
| <u>AB</u> | | | <u>100MG</u> | <u>A201722</u> | <u>003</u> | Feb 16, 2016 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>50MG</u> | <u>A205005</u> | <u>001</u> | Dec 12, 2017 |
| <u>AB</u> | | | <u>100MG</u> | <u>A205005</u> | <u>002</u> | Dec 12, 2017 |

NITROFURANTOIN; NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACROBID

| | | | | | | | |
|-----------|---|---|---------------|-------------------|----------------|------------|--------------|
| <u>AB</u> | + | ! | ALVOGEN MALTA | <u>75MG; 25MG</u> | <u>N020064</u> | <u>001</u> | Dec 24, 1991 |
|-----------|---|---|---------------|-------------------|----------------|------------|--------------|

NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS)

| | | | | | | |
|-----------|--|---------------------|-------------------|----------------|------------|--------------|
| <u>AB</u> | | AMNEAL PHARMS | <u>75MG; 25MG</u> | <u>A207372</u> | <u>001</u> | May 15, 2017 |
| <u>AB</u> | | MYLAN | <u>75MG; 25MG</u> | <u>A076648</u> | <u>001</u> | Mar 22, 2004 |
| <u>AB</u> | | SANDOZ | <u>75MG; 25MG</u> | <u>A077066</u> | <u>001</u> | Apr 05, 2005 |
| <u>AB</u> | | SUNNY PHARMTECH INC | <u>75MG; 25MG</u> | <u>A208516</u> | <u>001</u> | May 24, 2018 |
| <u>AB</u> | | WATSON LABS INC | <u>75MG; 25MG</u> | <u>A202250</u> | <u>001</u> | Jul 08, 2015 |

NITROGLYCERIN

AEROSOL, METERED; SUBLINGUAL

NITROMIST

| | | | | | | | |
|--|---|---|-----------------|-------------|---------|-----|--------------|
| | + | ! | MIST PHARMS LLC | 0.4MG/SPRAY | N021780 | 001 | Nov 02, 2006 |
|--|---|---|-----------------|-------------|---------|-----|--------------|

FILM, EXTENDED RELEASE; TRANSDERMAL

MINITRAN

| | | | | | | |
|------------|--|-------------------------|-----------------|----------------|------------|--------------|
| <u>AB1</u> | | MEDICIS | <u>0.4MG/HR</u> | <u>A089773</u> | <u>001</u> | Aug 30, 1996 |
| <u>AB1</u> | | VALEANT PHARMS | <u>0.1MG/HR</u> | <u>A089771</u> | <u>001</u> | Aug 30, 1996 |
| <u>AB1</u> | | | <u>0.6MG/HR</u> | <u>A089774</u> | <u>001</u> | Aug 30, 1996 |
| <u>AB1</u> | | VALEANT PHARMS NORTH | <u>0.2MG/HR</u> | <u>A089772</u> | <u>001</u> | Aug 30, 1996 |

NITRO-DUR

| | | | | | | | |
|------------|---|---|----------|-----------------|----------------|------------|--------------|
| <u>AB1</u> | + | ! | USPHARMA | <u>0.1MG/HR</u> | <u>N020145</u> | <u>001</u> | Apr 04, 1995 |
| <u>AB1</u> | + | ! | | <u>0.2MG/HR</u> | <u>N020145</u> | <u>002</u> | Apr 04, 1995 |
| <u>AB1</u> | + | ! | | <u>0.4MG/HR</u> | <u>N020145</u> | <u>004</u> | Apr 04, 1995 |
| <u>AB1</u> | + | ! | | <u>0.6MG/HR</u> | <u>N020145</u> | <u>005</u> | Apr 04, 1995 |

NITROGLYCERIN

| | | | | | | |
|------------|---|--------------------|-----------------|----------------|------------|--------------|
| <u>AB2</u> | | HERCON PHARM | <u>0.2MG/HR</u> | <u>A089884</u> | <u>001</u> | Oct 30, 1998 |
| <u>AB2</u> | | | <u>0.4MG/HR</u> | <u>A089885</u> | <u>001</u> | Oct 30, 1998 |
| <u>AB2</u> | | | <u>0.6MG/HR</u> | <u>A089886</u> | <u>001</u> | Oct 30, 1998 |
| <u>AB2</u> | ! | MYLAN TECHNOLOGIES | <u>0.2MG/HR</u> | <u>A074559</u> | <u>003</u> | Aug 30, 1996 |
| <u>AB2</u> | ! | | <u>0.4MG/HR</u> | <u>A074559</u> | <u>002</u> | Aug 30, 1996 |
| <u>AB2</u> | ! | | <u>0.6MG/HR</u> | <u>A074559</u> | <u>001</u> | Aug 30, 1996 |

NITRO-DUR

| | | | | | | | |
|--|---|---|----------|----------|---------|-----|--------------|
| | + | ! | USPHARMA | 0.3MG/HR | N020145 | 003 | Apr 04, 1995 |
| | + | ! | | 0.8MG/HR | N020145 | 006 | Apr 04, 1995 |

NITROGLYCERIN

| | | | | | | |
|--|---|--------------------|----------|---------|-----|--------------|
| | ! | MYLAN TECHNOLOGIES | 0.1MG/HR | A074559 | 004 | Feb 06, 1998 |
|--|---|--------------------|----------|---------|-----|--------------|

INJECTABLE; INJECTION

NITROGLYCERIN IN DEXTROSE 5%

| | | | | | | | |
|-----------|---|---|-----------------|-------------------|----------------|------------|--------------|
| <u>AP</u> | + | ! | BAXTER HLTHCARE | <u>10MG/100ML</u> | <u>N019970</u> | <u>001</u> | Dec 29, 1989 |
| <u>AP</u> | + | ! | | <u>20MG/100ML</u> | <u>N019970</u> | <u>002</u> | Dec 29, 1989 |
| <u>AP</u> | + | ! | | <u>40MG/100ML</u> | <u>N019970</u> | <u>003</u> | Dec 29, 1989 |

NITROGLYCERIN

| | | | | | | |
|--|---|----------|--------|---------|-----|--------------|
| | ! | LUITPOLD | 5MG/ML | A072034 | 001 | May 24, 1988 |
|--|---|----------|--------|---------|-----|--------------|

OINTMENT; INTRA-ANAL

RECTIV

| | | | | | | | |
|--|---|---|--------------------|------|---------|-----|--------------|
| | + | ! | ALLERGAN SALES LLC | 0.4% | N021359 | 001 | Jun 21, 2011 |
|--|---|---|--------------------|------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

NITROGLYCERIN

OINTMENT; TRANSDERMAL

NITROGLYCERIN

! FOUGERA PHARMS INC 2%

A087355 001 Jul 08, 1988

POWDER; SUBLINGUAL

GONITRO

+! POHL BOSKAMP 0.4MG/PACKET

N208424 001 Jun 08, 2016

SPRAY, METERED; SUBLINGUAL

NITROGLYCERIN**AB** PERRIGO ISRAEL 0.4MG/SPRAY **A091496 001** Sep 20, 2013NITROLINGUAL PUMPSPRAY**AB** +! POHL BOSKAMP 0.4MG/SPRAY **N018705 002** Jan 10, 1997

TABLET; SUBLINGUAL

NITROGLYCERIN**AB** ACTAVIS LABS FL INC 0.3MG **A203693 001** Oct 16, 2017**AB** 0.4MG **A203693 002** Oct 16, 2017**AB** 0.6MG **A203693 003** Oct 16, 2017**AB** DR REDDYS LABS INC 0.3MG **A208191 001** Aug 26, 2016**AB** 0.4MG **A208191 002** Aug 26, 2016**AB** 0.6MG **A208191 003** Aug 26, 2016**AB** GLENMARK PHARMS SA 0.3MG **A206391 001** Sep 19, 2017**AB** 0.4MG **A206391 002** Sep 19, 2017**AB** 0.6MG **A206391 003** Sep 19, 2017**AB** SIGMAPHARM LABS LLC 0.3MG **A207745 001** May 07, 2018**AB** 0.4MG **A207745 002** May 07, 2018**AB** 0.6MG **A207745 003** May 07, 2018NITROSTAT**AB** + PFIZER PHARMS 0.3MG **N021134 001** May 01, 2000**AB** + 0.4MG **N021134 002** May 01, 2000**AB** +! 0.6MG **N021134 003** May 01, 2000NIZATIDINE

CAPSULE; ORAL

NIZATIDINE**AB** ANI PHARMS INC 150MG **A075668 001** Sep 12, 2002**AB** 300MG **A075668 002** Sep 12, 2002**AB** DR REDDYS LABS LTD 150MG **A077314 001** Sep 15, 2005**AB** 300MG **A077314 002** Sep 15, 2005**AB** GLENMARK GENERICS 150MG **A090618 001** Jul 15, 2011**AB** 300MG **A090618 002** Jul 15, 2011**AB** MYLAN PHARMS INC 150MG **A075806 001** Jul 05, 2002**AB** ! 300MG **A075806 002** Jul 05, 2002**AB** SANDOZ 150MG **A076178 001** Jul 05, 2002**AB** 300MG **A076178 002** Jul 05, 2002**AB** WATSON LABS 150MG **A075616 001** Jul 09, 2002**AB** 300MG **A075616 002** Jul 09, 2002

SOLUTION; ORAL

NIZATIDINE

! AMNEAL PHARMS 15MG/ML

A090576 001 Nov 18, 2009

NOREPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

LEVOPHED**AP** +! HOSPIRA EQ 1MG BASE/ML **N007513 001**NOREPINEPHRINE BITARTRATE**AP** AMNEAL PHARMS CO EQ 1MG BASE/ML **A210839 001** Dec 17, 2018**AP** BAXTER HLTHCARE EQ 1MG BASE/ML **A040859 001** Mar 27, 2012

CORP

AP HIKMA FARMACEUTICA EQ 1MG BASE/ML **A203662 001** Nov 07, 2018**AP** MYLAN LABS LTD EQ 1MG BASE/ML **A211242 001** Oct 04, 2018**AP** SANDOZ INC EQ 1MG BASE/ML **A211359 001** Oct 18, 2018**AP** TEVA PHARMS USA EQ 1MG BASE/ML **A040455 001** Mar 03, 2003**AP** WEST-WARD PHARMS EQ 1MG BASE/ML **A040462 001** Oct 31, 2003

INT

NORETHINDRONE

TABLET; ORAL-28

CAMILA**AB1** MAYNE PHARMA 0.35MG **A076177 001** Oct 21, 2002HEATHER**AB1** GLENMARK GENERICS 0.35MG **A090454 001** Apr 23, 2010INCASSIA**AB1** AUROBINDO PHARMA 0.35MG **A207304 001** Sep 23, 2016

LTD

PRESCRIPTION DRUG PRODUCT LIST

NORETHINDRONE

TABLET; ORAL-28

NOR-QD

| | | | | | | |
|------------|------------|------|---------------|----------------|------------|--|
| AB1 | + ! | APIL | 0.35MG | N017060 | 001 | |
|------------|------------|------|---------------|----------------|------------|--|

NORETHINDRONE

| | | | | | | |
|------------|--|-----------------|---------------|----------------|------------|--------------|
| AB1 | | ACCORD HLTHCARE | 0.35MG | A206807 | 001 | Dec 13, 2016 |
| AB1 | | AMNEAL PHARMS | 0.35MG | A202260 | 001 | Aug 01, 2013 |
| AB1 | | LUPIN LTD | 0.35MG | A091325 | 001 | Sep 19, 2011 |
| AB1 | | MYLAN LABS LTD | 0.35MG | A201483 | 001 | Jun 24, 2013 |
| AB1 | | NOVAST LABS | 0.35MG | A202014 | 001 | Sep 13, 2013 |

ERRIN

| | | | | | | |
|------------|--|--------------|---------------|----------------|------------|--------------|
| AB2 | | MAYNE PHARMA | 0.35MG | A076225 | 001 | Oct 21, 2002 |
|------------|--|--------------|---------------|----------------|------------|--------------|

JENCYCLA

| | | | | | | |
|------------|--|-----------|---------------|----------------|------------|--------------|
| AB2 | | LUPIN LTD | 0.35MG | A091323 | 001 | Mar 28, 2013 |
|------------|--|-----------|---------------|----------------|------------|--------------|

MICRONOR

| | | | | | | |
|------------|------------|----------------|---------------|----------------|------------|--|
| AB2 | + ! | JANSSEN PHARMS | 0.35MG | N016954 | 001 | |
|------------|------------|----------------|---------------|----------------|------------|--|

NORETHINDRONE

| | | | | | | |
|------------|--|-------------------|---------------|----------------|------------|--------------|
| AB2 | | GLENMARK GENERICS | 0.35MG | A091209 | 001 | Jul 22, 2010 |
| AB2 | | MYLAN LABS LTD | 0.35MG | A200980 | 001 | Jun 12, 2013 |
| AB2 | | NOVAST LABS | 0.35MG | A200961 | 001 | Sep 13, 2013 |

NORETHINDRONE ACETATE

TABLET; ORAL

NORETHINDRONE ACETATE

| | | | | | | |
|-----------|----------|----------------------|------------|----------------|------------|--------------|
| AB | | AMNEAL PHARMS | 5MG | A200275 | 001 | Jul 30, 2012 |
| AB | | AUROBINDO PHARMA LTD | 5MG | A204236 | 001 | Jan 08, 2016 |
| AB | ! | BARR | 5MG | A075951 | 001 | May 25, 2001 |
| AB | | GLENMARK GENERICS | 5MG | A091090 | 001 | Jul 21, 2010 |
| AB | | MYLAN LABS LTD | 5MG | A205278 | 001 | Nov 10, 2016 |
| AB | | PACK PHARMS LLC | 5MG | A206490 | 001 | Nov 05, 2018 |

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL

NORTRIPTYLINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|--------------|---------------------|----------------|------------|--------------|
| AB | | MAYNE PHARMA | EQ 10MG BASE | A073556 | 002 | Mar 30, 1992 |
| AB | | | EQ 25MG BASE | A073556 | 003 | Mar 30, 1992 |
| AB | | | EQ 50MG BASE | A073556 | 004 | Mar 30, 1992 |
| AB | | | EQ 75MG BASE | A073556 | 001 | Mar 30, 1992 |
| AB | | TARO PHARM | EQ 10MG BASE | A075520 | 004 | May 08, 2000 |
| AB | | | EQ 25MG BASE | A075520 | 003 | May 08, 2000 |
| AB | | | EQ 50MG BASE | A075520 | 001 | May 08, 2000 |
| AB | | | EQ 75MG BASE | A075520 | 002 | May 08, 2000 |
| AB | | TEVA | EQ 10MG BASE | A074132 | 001 | Mar 27, 1995 |
| AB | | | EQ 25MG BASE | A074132 | 002 | Mar 27, 1995 |
| AB | | | EQ 50MG BASE | A074132 | 003 | Mar 27, 1995 |
| AB | | | EQ 75MG BASE | A074132 | 004 | Mar 27, 1995 |

PAMELOR

| | | | | | | |
|-----------|------------|------------|---------------------|----------------|------------|--|
| AB | + | SPECGX LLC | EQ 10MG BASE | N018013 | 001 | |
| AB | + | | EQ 25MG BASE | N018013 | 002 | |
| AB | + | | EQ 50MG BASE | N018013 | 004 | |
| AB | + ! | | EQ 75MG BASE | N018013 | 003 | |

SOLUTION; ORAL

NORTRIPTYLINE HYDROCHLORIDE

| | | | | | | |
|-----------|----------|-------------|-------------------------|----------------|------------|--------------|
| AA | ! | PHARM ASSOC | EQ 10MG BASE/5ML | A075606 | 001 | Aug 28, 2000 |
| AA | | TARO | EQ 10MG BASE/5ML | A077965 | 001 | Jun 20, 2006 |

NUSINERSEN SODIUM

SOLUTION; INTRATHECAL

SPINRAZA

| | | | | | |
|------------|-------------|---------------------|----------------|------------|--------------|
| + ! | BIOGEN IDEC | 12MG/5ML (2.4MG/ML) | N209531 | 001 | Dec 23, 2016 |
|------------|-------------|---------------------|----------------|------------|--------------|

NYSTATIN

CREAM; TOPICAL

NYSTATIN

| | | | | | | |
|-----------|----------|----------------------|-------------------------|----------------|------------|--------------|
| AT | | ACTAVIS MID ATLANTIC | 100,000 UNITS/GM | A062949 | 001 | Jun 13, 1988 |
| AT | | CROWN LABS INC | 100,000 UNITS/GM | A207733 | 001 | Sep 26, 2017 |
| AT | | FOUGERA PHARMS | 100,000 UNITS/GM | A062129 | 001 | |
| AT | | G AND W LABS INC | 100,000 UNITS/GM | A061966 | 001 | |
| AT | | PERRIGO NEW YORK | 100,000 UNITS/GM | A062225 | 001 | |
| AT | ! | TARO | 100,000 UNITS/GM | A064022 | 001 | Jan 29, 1993 |
| AT | | VINTAGE | 100,000 UNITS/GM | A065315 | 001 | May 31, 2006 |

PRESCRIPTION DRUG PRODUCT LIST

NYSTATIN

OINTMENT; TOPICAL

NYSTATIN

| | | | | |
|-----------|----------------------|-------------------------|--------------------|--------------|
| <u>AT</u> | ACTAVIS MID ATLANTIC | <u>100,000 UNITS/GM</u> | <u>A062840 001</u> | Nov 13, 1987 |
| <u>AT</u> | ! FOUGERA PHARMS | <u>100,000 UNITS/GM</u> | <u>A062124 002</u> | Sep 23, 1982 |
| <u>AT</u> | G AND W LABS INC | <u>100,000 UNITS/GM</u> | <u>A209114 001</u> | Oct 06, 2017 |
| <u>AT</u> | LYNE | <u>100,000 UNITS/GM</u> | <u>A209082 001</u> | May 21, 2018 |
| <u>AT</u> | PERRIGO NEW YORK | <u>100,000 UNITS/GM</u> | <u>A062472 001</u> | Feb 13, 1984 |
| <u>AT</u> | ZYDUS PHARMS USA INC | <u>100,000 UNITS/GM</u> | <u>A207767 001</u> | May 25, 2018 |

POWDER; TOPICAL

NYSTATIN

| | | | | |
|-----------|--------------------|-------------------------|--------------------|--------------|
| <u>AT</u> | EPIC PHARMA LLC | <u>100,000 UNITS/GM</u> | <u>A210532 001</u> | Apr 30, 2018 |
| <u>AT</u> | GAVIS PHARMS | <u>100,000 UNITS/GM</u> | <u>A065138 001</u> | Jul 23, 2004 |
| <u>AT</u> | LYNE | <u>100,000 UNITS/GM</u> | <u>A208838 001</u> | May 30, 2017 |
| <u>AT</u> | ! MAYNE PHARMA INC | <u>100,000 UNITS/GM</u> | <u>A065203 001</u> | Jul 15, 2004 |
| <u>AT</u> | NESHER PHARMS | <u>100,000 UNITS/GM</u> | <u>A208581 001</u> | Jun 08, 2017 |
| <u>AT</u> | UPSHER SMITH LABS | <u>100,000 UNITS/GM</u> | <u>A065183 001</u> | May 03, 2005 |
| <u>AT</u> | X GEN PHARMS | <u>100,000 UNITS/GM</u> | <u>A065175 001</u> | Dec 17, 2004 |

NYSTOP

| | | | | |
|-----------|-------------|-------------------------|--------------------|--------------|
| <u>AT</u> | PADDOCK LLC | <u>100,000 UNITS/GM</u> | <u>A064118 001</u> | Aug 16, 1996 |
|-----------|-------------|-------------------------|--------------------|--------------|

SUSPENSION; ORAL

NYSTATIN

| | | | | |
|-----------|---------------------|-------------------------|--------------------|--------------|
| <u>AA</u> | FOUGERA PHARMS INC | <u>100,000 UNITS/ML</u> | <u>A062517 001</u> | Jun 07, 1984 |
| <u>AA</u> | HI TECH PHARMA | <u>100,000 UNITS/ML</u> | <u>A064042 001</u> | Feb 28, 1994 |
| <u>AA</u> | LANNETT CO INC | <u>100,000 UNITS/ML</u> | <u>A065148 001</u> | Jun 28, 2005 |
| <u>AA</u> | PHARM ASSOC | <u>100,000 UNITS/ML</u> | <u>A203621 001</u> | Jan 07, 2016 |
| <u>AA</u> | TARO PHARM | <u>100,000 UNITS/ML</u> | <u>A062876 001</u> | Feb 29, 1988 |
| <u>AA</u> | VISTAPHARM | <u>100,000 UNITS/ML</u> | <u>A064142 001</u> | Jun 25, 1998 |
| <u>AA</u> | ! WOCHKHARDT BIO AG | <u>100,000 UNITS/ML</u> | <u>A065422 001</u> | Mar 07, 2011 |
| <u>AA</u> | | | <u>A062512 001</u> | Oct 29, 1984 |

TABLET; ORAL

NYSTATIN

| | | | | |
|-----------|----------------------|----------------------|--------------------|--------------|
| <u>AA</u> | HERITAGE PHARMS INC | <u>500,000 UNITS</u> | <u>A062474 001</u> | Dec 22, 1983 |
| <u>AA</u> | SUN PHARM INDUSTRIES | <u>500,000 UNITS</u> | <u>A062838 001</u> | Dec 22, 1988 |
| <u>AA</u> | ! TEVA | <u>500,000 UNITS</u> | <u>A062506 001</u> | Jan 16, 1984 |

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

MYKACET

| | | | | |
|-----------|------------------|-------------------------------|--------------------|--------------|
| <u>AT</u> | G AND W LABS INC | <u>100,000 UNITS/GM; 0.1%</u> | <u>A062367 001</u> | May 28, 1985 |
|-----------|------------------|-------------------------------|--------------------|--------------|

NYSTATIN AND TRIAMCINOLONE ACETONIDE

| | | | | |
|-----------|---------------------|-------------------------------|--------------------|--------------|
| <u>AT</u> | AMNEAL PHARMS LLC | <u>100,000 UNITS/GM; 0.1%</u> | <u>A209990 001</u> | Feb 15, 2018 |
| <u>AT</u> | CROWN LABS INC | <u>100,000 UNITS/GM; 0.1%</u> | <u>A207730 001</u> | Dec 26, 2017 |
| <u>AT</u> | DR REDDYS LABS LTD | <u>100,000 UNITS/GM; 0.1%</u> | <u>A208326 001</u> | Oct 26, 2016 |
| <u>AT</u> | FOUGERA PHARMS INC | <u>100,000 UNITS/GM; 0.1%</u> | <u>A062599 001</u> | Oct 08, 1985 |
| <u>AT</u> | GLENMARK PHARMS LTD | <u>100,000 UNITS/GM; 0.1%</u> | <u>A208136 001</u> | Oct 24, 2016 |
| <u>AT</u> | LUPIN LTD | <u>100,000 UNITS/GM; 0.1%</u> | <u>A208205 001</u> | May 31, 2018 |
| <u>AT</u> | PERRIGO UK FINCO | <u>100,000 UNITS/GM; 0.1%</u> | <u>A208479 001</u> | Aug 14, 2017 |
| <u>AT</u> | ! TARO | <u>100,000 UNITS/GM; 0.1%</u> | <u>A062364 001</u> | Dec 22, 1987 |

OINTMENT; TOPICAL

MYKACET

| | | | | |
|-----------|------------------|-------------------------------|--------------------|--------------|
| <u>AT</u> | G AND W LABS INC | <u>100,000 UNITS/GM; 0.1%</u> | <u>A062733 001</u> | Mar 06, 1987 |
|-----------|------------------|-------------------------------|--------------------|--------------|

NYSTATIN AND TRIAMCINOLONE ACETONIDE

| | | | | |
|-----------|----------------------|-------------------------------|--------------------|--------------|
| <u>AT</u> | AKORN | <u>100,000 UNITS/GM; 0.1%</u> | <u>A207217 001</u> | Aug 04, 2017 |
| <u>AT</u> | CROWN LABS INC | <u>100,000 UNITS/GM; 0.1%</u> | <u>A207731 001</u> | Dec 26, 2017 |
| <u>AT</u> | DR REDDYS LABS LTD | <u>100,000 UNITS/GM; 0.1%</u> | <u>A207741 001</u> | Jan 31, 2017 |
| <u>AT</u> | FOUGERA PHARMS INC | <u>100,000 UNITS/GM; 0.1%</u> | <u>A062602 001</u> | Oct 09, 1985 |
| <u>AT</u> | GLENMARK PHARMS LTD | <u>100,000 UNITS/GM; 0.1%</u> | <u>A208300 001</u> | Jun 23, 2016 |
| <u>AT</u> | PERRIGO UK FINCO | <u>100,000 UNITS/GM; 0.1%</u> | <u>A207380 001</u> | Dec 20, 2016 |
| <u>AT</u> | RISING PHARMS | <u>100,000 UNITS/GM; 0.1%</u> | <u>A206785 001</u> | Dec 29, 2016 |
| <u>AT</u> | STRIDES PHARMA | <u>100,000 UNITS/GM; 0.1%</u> | <u>A210077 001</u> | Jan 29, 2018 |
| <u>AT</u> | ! TARO | <u>100,000 UNITS/GM; 0.1%</u> | <u>A063305 001</u> | Mar 29, 1993 |
| <u>AT</u> | TELEGENT PHARMA INC | <u>100,000 UNITS/GM; 0.1%</u> | <u>A208287 001</u> | Dec 30, 2016 |
| <u>AT</u> | ZYDUS PHARMS USA INC | <u>100,000 UNITS/GM; 0.1%</u> | <u>A207764 001</u> | Nov 08, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

OBETICHOLIC ACID

TABLET; ORAL

OCALIVA

| | | | | | |
|---|-------------------------|------|---------|-----|--------------|
| + | INTERCEPT PHARMS INC | 5MG | N207999 | 001 | May 27, 2016 |
| + | ! | 10MG | N207999 | 002 | May 27, 2016 |

OCTREOTIDE ACETATE

INJECTABLE; INJECTION

OCTREOTIDE ACETATE

| | | | | | |
|-----------|---------------------------|--------------------------|----------------|------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 0.2MG BASE/ML</u> | <u>A077450</u> | <u>001</u> | Feb 10, 2006 |
| <u>AP</u> | | <u>EQ 1MG BASE/ML</u> | <u>A077450</u> | <u>002</u> | Feb 10, 2006 |
| <u>AP</u> | SAGENT PHARMS | <u>EQ 0.2MG BASE/ML</u> | <u>A091041</u> | <u>001</u> | Nov 12, 2013 |
| <u>AP</u> | | <u>EQ 1MG BASE/ML</u> | <u>A091041</u> | <u>002</u> | Nov 12, 2013 |
| <u>AP</u> | SUN PHARM INDS | <u>EQ 0.05MG BASE/ML</u> | <u>A077372</u> | <u>001</u> | Aug 14, 2007 |
| <u>AP</u> | | <u>EQ 0.1MG BASE/ML</u> | <u>A077372</u> | <u>002</u> | Aug 14, 2007 |
| <u>AP</u> | | <u>EQ 0.2MG BASE/ML</u> | <u>A077373</u> | <u>001</u> | Aug 14, 2007 |
| <u>AP</u> | | <u>EQ 0.5MG BASE/ML</u> | <u>A077372</u> | <u>003</u> | Aug 14, 2007 |
| <u>AP</u> | | <u>EQ 1MG BASE/ML</u> | <u>A077373</u> | <u>002</u> | Aug 14, 2007 |
| <u>AP</u> | TEVA PHARMS USA | <u>EQ 0.05MG BASE/ML</u> | <u>A075957</u> | <u>001</u> | Oct 03, 2005 |
| <u>AP</u> | | <u>EQ 0.1MG BASE/ML</u> | <u>A075957</u> | <u>002</u> | Oct 03, 2005 |
| <u>AP</u> | | <u>EQ 0.2MG BASE/ML</u> | <u>A075959</u> | <u>001</u> | Nov 21, 2005 |
| <u>AP</u> | | <u>EQ 0.5MG BASE/ML</u> | <u>A075957</u> | <u>003</u> | Oct 03, 2005 |
| <u>AP</u> | | <u>EQ 1MG BASE/ML</u> | <u>A075959</u> | <u>002</u> | Nov 21, 2005 |
| <u>AP</u> | USV NORTH AMERICA | <u>EQ 0.05MG BASE/ML</u> | <u>A204669</u> | <u>001</u> | Dec 27, 2018 |
| <u>AP</u> | | <u>EQ 0.1MG BASE/ML</u> | <u>A204669</u> | <u>002</u> | Dec 27, 2018 |
| <u>AP</u> | | <u>EQ 0.2MG BASE/ML</u> | <u>A203765</u> | <u>001</u> | Sep 07, 2018 |
| <u>AP</u> | | <u>EQ 0.5MG BASE/ML</u> | <u>A204669</u> | <u>003</u> | Dec 27, 2018 |
| <u>AP</u> | | <u>EQ 1MG BASE/ML</u> | <u>A203765</u> | <u>002</u> | Sep 07, 2018 |
| <u>AP</u> | ! WEST-WARD PHARMS INT | <u>EQ 0.2MG BASE/ML</u> | <u>A076330</u> | <u>001</u> | Apr 08, 2005 |
| <u>AP</u> | ! | <u>EQ 1MG BASE/ML</u> | <u>A076330</u> | <u>002</u> | Apr 08, 2005 |

OCTREOTIDE ACETATE (PRESERVATIVE FREE)

| | | | | | |
|-----------|---------------------------|--------------------------|----------------|------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 0.05MG BASE/ML</u> | <u>A077457</u> | <u>001</u> | Feb 10, 2006 |
| <u>AP</u> | | <u>EQ 0.1MG BASE/ML</u> | <u>A077457</u> | <u>002</u> | Feb 10, 2006 |
| <u>AP</u> | | <u>EQ 0.7MG BASE/ML</u> | <u>A077457</u> | <u>003</u> | Feb 10, 2006 |
| <u>AP</u> | MYLAN INSTITUTIONAL | <u>EQ 0.05MG BASE/ML</u> | <u>A079198</u> | <u>001</u> | Feb 10, 2011 |
| <u>AP</u> | | <u>EQ 0.1MG BASE/ML</u> | <u>A079198</u> | <u>002</u> | Feb 10, 2011 |
| <u>AP</u> | | <u>EQ 0.5MG BASE/ML</u> | <u>A079198</u> | <u>003</u> | Feb 10, 2011 |
| <u>AP</u> | SAGENT PHARMS | <u>EQ 0.05MG BASE/ML</u> | <u>A090834</u> | <u>001</u> | Nov 12, 2013 |
| <u>AP</u> | | <u>EQ 0.1MG BASE/ML</u> | <u>A090834</u> | <u>002</u> | Nov 12, 2013 |
| <u>AP</u> | | <u>EQ 0.5MG BASE/ML</u> | <u>A090834</u> | <u>003</u> | Nov 12, 2013 |
| <u>AP</u> | ! WEST-WARD PHARMS INT | <u>EQ 0.05MG BASE/ML</u> | <u>A076313</u> | <u>001</u> | Mar 28, 2005 |
| <u>AP</u> | ! | <u>EQ 0.1MG BASE/ML</u> | <u>A076313</u> | <u>003</u> | Mar 28, 2005 |
| <u>AP</u> | ! | <u>EQ 0.5MG BASE/ML</u> | <u>A076313</u> | <u>002</u> | Mar 28, 2005 |

SANDOSTATIN

| | | | | | | |
|-----------|---|-----------------|--------------------------|----------------|------------|--------------|
| <u>AP</u> | + | NOVARTIS | <u>EQ 0.05MG BASE/ML</u> | <u>N019667</u> | <u>001</u> | Oct 21, 1988 |
| <u>AP</u> | + | | <u>EQ 0.1MG BASE/ML</u> | <u>N019667</u> | <u>002</u> | Oct 21, 1988 |
| <u>AP</u> | + | | <u>EQ 0.2MG BASE/ML</u> | <u>N019667</u> | <u>004</u> | Jun 12, 1991 |
| <u>AP</u> | + | | <u>EQ 0.5MG BASE/ML</u> | <u>N019667</u> | <u>003</u> | Oct 21, 1988 |
| <u>AP</u> | + | | <u>EQ 1MG BASE/ML</u> | <u>N019667</u> | <u>005</u> | Jun 12, 1991 |
| | | SANDOSTATIN LAR | | | | |
| | + | NOVARTIS | EQ 10MG BASE/VIAL | N021008 | 001 | Nov 25, 1998 |
| | + | | EQ 20MG BASE/VIAL | N021008 | 002 | Nov 25, 1998 |
| | + | | EQ 30MG BASE/VIAL | N021008 | 003 | Nov 25, 1998 |

OFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

OCUFLOX

| | | | | | | |
|-----------|---|----------|-------------|----------------|------------|--------------|
| <u>AT</u> | + | ALLERGAN | <u>0.3%</u> | <u>N019921</u> | <u>001</u> | Jul 30, 1993 |
|-----------|---|----------|-------------|----------------|------------|--------------|

OFLOXACIN

| | | | | | | |
|-----------|--|--------------------|-------------|----------------|------------|--------------|
| <u>AT</u> | | AKORN | <u>0.3%</u> | <u>A076407</u> | <u>001</u> | Apr 15, 2008 |
| <u>AT</u> | | ALTAIRE PHARMS INC | <u>0.3%</u> | <u>A202692</u> | <u>001</u> | Apr 29, 2013 |
| <u>AT</u> | | ALVOGEN | <u>0.3%</u> | <u>A076830</u> | <u>001</u> | Aug 31, 2004 |
| <u>AT</u> | | BAUSCH AND LOMB | <u>0.3%</u> | <u>A076622</u> | <u>001</u> | May 14, 2004 |
| <u>AT</u> | | FDC LTD | <u>0.3%</u> | <u>A078559</u> | <u>001</u> | Feb 25, 2009 |
| <u>AT</u> | | HI TECH PHARMA | <u>0.3%</u> | <u>A076615</u> | <u>001</u> | May 14, 2004 |
| <u>AT</u> | | SANDOZ INC | <u>0.3%</u> | <u>A076231</u> | <u>001</u> | May 14, 2004 |

SOLUTION/DROPS; OTIC

OFLOXACIN

| | | | | | | |
|-----------|---|-----------------|-------------|----------------|------------|--------------|
| <u>AT</u> | | ALVOGEN | <u>0.3%</u> | <u>A090395</u> | <u>001</u> | Aug 11, 2009 |
| <u>AT</u> | | APOTEX INC | <u>0.3%</u> | <u>A076527</u> | <u>001</u> | Sep 28, 2007 |
| <u>AT</u> | ! | BAUSCH AND LOMB | <u>0.3%</u> | <u>A076128</u> | <u>001</u> | Mar 17, 2008 |

PRESCRIPTION DRUG PRODUCT LIST

OFLOXACIN

SOLUTION/DROPS;OTIC

OFLOXACIN

| | | | | | |
|-----------|----------------|-------------|----------------|------------|--------------|
| <u>AT</u> | HI TECH PHARMA | <u>0.3%</u> | <u>A076616</u> | <u>001</u> | Mar 17, 2008 |
| <u>AT</u> | SANDOZ INC | <u>0.3%</u> | <u>A078222</u> | <u>001</u> | Mar 17, 2008 |

TABLET;ORAL

OFLOXACIN

| | | | | | |
|-----------|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | CADILA PHARMS LTD | <u>200MG</u> | <u>A091656</u> | <u>001</u> | Sep 18, 2014 |
| <u>AB</u> | | <u>300MG</u> | <u>A091656</u> | <u>002</u> | Sep 18, 2014 |
| <u>AB</u> | | <u>400MG</u> | <u>A091656</u> | <u>003</u> | Sep 18, 2014 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>200MG</u> | <u>A077098</u> | <u>001</u> | Feb 10, 2006 |
| <u>AB</u> | | <u>300MG</u> | <u>A077098</u> | <u>002</u> | Feb 10, 2006 |
| <u>AB</u> | | <u>400MG</u> | <u>A077098</u> | <u>003</u> | Feb 10, 2006 |
| <u>AB</u> | LARKEN LABS | <u>400MG</u> | <u>A076093</u> | <u>003</u> | Sep 02, 2003 |
| <u>AB</u> | TEVA | <u>200MG</u> | <u>A076182</u> | <u>001</u> | Sep 02, 2003 |
| <u>AB</u> | | <u>300MG</u> | <u>A076182</u> | <u>002</u> | Sep 02, 2003 |
| <u>AB</u> | ! | <u>400MG</u> | <u>A076182</u> | <u>003</u> | Sep 02, 2003 |

OLANZAPINE

INJECTABLE;INTRAMUSCULAR

OLANZAPINE

| | | | | | |
|-----------|------------|------------------|----------------|------------|--------------|
| <u>AP</u> | LUITPOLD | <u>10MG/VIAL</u> | <u>A201741</u> | <u>001</u> | Mar 20, 2012 |
| <u>AP</u> | SANDOZ INC | <u>10MG/VIAL</u> | <u>A201588</u> | <u>001</u> | Oct 24, 2011 |

ZYPREXA

| | | | | | |
|-----------|----------|------------------|----------------|------------|--------------|
| <u>AP</u> | +! LILLY | <u>10MG/VIAL</u> | <u>N021253</u> | <u>001</u> | Mar 29, 2004 |
|-----------|----------|------------------|----------------|------------|--------------|

TABLET;ORAL

OLANZAPINE

| | | | | | |
|-----------|----------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ALKEM LABS LTD | <u>2.5MG</u> | <u>A202295</u> | <u>001</u> | Oct 20, 2015 |
| <u>AB</u> | | <u>5MG</u> | <u>A202295</u> | <u>002</u> | Oct 20, 2015 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A202295</u> | <u>003</u> | Oct 20, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A202295</u> | <u>004</u> | Oct 20, 2015 |
| <u>AB</u> | | <u>15MG</u> | <u>A202295</u> | <u>005</u> | Oct 20, 2015 |
| <u>AB</u> | | <u>20MG</u> | <u>A202295</u> | <u>006</u> | Oct 20, 2015 |
| <u>AB</u> | APOTEX INC | <u>2.5MG</u> | <u>A090798</u> | <u>001</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>5MG</u> | <u>A090798</u> | <u>002</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A090798</u> | <u>003</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>10MG</u> | <u>A090798</u> | <u>004</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>15MG</u> | <u>A090798</u> | <u>005</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>20MG</u> | <u>A090798</u> | <u>006</u> | Apr 23, 2012 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>2.5MG</u> | <u>A202050</u> | <u>001</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>5MG</u> | <u>A202050</u> | <u>002</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A202050</u> | <u>003</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>10MG</u> | <u>A202050</u> | <u>004</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>15MG</u> | <u>A202050</u> | <u>005</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>20MG</u> | <u>A202050</u> | <u>006</u> | Apr 23, 2012 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>2.5MG</u> | <u>A076255</u> | <u>001</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>5MG</u> | <u>A076255</u> | <u>002</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A076255</u> | <u>003</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>10MG</u> | <u>A076255</u> | <u>004</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>15MG</u> | <u>A076133</u> | <u>001</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>20MG</u> | <u>A076133</u> | <u>002</u> | Oct 24, 2011 |
| <u>AB</u> | HIKMA PHARMS | <u>2.5MG</u> | <u>A204866</u> | <u>001</u> | Jun 16, 2017 |
| <u>AB</u> | | <u>5MG</u> | <u>A204866</u> | <u>002</u> | Jun 16, 2017 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A204866</u> | <u>003</u> | Jun 16, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A204866</u> | <u>004</u> | Jun 16, 2017 |
| <u>AB</u> | | <u>15MG</u> | <u>A204866</u> | <u>005</u> | Jun 16, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A204866</u> | <u>006</u> | Jun 16, 2017 |
| <u>AB</u> | INVAGEN PHARMS | <u>2.5MG</u> | <u>A203333</u> | <u>001</u> | Mar 15, 2016 |
| <u>AB</u> | | <u>5MG</u> | <u>A203333</u> | <u>002</u> | Mar 15, 2016 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A203333</u> | <u>003</u> | Mar 15, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A203333</u> | <u>004</u> | Mar 15, 2016 |
| <u>AB</u> | | <u>15MG</u> | <u>A203333</u> | <u>005</u> | Mar 15, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A203333</u> | <u>006</u> | Mar 15, 2016 |
| <u>AB</u> | IVAX PHARMS INC | <u>20MG</u> | <u>A077301</u> | <u>001</u> | Apr 29, 2015 |
| <u>AB</u> | JIANGSU HANSOH PHARM | <u>2.5MG</u> | <u>A209399</u> | <u>001</u> | Sep 24, 2018 |
| <u>AB</u> | | <u>5MG</u> | <u>A209399</u> | <u>002</u> | Sep 24, 2018 |
| <u>AB</u> | | <u>10MG</u> | <u>A209399</u> | <u>003</u> | Sep 24, 2018 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>2.5MG</u> | <u>A202862</u> | <u>001</u> | Aug 15, 2014 |
| <u>AB</u> | | <u>5MG</u> | <u>A202862</u> | <u>002</u> | Aug 15, 2014 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A202862</u> | <u>003</u> | Aug 15, 2014 |
| <u>AB</u> | | <u>10MG</u> | <u>A202862</u> | <u>004</u> | Aug 15, 2014 |

PRESCRIPTION DRUG PRODUCT LIST

OLANZAPINE

TABLET; ORAL

OLANZAPINE

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | <u>15MG</u> | <u>A202862</u> | <u>005</u> | Aug 15, 2014 |
| <u>AB</u> | | <u>20MG</u> | <u>A202862</u> | <u>006</u> | Aug 15, 2014 |
| <u>AB</u> | ORCHID HLTHCARE | <u>2.5MG</u> | <u>A202287</u> | <u>001</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>5MG</u> | <u>A202287</u> | <u>002</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A202287</u> | <u>003</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>10MG</u> | <u>A202287</u> | <u>004</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>15MG</u> | <u>A202287</u> | <u>005</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>20MG</u> | <u>A202287</u> | <u>006</u> | Apr 23, 2012 |
| <u>AB</u> | QILU PHARM CO LTD | <u>2.5MG</u> | <u>A204319</u> | <u>001</u> | Jan 27, 2016 |
| <u>AB</u> | | <u>5MG</u> | <u>A204319</u> | <u>002</u> | Jan 27, 2016 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A204319</u> | <u>003</u> | Jan 27, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A204319</u> | <u>004</u> | Jan 27, 2016 |
| <u>AB</u> | | <u>15MG</u> | <u>A204319</u> | <u>005</u> | Jan 27, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A204319</u> | <u>006</u> | Jan 27, 2016 |
| <u>AB</u> | SUN PHARM INDS | <u>2.5MG</u> | <u>A091038</u> | <u>001</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>5MG</u> | <u>A091038</u> | <u>002</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A091038</u> | <u>003</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>10MG</u> | <u>A091038</u> | <u>004</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>15MG</u> | <u>A091038</u> | <u>005</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>20MG</u> | <u>A091038</u> | <u>006</u> | Apr 23, 2012 |
| <u>AB</u> | SUNSHINE LAKE | <u>2.5MG</u> | <u>A206238</u> | <u>001</u> | Nov 19, 2018 |
| <u>AB</u> | | <u>5MG</u> | <u>A206238</u> | <u>002</u> | Nov 19, 2018 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A206238</u> | <u>003</u> | Nov 19, 2018 |
| <u>AB</u> | | <u>10MG</u> | <u>A206238</u> | <u>004</u> | Nov 19, 2018 |
| <u>AB</u> | | <u>15MG</u> | <u>A206238</u> | <u>005</u> | Nov 19, 2018 |
| <u>AB</u> | | <u>20MG</u> | <u>A206238</u> | <u>006</u> | Nov 19, 2018 |
| <u>AB</u> | TEVA PHARMS | <u>2.5MG</u> | <u>A076000</u> | <u>001</u> | Oct 24, 2011 |
| <u>AB</u> | | <u>5MG</u> | <u>A076000</u> | <u>002</u> | Oct 24, 2011 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A076000</u> | <u>003</u> | Oct 24, 2011 |
| <u>AB</u> | | <u>10MG</u> | <u>A076000</u> | <u>004</u> | Oct 24, 2011 |
| <u>AB</u> | | <u>15MG</u> | <u>A076000</u> | <u>005</u> | Oct 24, 2011 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>2.5MG</u> | <u>A091434</u> | <u>001</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>5MG</u> | <u>A091434</u> | <u>002</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A091434</u> | <u>003</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>10MG</u> | <u>A091434</u> | <u>004</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>15MG</u> | <u>A091434</u> | <u>005</u> | Apr 23, 2012 |
| <u>AB</u> | | <u>20MG</u> | <u>A091434</u> | <u>006</u> | Apr 23, 2012 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>2.5MG</u> | <u>A090459</u> | <u>001</u> | Jul 16, 2018 |
| <u>AB</u> | | <u>5MG</u> | <u>A090459</u> | <u>002</u> | Jul 16, 2018 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A090459</u> | <u>003</u> | Jul 16, 2018 |
| <u>AB</u> | | <u>10MG</u> | <u>A090459</u> | <u>004</u> | Jul 16, 2018 |
| <u>AB</u> | | <u>15MG</u> | <u>A090459</u> | <u>005</u> | Jul 16, 2018 |
| <u>AB</u> | | <u>20MG</u> | <u>A090459</u> | <u>006</u> | Jul 16, 2018 |

ZYPREXA

| | | | | | | |
|-----------|---|-------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | LILLY | <u>2.5MG</u> | <u>N020592</u> | <u>001</u> | Sep 30, 1996 |
| <u>AB</u> | + | ! | <u>5MG</u> | <u>N020592</u> | <u>002</u> | Sep 30, 1996 |
| <u>AB</u> | + | | <u>7.5MG</u> | <u>N020592</u> | <u>003</u> | Sep 30, 1996 |
| <u>AB</u> | + | | <u>10MG</u> | <u>N020592</u> | <u>004</u> | Sep 30, 1996 |
| <u>AB</u> | + | | <u>15MG</u> | <u>N020592</u> | <u>005</u> | Sep 09, 1997 |
| <u>AB</u> | + | | <u>20MG</u> | <u>N020592</u> | <u>006</u> | Sep 09, 1997 |

TABLET, ORALLY DISINTEGRATING; ORAL

OLANZAPINE

| | | | | | |
|-----------|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | APOTEX INC | <u>5MG</u> | <u>A091265</u> | <u>001</u> | Oct 24, 2011 |
| <u>AB</u> | | <u>10MG</u> | <u>A091265</u> | <u>002</u> | Oct 24, 2011 |
| <u>AB</u> | | <u>15MG</u> | <u>A091265</u> | <u>003</u> | Oct 24, 2011 |
| <u>AB</u> | | <u>20MG</u> | <u>A091265</u> | <u>004</u> | Oct 24, 2011 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>5MG</u> | <u>A203708</u> | <u>001</u> | May 15, 2014 |
| <u>AB</u> | | <u>10MG</u> | <u>A203708</u> | <u>002</u> | May 15, 2014 |
| <u>AB</u> | | <u>15MG</u> | <u>A203708</u> | <u>003</u> | May 15, 2014 |
| <u>AB</u> | | <u>20MG</u> | <u>A203708</u> | <u>004</u> | May 15, 2014 |
| <u>AB</u> | BARR LABS INC | <u>5MG</u> | <u>A077243</u> | <u>001</u> | Jan 30, 2012 |
| <u>AB</u> | | <u>10MG</u> | <u>A077243</u> | <u>002</u> | Jan 30, 2012 |
| <u>AB</u> | | <u>15MG</u> | <u>A077243</u> | <u>003</u> | Jan 30, 2012 |
| <u>AB</u> | | <u>20MG</u> | <u>A077243</u> | <u>004</u> | Jan 30, 2012 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>5MG</u> | <u>A076534</u> | <u>001</u> | Oct 24, 2011 |
| <u>AB</u> | | <u>10MG</u> | <u>A076534</u> | <u>002</u> | Oct 24, 2011 |
| <u>AB</u> | | <u>15MG</u> | <u>A076534</u> | <u>003</u> | Oct 24, 2011 |
| <u>AB</u> | | <u>20MG</u> | <u>A076534</u> | <u>004</u> | Oct 24, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

OLANZAPINE

TABLET, ORALLY DISINTEGRATING;ORAL

OLANZAPINE

| | | | | |
|----------------------|---------------------|-------------|--------------------|--------------|
| <u>AB</u> | HEC PHARM | <u>5MG</u> | <u>A208146 001</u> | Jul 02, 2018 |
| <u>AB</u> | | <u>10MG</u> | <u>A208146 002</u> | Jul 02, 2018 |
| <u>AB</u> | | <u>15MG</u> | <u>A208146 003</u> | Jul 02, 2018 |
| <u>AB</u> | | <u>20MG</u> | <u>A208146 004</u> | Jul 02, 2018 |
| <u>AB</u> | INVAGEN PHARMS | <u>5MG</u> | <u>A203456 001</u> | Mar 16, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A203456 002</u> | Mar 16, 2016 |
| <u>AB</u> | | <u>15MG</u> | <u>A203456 003</u> | Mar 16, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A203456 004</u> | Mar 16, 2016 |
| <u>AB</u> | JUBILANT GENERICS | <u>5MG</u> | <u>A200221 001</u> | Sep 12, 2012 |
| <u>AB</u> | | <u>10MG</u> | <u>A200221 002</u> | Sep 12, 2012 |
| <u>AB</u> | | <u>15MG</u> | <u>A200221 003</u> | Sep 12, 2012 |
| <u>AB</u> | | <u>20MG</u> | <u>A200221 004</u> | Sep 12, 2012 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>5MG</u> | <u>A203044 001</u> | Feb 20, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A203044 002</u> | Feb 20, 2015 |
| <u>AB</u> | | <u>15MG</u> | <u>A203044 003</u> | Feb 20, 2015 |
| <u>AB</u> | | <u>20MG</u> | <u>A203044 004</u> | Feb 20, 2015 |
| <u>AB</u> | MYLAN PHARMS INC | <u>5MG</u> | <u>A202285 001</u> | May 12, 2014 |
| <u>AB</u> | | <u>10MG</u> | <u>A202285 002</u> | May 12, 2014 |
| <u>AB</u> | | <u>15MG</u> | <u>A202285 003</u> | May 12, 2014 |
| <u>AB</u> | | <u>20MG</u> | <u>A202285 004</u> | May 12, 2014 |
| <u>AB</u> | ORCHID HLTHCARE | <u>5MG</u> | <u>A202937 001</u> | Mar 02, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A202937 002</u> | Mar 02, 2015 |
| <u>AB</u> | | <u>15MG</u> | <u>A202937 003</u> | Mar 02, 2015 |
| <u>AB</u> | | <u>20MG</u> | <u>A202937 004</u> | Mar 02, 2015 |
| <u>AB</u> | PAR PHARM | <u>5MG</u> | <u>A078109 001</u> | Oct 24, 2011 |
| <u>AB</u> | | <u>10MG</u> | <u>A078109 002</u> | Oct 24, 2011 |
| <u>AB</u> | | <u>15MG</u> | <u>A078109 003</u> | Oct 24, 2011 |
| <u>AB</u> | | <u>20MG</u> | <u>A078109 004</u> | Oct 24, 2011 |
| <u>AB</u> | SUN PHARM INDS | <u>5MG</u> | <u>A090881 001</u> | Feb 28, 2012 |
| <u>AB</u> | | <u>10MG</u> | <u>A090881 002</u> | Feb 28, 2012 |
| <u>AB</u> | | <u>15MG</u> | <u>A090881 003</u> | Feb 28, 2012 |
| <u>AB</u> | | <u>20MG</u> | <u>A090881 004</u> | Feb 28, 2012 |
| <u>AB</u> | TORRENT PHARMS LLC | <u>5MG</u> | <u>A091415 001</u> | Oct 25, 2011 |
| <u>AB</u> | | <u>10MG</u> | <u>A091415 002</u> | Oct 25, 2011 |
| <u>AB</u> | | <u>15MG</u> | <u>A091415 003</u> | Oct 25, 2011 |
| <u>AB</u> | | <u>20MG</u> | <u>A091415 004</u> | Oct 25, 2011 |
| <u>ZYPREXA ZYDIS</u> | | | | |
| <u>AB</u> | +! LILLY | <u>5MG</u> | <u>N021086 001</u> | Apr 06, 2000 |
| <u>AB</u> | + | <u>10MG</u> | <u>N021086 002</u> | Apr 06, 2000 |
| <u>AB</u> | + | <u>15MG</u> | <u>N021086 003</u> | Apr 06, 2000 |
| <u>AB</u> | + | <u>20MG</u> | <u>N021086 004</u> | Apr 06, 2000 |

OLANZAPINE PAMOATE

SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

ZYPREXA RELPREV

| | | | | |
|----|--------------|--------------------|-------------|--------------|
| + | ELI LILLY CO | EQ 210MG BASE/VIAL | N022173 001 | Dec 11, 2009 |
| + | | EQ 300MG BASE/VIAL | N022173 002 | Dec 11, 2009 |
| +! | | EQ 405MG BASE/VIAL | N022173 003 | Dec 11, 2009 |

OLAPARIB

CAPSULE;ORAL

LYNPARZA

| | | | | |
|----|--------------------|------|-------------|--------------|
| +! | ASTRAZENECA PHARMS | 50MG | N206162 001 | Dec 19, 2014 |
|----|--------------------|------|-------------|--------------|

TABLET;ORAL

LYNPARZA

| | | | | |
|----|--------------------|-------|-------------|--------------|
| + | ASTRAZENECA PHARMS | 100MG | N208558 001 | Aug 17, 2017 |
| +! | | 150MG | N208558 002 | Aug 17, 2017 |

OLMESARTAN MEDOXOMIL

TABLET;ORAL

BENICAR

| | | | | | |
|-----------|----|----------------|-------------|--------------------|--------------|
| <u>AB</u> | + | DAIICHI SANKYO | <u>5MG</u> | <u>N021286 001</u> | Apr 25, 2002 |
| <u>AB</u> | + | | <u>20MG</u> | <u>N021286 003</u> | Apr 25, 2002 |
| <u>AB</u> | +! | | <u>40MG</u> | <u>N021286 004</u> | Apr 25, 2002 |

OLMESARTAN MEDOXOMIL

| | | | | |
|-----------|--------------------|-------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>5MG</u> | <u>A207662 001</u> | Apr 24, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A207662 002</u> | Apr 24, 2017 |
| <u>AB</u> | | <u>40MG</u> | <u>A207662 003</u> | Apr 24, 2017 |
| <u>AB</u> | ALEMBIC PHARMS LTD | <u>5MG</u> | <u>A203012 001</u> | Apr 24, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A203012 002</u> | Apr 24, 2017 |
| <u>AB</u> | | <u>40MG</u> | <u>A203012 003</u> | Apr 24, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

OLMESARTAN MEDOXOMIL

TABLET; ORAL

OLMESARTAN MEDOXOMIL

| | | | | |
|-----------|----------------------|-------------|--------------------|--------------|
| AB | ALKEM LABS LTD | 5MG | A206763 001 | Apr 24, 2017 |
| AB | | 20MG | A206763 002 | Apr 24, 2017 |
| AB | | 40MG | A206763 003 | Apr 24, 2017 |
| AB | AUROBINDO PHARMA LTD | 5MG | A204798 001 | Apr 24, 2017 |
| AB | | 20MG | A204798 002 | Apr 24, 2017 |
| AB | | 40MG | A204798 003 | Apr 24, 2017 |
| AB | GLENMARK PHARMS LTD | 5MG | A203281 001 | May 25, 2017 |
| AB | | 20MG | A203281 002 | May 25, 2017 |
| AB | | 40MG | A203281 003 | May 25, 2017 |
| AB | JUBILANT GENERICS | 5MG | A205482 001 | Apr 24, 2017 |
| AB | | 20MG | A205482 002 | Apr 24, 2017 |
| AB | | 40MG | A205482 003 | Apr 24, 2017 |
| AB | LUPIN LTD | 5MG | A206631 001 | Apr 27, 2017 |
| AB | | 20MG | A206631 002 | Apr 27, 2017 |
| AB | | 40MG | A206631 003 | Apr 27, 2017 |
| AB | MACLEODS PHARMS LTD | 5MG | A204814 001 | Apr 24, 2017 |
| AB | | 20MG | A204814 002 | Apr 24, 2017 |
| AB | | 40MG | A204814 003 | Apr 24, 2017 |
| AB | MYLAN PHARMS INC | 5MG | A078276 001 | Oct 26, 2016 |
| AB | | 20MG | A078276 002 | Oct 26, 2016 |
| AB | | 40MG | A078276 003 | Oct 26, 2016 |
| AB | QILU PHARM CO LTD | 5MG | A210552 001 | Jan 10, 2019 |
| AB | | 20MG | A210552 002 | Jan 10, 2019 |
| AB | | 40MG | A210552 003 | Jan 10, 2019 |
| AB | SCIEGEN PHARMS INC | 5MG | A208130 001 | Jun 29, 2018 |
| AB | | 20MG | A208130 002 | Jun 29, 2018 |
| AB | | 40MG | A208130 003 | Jun 29, 2018 |
| AB | TEVA PHARMS USA | 5MG | A091079 001 | Apr 24, 2017 |
| AB | | 20MG | A091079 002 | Apr 24, 2017 |
| AB | | 40MG | A091079 003 | Apr 24, 2017 |
| AB | TORRENT PHARMS LTD | 5MG | A202375 001 | Apr 24, 2017 |
| AB | | 20MG | A202375 002 | Apr 24, 2017 |
| AB | | 40MG | A202375 003 | Apr 24, 2017 |
| AB | ZYDUS PHARMS USA INC | 5MG | A205192 001 | Apr 24, 2017 |
| AB | | 20MG | A205192 002 | Apr 24, 2017 |
| AB | | 40MG | A205192 003 | Apr 24, 2017 |

OLODATEROL HYDROCHLORIDE

SPRAY, METERED; INHALATION

STRIVERDI RESPIMAT

| | | | | |
|---|----------------------|----------------------|-------------|--------------|
| + | BOEHRINGER INGELHEIM | EQ 0.0025MG BASE/INH | N203108 001 | Jul 31, 2014 |
|---|----------------------|----------------------|-------------|--------------|

OLODATEROL HYDROCHLORIDE; TIOTROPIUM BROMIDE

SPRAY, METERED; INHALATION

STIOLTO RESPIMAT

| | | | | |
|---|----------------------|--------------------------------------------|-------------|--------------|
| + | BOEHRINGER INGELHEIM | EQ 0.0025MG BASE/INH; EQ 0.0025MG BASE/INH | N206756 001 | May 21, 2015 |
|---|----------------------|--------------------------------------------|-------------|--------------|

OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

OLOPATADINE HYDROCHLORIDE

| | | | | |
|-----------------------|----------------------|----------------------|---------------------|---------------------------------|
| AT | AKORN | EQ 0.2% BASE | A204723 001 | Dec 05, 2017 |
| AT | AKORN INC | EQ 0.1% BASE | A204532 001 | Jan 10, 2017 |
| AT | ALEMBIC PHARMS LTD | EQ 0.1% BASE | A209919 001 | Dec 07, 2018 |
| AT | APOTEX INC | EQ 0.1% BASE | A078350 001 | Dec 07, 2015 |
| AT | | EQ 0.2% BASE | A090918 001 | Dec 05, 2017 |
| AT | AUROBINDO PHARMA LTD | EQ 0.1% BASE | A204812 001 | Dec 18, 2015 |
| AT | BARR LABS INC | EQ 0.2% BASE | A090848 001 | Jul 13, 2015 |
| AT | CIPLA | EQ 0.1% BASE | A206046 001 | Jul 26, 2017 |
| AT | | EQ 0.2% BASE | A206087 001 | Dec 05, 2017 |
| AT | MYLAN PHARMS INC | EQ 0.1% BASE | A204392 001 | Mar 21, 2018 |
| AT | SOMERSET THERAPS LLC | EQ 0.1% BASE | A206306 001 | Dec 07, 2015 |
| AT | USV NORTH AMERICA | EQ 0.1% BASE | A203152 001 | Dec 07, 2015 |
| AT | WOCKHARDT LTD | EQ 0.1% BASE | A200810 001 | Jun 28, 2017 |
| <u>PATADAY</u> | | | | |
| AT | + | NOVARTIS PHARMS CORP | EQ 0.2% BASE | N021545 001 Dec 22, 2004 |

PRESCRIPTION DRUG PRODUCT LIST

OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

PATANOL

| | | | | | |
|-----------|------------|-------------------------|---------------------|--------------------|--------------|
| AT | + ! | NOVARTIS PHARMS CORP | EQ 0.1% BASE | N020688 001 | Dec 18, 1996 |
|-----------|------------|-------------------------|---------------------|--------------------|--------------|

PAZEO

| | | | | | |
|--|------------|-------------------------|--------------|-------------|--------------|
| | + ! | NOVARTIS PHARMS CORP | EQ 0.7% BASE | N206276 001 | Jan 30, 2015 |
|--|------------|-------------------------|--------------|-------------|--------------|

SPRAY, METERED;NASAL

OLOPATADINE HYDROCHLORIDE

| | | | | | |
|-----------|--|------------|----------------------|--------------------|--------------|
| AB | | APOTEX INC | 0.665MG/SPRAY | A091572 001 | Oct 08, 2014 |
|-----------|--|------------|----------------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------|----------------------|--------------------|--------------|
| AB | | PERRIGO ISRAEL | 0.665MG/SPRAY | A202853 001 | Jan 31, 2017 |
|-----------|--|----------------|----------------------|--------------------|--------------|

PATANASE

| | | | | | |
|-----------|------------|-------------------------|----------------------|--------------------|--------------|
| AB | + ! | NOVARTIS PHARMS CORP | 0.665MG/SPRAY | N021861 001 | Apr 15, 2008 |
|-----------|------------|-------------------------|----------------------|--------------------|--------------|

OLSALAZINE SODIUM

CAPSULE;ORAL

DIPENTUM

| | | | | | |
|--|------------|---------------------|-------|-------------|--------------|
| | + ! | MYLAN SPECIALITY LP | 250MG | N019715 001 | Jul 31, 1990 |
|--|------------|---------------------|-------|-------------|--------------|

OMACETAXINE MEPESUCCINATE

POWDER;SUBCUTANEOUS

SYNRIBO

| | | | | | |
|--|------------|------------------|------------|-------------|--------------|
| | + ! | TEVA PHARMS INTL | 3.5MG/VIAL | N203585 001 | Oct 26, 2012 |
|--|------------|------------------|------------|-------------|--------------|

OMADACYCLINE TOSYLATE

POWDER;INTRAVENOUS

NUZYRA

| | | | | | |
|--|------------|--------------------|--------------------|-------------|--------------|
| | + ! | PARATEK PHARMS INC | EQ 100MG BASE/VIAL | N209817 001 | Oct 02, 2018 |
|--|------------|--------------------|--------------------|-------------|--------------|

TABLET;ORAL

NUZYRA

| | | | | | |
|--|------------|--------------------|---------------|-------------|--------------|
| | + ! | PARATEK PHARMS INC | EQ 150MG BASE | N209816 001 | Oct 02, 2018 |
|--|------------|--------------------|---------------|-------------|--------------|

OMBITASVIR; PARITAPREVIR; RITONAVIR

TABLET;ORAL

TECHNIVIE

| | | | | | |
|--|------------|------------|------------------|-------------|--------------|
| | + ! | ABEVIE INC | 12.5MG;75MG;50MG | N207931 001 | Jul 24, 2015 |
|--|------------|------------|------------------|-------------|--------------|

OMEGA-3-ACID ETHYL ESTERS

CAPSULE;ORAL

LOVAZA

| | | | | | |
|-----------|------------|--------------------|-------------------------------------------------------------------------------|--------------------|--------------|
| AB | + ! | SMITHKLINE BEECHAM | 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS | N021654 001 | Nov 10, 2004 |
|-----------|------------|--------------------|-------------------------------------------------------------------------------|--------------------|--------------|

OMEGA-3-ACID ETHYL ESTERS

| | | | | | |
|-----------|--|---------------|-------------------------------------------------------------------------------|--------------------|--------------|
| AB | | AMNEAL PHARMS | 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS | A204940 001 | Nov 27, 2015 |
|-----------|--|---------------|-------------------------------------------------------------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------|-------------------------------------------------------------------------------|--------------------|--------------|
| AB | | APOTEX INC | 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS | A090973 001 | Sep 30, 2014 |
|-----------|--|------------|-------------------------------------------------------------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|---------------|-------------------------------------------------------------------------------|--------------------|--------------|
| AB | | PAR PHARM INC | 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS | A091018 001 | Jun 24, 2014 |
|-----------|--|---------------|-------------------------------------------------------------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------|-------------------------------------------------------------------------------|--------------------|--------------|
| AB | | STRIDES PHARMA | 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS | A203893 001 | Sep 19, 2017 |
|-----------|--|----------------|-------------------------------------------------------------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-----------------|-------------------------------------------------------------------------------|--------------------|--------------|
| AB | | TEVA PHARMS USA | 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS | A091028 001 | Apr 07, 2014 |
|-----------|--|-----------------|-------------------------------------------------------------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-----------------|-------------------------------------------------------------------------------|--------------------|--------------|
| AB | | TEVA PHARMS USA | 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS | A091028 001 | Apr 07, 2014 |
|-----------|--|-----------------|-------------------------------------------------------------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-----------------|-------------------------------------------------------------------------------|--------------------|--------------|
| AB | | TEVA PHARMS USA | 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS | A091028 001 | Apr 07, 2014 |
|-----------|--|-----------------|-------------------------------------------------------------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-----------------|-------------------------------------------------------------------------------|--------------------|--------------|
| AB | | TEVA PHARMS USA | 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS | A091028 001 | Apr 07, 2014 |
|-----------|--|-----------------|-------------------------------------------------------------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-----------------|-------------------------------------------------------------------------------|--------------------|--------------|
| AB | | TEVA PHARMS USA | 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS | A091028 001 | Apr 07, 2014 |
|-----------|--|-----------------|-------------------------------------------------------------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-----------------|-------------------------------------------------------------------------------|--------------------|--------------|
| AB | | TEVA PHARMS USA | 1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS | A091028 001 | Apr 07, 2014 |
|-----------|--|-----------------|-------------------------------------------------------------------------------|--------------------|--------------|

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

OMEPRAZOLE

| | | | | | |
|-----------|--|---------------------|-------------|--------------------|--------------|
| AB | | ACTAVIS LABS FL INC | 10MG | A075347 001 | May 30, 2008 |
|-----------|--|---------------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 20MG | A075347 002 | May 30, 2008 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 40MG | A075347 003 | May 30, 2008 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------|-------------|--------------------|--------------|
| AB | | APOTEX | 10MG | A076048 001 | Oct 22, 2007 |
|-----------|--|--------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 20MG | A076048 002 | Oct 22, 2007 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 40MG | A076048 003 | Jan 21, 2009 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------------|-------------|--------------------|--------------|
| AB | | AUROBINDO PHARMA LTD | 10MG | A203270 001 | Aug 19, 2015 |
|-----------|--|----------------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 20MG | A203270 002 | Aug 19, 2015 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 40MG | A203270 003 | Aug 19, 2015 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------------|-------------|--------------------|--------------|
| AB | | BRECKENRIDGE PHARM | 10MG | A203481 001 | Jul 03, 2017 |
|-----------|--|--------------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 20MG | A203481 002 | Jul 03, 2017 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 40MG | A203481 003 | Jul 03, 2017 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------------------|-------------|--------------------|--------------|
| AB | | DR REDDYS LABS LTD | 10MG | A075576 003 | Oct 22, 2007 |
|-----------|--|--------------------|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 10MG | A078490 002 | Mar 16, 2009 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 20MG | A075576 002 | Oct 22, 2007 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 20MG | A078490 003 | Mar 16, 2009 |
|-----------|--|--|-------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|-------------|--------------------|--------------|
| AB | | | 40MG | A075576 001 | Jan 21, 2009 |
|-----------|--|--|-------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

OMEPRAZOLE

| | | | | | |
|-----------|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | <u>40MG</u> | <u>A078490</u> | <u>001</u> | Apr 17, 2009 |
| <u>AB</u> | GLENMARK GENERICS | <u>10MG</u> | <u>A091672</u> | <u>001</u> | Oct 31, 2014 |
| <u>AB</u> | | <u>20MG</u> | <u>A091672</u> | <u>002</u> | Oct 31, 2014 |
| <u>AB</u> | | <u>40MG</u> | <u>A091672</u> | <u>003</u> | Oct 31, 2014 |
| <u>AB</u> | IMPAX LABS | <u>10MG</u> | <u>A075785</u> | <u>001</u> | Oct 22, 2007 |
| <u>AB</u> | | <u>20MG</u> | <u>A075785</u> | <u>002</u> | Oct 22, 2007 |
| <u>AB</u> | | <u>40MG</u> | <u>A075785</u> | <u>003</u> | Jan 21, 2009 |
| <u>AB</u> | LANNETT CO INC | <u>10MG</u> | <u>A075410</u> | <u>001</u> | Nov 01, 2002 |
| <u>AB</u> | | <u>20MG</u> | <u>A075410</u> | <u>002</u> | Nov 01, 2002 |
| <u>AB</u> | | <u>40MG</u> | <u>A075410</u> | <u>003</u> | Jan 23, 2009 |
| <u>AB</u> | LUPIN LTD | <u>40MG</u> | <u>A202384</u> | <u>001</u> | Aug 25, 2015 |
| <u>AB</u> | MYLAN | <u>10MG</u> | <u>A075876</u> | <u>001</u> | May 29, 2003 |
| <u>AB</u> | | <u>20MG</u> | <u>A075876</u> | <u>002</u> | May 29, 2003 |
| <u>AB</u> | | <u>40MG</u> | <u>A075876</u> | <u>003</u> | Jan 21, 2009 |
| <u>AB</u> | MYLAN PHARMS INC | <u>10MG</u> | <u>A205070</u> | <u>001</u> | Jun 29, 2018 |
| <u>AB</u> | | <u>20MG</u> | <u>A205070</u> | <u>002</u> | Jun 29, 2018 |
| <u>AB</u> | | <u>40MG</u> | <u>A205070</u> | <u>003</u> | Jun 29, 2018 |
| <u>AB</u> | SANDOZ | <u>10MG</u> | <u>A075757</u> | <u>001</u> | Jan 28, 2003 |
| <u>AB</u> | ! | <u>20MG</u> | <u>A075757</u> | <u>002</u> | Jan 28, 2003 |
| <u>AB</u> | ! | <u>40MG</u> | <u>A076515</u> | <u>001</u> | Jan 21, 2009 |
| <u>AB</u> | TEVA PHARMS USA | <u>20MG</u> | <u>A204661</u> | <u>001</u> | Jun 13, 2017 |
| <u>AB</u> | | <u>40MG</u> | <u>A204661</u> | <u>002</u> | Jun 13, 2017 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>10MG</u> | <u>A091352</u> | <u>001</u> | Nov 19, 2012 |
| <u>AB</u> | | <u>20MG</u> | <u>A091352</u> | <u>002</u> | Nov 19, 2012 |
| <u>AB</u> | | <u>40MG</u> | <u>A091352</u> | <u>003</u> | Nov 19, 2012 |

OMEPRAZOLE MAGNESIUM

FOR SUSPENSION, DELAYED RELEASE;ORAL

PRILOSEC

| | | | | | |
|---|-----------------|----------------------|---------|-----|--------------|
| + | COVIS PHARMA BV | EQ 2.5MG BASE/PACKET | N022056 | 001 | Mar 20, 2008 |
| + | ! | EQ 10MG BASE/PACKET | N022056 | 002 | Mar 20, 2008 |

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

| | | | | | |
|-----------|-------------------------|-------------------|----------------|------------|--------------|
| <u>AB</u> | AJANTA PHARMA LTD | <u>20MG;1.1GM</u> | <u>A204228</u> | <u>001</u> | Jul 15, 2016 |
| <u>AB</u> | | <u>40MG;1.1GM</u> | <u>A204228</u> | <u>002</u> | Jul 15, 2016 |
| <u>AB</u> | AUROLIFE PHARMA LLC | <u>20MG;1.1GM</u> | <u>A204922</u> | <u>001</u> | Aug 19, 2016 |
| <u>AB</u> | | <u>40MG;1.1GM</u> | <u>A204922</u> | <u>002</u> | Aug 19, 2016 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>20MG;1.1GM</u> | <u>A204068</u> | <u>001</u> | Jul 15, 2016 |
| <u>AB</u> | | <u>40MG;1.1GM</u> | <u>A204068</u> | <u>002</u> | Jul 15, 2016 |
| <u>AB</u> | PAR PHARM | <u>20MG;1.1GM</u> | <u>A078966</u> | <u>001</u> | May 25, 2010 |
| <u>AB</u> | | <u>40MG;1.1GM</u> | <u>A078966</u> | <u>002</u> | May 25, 2010 |
| <u>AB</u> | SCIEGEN PHARMS INC | <u>20MG;1.1GM</u> | <u>A207476</u> | <u>001</u> | Dec 06, 2016 |
| <u>AB</u> | | <u>40MG;1.1GM</u> | <u>A207476</u> | <u>002</u> | Dec 06, 2016 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>20MG;1.1GM</u> | <u>A203290</u> | <u>001</u> | May 25, 2018 |
| <u>AB</u> | | <u>40MG;1.1GM</u> | <u>A203290</u> | <u>002</u> | May 25, 2018 |
| <u>AB</u> | <u>ZEGERID</u> | | | | |
| <u>AB</u> | + SANTARUS INC | <u>20MG;1.1GM</u> | <u>N021849</u> | <u>001</u> | Feb 27, 2006 |
| <u>AB</u> | + | <u>40MG;1.1GM</u> | <u>N021849</u> | <u>002</u> | Feb 27, 2006 |

FOR SUSPENSION;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

| | | | | | |
|-----------|-------------------|----------------------------------|----------------|------------|--------------|
| <u>AB</u> | AJANTA PHARMA LTD | <u>20MG/PACKET;1.68GM/PACKET</u> | <u>A205545</u> | <u>001</u> | Jul 27, 2016 |
| <u>AB</u> | | <u>40MG/PACKET;1.68GM/PACKET</u> | <u>A205545</u> | <u>002</u> | Jul 27, 2016 |
| <u>AB</u> | PAR PHARM | <u>20MG/PACKET;1.68GM/PACKET</u> | <u>A079182</u> | <u>001</u> | Apr 19, 2013 |
| <u>AB</u> | | <u>40MG/PACKET;1.68GM/PACKET</u> | <u>A079182</u> | <u>002</u> | Apr 19, 2013 |
| <u>AB</u> | <u>ZEGERID</u> | | | | |
| <u>AB</u> | + SANTARUS INC | <u>20MG/PACKET;1.68GM/PACKET</u> | <u>N021636</u> | <u>001</u> | Jun 15, 2004 |
| <u>AB</u> | + | <u>40MG/PACKET;1.68GM/PACKET</u> | <u>N021636</u> | <u>002</u> | Dec 21, 2004 |

ONDANSETRON

FILM;ORAL

ZUPLENZ

| | | | | | |
|---|--------------------|-----|---------|-----|--------------|
| + | MIDATECH PHARMA US | 4MG | N022524 | 001 | Jul 02, 2010 |
| + | ! | 8MG | N022524 | 002 | Jul 02, 2010 |

TABLET, ORALLY DISINTEGRATING;ORAL

ONDANSETRON

| | | | | | |
|-----------|------------------|------------|----------------|------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA | <u>4MG</u> | <u>A090469</u> | <u>001</u> | Apr 12, 2010 |
| <u>AB</u> | | <u>8MG</u> | <u>A090469</u> | <u>002</u> | Apr 12, 2010 |

PRESCRIPTION DRUG PRODUCT LIST

ONDANSETRON

TABLET, ORALLY DISINTEGRATING;ORAL

ONDANSETRON

| | | | | | |
|-----------|--------------------|------------|----------------|------------|--------------|
| <u>AB</u> | BARR | <u>4MG</u> | <u>A076693</u> | <u>001</u> | Jun 25, 2007 |
| <u>AB</u> | | <u>8MG</u> | <u>A076693</u> | <u>002</u> | Jun 25, 2007 |
| <u>AB</u> | GLENMARK GENERICS | <u>4MG</u> | <u>A078152</u> | <u>001</u> | Jun 27, 2007 |
| <u>AB</u> | | <u>8MG</u> | <u>A078152</u> | <u>002</u> | Jun 27, 2007 |
| <u>AB</u> | MYLAN | <u>4MG</u> | <u>A078139</u> | <u>001</u> | Jun 25, 2007 |
| <u>AB</u> | | <u>8MG</u> | <u>A078139</u> | <u>002</u> | Jun 25, 2007 |
| <u>AB</u> | SANDOZ | <u>4MG</u> | <u>A078050</u> | <u>001</u> | Aug 13, 2007 |
| <u>AB</u> | | <u>8MG</u> | <u>A078050</u> | <u>002</u> | Aug 13, 2007 |
| <u>AB</u> | SUN PHARM INDS | <u>4MG</u> | <u>A077557</u> | <u>001</u> | Aug 02, 2007 |
| <u>AB</u> | | <u>8MG</u> | <u>A077557</u> | <u>002</u> | Aug 02, 2007 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>4MG</u> | <u>A078602</u> | <u>001</u> | Feb 24, 2011 |
| <u>AB</u> | | <u>8MG</u> | <u>A078602</u> | <u>002</u> | Feb 24, 2011 |
| <u>AB</u> | TEVA | <u>4MG</u> | <u>A076810</u> | <u>001</u> | Jun 25, 2007 |
| <u>AB</u> | | <u>8MG</u> | <u>A076810</u> | <u>002</u> | Jun 25, 2007 |

ZOFRAN ODT

| | | | | | |
|-----------|---------------------------|------------|----------------|------------|--------------|
| <u>AB</u> | + NOVARTIS PHARMS CORP | <u>4MG</u> | <u>N020781</u> | <u>001</u> | Jan 27, 1999 |
| <u>AB</u> | +! | <u>8MG</u> | <u>N020781</u> | <u>002</u> | Jan 27, 1999 |

ONDANSETRON HYDROCHLORIDE

INJECTABLE; INJECTION

ONDANSETRON HYDROCHLORIDE

| | | | | | |
|-----------|---------------------------|-----------------------|----------------|------------|--------------|
| <u>AP</u> | ACCORD HLTHCARE | <u>EQ 2MG BASE/ML</u> | <u>A206846</u> | <u>001</u> | Jul 13, 2015 |
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>EQ 2MG BASE/ML</u> | <u>A202599</u> | <u>001</u> | Dec 21, 2012 |
| <u>AP</u> | ! BAXTER HLTHCARE CORP | <u>EQ 2MG BASE/ML</u> | <u>A078288</u> | <u>001</u> | Feb 22, 2013 |
| <u>AP</u> | EMCURE PHARMS | <u>EQ 2MG BASE/ML</u> | <u>A090424</u> | <u>001</u> | Apr 16, 2010 |
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 2MG BASE/ML</u> | <u>A076974</u> | <u>001</u> | Dec 26, 2006 |
| <u>AP</u> | GLAND PHARMA LTD | <u>EQ 2MG BASE/ML</u> | <u>A079224</u> | <u>001</u> | Sep 25, 2009 |
| <u>AP</u> | | <u>EQ 2MG BASE/ML</u> | <u>A090648</u> | <u>001</u> | Jun 15, 2012 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>EQ 2MG BASE/ML</u> | <u>A076781</u> | <u>001</u> | Dec 26, 2006 |
| <u>AP</u> | HOSPIRA | <u>EQ 2MG BASE/ML</u> | <u>A077473</u> | <u>001</u> | Dec 26, 2006 |
| <u>AP</u> | | <u>EQ 2MG BASE/ML</u> | <u>A077840</u> | <u>001</u> | Jan 19, 2007 |
| <u>AP</u> | LUITPOLD | <u>EQ 2MG BASE/ML</u> | <u>A079039</u> | <u>001</u> | Nov 18, 2008 |
| <u>AP</u> | MYLAN LABS LTD | <u>EQ 2MG BASE/ML</u> | <u>A204906</u> | <u>001</u> | Jul 31, 2017 |
| <u>AP</u> | QILU PHARM CO LTD | <u>EQ 2MG BASE/ML</u> | <u>A203711</u> | <u>001</u> | Sep 08, 2014 |
| <u>AP</u> | SANDOZ INC | <u>EQ 2MG BASE/ML</u> | <u>A077430</u> | <u>001</u> | Jun 27, 2007 |
| <u>AP</u> | TEVA | <u>EQ 2MG BASE/ML</u> | <u>A076876</u> | <u>001</u> | Nov 22, 2006 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>EQ 2MG BASE/ML</u> | <u>A076967</u> | <u>001</u> | Dec 26, 2006 |
| <u>AP</u> | | <u>EQ 2MG BASE/ML</u> | <u>A077365</u> | <u>001</u> | Dec 26, 2006 |
| <u>AP</u> | WOCKHARDT | <u>EQ 2MG BASE/ML</u> | <u>A077577</u> | <u>001</u> | Dec 26, 2006 |

ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

| | | | | | |
|-----------|---------------------------|-----------------------|----------------|------------|--------------|
| <u>AP</u> | ACCORD HLTHCARE | <u>EQ 2MG BASE/ML</u> | <u>A206845</u> | <u>001</u> | Mar 10, 2016 |
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>EQ 2MG BASE/ML</u> | <u>A202600</u> | <u>001</u> | Dec 21, 2012 |
| <u>AP</u> | ! BAXTER HLTHCARE CORP | <u>EQ 2MG BASE/ML</u> | <u>A078287</u> | <u>001</u> | Feb 22, 2013 |
| <u>AP</u> | EMCURE PHARMS LTD | <u>EQ 2MG BASE/ML</u> | <u>A078945</u> | <u>001</u> | Jan 03, 2013 |
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 2MG BASE/ML</u> | <u>A076972</u> | <u>001</u> | Dec 26, 2006 |
| <u>AP</u> | | <u>EQ 2MG BASE/ML</u> | <u>A202253</u> | <u>001</u> | Jul 19, 2013 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>EQ 2MG BASE/ML</u> | <u>A076780</u> | <u>001</u> | Dec 26, 2006 |
| <u>AP</u> | HOSPIRA | <u>EQ 2MG BASE/ML</u> | <u>A077548</u> | <u>001</u> | Dec 26, 2006 |
| <u>AP</u> | LUITPOLD | <u>EQ 2MG BASE/ML</u> | <u>A079032</u> | <u>001</u> | Nov 18, 2008 |
| <u>AP</u> | SANDOZ INC | <u>EQ 2MG BASE/ML</u> | <u>A077551</u> | <u>001</u> | Jun 27, 2007 |
| <u>AP</u> | TEVA | <u>EQ 2MG BASE/ML</u> | <u>A076759</u> | <u>001</u> | Nov 22, 2006 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>EQ 2MG BASE/ML</u> | <u>A077011</u> | <u>001</u> | Dec 26, 2006 |
| <u>AP</u> | | <u>EQ 2MG BASE/ML</u> | <u>A077541</u> | <u>001</u> | Dec 26, 2006 |
| <u>AP</u> | WOCKHARDT | <u>EQ 2MG BASE/ML</u> | <u>A077716</u> | <u>001</u> | Dec 26, 2006 |

SOLUTION;ORAL

ONDANSETRON HYDROCHLORIDE

| | | | | | |
|-----------|-------------------------|------------------------|----------------|------------|--------------|
| <u>AA</u> | AMNEAL PHARMS | <u>EQ 4MG BASE/5ML</u> | <u>A091483</u> | <u>001</u> | Jan 31, 2011 |
| <u>AA</u> | APOTEX | <u>EQ 4MG BASE/5ML</u> | <u>A078127</u> | <u>001</u> | Jun 25, 2007 |
| <u>AA</u> | AUROBINDO PHARMA | <u>EQ 4MG BASE/5ML</u> | <u>A078776</u> | <u>001</u> | Nov 28, 2007 |
| <u>AA</u> | LANNETT CO INC | <u>EQ 4MG BASE/5ML</u> | <u>A091342</u> | <u>001</u> | Jan 27, 2011 |
| <u>AA</u> | TARO | <u>EQ 4MG BASE/5ML</u> | <u>A077009</u> | <u>001</u> | Nov 30, 2007 |
| <u>AA</u> | WEST-WARD PHARMS INT | <u>EQ 4MG BASE/5ML</u> | <u>A076960</u> | <u>001</u> | Dec 26, 2006 |

PRESCRIPTION DRUG PRODUCT LIST

ONDANSETRON HYDROCHLORIDE

SOLUTION; ORAL

ZOFRAN

| | | | | | | |
|-----------|------------|-------------------------|-------------------------------|-----------------------|-------------------|--------------|
| AA | + ! | NOVARTIS PHARMS CORP | <u>EQ 4MG BASE/5ML</u> | <u>N020605</u> | <u>001</u> | Jan 24, 1997 |
|-----------|------------|-------------------------|-------------------------------|-----------------------|-------------------|--------------|

TABLET; ORAL

ONDANSETRON HYDROCHLORIDE

| | | | | | | |
|-----------|--|---------------------|----------------------------|-----------------------|-------------------|--------------|
| AB | | APOTEX | <u>EQ 4MG BASE</u> | <u>A077306</u> | <u>001</u> | Jun 25, 2007 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A077306</u> | <u>002</u> | Jun 25, 2007 |
| AB | | AUROBINDO PHARMA | <u>EQ 4MG BASE</u> | <u>A078539</u> | <u>001</u> | Jul 31, 2007 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A078539</u> | <u>002</u> | Jul 31, 2007 |
| AB | | | <u>EQ 24MG BASE</u> | <u>A078539</u> | <u>003</u> | Jul 31, 2007 |
| AB | | CASI PHARMS INC | <u>EQ 4MG BASE</u> | <u>A077517</u> | <u>001</u> | Jun 25, 2007 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A077517</u> | <u>002</u> | Jun 25, 2007 |
| AB | | | <u>EQ 24MG BASE</u> | <u>A077517</u> | <u>003</u> | Jun 25, 2007 |
| AB | | DR REDDYS LABS LTD | <u>EQ 4MG BASE</u> | <u>A076183</u> | <u>003</u> | Dec 26, 2006 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A076183</u> | <u>002</u> | Dec 26, 2006 |
| AB | | | <u>EQ 24MG BASE</u> | <u>A076183</u> | <u>001</u> | Dec 26, 2006 |
| AB | | GLENMARK GENERICS | <u>EQ 4MG BASE</u> | <u>A077535</u> | <u>001</u> | Jun 25, 2007 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A077535</u> | <u>002</u> | Jun 25, 2007 |
| AB | | | <u>EQ 24MG BASE</u> | <u>A077535</u> | <u>003</u> | Jun 25, 2007 |
| AB | | IPCA LABS LTD | <u>EQ 4MG BASE</u> | <u>A203761</u> | <u>001</u> | Jan 23, 2014 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A203761</u> | <u>002</u> | Jan 23, 2014 |
| AB | | MYLAN | <u>EQ 4MG BASE</u> | <u>A076930</u> | <u>001</u> | Jun 25, 2007 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A076930</u> | <u>002</u> | Jun 25, 2007 |
| AB | | | <u>EQ 24MG BASE</u> | <u>A076930</u> | <u>004</u> | Jun 25, 2007 |
| AB | | NATCO PHARMA LTD | <u>EQ 4MG BASE</u> | <u>A077851</u> | <u>001</u> | Jun 25, 2007 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A077851</u> | <u>002</u> | Jun 25, 2007 |
| AB | | PLIVA HRVATSKA DOO | <u>EQ 4MG BASE</u> | <u>A077112</u> | <u>001</u> | Jun 25, 2007 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A077112</u> | <u>002</u> | Jun 25, 2007 |
| AB | | | <u>EQ 24MG BASE</u> | <u>A077112</u> | <u>003</u> | Jun 25, 2007 |
| AB | | SUN PHARM INDS (IN) | <u>EQ 4MG BASE</u> | <u>A077050</u> | <u>001</u> | Jun 25, 2007 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A077050</u> | <u>002</u> | Jun 25, 2007 |
| AB | | TEVA | <u>EQ 4MG BASE</u> | <u>A076252</u> | <u>001</u> | Jun 25, 2007 |
| AB | | | <u>EQ 8MG BASE</u> | <u>A076252</u> | <u>002</u> | Jun 25, 2007 |
| AB | | | <u>EQ 24MG BASE</u> | <u>A076252</u> | <u>003</u> | Jun 25, 2007 |

ZOFRAN

| | | | | | | |
|-----------|----------|-------------------------|---------------------------|-----------------------|-------------------|--------------|
| AB | + | NOVARTIS PHARMS CORP | <u>EQ 4MG BASE</u> | <u>N020103</u> | <u>001</u> | Dec 31, 1992 |
|-----------|----------|-------------------------|---------------------------|-----------------------|-------------------|--------------|

| | | | | | | |
|-----------|----------|--|---------------------------|-----------------------|-------------------|--------------|
| AB | + | | <u>EQ 8MG BASE</u> | <u>N020103</u> | <u>002</u> | Dec 31, 1992 |
|-----------|----------|--|---------------------------|-----------------------|-------------------|--------------|

| | | | | | | |
|-----------|------------|--|----------------------------|-----------------------|-------------------|--------------|
| AB | + ! | | <u>EQ 24MG BASE</u> | <u>N020103</u> | <u>003</u> | Aug 27, 1999 |
|-----------|------------|--|----------------------------|-----------------------|-------------------|--------------|

ONDANSETRON HYDROCHLORIDE

| | | | | | | |
|--|--|--------------------|----------------------------|-----------------------|-------------------|--------------|
| | | DR REDDYS LABS LTD | <u>EQ 16MG BASE</u> | <u>A076183</u> | <u>004</u> | Dec 26, 2006 |
|--|--|--------------------|----------------------------|-----------------------|-------------------|--------------|

ORITAVANCIN DIPHOSPHATE

POWDER; INTRAVENOUS

ORBACTIV

| | | | | | |
|------------|----------------|----------------------------------|-----------------------|-------------------|--------------|
| + ! | MELINTA THERAP | <u>EQ 400MG BASE/VIAL</u> | <u>N206334</u> | <u>001</u> | Aug 06, 2014 |
|------------|----------------|----------------------------------|-----------------------|-------------------|--------------|

ORLISTAT

CAPSULE; ORAL

XENICAL

| | | | | | |
|------------|-------------|---------------------|-----------------------|-------------------|--------------|
| + ! | CHEPLAPHARM | <u>120MG</u> | <u>N020766</u> | <u>001</u> | Apr 23, 1999 |
|------------|-------------|---------------------|-----------------------|-------------------|--------------|

ORPHENADRINE CITRATE

INJECTABLE; INJECTION

ORPHENADRINE CITRATE

| | | | | | | |
|-----------|----------|-------------------------|-----------------------|-----------------------|-------------------|--------------|
| AP | ! | AKORN | <u>30MG/ML</u> | <u>A040484</u> | <u>001</u> | May 24, 2006 |
| AP | | SAGENT PHARMS | <u>30MG/ML</u> | <u>A090585</u> | <u>001</u> | Aug 30, 2011 |
| AP | | WATSON LABS | <u>30MG/ML</u> | <u>A084779</u> | <u>001</u> | Mar 15, 1982 |
| AP | | WEST-WARD PHARMS INT | <u>30MG/ML</u> | <u>A040463</u> | <u>001</u> | Mar 04, 2003 |

TABLET, EXTENDED RELEASE; ORAL

ORPHENADRINE CITRATE

| | | | | | | |
|-----------|----------|-----------------|---------------------|-----------------------|-------------------|--------------|
| AB | | ANDA REPOSITORY | <u>100MG</u> | <u>A040249</u> | <u>001</u> | Jan 29, 1999 |
| AB | | GAVIS PHARMS | <u>100MG</u> | <u>A040284</u> | <u>001</u> | Jun 19, 1998 |
| AB | | IMPAX PHARMS | <u>100MG</u> | <u>A040368</u> | <u>001</u> | Jun 23, 2000 |
| AB | | INVAGEN PHARMS | <u>100MG</u> | <u>A091158</u> | <u>001</u> | Jul 27, 2012 |
| AB | ! | SANDOZ | <u>100MG</u> | <u>A040327</u> | <u>001</u> | Feb 15, 2000 |

PRESCRIPTION DRUG PRODUCT LIST

OSELTAMIVIR PHOSPHATE

CAPSULE; ORAL

OSELTAMIVIR PHOSPHATE

| | | | | |
|-----------|---------------------|---------------------|--------------------|--------------|
| <u>AB</u> | AMNEAL PHARMS | <u>EQ 30MG BASE</u> | <u>A209093 001</u> | May 17, 2017 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A209093 002</u> | May 17, 2017 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A209093 003</u> | May 17, 2017 |
| <u>AB</u> | HETERO LABS LTD III | <u>EQ 30MG BASE</u> | <u>A209438 001</u> | Feb 23, 2018 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A209438 002</u> | Feb 23, 2018 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A209438 003</u> | Feb 23, 2018 |
| <u>AB</u> | LUPIN ATLANTIS | <u>EQ 30MG BASE</u> | <u>A208348 001</u> | Jan 09, 2018 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A208348 002</u> | Jan 09, 2018 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A208348 003</u> | Jan 09, 2018 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>EQ 30MG BASE</u> | <u>A207211 001</u> | Sep 14, 2017 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A207211 002</u> | Sep 14, 2017 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A207211 003</u> | Sep 14, 2017 |
| <u>AB</u> | NATCO PHARMA LTD | <u>EQ 30MG BASE</u> | <u>A202595 001</u> | Aug 03, 2016 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A202595 002</u> | Aug 03, 2016 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A202595 003</u> | Aug 03, 2016 |
| <u>AB</u> | NESHER PHARMS | <u>EQ 30MG BASE</u> | <u>A208578 001</u> | Feb 24, 2017 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A208578 002</u> | Feb 24, 2017 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A208578 003</u> | Feb 24, 2017 |
| <u>AB</u> | STRIDES PHARMA | <u>EQ 30MG BASE</u> | <u>A209421 001</u> | Jun 08, 2018 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A209421 002</u> | Jun 08, 2018 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A209421 003</u> | Jun 08, 2018 |

TAMIFLU

| | | | | |
|-----------|---------|---------------------|--------------------|--------------|
| <u>AB</u> | + ROCHE | <u>EQ 30MG BASE</u> | <u>N021087 003</u> | Jul 02, 2007 |
| <u>AB</u> | + | <u>EQ 45MG BASE</u> | <u>N021087 002</u> | Jul 02, 2007 |
| <u>AB</u> | +! | <u>EQ 75MG BASE</u> | <u>N021087 001</u> | Oct 27, 1999 |

FOR SUSPENSION; ORAL

OSELTAMIVIR PHOSPHATE

| | | | | |
|-----------|--------------------|-----------------------|--------------------|--------------|
| <u>AB</u> | ALVOGEN PINE BROOK | <u>EQ 6MG BASE/ML</u> | <u>A208823 001</u> | Oct 31, 2017 |
| <u>AB</u> | AMNEAL PHARMS NY | <u>EQ 6MG BASE/ML</u> | <u>A210186 001</u> | Feb 27, 2018 |
| <u>AB</u> | LUPIN ATLANTIS | <u>EQ 6MG BASE/ML</u> | <u>A208347 001</u> | Feb 20, 2018 |
| <u>AB</u> | NESHER PHARMS | <u>EQ 6MG BASE/ML</u> | <u>A209113 001</u> | Sep 14, 2017 |

TAMIFLU

| | | | | |
|-----------|----------|-----------------------|--------------------|--------------|
| <u>AB</u> | +! ROCHE | <u>EQ 6MG BASE/ML</u> | <u>N021246 002</u> | Mar 21, 2011 |
|-----------|----------|-----------------------|--------------------|--------------|

OSIMERTINIB MESYLATE

TABLET; ORAL

TAGRISSO

| | | | | |
|---|--------------------|--------------|-------------|--------------|
| + | ASTRAZENECA PHARMS | EQ 40MG BASE | N208065 001 | Nov 13, 2015 |
| + | ! | EQ 80MG BASE | N208065 002 | Nov 13, 2015 |

OSPEMIFENE

TABLET; ORAL

OSPHENA

| | | | | | |
|---|---|-----------|------|-------------|--------------|
| + | ! | DUCHESNAY | 60MG | N203505 001 | Feb 26, 2013 |
|---|---|-----------|------|-------------|--------------|

OXACILLIN SODIUM

INJECTABLE; INJECTION

OXACILLIN SODIUM

| | | | | | |
|--------------------------------|----------------------|--------------------------|--------------------|--------------|--------------|
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>EQ 1GM BASE/VIAL</u> | <u>A201539 001</u> | Jan 18, 2013 | |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A201539 002</u> | Jan 18, 2013 | |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A201538 001</u> | Jan 18, 2013 | |
| <u>AP</u> | HOSPIRA INC | <u>EQ 1GM BASE/VIAL</u> | <u>A203950 001</u> | Dec 11, 2015 | |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A203950 002</u> | Dec 11, 2015 | |
| <u>AP</u> | RENAISSANCE SSA LLC | <u>EQ 1GM BASE/VIAL</u> | <u>A206681 001</u> | Sep 11, 2017 | |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A206681 002</u> | Sep 11, 2017 | |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A206760 001</u> | Oct 26, 2017 | |
| <u>AP</u> | ! SAGENT PHARMS | <u>EQ 1GM BASE/VIAL</u> | <u>A091246 001</u> | Mar 30, 2012 | |
| <u>AP</u> | ! | <u>EQ 2GM BASE/VIAL</u> | <u>A091246 002</u> | Mar 30, 2012 | |
| <u>AP</u> | ! | <u>EQ 10GM BASE/VIAL</u> | <u>A091245 001</u> | Mar 30, 2012 | |
| <u>AP</u> | WOCKHARDT BIO AG | <u>EQ 1GM BASE/VIAL</u> | <u>A207147 001</u> | Jul 31, 2017 | |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL</u> | <u>A207147 002</u> | Jul 31, 2017 | |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A207148 001</u> | Nov 24, 2017 | |
| BACTOCILL IN PLASTIC CONTAINER | | | | | |
| + | ! | BAXTER HLTHCARE | EQ 20MG BASE/ML | N050640 001 | Oct 26, 1989 |
| + | ! | | EQ 40MG BASE/ML | N050640 002 | Oct 26, 1989 |

PRESCRIPTION DRUG PRODUCT LIST

OXALIPLATIN

INJECTABLE; IV (INFUSION)

ELOXATIN

| | | | | | |
|-----------|------------|-------------------|----------------------------|--------------------|--------------|
| <u>AP</u> | <u>+</u> ! | SANOFI AVENTIS US | <u>50MG/10ML (5MG/ML)</u> | <u>N021759 001</u> | Jan 31, 2005 |
| <u>AP</u> | <u>+</u> ! | | <u>100MG/20ML (5MG/ML)</u> | <u>N021759 002</u> | Jan 31, 2005 |

OXALIPLATIN

| | | | | | |
|-----------|------------|---------------------|----------------------------|--------------------|--------------|
| <u>AP</u> | | ACCORD HLTHCARE | <u>50MG/10ML (5MG/ML)</u> | <u>A207474 001</u> | Mar 21, 2017 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A207474 002</u> | Mar 21, 2017 |
| <u>AP</u> | | | <u>200MG/40ML (5MG/ML)</u> | <u>A207474 003</u> | Mar 21, 2017 |
| <u>AP</u> | | ACTAVIS LLC | <u>50MG/10ML (5MG/ML)</u> | <u>A204880 001</u> | Mar 05, 2018 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A204880 002</u> | Mar 05, 2018 |
| <u>AP</u> | | ACTAVIS TOTOWA | <u>50MG/VIAL</u> | <u>A078803 001</u> | Aug 08, 2012 |
| <u>AP</u> | | | <u>100MG/VIAL</u> | <u>A078803 002</u> | Aug 08, 2012 |
| <u>AP</u> | | CIPLA | <u>50MG/10ML (5MG/ML)</u> | <u>A208523 001</u> | Feb 10, 2017 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A208523 002</u> | Feb 10, 2017 |
| <u>AP</u> | | EUGIA PHARMA | <u>50MG/10ML (5MG/ML)</u> | <u>A205529 001</u> | Sep 06, 2017 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A205529 002</u> | Sep 06, 2017 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>50MG/10ML (5MG/ML)</u> | <u>A078811 001</u> | Jun 10, 2010 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A078811 002</u> | Jun 10, 2010 |
| <u>AP</u> | | | <u>50MG/VIAL</u> | <u>A078819 001</u> | Jun 02, 2010 |
| <u>AP</u> | | | <u>50MG/10ML (5MG/ML)</u> | <u>A090030 001</u> | Jan 31, 2017 |
| <u>AP</u> | | | <u>100MG/VIAL</u> | <u>A078819 002</u> | Jun 02, 2010 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A090030 002</u> | Jan 31, 2017 |
| <u>AP</u> | | | <u>200MG/40ML (5MG/ML)</u> | <u>A090030 003</u> | Jan 31, 2017 |
| <u>AP</u> | | GLAND PHARMA LTD | <u>50MG/10ML (5MG/ML)</u> | <u>A207325 001</u> | Feb 10, 2017 |
| <u>AP</u> | | | <u>50MG/VIAL</u> | <u>A207385 001</u> | May 23, 2017 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A207325 002</u> | Feb 10, 2017 |
| <u>AP</u> | | | <u>100MG/VIAL</u> | <u>A207385 002</u> | May 23, 2017 |
| <u>AP</u> | | | <u>200MG/40ML (5MG/ML)</u> | <u>A207325 003</u> | Oct 18, 2017 |
| <u>AP</u> | | HOSPIRA INC | <u>50MG/VIAL</u> | <u>A078815 001</u> | Sep 30, 2009 |
| <u>AP</u> | | | <u>100MG/VIAL</u> | <u>A078815 002</u> | Sep 30, 2009 |
| <u>AP</u> | | HOSPIRA WORLDWIDE | <u>50MG/10ML (5MG/ML)</u> | <u>A078813 001</u> | Aug 07, 2009 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A078813 002</u> | Aug 07, 2009 |
| <u>AP</u> | | INGENUS PHARMS LLC | <u>50MG/10ML (5MG/ML)</u> | <u>A207562 001</u> | Oct 16, 2018 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A207562 002</u> | Oct 16, 2018 |
| <u>AP</u> | | JIANGSU HENGRUI MED | <u>50MG/10ML (5MG/ML)</u> | <u>A203869 001</u> | Jun 18, 2014 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A203869 002</u> | Jun 18, 2014 |
| <u>AP</u> | | LUITPOLD | <u>50MG/10ML (5MG/ML)</u> | <u>A204378 001</u> | May 12, 2017 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A204378 002</u> | May 12, 2017 |
| <u>AP</u> | | MYLAN LABS LTD | <u>50MG/10ML (5MG/ML)</u> | <u>A091358 001</u> | Aug 07, 2012 |
| <u>AP</u> | | | <u>50MG/VIAL</u> | <u>A200979 001</u> | Aug 08, 2012 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A091358 002</u> | Aug 07, 2012 |
| <u>AP</u> | | | <u>100MG/VIAL</u> | <u>A200979 002</u> | Aug 08, 2012 |
| <u>AP</u> | | | <u>200MG/40ML (5MG/ML)</u> | <u>A091358 003</u> | Nov 14, 2017 |
| <u>AP</u> | | QILU PHARM CO LTD | <u>50MG/10ML (5MG/ML)</u> | <u>A204368 001</u> | Jun 07, 2016 |
| <u>AP</u> | | | <u>50MG/VIAL</u> | <u>A204616 001</u> | May 11, 2016 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A204368 002</u> | Jun 07, 2016 |
| <u>AP</u> | | | <u>100MG/VIAL</u> | <u>A204616 002</u> | May 11, 2016 |
| <u>AP</u> | <u>!</u> | | <u>200MG/40ML (5MG/ML)</u> | <u>A204368 003</u> | Jun 07, 2016 |
| <u>AP</u> | | SANDOZ | <u>50MG/10ML (5MG/ML)</u> | <u>A078817 001</u> | Jan 24, 2011 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A078817 002</u> | Jan 24, 2011 |
| <u>AP</u> | <u>!</u> | SUN PHARMA GLOBAL | <u>50MG/VIAL</u> | <u>A078818 001</u> | Aug 07, 2009 |
| <u>AP</u> | | | <u>50MG/10ML (5MG/ML)</u> | <u>A202922 001</u> | Apr 08, 2014 |
| <u>AP</u> | <u>!</u> | | <u>100MG/VIAL</u> | <u>A078818 002</u> | Aug 07, 2009 |
| <u>AP</u> | | | <u>100MG/20ML (5MG/ML)</u> | <u>A202922 002</u> | Apr 08, 2014 |
| <u>AP</u> | <u>+</u> ! | TEVA PHARMS | <u>50MG/10ML (5MG/ML)</u> | <u>N022160 001</u> | Aug 07, 2009 |
| <u>AP</u> | <u>+</u> ! | | <u>100MG/20ML (5MG/ML)</u> | <u>N022160 002</u> | Aug 07, 2009 |

OXANDROLONE

TABLET; ORAL

OXANDROLONE

| | | | | | |
|-----------|----------|-------------------|--------------|--------------------|--------------|
| <u>AB</u> | | PAR PHARM | <u>2.5MG</u> | <u>A077827 001</u> | Jun 22, 2007 |
| <u>AB</u> | <u>!</u> | | <u>10MG</u> | <u>A077827 002</u> | Jun 22, 2007 |
| <u>AB</u> | | UPSHER SMITH LABS | <u>2.5MG</u> | <u>A076761 001</u> | Dec 01, 2006 |
| <u>AB</u> | | | <u>10MG</u> | <u>A078033 001</u> | Mar 22, 2007 |

OXAPROZIN

TABLET; ORAL

DAYPRO

| | | | | | |
|-----------|------------|-----------|--------------|--------------------|--------------|
| <u>AB</u> | <u>+</u> ! | GD SEARLE | <u>600MG</u> | <u>N018841 004</u> | Oct 29, 1992 |
|-----------|------------|-----------|--------------|--------------------|--------------|

OXAPROZIN

| | | | | | |
|-----------|--|------------------|--------------|--------------------|--------------|
| <u>AB</u> | | AMNEAL PHARMS CO | <u>600MG</u> | <u>A208633 001</u> | May 04, 2017 |
| <u>AB</u> | | APOTEX INC | <u>600MG</u> | <u>A075987 001</u> | Sep 02, 2004 |

PRESCRIPTION DRUG PRODUCT LIST

OXAPROZIN

TABLET; ORAL

OXAPROZIN

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | DR REDDYS LABS LTD | <u>600MG</u> | <u>A075855</u> | <u>001</u> | Jan 31, 2001 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>600MG</u> | <u>A075846</u> | <u>001</u> | May 13, 2002 |
| <u>AB</u> | SANDOZ | <u>600MG</u> | <u>A075845</u> | <u>001</u> | Jan 31, 2001 |
| <u>AB</u> | SUN PHARM INDS INC | <u>600MG</u> | <u>A075844</u> | <u>001</u> | Jan 03, 2002 |
| <u>AB</u> | TEVA | <u>600MG</u> | <u>A075849</u> | <u>001</u> | Jul 03, 2002 |

OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

| | | | | | |
|-----------|-------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | ACTAVIS ELIZABETH | <u>10MG</u> | <u>A072253</u> | <u>002</u> | Apr 14, 1988 |
| <u>AB</u> | | <u>15MG</u> | <u>A072253</u> | <u>003</u> | Apr 14, 1988 |
| <u>AB</u> | ! | <u>30MG</u> | <u>A072253</u> | <u>001</u> | Apr 14, 1988 |
| <u>AB</u> | SANDOZ | <u>10MG</u> | <u>A071813</u> | <u>001</u> | Apr 19, 1988 |
| <u>AB</u> | | <u>15MG</u> | <u>A071756</u> | <u>001</u> | Apr 19, 1988 |
| <u>AB</u> | | <u>30MG</u> | <u>A071814</u> | <u>001</u> | Apr 19, 1988 |

OXCARBAZEPINE

SUSPENSION; ORAL

OXCARBAZEPINE

| | | | | | |
|-----------|-------------------------|------------------|----------------|------------|--------------|
| <u>AB</u> | AMNEAL PHARMS | <u>300MG/5ML</u> | <u>A202961</u> | <u>001</u> | Sep 17, 2012 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>300MG/5ML</u> | <u>A078734</u> | <u>001</u> | Jun 26, 2009 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>300MG/5ML</u> | <u>A201193</u> | <u>001</u> | Oct 03, 2012 |

TRILEPTAL

| | | | | | | |
|-----------|----|----------|------------------|----------------|------------|--------------|
| <u>AB</u> | +! | NOVARTIS | <u>300MG/5ML</u> | <u>N021285</u> | <u>001</u> | May 25, 2001 |
|-----------|----|----------|------------------|----------------|------------|--------------|

TABLET; ORAL

OXCARBAZEPINE

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ANI PHARMS INC | <u>150MG</u> | <u>A078005</u> | <u>001</u> | Dec 11, 2007 |
| <u>AB</u> | | <u>300MG</u> | <u>A078005</u> | <u>002</u> | Dec 11, 2007 |
| <u>AB</u> | | <u>600MG</u> | <u>A078005</u> | <u>003</u> | Dec 11, 2007 |
| <u>AB</u> | APOTEX INC | <u>150MG</u> | <u>A077747</u> | <u>001</u> | Apr 09, 2008 |
| <u>AB</u> | | <u>300MG</u> | <u>A077747</u> | <u>002</u> | Apr 09, 2008 |
| <u>AB</u> | | <u>600MG</u> | <u>A077747</u> | <u>003</u> | Apr 09, 2008 |
| <u>AB</u> | BRECKENRIDGE PHARM | <u>150MG</u> | <u>A078069</u> | <u>001</u> | Jan 11, 2008 |
| <u>AB</u> | | <u>300MG</u> | <u>A078069</u> | <u>002</u> | Jan 11, 2008 |
| <u>AB</u> | | <u>600MG</u> | <u>A078069</u> | <u>003</u> | Jan 11, 2008 |
| <u>AB</u> | GLENMARK PHARMS LTD | <u>150MG</u> | <u>A077802</u> | <u>001</u> | Oct 09, 2007 |
| <u>AB</u> | | <u>300MG</u> | <u>A077802</u> | <u>002</u> | Oct 09, 2007 |
| <u>AB</u> | | <u>600MG</u> | <u>A077802</u> | <u>003</u> | Oct 09, 2007 |
| <u>AB</u> | SUN PHARM INDS | <u>150MG</u> | <u>A077794</u> | <u>001</u> | Oct 09, 2007 |
| <u>AB</u> | | <u>300MG</u> | <u>A077794</u> | <u>002</u> | Oct 09, 2007 |
| <u>AB</u> | | <u>600MG</u> | <u>A077794</u> | <u>003</u> | Oct 09, 2007 |
| <u>AB</u> | TARO | <u>150MG</u> | <u>A077801</u> | <u>001</u> | Nov 15, 2007 |
| <u>AB</u> | | <u>300MG</u> | <u>A077801</u> | <u>002</u> | Nov 15, 2007 |
| <u>AB</u> | | <u>600MG</u> | <u>A077801</u> | <u>003</u> | Nov 15, 2007 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>150MG</u> | <u>A077795</u> | <u>001</u> | Oct 09, 2007 |
| <u>AB</u> | | <u>300MG</u> | <u>A077795</u> | <u>002</u> | Oct 09, 2007 |
| <u>AB</u> | | <u>600MG</u> | <u>A077795</u> | <u>003</u> | Oct 09, 2007 |

TRILEPTAL

| | | | | | | |
|-----------|---|----------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | NOVARTIS | <u>150MG</u> | <u>N021014</u> | <u>001</u> | Jan 14, 2000 |
| <u>AB</u> | + | | <u>300MG</u> | <u>N021014</u> | <u>002</u> | Jan 14, 2000 |
| <u>AB</u> | + | ! | <u>600MG</u> | <u>N021014</u> | <u>003</u> | Jan 14, 2000 |

TABLET, EXTENDED RELEASE; ORAL

OXTELLAR XR

| | | | | | |
|---|-----------------|-------|---------|-----|--------------|
| + | SUPERNUS PHARMS | 150MG | N202810 | 001 | Oct 19, 2012 |
| + | | 300MG | N202810 | 002 | Oct 19, 2012 |
| + | ! | 600MG | N202810 | 003 | Oct 19, 2012 |

OXICONAZOLE NITRATE

CREAM; TOPICAL

OXICONAZOLE NITRATE

| | | | | | |
|-----------|-------------|-------------------|----------------|------------|--------------|
| <u>AB</u> | TARO PHARMS | <u>EQ 1% BASE</u> | <u>A205076</u> | <u>001</u> | Mar 07, 2016 |
|-----------|-------------|-------------------|----------------|------------|--------------|

OXISTAT

| | | | | | | | |
|-----------|---|---|----------------|-------------------|----------------|------------|--------------|
| <u>AB</u> | + | ! | FOUGERA PHARMS | <u>EQ 1% BASE</u> | <u>N019828</u> | <u>001</u> | Dec 30, 1988 |
|-----------|---|---|----------------|-------------------|----------------|------------|--------------|

LOTION; TOPICAL

OXISTAT

| | | | | | | |
|---|---|----------------|------------|---------|-----|--------------|
| + | ! | FOUGERA PHARMS | EQ 1% BASE | N020209 | 001 | Sep 30, 1992 |
|---|---|----------------|------------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

OXYBUTYNIN

FILM, EXTENDED RELEASE;TRANSDERMAL

OXYTROL

+! ALLERGAN SALES LLC 3.9MG/24HR N021351 002 Feb 26, 2003

OXYBUTYNIN CHLORIDE

GEL;TRANSDERMAL

GELNIQUE**AB** +! ALLERGAN SALES LLC 10% (100MG/PACKET) **N022204 001** Jan 27, 2009OXYBUTYNIN CHLORIDE**AB** PAR PHARM INC 10% (100MG/PACKET) **A207329 001** May 31, 2018

SYRUP;ORAL

OXYBUTYNIN CHLORIDE**AA** LANNETT CO INC 5MG/5ML **A074520 001** Mar 29, 1996**AA** 5MG/5ML **A076682 001** Dec 28, 2004**AA** PHARM ASSOC 5MG/5ML **A075137 001** Dec 18, 1998**AA** ! WOCKHARDT BIO AG 5MG/5ML **A074868 001** Feb 12, 1997

TABLET;ORAL

OXYBUTYNIN CHLORIDE**AB** ABHAI LLC 5MG **A209335 001** Dec 22, 2017**AB** APPCO PHARMA LLC 5MG **A209025 001** Dec 21, 2017**AB** NOVITIUM PHARMA 5MG **A209823 001** Oct 23, 2017**AB** TEVA PHARMS USA 5MG **A071655 001** Nov 14, 1988**AB** TULEX PHARMS INC 5MG **A210125 001** Sep 06, 2018**AB** UPSHER SMITH LABS 5MG **A074625 001** Jul 31, 1996**AB** ! VINTAGE PHARMS 5MG **A075079 001** Oct 31, 1997

TABLET, EXTENDED RELEASE;ORAL

DITROPAN XL**AB** + JANSSEN PHARMS 5MG **N020897 001** Dec 16, 1998**AB** + 10MG **N020897 002** Dec 16, 1998OXYBUTYNIN CHLORIDE**AB** ACCORD HLTHCARE 5MG **A207138 001** Feb 29, 2016**AB** 10MG **A207138 002** Feb 29, 2016**AB** 15MG **A207138 003** Feb 29, 2016**AB** AMNEAL PHARMS 5MG **A204010 001** Nov 23, 2015**AB** 10MG **A204010 002** Nov 23, 2015**AB** 15MG **A204010 003** Nov 23, 2015**AB** IMPAX PHARMS 5MG **A076745 002** May 09, 2007**AB** 10MG **A076745 003** May 09, 2007**AB** 15MG **A076745 001** Nov 09, 2006**AB** MYLAN 5MG **A076702 001** Nov 09, 2006**AB** MYLAN PHARMS INC 10MG **A076644 001** Nov 09, 2006**AB** ! 15MG **A076644 002** May 10, 2007**AB** OSMOTICA PHARM US 5MG **A078503 001** Feb 04, 2009**AB** 10MG **A078503 002** Feb 04, 2009**AB** 15MG **A078503 003** Feb 04, 2009**AB** UNIQUE PHARM LABS 5MG **A206121 001** May 27, 2016**AB** 10MG **A206121 002** May 27, 2016**AB** 15MG **A206121 003** May 27, 2016**AB** ZYDUS PHARMS USA 5MG **A202332 001** Jun 26, 2017**AB** 10MG **A202332 002** Jun 26, 2017**AB** 15MG **A202332 003** Jun 26, 2017OXYCODONE

CAPSULE, EXTENDED RELEASE;ORAL

XTAMPZA ER

+ COLLEGIUM PHARM INC 9MG N208090 001 Apr 26, 2016

+ 13.5MG N208090 002 Apr 26, 2016

+ 18MG N208090 003 Apr 26, 2016

+ 27MG N208090 004 Apr 26, 2016

+! 36MG N208090 005 Apr 26, 2016

OXYCODONE HYDROCHLORIDE

CAPSULE;ORAL

OXYCODONE HYDROCHLORIDE**AB** ANI PHARMS INC 5MG **A205177 001** Mar 31, 2016**AB** AVANTHI INC 5MG **A202773 001** Aug 17, 2015**AB** +! GENUS LIFESCIENCES 5MG **N200534 001** Oct 20, 2010**AB** LANNETT CO INC 5MG **A203823 001** Aug 01, 2014**AB** MAYNE PHARMA INC 5MG **A203107 001** Jul 26, 2012**AB** NOVEL LABS INC 5MG **A204752 001** Aug 24, 2015

PRESCRIPTION DRUG PRODUCT LIST

OXYCODONE HYDROCHLORIDE

SOLUTION;ORAL

OXYCODONE HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|------------------|--------------------|--------------|
| <u>AA</u> | ABHAI LLC | <u>5MG/5ML</u> | <u>A208593 001</u> | Jul 21, 2017 |
| <u>AA</u> | | <u>100MG/5ML</u> | <u>A208593 002</u> | Jul 21, 2017 |
| <u>AA</u> | ANI PHARMS INC | <u>5MG/5ML</u> | <u>A204979 001</u> | Jun 01, 2015 |
| <u>AA</u> | | <u>100MG/5ML</u> | <u>A203447 001</u> | Aug 30, 2017 |
| <u>AA</u> | ASCENT PHARMS INC | <u>5MG/5ML</u> | <u>A209021 001</u> | Nov 09, 2017 |
| <u>AA</u> | | <u>100MG/5ML</u> | <u>A209021 002</u> | Nov 09, 2017 |
| <u>AA</u> | + GENUS LIFESCIENCES | <u>5MG/5ML</u> | <u>N200535 002</u> | Aug 22, 2013 |
| <u>AA</u> | +! | <u>100MG/5ML</u> | <u>N200535 001</u> | Oct 20, 2010 |
| <u>AA</u> | HI-TECH PHARMACAL | <u>5MG/5ML</u> | <u>A208817 001</u> | Aug 10, 2017 |
| <u>AA</u> | | <u>100MG/5ML</u> | <u>A208795 001</u> | Aug 07, 2017 |
| <u>AA</u> | LANNETT CO INC | <u>100MG/5ML</u> | <u>A204085 001</u> | Sep 09, 2014 |
| <u>AA</u> | MAYNE PHARMA INC | <u>100MG/5ML</u> | <u>A204092 001</u> | Jun 05, 2014 |
| <u>AA</u> | NOVEL LABS INC | <u>100MG/5ML</u> | <u>A204603 001</u> | Apr 29, 2015 |
| <u>AA</u> | PHARM ASSOC | <u>100MG/5ML</u> | <u>A206822 001</u> | Aug 15, 2017 |
| <u>AA</u> | SPECGX LLC | <u>5MG/5ML</u> | <u>A210758 001</u> | Apr 30, 2018 |
| <u>AA</u> | | <u>100MG/5ML</u> | <u>A210758 002</u> | Apr 30, 2018 |
| <u>AA</u> | +! VISTAPHARM | <u>5MG/5ML</u> | <u>N201194 001</u> | Jan 12, 2012 |
| <u>AA</u> | | <u>100MG/5ML</u> | <u>A202537 001</u> | Jul 30, 2012 |
| <u>AA</u> | WES PHARMA INC | <u>5MG/5ML</u> | <u>A207511 001</u> | Nov 23, 2016 |
| <u>AA</u> | | <u>100MG/5ML</u> | <u>A209897 001</u> | Sep 06, 2017 |
| <u>AA</u> | WEST-WARD PHARMS INT | <u>5MG/5ML</u> | <u>A204037 001</u> | Jul 15, 2013 |
| <u>AA</u> | | <u>100MG/5ML</u> | <u>A203208 001</u> | Jul 12, 2013 |
| <u>AA</u> | WOCKHARDT BIO AG | <u>5MG/5ML</u> | <u>A206456 001</u> | Jun 16, 2015 |

TABLET;ORAL

OXYCODONE HYDROCHLORIDE

| | | | | |
|-----------|---------------------|-------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS ELIZABETH | <u>5MG</u> | <u>A076636 003</u> | Apr 07, 2015 |
| <u>AB</u> | | <u>15MG</u> | <u>A076636 001</u> | Feb 06, 2004 |
| <u>AB</u> | | <u>30MG</u> | <u>A076636 002</u> | Feb 06, 2004 |
| <u>AB</u> | ALVOGEN MALTA | <u>5MG</u> | <u>A202116 001</u> | Dec 30, 2011 |
| <u>AB</u> | | <u>15MG</u> | <u>A202116 002</u> | Dec 30, 2011 |
| <u>AB</u> | | <u>30MG</u> | <u>A202116 003</u> | Dec 30, 2011 |
| <u>AB</u> | AMNEAL PHARMS | <u>5MG</u> | <u>A203638 001</u> | Jun 03, 2014 |
| <u>AB</u> | | <u>10MG</u> | <u>A203638 002</u> | Jun 03, 2014 |
| <u>AB</u> | | <u>15MG</u> | <u>A203638 003</u> | Jun 03, 2014 |
| <u>AB</u> | | <u>20MG</u> | <u>A203638 004</u> | Jun 03, 2014 |
| <u>AB</u> | | <u>30MG</u> | <u>A203638 005</u> | Jun 03, 2014 |
| <u>AB</u> | ASCENT PHARMS INC | <u>15MG</u> | <u>A207418 001</u> | Aug 07, 2017 |
| <u>AB</u> | | <u>30MG</u> | <u>A207418 002</u> | Aug 07, 2017 |
| <u>AB</u> | AUROLIFE PHARMA LLC | <u>5MG</u> | <u>A202160 001</u> | Nov 19, 2012 |
| <u>AB</u> | | <u>15MG</u> | <u>A202160 002</u> | Nov 19, 2012 |
| <u>AB</u> | | <u>30MG</u> | <u>A202160 003</u> | Nov 19, 2012 |
| <u>AB</u> | AVANTHI INC | <u>5MG</u> | <u>A091393 001</u> | Aug 31, 2009 |
| <u>AB</u> | | <u>10MG</u> | <u>A091393 002</u> | Aug 31, 2009 |
| <u>AB</u> | | <u>15MG</u> | <u>A091393 003</u> | Aug 31, 2009 |
| <u>AB</u> | | <u>20MG</u> | <u>A091393 004</u> | Aug 31, 2009 |
| <u>AB</u> | | <u>30MG</u> | <u>A091393 005</u> | Aug 31, 2009 |
| <u>AB</u> | EPIC PHARMA LLC | <u>5MG</u> | <u>A090895 001</u> | Aug 24, 2009 |
| <u>AB</u> | | <u>5MG</u> | <u>A202662 001</u> | Sep 22, 2015 |
| <u>AB</u> | | <u>10MG</u> | <u>A202662 002</u> | Sep 22, 2015 |
| <u>AB</u> | | <u>15MG</u> | <u>A090895 002</u> | Aug 24, 2009 |
| <u>AB</u> | | <u>15MG</u> | <u>A202662 003</u> | Sep 22, 2015 |
| <u>AB</u> | | <u>20MG</u> | <u>A202662 005</u> | Apr 27, 2017 |
| <u>AB</u> | | <u>30MG</u> | <u>A090895 003</u> | Aug 24, 2009 |
| <u>AB</u> | | <u>30MG</u> | <u>A202662 004</u> | Sep 22, 2015 |
| <u>AB</u> | MAYNE PHARMA INC | <u>5MG</u> | <u>A091313 001</u> | Feb 18, 2011 |
| <u>AB</u> | | <u>10MG</u> | <u>A091313 004</u> | Apr 29, 2016 |
| <u>AB</u> | | <u>15MG</u> | <u>A091313 002</u> | Feb 18, 2011 |
| <u>AB</u> | | <u>20MG</u> | <u>A091313 005</u> | Apr 29, 2016 |
| <u>AB</u> | | <u>30MG</u> | <u>A091313 003</u> | Feb 18, 2011 |
| <u>AB</u> | NESHER PHARMS | <u>5MG</u> | <u>A077290 001</u> | Dec 08, 2005 |
| <u>AB</u> | | <u>10MG</u> | <u>A077290 002</u> | Dec 08, 2005 |
| <u>AB</u> | | <u>15MG</u> | <u>A077290 003</u> | Dec 08, 2005 |
| <u>AB</u> | | <u>20MG</u> | <u>A077290 004</u> | Dec 08, 2005 |
| <u>AB</u> | | <u>30MG</u> | <u>A077290 005</u> | Dec 08, 2005 |
| <u>AB</u> | NOVEL LABS INC | <u>5MG</u> | <u>A204021 001</u> | Jun 12, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A204021 002</u> | Jun 12, 2017 |
| <u>AB</u> | | <u>15MG</u> | <u>A204021 003</u> | Jun 12, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A204021 004</u> | Jun 12, 2017 |
| <u>AB</u> | | <u>30MG</u> | <u>A204021 005</u> | Jun 12, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE HYDROCHLORIDE

| | | | | |
|-----------|--------------------|-------------|--------------------|--------------|
| <u>AB</u> | NUVO PHARM | <u>5MG</u> | <u>A207119 001</u> | Apr 12, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A207119 002</u> | Apr 12, 2016 |
| <u>AB</u> | | <u>15MG</u> | <u>A207119 003</u> | Apr 12, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A207119 004</u> | Apr 12, 2016 |
| <u>AB</u> | | <u>30MG</u> | <u>A207119 005</u> | Apr 12, 2016 |
| <u>AB</u> | RHODES PHARMS | <u>5MG</u> | <u>A091490 001</u> | Mar 09, 2011 |
| <u>AB</u> | | <u>10MG</u> | <u>A091490 002</u> | Mar 09, 2011 |
| <u>AB</u> | | <u>15MG</u> | <u>A091490 003</u> | Mar 09, 2011 |
| <u>AB</u> | | <u>20MG</u> | <u>A091490 004</u> | Mar 09, 2011 |
| <u>AB</u> | | <u>30MG</u> | <u>A091490 005</u> | Mar 09, 2011 |
| <u>AB</u> | SPECGX LLC | <u>5MG</u> | <u>A076758 003</u> | Mar 19, 2007 |
| <u>AB</u> | | <u>15MG</u> | <u>A076758 001</u> | Jun 30, 2004 |
| <u>AB</u> | | <u>30MG</u> | <u>A076758 002</u> | Jun 30, 2004 |
| <u>AB</u> | SUN PHARM INDS INC | <u>5MG</u> | <u>A090659 001</u> | Apr 10, 2009 |
| <u>AB</u> | | <u>10MG</u> | <u>A090659 005</u> | Nov 06, 2012 |
| <u>AB</u> | | <u>15MG</u> | <u>A090659 002</u> | Apr 10, 2009 |
| <u>AB</u> | | <u>20MG</u> | <u>A090659 004</u> | Nov 06, 2012 |
| <u>AB</u> | | <u>30MG</u> | <u>A090659 003</u> | Apr 10, 2009 |
| <u>AB</u> | VINTAGE PHARMS | <u>5MG</u> | <u>A077712 003</u> | Mar 02, 2009 |
| <u>AB</u> | | <u>10MG</u> | <u>A077712 004</u> | Apr 13, 2015 |
| <u>AB</u> | | <u>15MG</u> | <u>A077712 001</u> | Jan 31, 2007 |
| <u>AB</u> | | <u>20MG</u> | <u>A077712 005</u> | Apr 13, 2015 |
| <u>AB</u> | | <u>30MG</u> | <u>A077712 002</u> | Jan 31, 2007 |

ROXICODONE

| | | | | |
|-----------|--------------|-------------|--------------------|--------------|
| <u>AB</u> | + SPECGX LLC | <u>5MG</u> | <u>N021011 003</u> | May 15, 2009 |
| <u>AB</u> | +! | <u>15MG</u> | <u>N021011 001</u> | Aug 31, 2000 |
| <u>AB</u> | + | <u>30MG</u> | <u>N021011 002</u> | Aug 31, 2000 |

OXAYDO

| | | | | |
|--|---------------|-------|-------------|--------------|
| | EGALET US INC | 5MG | N202080 001 | Jun 17, 2011 |
| | | 7.5MG | N202080 002 | Jun 17, 2011 |

TABLET, EXTENDED RELEASE; ORAL

OXYCONTIN

| | | | | |
|---|------------------|------|-------------|--------------|
| + | PURDUE PHARMA LP | 10MG | N022272 001 | Apr 05, 2010 |
| + | | 15MG | N022272 002 | Apr 05, 2010 |
| + | | 20MG | N022272 003 | Apr 05, 2010 |
| + | | 30MG | N022272 004 | Apr 05, 2010 |
| + | ! | 40MG | N022272 005 | Apr 05, 2010 |
| + | | 60MG | N022272 006 | Apr 05, 2010 |
| + | | 80MG | N022272 007 | Apr 05, 2010 |

OXYMETAZOLINE HYDROCHLORIDE

CREAM; TOPICAL

RHOFADE

| | | | | |
|---|-----------|----|-------------|--------------|
| + | ! ACLARIS | 1% | N208552 001 | Jan 18, 2017 |
|---|-----------|----|-------------|--------------|

OXYMETAZOLINE HYDROCHLORIDE; TETRACAINE HYDROCHLORIDE

SPRAY, METERED; NASAL

KOVANAZE

| | | | | |
|---|--------------|------------------------|-------------|--------------|
| + | ! ST RENATUS | 0.1MG/SPRAY; 6MG/SPRAY | N208032 001 | Jun 29, 2016 |
|---|--------------|------------------------|-------------|--------------|

OXYMETHOLONE

TABLET; ORAL

ANADROL-50

| | | | | |
|---|-----------------------|------|-------------|--|
| + | ! MYLAN SPECIALITY LP | 50MG | N016848 001 | |
|---|-----------------------|------|-------------|--|

OXYMORPHONE HYDROCHLORIDE

TABLET; ORAL

OPANA

| | | | | |
|-----------|---------------|-------------|--------------------|--------------|
| <u>AB</u> | + ENDO PHARMS | <u>5MG</u> | <u>N021611 001</u> | Jun 22, 2006 |
| <u>AB</u> | +! | <u>10MG</u> | <u>N021611 002</u> | Jun 22, 2006 |

OXYMORPHONE HYDROCHLORIDE

| | | | | |
|-----------|---------------------|-------------|--------------------|--------------|
| <u>AB</u> | ASCENT PHARMS INC | <u>5MG</u> | <u>A210175 001</u> | Feb 02, 2018 |
| <u>AB</u> | | <u>10MG</u> | <u>A210175 002</u> | Feb 02, 2018 |
| <u>AB</u> | AUROLIFE PHARMA LLC | <u>5MG</u> | <u>A204459 001</u> | Apr 26, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A204459 002</u> | Apr 26, 2016 |
| <u>AB</u> | AVANTHI INC | <u>5MG</u> | <u>A203601 001</u> | Jan 30, 2013 |
| <u>AB</u> | | <u>10MG</u> | <u>A203601 002</u> | Jan 30, 2013 |
| <u>AB</u> | EPIC PHARMA LLC | <u>5MG</u> | <u>A201187 001</u> | Dec 15, 2014 |
| <u>AB</u> | | <u>10MG</u> | <u>A201187 002</u> | Dec 15, 2014 |
| <u>AB</u> | SPECGX LLC | <u>5MG</u> | <u>A202321 001</u> | Apr 25, 2013 |
| <u>AB</u> | | <u>10MG</u> | <u>A202321 002</u> | Apr 25, 2013 |

PRESCRIPTION DRUG PRODUCT LIST

OXYMORPHONE HYDROCHLORIDE

TABLET; ORAL

OXYMORPHONE HYDROCHLORIDE

| | | | | |
|-----------|------------------|-------------|--------------------|--------------|
| AB | TEVA | 5MG | A091443 002 | Feb 15, 2011 |
| AB | | 10MG | A091443 001 | Feb 15, 2011 |
| AB | WEST-WARD PHARMS | 5MG | A090964 001 | Sep 27, 2010 |
| | INT | | | |
| AB | | 10MG | A090964 002 | Sep 27, 2010 |

TABLET, EXTENDED RELEASE; ORAL

OXYMORPHONE HYDROCHLORIDE

| | | | | |
|-----------|-------------------|--------------|--------------------|--------------|
| AB | ACTAVIS ELIZABETH | 5MG | A079046 003 | Jul 11, 2013 |
| AB | | 7.5MG | A079046 001 | Dec 13, 2010 |
| AB | | 10MG | A079046 004 | Jul 11, 2013 |
| AB | | 15MG | A079046 002 | Dec 13, 2010 |
| AB | | 20MG | A079046 005 | Jul 11, 2013 |
| AB | | 30MG | A079046 006 | Jul 11, 2013 |
| AB | | 40MG | A079046 007 | Jul 11, 2013 |
| AB | IMPAX LABS | 5MG | A079087 001 | Jun 14, 2010 |
| AB | | 7.5MG | A079087 002 | Dec 21, 2010 |
| AB | | 10MG | A079087 003 | Jun 14, 2010 |
| AB | | 15MG | A079087 004 | Dec 21, 2010 |
| AB | | 20MG | A079087 005 | Jun 14, 2010 |
| AB | | 30MG | A079087 006 | Jul 22, 2010 |
| AB | SPECGX LLC | 5MG | A202946 001 | Jun 27, 2014 |
| AB | | 7.5MG | A202946 002 | Jun 27, 2014 |
| AB | | 10MG | A202946 003 | Jun 27, 2014 |
| AB | | 15MG | A202946 004 | Jun 27, 2014 |
| AB | | 20MG | A202946 005 | Jun 27, 2014 |
| AB | | 30MG | A202946 006 | Jun 27, 2014 |
| AB | | 40MG | A202946 007 | Jun 27, 2014 |
| AB | WEST-WARD PHARMS | 5MG | A200822 002 | Jul 15, 2013 |
| | INT | | | |
| AB | | 7.5MG | A200822 003 | Jul 15, 2013 |
| AB | | 10MG | A200822 004 | Jul 15, 2013 |
| AB | | 15MG | A200822 005 | Jul 15, 2013 |
| AB | | 20MG | A200822 006 | Jul 15, 2013 |
| AB | | 30MG | A200822 007 | Jul 15, 2013 |
| AB | | 40MG | A200822 001 | Jul 15, 2013 |
| ! | IMPAX LABS | 40MG | A079087 007 | Jun 14, 2010 |

OXYTOCIN

INJECTABLE; INJECTION

OXYTOCIN

| | | | | |
|-----------|------------|--------------------|-------------------------------------------|--------------------|
| AP | + ! | FRESENIUS KABI USA | 10USP UNITS/ML (10USP UNITS/ML) | N018248 001 |
| AP | + ! | | 100USP UNITS/10ML (10USP UNITS/ML) | N018248 002 |
| AP | | HIKMA FARMACEUTICA | 10USP UNITS/ML (10USP UNITS/ML) | A200219 001 |
| AP | | SAGENT PHARMS | 10USP UNITS/ML (10USP UNITS/ML) | A091676 001 |
| AP | | | 100USP UNITS/10ML (10USP UNITS/ML) | A091676 002 |
| AP | + ! | WEST-WARD PHARMS | 10USP UNITS/ML (10USP UNITS/ML) | N018243 001 |
| | | INT | | |
| AP | + ! | | 100USP UNITS/10ML (10USP UNITS/ML) | N018243 002 |

PITOCIN

| | | | | |
|-----------|------------|----------------------|-------------------------------------------|--------------------|
| AP | + ! | PAR STERILE PRODUCTS | 10USP UNITS/ML (10USP UNITS/ML) | N018261 001 |
| AP | + | | 100USP UNITS/10ML (10USP UNITS/ML) | N018261 002 |
| | | OXYTOCIN | | |
| | + ! | FRESENIUS KABI USA | 300USP UNITS/30ML (10USP UNITS/ML) | N018248 003 |
| | | PITOCIN | | |
| | + | PAR STERILE PRODUCTS | 500USP UNITS/50ML (10USP UNITS/ML) | N018261 003 |

OZENOXACIN

CREAM; TOPICAL

XEPI

| | | | | |
|------------|----------------------|----|-------------|--------------|
| + ! | FERRER INTERNACIONAL | 1% | N208945 001 | Dec 11, 2017 |
|------------|----------------------|----|-------------|--------------|

PACLITAXEL

FOR SUSPENSION; IV (INFUSION)

ABRAXANE

| | | | | |
|------------|--------------------|------------|-------------|--------------|
| + ! | ABRAXIS BIOSCIENCE | 100MG/VIAL | N021660 001 | Jan 07, 2005 |
|------------|--------------------|------------|-------------|--------------|

INJECTABLE; INJECTION

PACITAXEL

| | | | | |
|-----------|------------------|---------------|--------------------|--------------|
| AP | GLAND PHARMA LTD | 6MG/ML | A207326 001 | Aug 23, 2016 |
|-----------|------------------|---------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

PACLITAXEL

INJECTABLE; INJECTION

PACLITAXEL

| | | | | | |
|-----------|---|-------------------------|---------------|--------------------|--------------|
| <u>AP</u> | | ACCORD HLTHCARE | <u>6MG/ML</u> | <u>A205720 001</u> | Aug 17, 2018 |
| <u>AP</u> | | ACTAVIS TOTOWA | <u>6MG/ML</u> | <u>A090130 001</u> | Dec 09, 2009 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>6MG/ML</u> | <u>A077574 001</u> | Nov 27, 2006 |
| <u>AP</u> | ! | HOSPIRA | <u>6MG/ML</u> | <u>A076131 001</u> | May 08, 2002 |
| <u>AP</u> | | MYLAN LABS LTD | <u>6MG/ML</u> | <u>A091540 001</u> | Sep 29, 2011 |
| <u>AP</u> | | SANDOZ INC | <u>6MG/ML</u> | <u>A078167 001</u> | Dec 26, 2007 |
| <u>AP</u> | | TEVA PHARMS | <u>6MG/ML</u> | <u>A075184 001</u> | Jan 25, 2002 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>6MG/ML</u> | <u>A075190 001</u> | Jan 28, 2002 |

TAXOL

| | | | | | |
|-----------|---|-----------------|---------------|--------------------|--------------|
| <u>AP</u> | + | HQ SPCLT PHARMA | <u>6MG/ML</u> | <u>N020262 001</u> | Dec 29, 1992 |
|-----------|---|-----------------|---------------|--------------------|--------------|

PALBOCICLIB

CAPSULE; ORAL

IBRANCE

| | | | | | |
|--|---|------------|-------|-------------|--------------|
| | + | PFIZER INC | 75MG | N207103 001 | Feb 03, 2015 |
| | + | | 100MG | N207103 002 | Feb 03, 2015 |
| | + | | 125MG | N207103 003 | Feb 03, 2015 |

PALIPERIDONE

TABLET, EXTENDED RELEASE; ORAL

INVEGA

| | | | | | |
|-----------|---|----------------|--------------|--------------------|--------------|
| <u>AB</u> | + | JANSSEN PHARMS | <u>1.5MG</u> | <u>N021999 006</u> | Aug 26, 2008 |
| <u>AB</u> | + | | <u>3MG</u> | <u>N021999 001</u> | Dec 19, 2006 |
| <u>AB</u> | + | | <u>6MG</u> | <u>N021999 002</u> | Dec 19, 2006 |
| <u>AB</u> | + | | <u>9MG</u> | <u>N021999 003</u> | Dec 19, 2006 |

PALIPERIDONE

| | | | | | |
|-----------|--|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS LABS FL INC | <u>1.5MG</u> | <u>A202645 001</u> | Aug 03, 2015 |
| <u>AB</u> | | | <u>3MG</u> | <u>A202645 002</u> | Aug 03, 2015 |
| <u>AB</u> | | | <u>6MG</u> | <u>A202645 003</u> | Aug 03, 2015 |
| <u>AB</u> | | | <u>9MG</u> | <u>A202645 004</u> | Aug 03, 2015 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>1.5MG</u> | <u>A203802 001</u> | Sep 24, 2015 |
| <u>AB</u> | | | <u>3MG</u> | <u>A203802 002</u> | Sep 24, 2015 |
| <u>AB</u> | | | <u>6MG</u> | <u>A203802 003</u> | Sep 24, 2015 |
| <u>AB</u> | | | <u>9MG</u> | <u>A203802 004</u> | Sep 24, 2015 |
| <u>AB</u> | | SUN PHARMA GLOBAL | <u>1.5mg</u> | <u>A205618 001</u> | Apr 06, 2018 |
| <u>AB</u> | | | <u>3MG</u> | <u>A205618 002</u> | Apr 06, 2018 |
| <u>AB</u> | | | <u>6MG</u> | <u>A205618 003</u> | Apr 06, 2018 |
| <u>AB</u> | | | <u>9MG</u> | <u>A205618 004</u> | Apr 06, 2018 |

PALIPERIDONE PALMITATE

SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

INVEGA SUSTENNA

| | | | | | |
|--|---|----------------|-----------------------------|-------------|--------------|
| | + | JANSSEN PHARMS | 39MG/0.25ML (39MG/0.25ML) | N022264 001 | Jul 31, 2009 |
| | + | | 78MG/0.5ML (78MG/0.5ML) | N022264 002 | Jul 31, 2009 |
| | + | | 117MG/0.75ML (117MG/0.75ML) | N022264 003 | Jul 31, 2009 |
| | + | | 156MG/ML (156MG/ML) | N022264 004 | Jul 31, 2009 |
| | + | | 234MG/1.5ML (156MG/ML) | N022264 005 | Jul 31, 2009 |

INVEGA TRINZA

| | | | | | |
|--|---|----------------|-------------------------------|-------------|--------------|
| | + | JANSSEN PHARMS | 273MG/0.875ML (273MG/0.875ML) | N207946 001 | May 18, 2015 |
| | + | | 410MG/1.315ML (311.79MG/ML) | N207946 002 | May 18, 2015 |
| | + | | 546MG/1.75ML (312MG/ML) | N207946 003 | May 18, 2015 |
| | + | | 819MG/2.625ML (312MG/ML) | N207946 004 | May 18, 2015 |

PALONOSETRON HYDROCHLORIDE

INJECTABLE; INTRAVENOUS

ALOXI

| | | | | | |
|-----------|---|------------------|--------------------------------------------------|--------------------|--------------|
| <u>AP</u> | + | HELFINN HLTHCARE | <u>EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)</u> | <u>N021372 002</u> | Feb 29, 2008 |
| <u>AP</u> | + | | <u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u> | <u>N021372 001</u> | Jul 25, 2003 |

PALONOSETRON HYDROCHLORIDE

| | | | | | |
|-----------|--|-------------------------|--------------------------------------------------|--------------------|--------------|
| <u>AP</u> | | AUROBINDO PHARMA LTD | <u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u> | <u>A204702 001</u> | Nov 06, 2018 |
| <u>AP</u> | | CIPLA | <u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u> | <u>A206396 001</u> | Sep 19, 2018 |
| <u>AP</u> | | DR REDDYS LABS LTD | <u>EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)</u> | <u>A201533 001</u> | Apr 21, 2016 |
| <u>AP</u> | | | <u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u> | <u>A201533 002</u> | Apr 21, 2016 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u> | <u>A206801 001</u> | Sep 19, 2018 |
| <u>AP</u> | | | <u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u> | <u>A206802 001</u> | Sep 19, 2018 |
| <u>AP</u> | | HOSPIRA INC | <u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u> | <u>A207005 001</u> | Sep 19, 2018 |
| <u>AP</u> | | | <u>EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)</u> | <u>A207005 002</u> | Sep 19, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

PALONOSETRON HYDROCHLORIDE

INJECTABLE; INTRAVENOUS

PALONOSETRON HYDROCHLORIDE

| | | | | |
|-----------|----------------------------|--------------------------------------------------|--------------------|--------------|
| <u>AP</u> | MYLAN INSTITUTIONAL | <u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u> | <u>A206416 001</u> | Sep 19, 2018 |
| <u>AP</u> | QILU PHARM CO LTD | <u>EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)</u> | <u>A205648 002</u> | Sep 19, 2018 |
| <u>AP</u> | | <u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u> | <u>A205648 001</u> | Sep 19, 2018 |
| <u>AP</u> | SAGENT PHARMS | <u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u> | <u>A204289 001</u> | Sep 19, 2018 |
| <u>AP</u> | | <u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u> | <u>A205870 001</u> | Sep 19, 2018 |
| <u>AP</u> | SANDOZ INC | <u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u> | <u>A202521 001</u> | Oct 13, 2015 |
| <u>AP</u> | TEVA PHARMS USA | <u>EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)</u> | <u>A090713 002</u> | Mar 23, 2018 |
| <u>AP</u> | | <u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u> | <u>A090713 001</u> | Mar 23, 2018 |
| <u>AP</u> | VIRTUS PHARM | <u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u> | <u>A209287 001</u> | Sep 19, 2018 |
| | SOLUTION; INTRAVENOUS | | | |
| | PALONOSETRON HYDROCHLORIDE | | | |
| | EXELA PHARMA | EQ 0.25MG BASE/2ML (EQ 0.125MG BASE/ML) | N207963 001 | Aug 22, 2016 |
| | SCIENCE | | | |
| | + FRESENIUS KABI USA | EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) | N208109 001 | Nov 21, 2017 |

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

PAMIDRONATE DISODIUM

| | | | | |
|-----------|--------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | AREVA PHARMS | <u>30MG/VIAL</u> | <u>A077433 001</u> | Nov 26, 2008 |
| <u>AP</u> | | <u>90MG/VIAL</u> | <u>A077433 003</u> | Nov 26, 2008 |
| <u>AP</u> | FRESENIUS KABI USA | <u>30MG/VIAL</u> | <u>A075773 001</u> | May 06, 2002 |
| <u>AP</u> | | <u>30MG/10ML (3MG/ML)</u> | <u>A076207 001</u> | May 17, 2002 |
| <u>AP</u> | | <u>90MG/VIAL</u> | <u>A075773 002</u> | May 06, 2002 |
| <u>AP</u> | | <u>90MG/10ML (9MG/ML)</u> | <u>A076207 002</u> | May 17, 2002 |
| <u>AP</u> | ! HOSPIRA | <u>30MG/10ML (3MG/ML)</u> | <u>A075841 001</u> | Jun 27, 2002 |
| <u>AP</u> | ! | <u>60MG/10ML (6MG/ML)</u> | <u>A075841 002</u> | Jun 27, 2002 |
| <u>AP</u> | ! | <u>90MG/10ML (9MG/ML)</u> | <u>A075841 003</u> | Jun 27, 2002 |
| <u>AP</u> | LUITPOLD | <u>30MG/10ML (3MG/ML)</u> | <u>A078942 001</u> | Jul 25, 2008 |
| <u>AP</u> | | <u>90MG/10ML (9MG/ML)</u> | <u>A078942 002</u> | Jul 25, 2008 |
| <u>AP</u> | MYLAN LABS LTD | <u>30MG/10ML (3MG/ML)</u> | <u>A078520 001</u> | Oct 31, 2008 |
| <u>AP</u> | | <u>90MG/10ML (9MG/ML)</u> | <u>A078520 002</u> | Oct 31, 2008 |
| <u>AP</u> | PLIVA LACHEMA | <u>30MG/10ML (3MG/ML)</u> | <u>A078156 001</u> | Aug 19, 2008 |
| <u>AP</u> | | <u>60MG/10ML (6MG/ML)</u> | <u>A078156 002</u> | Aug 19, 2008 |
| <u>AP</u> | | <u>90MG/10ML (9MG/ML)</u> | <u>A078156 003</u> | Aug 19, 2008 |
| <u>AP</u> | SAGENT PHARMS | <u>30MG/10ML (3MG/ML)</u> | <u>A078373 001</u> | Dec 23, 2008 |
| <u>AP</u> | | <u>90MG/10ML (9MG/ML)</u> | <u>A078373 002</u> | Dec 23, 2008 |
| <u>AP</u> | SUN PHARMA GLOBAL | <u>30MG/VIAL</u> | <u>A077703 001</u> | Dec 24, 2008 |
| <u>AP</u> | | <u>90MG/VIAL</u> | <u>A077703 002</u> | Dec 24, 2008 |
| <u>AP</u> | TEVA PHARMS USA | <u>30MG/10ML (3MG/ML)</u> | <u>A076153 001</u> | Mar 27, 2002 |
| <u>AP</u> | | <u>90MG/10ML (9MG/ML)</u> | <u>A076153 002</u> | Mar 27, 2002 |
| <u>AP</u> | WEST-WARD PHARMS | <u>30MG/VIAL</u> | <u>A075290 001</u> | Apr 30, 2001 |
| <u>AP</u> | INT | | | |
| <u>AP</u> | +! | <u>30MG/10ML (3MG/ML)</u> | <u>N021113 001</u> | Mar 04, 2002 |
| <u>AP</u> | | <u>90MG/VIAL</u> | <u>A075290 003</u> | Apr 30, 2001 |
| <u>AP</u> | +! | <u>90MG/10ML (9MG/ML)</u> | <u>N021113 002</u> | Mar 04, 2002 |
| | AREVA PHARMS | 60MG/VIAL | A077433 002 | Nov 26, 2008 |

PANCRELIPASE (AMYLASE; LIPASE; PROTEASE)

CAPSULE, DELAYED RELEASE; ORAL

CREON

| | | | | |
|--|----------|-----------------------------------------------------|-------------|--------------|
| | + ABBVIE | 60,000USP UNITS; 12,000USP UNITS; 38,000USP UNITS | N020725 002 | Apr 30, 2009 |
| | + | 15,000USP UNITS; 3,000USP UNITS; 9,500USP UNITS | N020725 004 | Jul 12, 2011 |
| | + | 30,000USP UNITS; 6,000USP UNITS; 19,000USP UNITS | N020725 001 | Apr 30, 2009 |
| | + | 180,000USP UNITS; 36,000USP UNITS; 114,000USP UNITS | N020725 005 | Mar 14, 2013 |
| | +! | 120,000USP UNITS; 24,000USP UNITS; 76,000USP UNITS | N020725 003 | Apr 30, 2009 |

PANCREAZE

| | | | | |
|--|-------------|---------------------------------------------------|-------------|--------------|
| | + VIVUS INC | 10,850USP UNITS; 2,600USP UNITS; 6,200USP UNITS | N022523 005 | Mar 07, 2014 |
| | + | 24,600USP UNITS; 4,200USP UNITS; 14,200USP UNITS | N022523 001 | Apr 12, 2010 |
| | + | 61,500USP UNITS; 10,500USP UNITS; 35,500USP UNITS | N022523 002 | Apr 12, 2010 |
| | + | 83,900USP UNITS; 21,000USP UNITS; 54,700USP UNITS | N022523 004 | Apr 12, 2010 |
| | +! | 98,400USP UNITS; 16,800USP UNITS; 56,800USP UNITS | N022523 003 | Apr 12, 2010 |

PRESCRIPTION DRUG PRODUCT LIST

PANCRELIPASE (AMYLASE;LIPASE;PROTEASE)

CAPSULE, DELAYED RELEASE;ORAL

PERTZYE

| | | | | |
|---|--------------------|-------------------------------------------------|-------------|--------------|
| + | DIGESTIVE CARE INC | 30,250USP UNITS;8,000USP UNITS;28,750USP UNITS | N022175 001 | May 17, 2012 |
| + | ! | 60,500USP UNITS;16,000USP UNITS;57,500USP UNITS | N022175 002 | May 17, 2012 |
| + | | 15,125USP UNITS;4,000USP UNITS;14,375USP UNITS | N022175 003 | Oct 06, 2016 |
| + | | 90,750USP UNITS;24,000USP UNITS;86,250USP UNITS | N022175 004 | Jul 13, 2017 |

ZENPEP

| | | | | |
|---|-----------------|---------------------------------------------------|-------------|--------------|
| + | FOREST LABS INC | 168,000USP UNITS;40,000USP UNITS;126,000USP UNITS | N022210 007 | Mar 25, 2014 |
| + | | 14,000USP UNITS;3,000USP UNITS;10,000USP UNITS | N022210 005 | Jun 15, 2011 |
| + | | 24,000USP UNITS;5,000USP UNITS;17,000USP UNITS | N022210 001 | Aug 27, 2009 |
| + | | 42,000USP UNITS;10,000USP UNITS;32,000USP UNITS | N022210 002 | Aug 27, 2009 |
| + | | 63,000USP UNITS;15,000USP UNITS;47,000USP UNITS | N022210 003 | Aug 27, 2009 |
| + | | 84,000USP UNITS;20,000USP UNITS;63,000USP UNITS | N022210 004 | Aug 27, 2009 |
| + | ! | 105,000USP UNITS;25,000USP UNITS;79,000USP UNITS | N022210 006 | Jul 13, 2011 |

TABLET;ORAL

VIOKACE

| | | | | |
|---|-----------------|-------------------------------------------------|-------------|--------------|
| + | FOREST LABS INC | 39,150USP UNITS;10,440USP UNITS;39,150USP UNITS | N022542 001 | Mar 01, 2012 |
| + | ! | 78,300USP UNITS;20,880USP UNITS;78,300USP UNITS | N022542 002 | Mar 01, 2012 |

PANCURONIUM BROMIDE

INJECTABLE;INJECTION

PANCURONIUM BROMIDE

| | | | | |
|-----------|---------|---------------|--------------------|--------------|
| <u>AP</u> | HOSPIRA | <u>1MG/ML</u> | <u>A072320 001</u> | Jan 19, 1989 |
| <u>AP</u> | ! | <u>1MG/ML</u> | <u>A072759 001</u> | Jul 31, 1990 |
| | ! | 2MG/ML | A072760 001 | Jul 31, 1990 |

PANOBINOSTAT LACTATE

CAPSULE;ORAL

FARYDAK

| | | | | |
|---|----------------------|--------------|-------------|--------------|
| + | NOVARTIS PHARMS CORP | EQ 10MG BASE | N205353 001 | Feb 23, 2015 |
| + | | EQ 15MG BASE | N205353 002 | Feb 23, 2015 |
| + | ! | EQ 20MG BASE | N205353 003 | Feb 23, 2015 |

PANTOPRAZOLE SODIUM

FOR SUSPENSION, DELAYED RELEASE;ORAL

PROTONIX

| | | | | | |
|---|---|--------------|--------------|-------------|--------------|
| + | ! | WYETH PHARMS | EQ 40MG BASE | N022020 001 | Nov 14, 2007 |
|---|---|--------------|--------------|-------------|--------------|

INJECTABLE;IV (INFUSION)

PANTOPRAZOLE SODIUM

| | | | | |
|-----------|----------------------|--------------------------|--------------------|--------------|
| <u>AP</u> | AKORN INC | <u>EQ 40MG BASE/VIAL</u> | <u>A079197 001</u> | Nov 08, 2012 |
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>EQ 40MG BASE/VIAL</u> | <u>A205675 001</u> | Mar 30, 2016 |
| <u>AP</u> | MYLAN LABS LTD | <u>EQ 40MG BASE/VIAL</u> | <u>A208580 001</u> | May 04, 2018 |
| <u>AP</u> | SANDOZ INC | <u>EQ 40MG BASE/VIAL</u> | <u>A090296 001</u> | Jul 14, 2015 |

PROTONIX IV

| | | | | | | |
|-----------|---|---|--------------|--------------------------|--------------------|--------------|
| <u>AP</u> | + | ! | WYETH PHARMS | <u>EQ 40MG BASE/VIAL</u> | <u>N020988 001</u> | Mar 22, 2001 |
|-----------|---|---|--------------|--------------------------|--------------------|--------------|

POWDER;IV (INFUSION)

PANTOPRAZOLE SODIUM

| | | | | | |
|--|--|----------------------|-------------------|-------------|--------------|
| | | EXELA PHARMA SCS LLC | EQ 40MG BASE/VIAL | N209463 001 | Jun 30, 2017 |
|--|--|----------------------|-------------------|-------------|--------------|

TABLET, DELAYED RELEASE;ORAL

PANTOPRAZOLE SODIUM

| | | | | |
|-----------|----------------------|---------------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS TOTOWA | <u>EQ 20MG BASE</u> | <u>A090797 001</u> | Feb 07, 2011 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A090797 002</u> | Feb 07, 2011 |
| <u>AB</u> | AMNEAL PHARMS | <u>EQ 20MG BASE</u> | <u>A205119 001</u> | Jan 26, 2016 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A205119 002</u> | Jan 26, 2016 |
| <u>AB</u> | APOTEX INC | <u>EQ 20MG BASE</u> | <u>A090807 001</u> | May 02, 2012 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A090807 002</u> | May 02, 2012 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 20MG BASE</u> | <u>A202038 001</u> | Sep 28, 2012 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A202038 002</u> | Sep 28, 2012 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 20MG BASE</u> | <u>A077619 001</u> | Jan 19, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

PANTOPRAZOLE SODIUM

TABLET, DELAYED RELEASE;ORAL

PANTOPRAZOLE SODIUM

| | | | | |
|-----------|---------------------|---------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077619 002</u> | Jan 19, 2011 |
| <u>AB</u> | HETERO LABS LTD V | <u>EQ 20MG BASE</u> | <u>A202882 001</u> | Sep 10, 2014 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A202882 002</u> | Sep 10, 2014 |
| <u>AB</u> | JUBILANT GENERICS | <u>EQ 20MG BASE</u> | <u>A090901 001</u> | Aug 30, 2011 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A090901 002</u> | Aug 30, 2011 |
| <u>AB</u> | LANNETT CO INC | <u>EQ 20MG BASE</u> | <u>A078281 001</u> | Jan 20, 2011 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A078281 002</u> | Jan 20, 2011 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>EQ 20MG BASE</u> | <u>A200821 001</u> | Feb 16, 2012 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A200821 002</u> | Feb 16, 2012 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 20MG BASE</u> | <u>A090970 001</u> | Jan 19, 2011 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A090970 002</u> | Jan 19, 2011 |
| <u>AB</u> | ORCHID HLTHCARE | <u>EQ 20MG BASE</u> | <u>A202052 001</u> | Dec 02, 2014 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A202052 002</u> | Dec 02, 2014 |
| <u>AB</u> | PERRIGO R AND D | <u>EQ 20MG BASE</u> | <u>A203024 001</u> | May 07, 2014 |
| <u>AB</u> | TEVA | <u>EQ 20MG BASE</u> | <u>A077056 001</u> | Aug 02, 2007 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077056 002</u> | Aug 02, 2007 |
| <u>AB</u> | TORRENT PHARMS | <u>EQ 20MG BASE</u> | <u>A090074 001</u> | Jan 19, 2011 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A090074 002</u> | Jan 19, 2011 |
| <u>AB</u> | WOCKHARDT | <u>EQ 20MG BASE</u> | <u>A091231 001</u> | Jan 19, 2011 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A091231 002</u> | Jan 19, 2011 |

PROTONIX

| | | | | |
|-----------|----------------|---------------------|--------------------|--------------|
| <u>AB</u> | + WYETH PHARMS | <u>EQ 20MG BASE</u> | <u>N020987 002</u> | Jun 12, 2001 |
| <u>AB</u> | +! | <u>EQ 40MG BASE</u> | <u>N020987 001</u> | Feb 02, 2000 |

PARICALCITOL

CAPSULE;ORAL

PARICALCITOL

| | | | | |
|-----------|----------------------|-------------|--------------------|--------------|
| <u>AB</u> | AMNEAL PHARMS | <u>1MCG</u> | <u>A204327 001</u> | Jan 13, 2016 |
| <u>AB</u> | | <u>2MCG</u> | <u>A204327 002</u> | Jan 13, 2016 |
| <u>AB</u> | | <u>4MCG</u> | <u>A204327 003</u> | Jan 13, 2016 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>1MCG</u> | <u>A207672 001</u> | Jan 14, 2016 |
| <u>AB</u> | | <u>2MCG</u> | <u>A207672 002</u> | Jan 14, 2016 |
| <u>AB</u> | | <u>4MCG</u> | <u>A207672 003</u> | Jan 14, 2016 |
| <u>AB</u> | BIONPHARMA INC | <u>1MCG</u> | <u>A202539 001</u> | Mar 27, 2014 |
| <u>AB</u> | | <u>2MCG</u> | <u>A202539 002</u> | Mar 27, 2014 |
| <u>AB</u> | | <u>4MCG</u> | <u>A202539 003</u> | Mar 27, 2014 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>1MCG</u> | <u>A091412 001</u> | Jun 24, 2014 |
| <u>AB</u> | | <u>2MCG</u> | <u>A091412 002</u> | Jun 24, 2014 |
| <u>AB</u> | | <u>4MCG</u> | <u>A091412 003</u> | Jun 24, 2014 |
| <u>AB</u> | LOTUS PHARM CO LTD | <u>1MCG</u> | <u>A206710 001</u> | Feb 24, 2016 |
| <u>AB</u> | | <u>2MCG</u> | <u>A206710 002</u> | Feb 24, 2016 |
| <u>AB</u> | | <u>4MCG</u> | <u>A206710 003</u> | Feb 24, 2016 |
| <u>AB</u> | MARKSANS PHARMA | <u>1MCG</u> | <u>A204948 001</u> | Oct 07, 2016 |
| <u>AB</u> | | <u>2MCG</u> | <u>A204948 002</u> | Oct 07, 2016 |
| <u>AB</u> | | <u>4MCG</u> | <u>A204948 003</u> | Oct 07, 2016 |
| <u>AB</u> | RISING PHARMS | <u>1MCG</u> | <u>A202124 001</u> | Jun 24, 2014 |
| <u>AB</u> | | <u>2MCG</u> | <u>A202124 002</u> | Jun 24, 2014 |
| <u>AB</u> | | <u>4MCG</u> | <u>A202124 003</u> | Jun 24, 2014 |
| <u>AB</u> | TEVA PHARMS USA | <u>1MCG</u> | <u>A090829 001</u> | Sep 27, 2013 |
| <u>AB</u> | | <u>2MCG</u> | <u>A090829 002</u> | Sep 27, 2013 |
| <u>AB</u> | ! | <u>4MCG</u> | <u>A090829 003</u> | Sep 27, 2013 |

ZEMPLAR

| | | | | |
|-----------|----------|-------------|--------------------|--------------|
| <u>AB</u> | + ABBVIE | <u>1MCG</u> | <u>N021606 001</u> | May 26, 2005 |
| <u>AB</u> | + | <u>2MCG</u> | <u>N021606 002</u> | May 26, 2005 |

SOLUTION;INTRAVENOUS

PARICALCITOL

| | | | | |
|-----------|----------------------|--------------------------------|--------------------|--------------|
| <u>AP</u> | ACCORD HLTHCARE | <u>0.002MG/ML (0.002MG/ML)</u> | <u>N207174 001</u> | Feb 04, 2016 |
| <u>AP</u> | | <u>0.005MG/ML (0.005MG/ML)</u> | <u>N207174 002</u> | Feb 04, 2016 |
| <u>AP</u> | | <u>0.01MG/2ML (0.005MG/ML)</u> | <u>N207174 003</u> | Feb 04, 2016 |
| <u>AP</u> | AKORN | <u>0.005MG/ML (0.005MG/ML)</u> | <u>A207692 001</u> | Oct 16, 2017 |
| <u>AP</u> | AMNEAL PHARMS CO | <u>0.002MG/ML (0.002MG/ML)</u> | <u>A206699 001</u> | Mar 09, 2017 |
| <u>AP</u> | | <u>0.005MG/ML (0.005MG/ML)</u> | <u>A206699 002</u> | Mar 09, 2017 |
| <u>AP</u> | | <u>0.01MG/2ML (0.005MG/ML)</u> | <u>A206699 003</u> | Mar 09, 2017 |
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>0.002MG/ML (0.002MG/ML)</u> | <u>A205982 001</u> | Oct 09, 2018 |
| <u>AP</u> | | <u>0.005MG/ML (0.005MG/ML)</u> | <u>A205982 002</u> | Oct 09, 2018 |
| <u>AP</u> | | <u>0.01MG/2ML (0.005MG/ML)</u> | <u>A205982 003</u> | Oct 09, 2018 |
| <u>AP</u> | DR REDDYS LABS LTD | <u>0.002MG/ML (0.002MG/ML)</u> | <u>A204910 001</u> | Aug 17, 2016 |
| <u>AP</u> | | <u>0.005MG/ML (0.005MG/ML)</u> | <u>A204910 002</u> | Aug 17, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

PARICALCITOL

SOLUTION; INTRAVENOUS

PARICALCITOL

| | | | | |
|-----------|----------------|--------------------------------|--------------------|--------------|
| <u>AP</u> | | <u>0.01MG/2ML (0.005MG/ML)</u> | <u>A204910 003</u> | Aug 17, 2016 |
| <u>AP</u> | HIKMA PHARMS | <u>0.002MG/ML (0.002MG/ML)</u> | <u>N205917 001</u> | Nov 18, 2014 |
| <u>AP</u> | | <u>0.005MG/ML (0.005MG/ML)</u> | <u>N205917 002</u> | Nov 18, 2014 |
| <u>AP</u> | | <u>0.01MG/2ML (0.005MG/ML)</u> | <u>N205917 003</u> | Nov 18, 2014 |
| <u>AP</u> | HOSPIRA INC | <u>0.002MG/ML (0.002MG/ML)</u> | <u>N201657 001</u> | Oct 21, 2014 |
| <u>AP</u> | | <u>0.005MG/ML (0.005MG/ML)</u> | <u>N201657 002</u> | Oct 21, 2014 |
| <u>AP</u> | | <u>0.01MG/2ML (0.005MG/ML)</u> | <u>N201657 003</u> | Oct 21, 2014 |
| <u>AP</u> | MYLAN LABS LTD | <u>0.002MG/ML (0.002MG/ML)</u> | <u>A203897 001</u> | Nov 02, 2017 |
| <u>AP</u> | | <u>0.005MG/ML (0.005MG/ML)</u> | <u>A203897 002</u> | Nov 02, 2017 |
| <u>AP</u> | | <u>0.01MG/2ML (0.005MG/ML)</u> | <u>A203897 003</u> | Nov 02, 2017 |
| <u>AP</u> | SANDOZ INC | <u>0.002MG/ML (0.002MG/ML)</u> | <u>A091108 001</u> | Jul 27, 2011 |
| <u>AP</u> | | <u>0.005MG/ML (0.005MG/ML)</u> | <u>A091108 002</u> | Jul 27, 2011 |
| <u>AP</u> | | <u>0.01MG/2ML (0.005MG/ML)</u> | <u>A091108 003</u> | Jul 27, 2011 |

ZEMPLAR

| | | | | | |
|-----------|----|--------|--------------------------------|--------------------|--------------|
| <u>AP</u> | +! | ABEVIE | <u>0.002MG/ML (0.002MG/ML)</u> | <u>N020819 002</u> | Feb 01, 2000 |
| <u>AP</u> | +! | | <u>0.005MG/ML (0.005MG/ML)</u> | <u>N020819 001</u> | Apr 17, 1998 |
| <u>AP</u> | +! | | <u>0.01MG/2ML (0.005MG/ML)</u> | <u>N020819 003</u> | Feb 01, 2000 |

PAROMOMYCIN SULFATE

CAPSULE; ORAL

PAROMOMYCIN SULFATE

| | | | | | |
|-----------|---|---------------------|----------------------|--------------------|--------------|
| <u>AA</u> | | HERITAGE PHARMS INC | <u>EQ 250MG BASE</u> | <u>A065173 001</u> | Dec 14, 2007 |
| <u>AA</u> | ! | SUN PHARM INDS INC | <u>EQ 250MG BASE</u> | <u>A064171 001</u> | Jun 30, 1997 |

PAROXETINE HYDROCHLORIDE

SUSPENSION; ORAL

PAXIL

| | | | | | |
|--|----|---------------------|-------------------------|--------------------|--------------|
| | +! | APOTEX TECHNOLOGIES | <u>EQ 10MG BASE/5ML</u> | <u>N020710 001</u> | Jun 25, 1997 |
|--|----|---------------------|-------------------------|--------------------|--------------|

TABLET; ORAL

PAROXETINE

| | | | | | |
|-----------|--|--------------|---------------------|--------------------|--------------|
| <u>AB</u> | | PRINSTON INC | <u>EQ 10MG BASE</u> | <u>A203854 001</u> | Oct 31, 2014 |
| <u>AB</u> | | | <u>EQ 20MG BASE</u> | <u>A203854 002</u> | Oct 31, 2014 |
| <u>AB</u> | | | <u>EQ 30MG BASE</u> | <u>A203854 003</u> | Oct 31, 2014 |
| <u>AB</u> | | | <u>EQ 40MG BASE</u> | <u>A203854 004</u> | Oct 31, 2014 |

PAROXETINE HYDROCHLORIDE

| | | | | | |
|-----------|--|--------------------|---------------------|--------------------|--------------|
| <u>AB</u> | | APOTEX | <u>EQ 10MG BASE</u> | <u>A075356 001</u> | Jul 30, 2003 |
| <u>AB</u> | | | <u>EQ 20MG BASE</u> | <u>A075356 002</u> | Jul 30, 2003 |
| <u>AB</u> | | | <u>EQ 30MG BASE</u> | <u>A075356 003</u> | Jul 30, 2003 |
| <u>AB</u> | | | <u>EQ 40MG BASE</u> | <u>A075356 004</u> | Jul 30, 2003 |
| <u>AB</u> | | AUROBINDO PHARMA | <u>EQ 10MG BASE</u> | <u>A078406 001</u> | Jul 25, 2007 |
| <u>AB</u> | | | <u>EQ 20MG BASE</u> | <u>A078406 002</u> | Jul 25, 2007 |
| <u>AB</u> | | | <u>EQ 30MG BASE</u> | <u>A078406 003</u> | Jul 25, 2007 |
| <u>AB</u> | | | <u>EQ 40MG BASE</u> | <u>A078406 004</u> | Jul 25, 2007 |
| <u>AB</u> | | JUBILANT GENERICS | <u>EQ 10MG BASE</u> | <u>A205528 001</u> | Nov 27, 2015 |
| <u>AB</u> | | | <u>EQ 20MG BASE</u> | <u>A205528 002</u> | Nov 27, 2015 |
| <u>AB</u> | | | <u>EQ 30MG BASE</u> | <u>A205528 003</u> | Nov 27, 2015 |
| <u>AB</u> | | | <u>EQ 40MG BASE</u> | <u>A205528 004</u> | Nov 27, 2015 |
| <u>AB</u> | | MYLAN | <u>EQ 10MG BASE</u> | <u>A078902 001</u> | Mar 13, 2008 |
| <u>AB</u> | | | <u>EQ 20MG BASE</u> | <u>A078902 002</u> | Mar 13, 2008 |
| <u>AB</u> | | | <u>EQ 30MG BASE</u> | <u>A078902 003</u> | Mar 13, 2008 |
| <u>AB</u> | | | <u>EQ 40MG BASE</u> | <u>A078902 004</u> | Mar 13, 2008 |
| <u>AB</u> | | OXFORD PHARMS | <u>EQ 10MG BASE</u> | <u>A076968 001</u> | Jun 21, 2010 |
| <u>AB</u> | | | <u>EQ 20MG BASE</u> | <u>A076968 002</u> | Jun 21, 2010 |
| <u>AB</u> | | | <u>EQ 30MG BASE</u> | <u>A076968 003</u> | Jun 21, 2010 |
| <u>AB</u> | | | <u>EQ 40MG BASE</u> | <u>A076968 004</u> | Jun 21, 2010 |
| <u>AB</u> | | SUN PHARM INDS INC | <u>EQ 10MG BASE</u> | <u>A078194 001</u> | Jun 29, 2007 |
| <u>AB</u> | | | <u>EQ 20MG BASE</u> | <u>A078194 002</u> | Jun 29, 2007 |
| <u>AB</u> | | | <u>EQ 30MG BASE</u> | <u>A078194 003</u> | Jun 29, 2007 |
| <u>AB</u> | | | <u>EQ 40MG BASE</u> | <u>A078194 004</u> | Jun 29, 2007 |
| <u>AB</u> | | TEVA | <u>EQ 10MG BASE</u> | <u>A076618 001</u> | Aug 15, 2005 |
| <u>AB</u> | | | <u>EQ 20MG BASE</u> | <u>A076618 002</u> | Aug 15, 2005 |
| <u>AB</u> | | | <u>EQ 30MG BASE</u> | <u>A076618 003</u> | Aug 15, 2005 |
| <u>AB</u> | | | <u>EQ 40MG BASE</u> | <u>A076618 004</u> | Aug 15, 2005 |
| <u>AB</u> | | ZYDUS PHARMS USA | <u>EQ 10MG BASE</u> | <u>A077584 001</u> | Mar 07, 2007 |
| <u>AB</u> | | | <u>EQ 20MG BASE</u> | <u>A077584 002</u> | Mar 07, 2007 |
| <u>AB</u> | | | <u>EQ 30MG BASE</u> | <u>A077584 003</u> | Mar 07, 2007 |
| <u>AB</u> | | | <u>EQ 40MG BASE</u> | <u>A077584 004</u> | Mar 07, 2007 |

PAXIL

| | | | | | |
|-----------|---|---------------------|---------------------|--------------------|--------------|
| <u>AB</u> | + | APOTEX TECHNOLOGIES | <u>EQ 10MG BASE</u> | <u>N020031 001</u> | Dec 29, 1992 |
| <u>AB</u> | + | | <u>EQ 20MG BASE</u> | <u>N020031 002</u> | Dec 29, 1992 |

PRESCRIPTION DRUG PRODUCT LIST

PAROXETINE HYDROCHLORIDE

TABLET; ORAL

PAXIL

| | | | | | |
|-----------|---|---|---------------------|--------------------|--------------|
| <u>AB</u> | + | | <u>EQ 30MG BASE</u> | <u>N020031 003</u> | Dec 29, 1992 |
| <u>AB</u> | + | ! | <u>EQ 40MG BASE</u> | <u>N020031 005</u> | Dec 29, 1992 |

TABLET, EXTENDED RELEASE; ORAL

PAROXETINE HYDROCHLORIDE

| | | | | | |
|-----------|--|---------------------|-----------------------|--------------------|--------------|
| <u>AB</u> | | LANNETT CO INC | <u>EQ 12.5MG BASE</u> | <u>A204744 001</u> | Jun 10, 2016 |
| <u>AB</u> | | | <u>EQ 25MG BASE</u> | <u>A204744 002</u> | Jun 10, 2016 |
| <u>AB</u> | | | <u>EQ 37.5MG BASE</u> | <u>A204744 003</u> | Jun 10, 2016 |
| <u>AB</u> | | LUPIN LTD | <u>EQ 12.5MG BASE</u> | <u>A204134 001</u> | Jan 20, 2017 |
| <u>AB</u> | | | <u>EQ 25MG BASE</u> | <u>A204134 002</u> | Jan 20, 2017 |
| <u>AB</u> | | | <u>EQ 37.5MG BASE</u> | <u>A204134 003</u> | Jan 20, 2017 |
| <u>AB</u> | | MYLAN | <u>EQ 12.5MG BASE</u> | <u>A077873 001</u> | Jun 29, 2007 |
| <u>AB</u> | | | <u>EQ 25MG BASE</u> | <u>A077873 002</u> | Jun 29, 2007 |
| <u>AB</u> | | | <u>EQ 37.5MG BASE</u> | <u>A091427 001</u> | Apr 14, 2011 |
| <u>AB</u> | | SCIECURE PHARMA INC | <u>EQ 12.5MG BASE</u> | <u>A209293 001</u> | Jun 12, 2018 |
| <u>AB</u> | | | <u>EQ 25MG BASE</u> | <u>A209293 002</u> | Jun 12, 2018 |
| <u>AB</u> | | | <u>EQ 37.5MG BASE</u> | <u>A209293 003</u> | Jun 12, 2018 |

PAXIL CR

| | | | | | |
|-----------|---|---------------------|-----------------------|--------------------|--------------|
| <u>AB</u> | + | APOTEX TECHNOLOGIES | <u>EQ 12.5MG BASE</u> | <u>N020936 001</u> | Feb 16, 1999 |
| <u>AB</u> | + | | <u>EQ 25MG BASE</u> | <u>N020936 002</u> | Feb 16, 1999 |
| <u>AB</u> | + | ! | <u>EQ 37.5MG BASE</u> | <u>N020936 003</u> | Dec 06, 2000 |

PAROXETINE MESYLATE

CAPSULE; ORAL

BRISDELLE

| | | | | | | |
|-----------|---|---|--------------------|----------------------|--------------------|--------------|
| <u>AB</u> | + | ! | SEBELA IRELAND LTD | <u>EQ 7.5MG BASE</u> | <u>N204516 001</u> | Jun 28, 2013 |
|-----------|---|---|--------------------|----------------------|--------------------|--------------|

PAROXETINE MESYLATE

| | | | | | |
|-----------|--|---------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS LABS FL INC | <u>EQ 7.5MG BASE</u> | <u>A207139 001</u> | Jun 20, 2017 |
| <u>AB</u> | | PRINSTON INC | <u>EQ 7.5MG BASE</u> | <u>A207188 001</u> | Aug 18, 2017 |

TABLET; ORAL

PEXEVA

| | | | | | |
|--|---|--------------------|--------------|-------------|--------------|
| | + | SEBELA IRELAND LTD | EQ 10MG BASE | N021299 001 | Jul 03, 2003 |
| | + | | EQ 20MG BASE | N021299 002 | Jul 03, 2003 |
| | + | | EQ 30MG BASE | N021299 003 | Jul 03, 2003 |
| | + | ! | EQ 40MG BASE | N021299 004 | Jul 03, 2003 |

PASIREOTIDE DIASPARTATE

SOLUTION; SUBCUTANEOUS

SIGNIFOR

| | | | | | |
|--|---|----------|-------------------------------------|-------------|--------------|
| | + | NOVARTIS | EQ 0.3MG BASE/ML (EQ 0.3MG BASE/ML) | N200677 001 | Dec 14, 2012 |
| | + | | EQ 0.6MG BASE/ML (EQ 0.6MG BASE/ML) | N200677 002 | Dec 14, 2012 |
| | + | ! | EQ 0.9MG BASE/ML (EQ 0.9MG BASE/ML) | N200677 003 | Dec 14, 2012 |

PASIREOTIDE PAMOATE

FOR SUSPENSION; INTRAMUSCULAR

SIGNIFOR LAR KIT

| | | | | | |
|--|---|----------------------|-------------------|-------------|--------------|
| | + | NOVARTIS PHARMS CORP | EQ 10MG BASE/VIAL | N203255 004 | Jun 29, 2018 |
| | + | | EQ 20MG BASE/VIAL | N203255 001 | Dec 15, 2014 |
| | + | | EQ 30MG BASE/VIAL | N203255 005 | Jun 29, 2018 |
| | + | | EQ 40MG BASE/VIAL | N203255 002 | Dec 15, 2014 |
| | + | ! | EQ 60MG BASE/VIAL | N203255 003 | Dec 15, 2014 |

PATIOMER SORBITE X CALCIUM

POWDER; ORAL

VELTASSA

| | | | | | |
|--|---|-------------|-----------------------|-------------|--------------|
| | + | RELYPSA INC | EQ 8.4GM BASE/PACKET | N205739 001 | Oct 21, 2015 |
| | + | | EQ 16.8GM BASE/PACKET | N205739 002 | Oct 21, 2015 |
| | + | ! | EQ 25.2GM BASE/PACKET | N205739 003 | Oct 21, 2015 |

PATISIRAN SODIUM

SOLUTION; INTRAVENOUS

ONPATRO

| | | | | | | |
|--|---|---|--------------------|-----------------------------------|-------------|--------------|
| | + | ! | ALNYLAM PHARMS INC | EQ 10MG BASE/5ML (EQ 2MG BASE/ML) | N210922 001 | Aug 10, 2018 |
|--|---|---|--------------------|-----------------------------------|-------------|--------------|

PAZOPANIB HYDROCHLORIDE

TABLET; ORAL

VOTRIENT

| | | | | | | |
|--|---|---|----------------------|---------------|-------------|--------------|
| | + | ! | NOVARTIS PHARMS CORP | EQ 200MG BASE | N022465 001 | Oct 19, 2009 |
|--|---|---|----------------------|---------------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

PEGADEMASE BOVINE

INJECTABLE; INJECTION

ADAGEN

+! LEADIANT BIOSCI INC 250 UNITS/ML N019818 001 Mar 21, 1990

PEGAPTANIB SODIUM

INJECTABLE; INTRAVITREAL

MACUGEN

+! VALEANT PHARMS LLC EQ 0.3MG ACID/0.09ML N021756 001 Dec 17, 2004

PEGVISOMANT

INJECTABLE; SUBCUTANEOUS

SOMAVERT

+! PHARMACIA AND 10MG/VIAL N021106 001 Mar 25, 2003

UPJOHN

+! 15MG/VIAL N021106 002 Mar 25, 2003

+! 20MG/VIAL N021106 003 Mar 25, 2003

+! 25MG/VIAL N021106 004 Jul 31, 2014

+! 30MG/VIAL N021106 005 Jul 31, 2014

PEMETREXED DISODIUM

POWDER; INTRAVENOUS

ALIMTA

+! LILLY EQ 100MG BASE/VIAL N021462 002 Sep 07, 2007

+! EQ 500MG BASE/VIAL N021462 001 Feb 04, 2004

PENCICLOVIR

CREAM; TOPICAL

DENA VIR

+! MYLAN PHARMS INC 1% N020629 001 Sep 24, 1996

PENICILLAMINE

CAPSULE; ORAL

CUPRIMINE

+! ATON 250MG N019853 001

TABLET; ORAL

DEPEN

+! MYLAN SPECIALITY LP 250MG N019854 001

PENICILLIN G BENZATHINE

INJECTABLE; INJECTION

BICILLIN L-A

BC +! KING PHARMS LLC 600,000 UNITS/ML N050141 001

PENICILLIN G BENZATHINE; PENICILLIN G PROCAINE

INJECTABLE; INJECTION

BICILLIN C-R

+! KING PHARMS LLC 300,000 UNITS/ML; 300,000 UNITS/ML N050138 001

BICILLIN C-R 900/300

+! KING PHARMS LLC 900,000 UNITS/2ML; 300,000 UNITS/2ML N050138 003

PENICILLIN G POTASSIUM

INJECTABLE; INJECTION

PENICILLIN G POTASSIUM**AP** ACS DOBFAR SPA **20,000,000 UNITS/VIAL** **A205043 002** Oct 26, 2018**AP** **5,000,000 UNITS/VIAL** **A205043 001** Oct 26, 2018**AP** HANFORD GC **5,000,000 UNITS/VIAL** **A065149 002** Jul 23, 2009**AP** **20,000,000 UNITS/VIAL** **A065149 003** Jul 23, 2009**AP** ISTITUTO BIO ITA SPA **5,000,000 UNITS/VIAL** **A065448 001** Aug 18, 2009**AP** **20,000,000 UNITS/VIAL** **A065448 002** Aug 18, 2009**AP** SANDOZ **5,000,000 UNITS/VIAL** **A065079 002** Aug 30, 2002**AP** **20,000,000 UNITS/VIAL** **A065079 003** Aug 30, 2002PFIZERPEN**AP** ! PFIZER **5,000,000 UNITS/VIAL** **A060657 002****AP** ! **20,000,000 UNITS/VIAL** **A060657 003**

PENICILLIN G POTASSIUM

HANFORD GC

1,000,000 UNITS/VIAL A065149 001 Jul 23, 2009

PENICILLIN G POTASSIUM IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 20,000 UNITS/ML N050638 001 Jun 25, 1990

+! 40,000 UNITS/ML N050638 002 Jun 25, 1990

+! 60,000 UNITS/ML N050638 003 Jun 25, 1990

PRESCRIPTION DRUG PRODUCT LIST

PENICILLIN G PROCAINE

INJECTABLE; INJECTION

PENICILLIN G PROCAINE

! KING PHARMS LLC

300,000 UNITS/ML

A060101 002

!

600,000 UNITS/ML

A060101 001

PENICILLIN G SODIUM

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

PENICILLIN G SODIUM

! SANDOZ

5,000,000 UNITS/VIAL

A065068 001 Feb 26, 2001

PENICILLIN V POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN V POTASSIUMAA DAVA PHARMS INCEQ 125MG BASE/5MLA062981 001 Feb 10, 1989AA !EQ 250MG BASE/5MLA062981 002 Feb 10, 1989PENICILLIN-VKAA TEVAEQ 125MG BASE/5MLA060456 001AA !EQ 250MG BASE/5MLA060456 002

TABLET; ORAL

PENICILLIN V POTASSIUMAB AUROBINDO PHARMAEQ 250MG BASEA065435 001 Apr 29, 2008AB !EQ 500MG BASEA065435 002 Apr 29, 2008AB DAVA PHARMS INCEQ 250MG BASEA062936 001 Nov 25, 1988AB !EQ 500MG BASEA062935 001 Nov 23, 1988AB HIKMA PHARMSEQ 250MG BASEA090549 001 Oct 11, 2013AB !EQ 500MG BASEA090549 002 Oct 11, 2013AB SANDOZEQ 250MG BASEA064071 001 Nov 30, 1995AB !EQ 500MG BASEA064071 002 Nov 30, 1995PENICILLIN-VKAB TEVAEQ 250MG BASEA060711 002AB !EQ 500MG BASEA060711 003PENTAMIDINE ISETHIONATE

FOR SOLUTION; INHALATION

NEBUPENT

+! FRESENIUS KABI USA

300MG/VIAL

N019887 001 Jun 15, 1989

INJECTABLE; INJECTION

PENTAMAP +! FRESENIUS KABI USA300MG/VIALN019264 001 Oct 16, 1984PENTAMIDINE ISETHIONATEAP SETON PHARMS300MG/VIALA206666 001 Sep 28, 2017PENTETATE CALCIUM TRISODIUM

SOLUTION; INHALATION, INTRAVENOUS

PENTETATE CALCIUM TRISODIUM

+! HAMELN PHARMA PLUS EQ 1GM BASE/5ML (EQ 200MG BASE/ML)

N021749 001 Aug 11, 2004

PENTETATE ZINC TRISODIUM

SOLUTION; INHALATION, INTRAVENOUS

PENTETATE ZINC TRISODIUM

+! HAMELN PHARMA PLUS EQ 1GM BASE/5ML (EQ 200MG BASE/ML)

N021751 001 Aug 11, 2004

PENTOBARBITAL SODIUM

INJECTABLE; INJECTION

NEMBUTAL SODIUMAP ! OAK PHARMS50MG/MLA083246 001PENTOBARBITAL SODIUMAP CUSTOPHARM INC50MG/MLA203619 001 Nov 13, 2017AP RENAISSANCE SSA LLC50MG/MLA206677 001 Nov 27, 2017AP SAGENT PHARMS50MG/MLA206404 001 May 23, 2016PENTOSAN POLYSULFATE SODIUM

CAPSULE; ORAL

ELMIRON

+! JANSSEN PHARMS

100MG

N020193 001 Sep 26, 1996

PENTOSTATIN

INJECTABLE; INJECTION

NIPENTAP +! HOSPIRA INC10MG/VIALN020122 001 Oct 11, 1991PENTOSTATINAP MYLAN INSTITUTIONAL10MG/VIALA203554 001 Sep 19, 2014AP WEST-WARD PHARMS10MG/VIALA077841 001 Aug 07, 2007

INT

PRESCRIPTION DRUG PRODUCT LIST

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE;ORAL

PENTOXIFYLLINE

| | | | | | | |
|-----------|---|----------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ! | APOTEX | <u>400MG</u> | <u>A075191</u> | <u>001</u> | Jun 09, 1999 |
| <u>AB</u> | | MYLAN | <u>400MG</u> | <u>A074425</u> | <u>001</u> | Jul 08, 1997 |
| <u>AB</u> | | VALEANT PHARMS | <u>400MG</u> | <u>A075028</u> | <u>001</u> | Jul 20, 1998 |

PENTOXIL

| | | | | | | |
|-----------|--|-------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | UPSHER SMITH LABS | <u>400MG</u> | <u>A074962</u> | <u>001</u> | Mar 31, 1999 |
|-----------|--|-------------------|--------------|----------------|------------|--------------|

PERAMIVIR

SOLUTION;INTRAVENOUS

RAPIVAB

| | | | | | | |
|---|---|----------|----------------------|---------|-----|--------------|
| + | ! | BIOCRIST | 200MG/20ML (10MG/ML) | N206426 | 001 | Dec 19, 2014 |
|---|---|----------|----------------------|---------|-----|--------------|

PERAMPANEL

SUSPENSION;ORAL

FYCOMPA

| | | | | | | |
|---|---|-----------|----------|---------|-----|--------------|
| + | ! | EISAI INC | 0.5MG/ML | N208277 | 001 | Apr 29, 2016 |
|---|---|-----------|----------|---------|-----|--------------|

TABLET;ORAL

FYCOMPA

| | | | | | | |
|---|---|-----------|------|---------|-----|--------------|
| + | | EISAI INC | 2MG | N202834 | 001 | Oct 22, 2012 |
| + | | | 4MG | N202834 | 002 | Oct 22, 2012 |
| + | | | 6MG | N202834 | 003 | Oct 22, 2012 |
| + | | | 8MG | N202834 | 004 | Oct 22, 2012 |
| + | | | 10MG | N202834 | 005 | Oct 22, 2012 |
| + | ! | | 12MG | N202834 | 006 | Oct 22, 2012 |

PERFLUTREN

INJECTABLE;INTRAVENOUS

DEFINITY

| | | | | | | |
|---|---|----------------|-----------|---------|-----|--------------|
| + | ! | LANTHEUS MEDCL | 6.52MG/ML | N021064 | 001 | Jul 31, 2001 |
|---|---|----------------|-----------|---------|-----|--------------|

PERINDOPRIL ERBUMINE

TABLET;ORAL

PERINDOPRIL ERBUMINE

| | | | | | | |
|-----------|---|-------------------------|------------|----------------|------------|--------------|
| <u>AB</u> | | ANI PHARMS INC | <u>2MG</u> | <u>A078138</u> | <u>001</u> | Nov 10, 2009 |
| <u>AB</u> | | | <u>4MG</u> | <u>A078138</u> | <u>002</u> | Nov 10, 2009 |
| <u>AB</u> | | | <u>8MG</u> | <u>A078138</u> | <u>003</u> | Nov 10, 2009 |
| <u>AB</u> | | AUROBINDO PHARMA | <u>2MG</u> | <u>A079070</u> | <u>001</u> | Nov 10, 2009 |
| <u>AB</u> | | | <u>4MG</u> | <u>A079070</u> | <u>002</u> | Nov 10, 2009 |
| <u>AB</u> | ! | | <u>8MG</u> | <u>A079070</u> | <u>003</u> | Nov 10, 2009 |
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>2MG</u> | <u>A090072</u> | <u>001</u> | Nov 10, 2009 |
| <u>AB</u> | | | <u>4MG</u> | <u>A090072</u> | <u>002</u> | Nov 10, 2009 |
| <u>AB</u> | | | <u>8MG</u> | <u>A090072</u> | <u>003</u> | Nov 10, 2009 |

PERMETHRIN

CREAM;TOPICAL

ELIMITE

| | | | | | | | |
|-----------|---|---|------------------|-----------|----------------|------------|--------------|
| <u>AB</u> | + | ! | MYLAN PHARMS INC | <u>5%</u> | <u>N019855</u> | <u>001</u> | Aug 25, 1989 |
|-----------|---|---|------------------|-----------|----------------|------------|--------------|

PERMETHRIN

| | | | | | | |
|-----------|--|------------------|-----------|----------------|------------|--------------|
| <u>AB</u> | | ACTAVIS LABS | <u>5%</u> | <u>A074806</u> | <u>001</u> | Jan 23, 1998 |
| <u>AB</u> | | PERRIGO NEW YORK | <u>5%</u> | <u>A076369</u> | <u>001</u> | Apr 21, 2003 |

PERPHENAZINE

TABLET;ORAL

PERPHENAZINE

| | | | | | | |
|-----------|---|---------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | MYLAN PHARMS INC | <u>2MG</u> | <u>A206691</u> | <u>001</u> | Apr 14, 2017 |
| <u>AB</u> | | | <u>4MG</u> | <u>A206691</u> | <u>002</u> | Apr 14, 2017 |
| <u>AB</u> | | | <u>8MG</u> | <u>A206691</u> | <u>003</u> | Apr 14, 2017 |
| <u>AB</u> | | | <u>16MG</u> | <u>A206691</u> | <u>004</u> | Apr 14, 2017 |
| <u>AB</u> | | SANDOZ | <u>2MG</u> | <u>A089685</u> | <u>002</u> | Dec 08, 1988 |
| <u>AB</u> | | | <u>4MG</u> | <u>A089685</u> | <u>003</u> | Dec 08, 1988 |
| <u>AB</u> | | | <u>8MG</u> | <u>A089685</u> | <u>001</u> | Dec 08, 1988 |
| <u>AB</u> | ! | | <u>16MG</u> | <u>A089685</u> | <u>004</u> | Dec 08, 1988 |
| <u>AB</u> | | VINTAGE PHARMS | <u>2MG</u> | <u>A040226</u> | <u>001</u> | Dec 31, 1998 |
| <u>AB</u> | | | <u>4MG</u> | <u>A040226</u> | <u>002</u> | Dec 31, 1998 |
| <u>AB</u> | | | <u>8MG</u> | <u>A040226</u> | <u>003</u> | Dec 31, 1998 |
| <u>AB</u> | | | <u>16MG</u> | <u>A040226</u> | <u>004</u> | Dec 31, 1998 |
| <u>AB</u> | | WATSON LABS INC | <u>2MG</u> | <u>A207582</u> | <u>001</u> | Oct 17, 2016 |
| <u>AB</u> | | | <u>4MG</u> | <u>A207582</u> | <u>002</u> | Oct 17, 2016 |
| <u>AB</u> | | | <u>8MG</u> | <u>A207582</u> | <u>003</u> | Oct 17, 2016 |
| <u>AB</u> | | | <u>16MG</u> | <u>A207582</u> | <u>004</u> | Oct 17, 2016 |
| <u>AB</u> | | WILSHIRE PHARMS INC | <u>2MG</u> | <u>A205973</u> | <u>001</u> | Dec 17, 2015 |
| <u>AB</u> | | | <u>4MG</u> | <u>A205973</u> | <u>002</u> | Dec 17, 2015 |
| <u>AB</u> | | | <u>8MG</u> | <u>A205973</u> | <u>003</u> | Dec 17, 2015 |

PRESCRIPTION DRUG PRODUCT LIST

PERPHENAZINE

TABLET; ORAL

PERPHENAZINE

| | | | | | |
|-----------|--|-------------|----------------|------------|--------------|
| <u>AB</u> | | <u>16MG</u> | <u>A205973</u> | <u>004</u> | Dec 17, 2015 |
|-----------|--|-------------|----------------|------------|--------------|

PHENDIMETRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

PHENDIMETRAZINE TARTRATE

| | | | | | |
|---|---------------|-------|---------|-----|--|
| + | VIRTUS PHARMS | 105MG | N018074 | 001 | |
|---|---------------|-------|---------|-----|--|

TABLET; ORAL

BONTRIL PDM

| | | | | | |
|-----------|---|---------|-------------|----------------|------------|
| <u>AA</u> | ! | VALEANT | <u>35MG</u> | <u>A085272</u> | <u>001</u> |
|-----------|---|---------|-------------|----------------|------------|

PHENDIMETRAZINE TARTRATE

| | | | | | | |
|-----------|--|----------------|-------------|----------------|------------|--------------|
| <u>AA</u> | | ELITE LABS INC | <u>35MG</u> | <u>A040762</u> | <u>001</u> | Jan 28, 2008 |
|-----------|--|----------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AA</u> | | | <u>35MG</u> | <u>A203600</u> | <u>001</u> | Dec 27, 2017 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|----------|-------------|----------------|------------|--------------|
| <u>AA</u> | | KVK TECH | <u>35MG</u> | <u>A091042</u> | <u>001</u> | Aug 31, 2010 |
|-----------|--|----------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--------|-------------|----------------|------------|--------------|
| <u>AA</u> | | MIKART | <u>35MG</u> | <u>A089452</u> | <u>001</u> | Oct 30, 1991 |
|-----------|--|--------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|---------------|-------------|----------------|------------|--|
| <u>AA</u> | | VIRTUS PHARMS | <u>35MG</u> | <u>A085588</u> | <u>001</u> | |
|-----------|--|---------------|-------------|----------------|------------|--|

PHENELZINE SULFATE

TABLET; ORAL

NARDIL

| | | | | | | |
|-----------|---|---|-------------|---------------------|----------------|------------|
| <u>AB</u> | + | ! | PARKE DAVIS | <u>EQ 15MG BASE</u> | <u>N011909</u> | <u>002</u> |
|-----------|---|---|-------------|---------------------|----------------|------------|

PHENELZINE SULFATE

| | | | | | | |
|-----------|--|----------------|---------------------|----------------|------------|--------------|
| <u>AB</u> | | NOVEL LABS INC | <u>EQ 15MG BASE</u> | <u>A200181</u> | <u>001</u> | Dec 08, 2010 |
|-----------|--|----------------|---------------------|----------------|------------|--------------|

PHENOXYBENZAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIBENZYLINE

| | | | | | | |
|-----------|---|---|-------------------------|-------------|----------------|------------|
| <u>AB</u> | + | ! | CONCORDIA PHARMS INC | <u>10MG</u> | <u>N008708</u> | <u>001</u> |
|-----------|---|---|-------------------------|-------------|----------------|------------|

PHENOXYBENZAMINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|---------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | PAR PHARM INC | <u>10MG</u> | <u>A204522</u> | <u>001</u> | Jan 24, 2017 |
|-----------|--|---------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>10MG</u> | <u>A201050</u> | <u>001</u> | Jul 16, 2012 |
|-----------|--|-------------------------|-------------|----------------|------------|--------------|

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

ADIPEX-P

| | | | | | | |
|-----------|---|------|---------------|----------------|------------|--------------|
| <u>AA</u> | ! | TEVA | <u>37.5MG</u> | <u>A088023</u> | <u>001</u> | Aug 02, 1983 |
|-----------|---|------|---------------|----------------|------------|--------------|

PHENTERMINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|---------------------|-------------|----------------|------------|--------------|
| <u>AA</u> | | AUROLIFE PHARMA LLC | <u>15MG</u> | <u>A204318</u> | <u>001</u> | Nov 09, 2016 |
|-----------|--|---------------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AA</u> | | | <u>30MG</u> | <u>A204318</u> | <u>002</u> | Nov 09, 2016 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|------|-------------|----------------|------------|--------------|
| <u>AA</u> | | BARR | <u>15MG</u> | <u>A090591</u> | <u>001</u> | Mar 18, 2010 |
|-----------|--|------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AA</u> | | | <u>30MG</u> | <u>A090591</u> | <u>002</u> | Mar 18, 2010 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|------------|-------------|----------------|------------|--------------|
| <u>AA</u> | | ELITE LABS | <u>15MG</u> | <u>A202248</u> | <u>001</u> | Sep 28, 2012 |
|-----------|--|------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AA</u> | | | <u>30MG</u> | <u>A202248</u> | <u>002</u> | Sep 28, 2012 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|----------------|---------------|----------------|------------|--------------|
| <u>AA</u> | | ELITE LABS INC | <u>37.5MG</u> | <u>A040228</u> | <u>001</u> | Jun 19, 1997 |
|-----------|--|----------------|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|----------------|-------------|----------------|------------|--------------|
| <u>AA</u> | | INVAGEN PHARMS | <u>15MG</u> | <u>A202858</u> | <u>001</u> | Feb 14, 2014 |
|-----------|--|----------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AA</u> | | | <u>30MG</u> | <u>A202858</u> | <u>002</u> | Feb 14, 2014 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AA</u> | | | <u>30MG</u> | <u>A204414</u> | <u>001</u> | May 05, 2014 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|---------------|----------------|------------|--------------|
| <u>AA</u> | | | <u>37.5MG</u> | <u>A202846</u> | <u>001</u> | Feb 05, 2014 |
|-----------|--|--|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|----------|-------------|----------------|------------|--------------|
| <u>AA</u> | | KVK TECH | <u>15MG</u> | <u>A040886</u> | <u>002</u> | Mar 31, 2008 |
|-----------|--|----------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AA</u> | | | <u>30MG</u> | <u>A040875</u> | <u>001</u> | Mar 21, 2008 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AA</u> | | | <u>30MG</u> | <u>A040886</u> | <u>001</u> | Mar 31, 2008 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|---------------|----------------|------------|--------------|
| <u>AA</u> | | | <u>37.5MG</u> | <u>A040887</u> | <u>001</u> | Apr 24, 2008 |
|-----------|--|--|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|---------|-------------|----------------|------------|--------------|
| <u>AA</u> | | LANNETT | <u>15MG</u> | <u>A087022</u> | <u>002</u> | Jan 20, 2012 |
|-----------|--|---------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AA</u> | | | <u>30MG</u> | <u>A087022</u> | <u>001</u> | Feb 03, 1983 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|----------------|-------------|----------------|------------|--------------|
| <u>AA</u> | | LANNETT CO INC | <u>30MG</u> | <u>A091359</u> | <u>001</u> | Jul 16, 2010 |
|-----------|--|----------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|---------------|----------------|------------|--------------|
| <u>AA</u> | | | <u>37.5MG</u> | <u>A201961</u> | <u>001</u> | Jul 20, 2011 |
|-----------|--|--|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|------------|-------------|----------------|------------|--------------|
| <u>AA</u> | | NUVO PHARM | <u>15MG</u> | <u>A205019</u> | <u>001</u> | Dec 05, 2014 |
|-----------|--|------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AA</u> | | | <u>30MG</u> | <u>A205019</u> | <u>002</u> | Dec 05, 2014 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|---------------|----------------|------------|--------------|
| <u>AA</u> | | | <u>37.5MG</u> | <u>A205017</u> | <u>001</u> | Sep 25, 2014 |
|-----------|--|--|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|--------|-------------|----------------|------------|--|
| <u>AA</u> | ! | SANDOZ | <u>15MG</u> | <u>A087190</u> | <u>002</u> | |
|-----------|---|--------|-------------|----------------|------------|--|

| | | | | | | |
|-----------|---|--|-------------|----------------|------------|--------------|
| <u>AA</u> | ! | | <u>30MG</u> | <u>A086945</u> | <u>001</u> | Jul 20, 1983 |
|-----------|---|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|--|-------------|----------------|------------|--|
| <u>AA</u> | ! | | <u>30MG</u> | <u>A087190</u> | <u>001</u> | |
|-----------|---|--|-------------|----------------|------------|--|

| | | | | | | |
|-----------|--|-------------------------|-------------|----------------|------------|--------------|
| <u>AA</u> | | SUN PHARM INDUSTRIES | <u>30MG</u> | <u>A040525</u> | <u>001</u> | Oct 23, 2003 |
|-----------|--|-------------------------|-------------|----------------|------------|--------------|

TABLET; ORAL

ADIPEX-P

| | | | | | | |
|-----------|---|------|---------------|----------------|------------|--|
| <u>AA</u> | ! | TEVA | <u>37.5MG</u> | <u>A085128</u> | <u>001</u> | |
|-----------|---|------|---------------|----------------|------------|--|

LOMAIRA

| | | | | | | |
|-----------|---|-------------|------------|----------------|------------|--------------|
| <u>AA</u> | ! | AVANTHI INC | <u>8MG</u> | <u>A203495</u> | <u>001</u> | Sep 12, 2016 |
|-----------|---|-------------|------------|----------------|------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

PHENTERMINE HYDROCHLORIDE

TABLET; ORAL

PHENTERMINE HYDROCHLORIDE

| | | | | | |
|-----------|-------------------------|---------------|----------------|------------|--------------|
| <u>AA</u> | AUROLIFE PHARMA LLC | <u>37.5MG</u> | <u>A203068</u> | <u>001</u> | Aug 06, 2014 |
| <u>AA</u> | BARR | <u>37.5MG</u> | <u>A090470</u> | <u>001</u> | Aug 31, 2009 |
| <u>AA</u> | ELITE LABS | <u>37.5MG</u> | <u>A200272</u> | <u>001</u> | Jan 31, 2011 |
| <u>AA</u> | ELITE LABS INC | <u>37.5MG</u> | <u>A040190</u> | <u>001</u> | May 30, 1997 |
| <u>AA</u> | INVAGEN PHARMS | <u>37.5MG</u> | <u>A202942</u> | <u>001</u> | Feb 05, 2014 |
| <u>AA</u> | KVK TECH | <u>37.5MG</u> | <u>A040876</u> | <u>001</u> | Mar 31, 2008 |
| <u>AA</u> | KVK TECH INC | <u>8MG</u> | <u>A203436</u> | <u>001</u> | Mar 17, 2017 |
| <u>AA</u> | LANNETT | <u>37.5MG</u> | <u>A040555</u> | <u>001</u> | Apr 15, 2005 |
| <u>AA</u> | NOVAST LABS | <u>37.5MG</u> | <u>A091451</u> | <u>001</u> | Sep 21, 2012 |
| <u>AA</u> | NUVO PHARM | <u>37.5MG</u> | <u>A205008</u> | <u>001</u> | Sep 25, 2014 |
| <u>AA</u> | POLYGEN PHARMS | <u>37.5MG</u> | <u>A206342</u> | <u>001</u> | Nov 18, 2016 |
| <u>AA</u> | PRINSTON INC | <u>37.5MG</u> | <u>A040377</u> | <u>001</u> | Jan 04, 2002 |
| <u>AA</u> | SUN PHARM INDS INC | <u>37.5MG</u> | <u>A040790</u> | <u>001</u> | Aug 21, 2007 |
| <u>AA</u> | SUN PHARM INDUSTRIES | <u>37.5MG</u> | <u>A040526</u> | <u>001</u> | Oct 23, 2003 |

TABLET, ORALLY DISINTEGRATING; ORAL

PHENTERMINE HYDROCHLORIDE

| | | | | |
|------------------|--------|---------|-----|--------------|
| ZYDUS PHARMS USA | 15MG | A204663 | 001 | Jun 28, 2017 |
| INC | 30MG | A204663 | 002 | Jun 28, 2017 |
| | 37.5MG | A204663 | 003 | Jun 28, 2017 |

PHENTERMINE HYDROCHLORIDE; TOPIRAMATE

CAPSULE, EXTENDED RELEASE; ORAL

QSYMIA

| | | | | | |
|---|-------|-----------------------|---------|-----|--------------|
| + | VIVUS | EQ 3.75MG BASE; 23MG | N022580 | 001 | Jul 17, 2012 |
| + | | EQ 7.5MG BASE; 46MG | N022580 | 002 | Jul 17, 2012 |
| + | | EQ 11.25MG BASE; 69MG | N022580 | 003 | Jul 17, 2012 |
| + | ! | EQ 15MG BASE; 92MG | N022580 | 004 | Jul 17, 2012 |

PHENTOLAMINE MESYLATE

INJECTABLE; INJECTION

PHENTOLAMINE MESYLATE

| | | | | | | |
|-----------|---------------------------|-------------------|----------------|------------|--------------|--------------|
| <u>AP</u> | PRECISION DOSE INC | <u>5MG/VIAL</u> | <u>A207686</u> | <u>001</u> | Jul 14, 2017 | |
| <u>AP</u> | ! WEST-WARD PHARMS INT | <u>5MG/VIAL</u> | <u>A040235</u> | <u>001</u> | Mar 11, 1998 | |
| | ORAVERSE | | | | | |
| + | ! | SEPTODONT HOLDING | 0.4MG/1.7ML | N022159 | 001 | May 09, 2008 |

PHENYLEPHRINE HYDROCHLORIDE

SOLUTION; INTRAVENOUS

PHENYLEPHRINE HYDROCHLORIDE

| | | | | | |
|-----------|----------------------------|-----------------------------|----------------|------------|--------------|
| <u>AP</u> | AMNEAL PHARMS CO | <u>10MG/ML (10MG/ML)</u> | <u>A211079</u> | <u>001</u> | Jul 05, 2018 |
| <u>AP</u> | | <u>50MG/5ML (10MG/ML)</u> | <u>A211078</u> | <u>001</u> | Jul 19, 2018 |
| <u>AP</u> | | <u>100MG/10ML (10MG/ML)</u> | <u>A211078</u> | <u>002</u> | Jul 19, 2018 |
| <u>AP</u> | CIPLA | <u>10MG/ML (10MG/ML)</u> | <u>A210334</u> | <u>001</u> | Apr 27, 2018 |
| <u>AP</u> | | <u>50MG/5ML (10MG/ML)</u> | <u>A210333</u> | <u>001</u> | Apr 27, 2018 |
| <u>AP</u> | | <u>100MG/10ML (10MG/ML)</u> | <u>A210333</u> | <u>002</u> | Apr 27, 2018 |
| <u>AP</u> | PAR STERILE PRODUCTS | <u>10MG/ML (10MG/ML)</u> | <u>A210025</u> | <u>001</u> | Dec 21, 2018 |
| <u>AP</u> | | <u>50MG/5ML (10MG/ML)</u> | <u>A210025</u> | <u>002</u> | Dec 21, 2018 |
| <u>AP</u> | | <u>100MG/10ML (10MG/ML)</u> | <u>A210025</u> | <u>003</u> | Dec 21, 2018 |
| <u>AP</u> | +! WEST WARD PHARM CORP | <u>10MG/ML (10MG/ML)</u> | <u>N203826</u> | <u>001</u> | Dec 20, 2012 |

VAZCULEP

| | | | | | | |
|-----------|---|---------------|-----------------------------|----------------|------------|--------------|
| <u>AP</u> | + | AVADEL LEGACY | <u>10MG/ML (10MG/ML)</u> | <u>N204300</u> | <u>001</u> | Jun 27, 2014 |
| <u>AP</u> | + | | <u>50MG/5ML (10MG/ML)</u> | <u>N204300</u> | <u>002</u> | Jun 27, 2014 |
| <u>AP</u> | + | ! | <u>100MG/10ML (10MG/ML)</u> | <u>N204300</u> | <u>003</u> | Jun 27, 2014 |

SOLUTION/DROPS; OPHTHALMIC

PHENYLEPHRINE HYDROCHLORIDE

| | | | | | | |
|---|---|-----------------|------|---------|-----|--------------|
| + | ! | AKORN INC | 2.5% | N207926 | 001 | Jan 15, 2015 |
| + | ! | | 10% | N207926 | 002 | Jan 15, 2015 |
| + | ! | PARAGON BIOTECK | 2.5% | N203510 | 001 | Mar 21, 2013 |
| + | ! | | 10% | N203510 | 002 | Mar 21, 2013 |

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE

| | | | | | |
|-----------|-------------------------------------------------------------------|----------------------------|----------------|------------|--------------|
| <u>AA</u> | HI-TECH PHARMACAL | <u>5MG/5ML; 6.25MG/5ML</u> | <u>A040675</u> | <u>001</u> | Dec 23, 2014 |
| <u>AA</u> | ! VINTAGE | <u>5MG/5ML; 6.25MG/5ML</u> | <u>A040654</u> | <u>001</u> | Dec 07, 2006 |
| | <u>PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE</u> | | | | |
| <u>AA</u> | AMNEAL PHARMS | <u>5MG/5ML; 6.25MG/5ML</u> | <u>A040902</u> | <u>001</u> | Aug 25, 2009 |

PRESCRIPTION DRUG PRODUCT LIST

PHENYTOIN

SUSPENSION; ORAL

DILANTIN-125

| | | | | | | |
|-----------|------------|-------------|------------------|----------------|------------|--|
| AB | + ! | PARKE DAVIS | 125MG/5ML | N008762 | 001 | |
|-----------|------------|-------------|------------------|----------------|------------|--|

PHENYTOIN

| | | | | | | |
|-----------|--|------------------|------------------|----------------|------------|--------------|
| AB | | TARO | 125MG/5ML | A040521 | 001 | Mar 08, 2004 |
| AB | | VISTAPHARM | 125MG/5ML | A040342 | 001 | Jan 31, 2001 |
| AB | | | 125MG/5ML | A040610 | 001 | Aug 18, 2005 |
| AB | | WOCKHARDT BIO AG | 125MG/5ML | A040420 | 001 | Apr 19, 2002 |

TABLET, CHEWABLE; ORAL

DILANTIN

| | | | | | | |
|-----------|----------|--------|-------------|----------------|------------|--|
| AB | ! | PFIZER | 50MG | A084427 | 001 | |
|-----------|----------|--------|-------------|----------------|------------|--|

PHENYTOIN

| | | | | | | |
|-----------|--|------------------|-------------|----------------|------------|--------------|
| AB | | EPIC PHARMA LLC | 50MG | A040884 | 001 | Nov 28, 2014 |
| AB | | MYLAN PHARMS INC | 50MG | A200691 | 001 | Dec 26, 2012 |
| AB | | TARO | 50MG | A200565 | 001 | Apr 17, 2014 |

PHENYTOIN SODIUM

CAPSULE; ORAL

DILANTIN

| | | | | | | |
|-----------|----------|-------------|-----------------------|----------------|------------|--|
| AB | ! | PARKE-DAVIS | 100MG EXTENDED | A084349 | 002 | |
|-----------|----------|-------------|-----------------------|----------------|------------|--|

EXTENDED PHENYTOIN SODIUM

| | | | | | | |
|-----------|--|---------------------|-----------------------|----------------|------------|--------------|
| AB | | AMNEAL PHARMS NY | 100MG EXTENDED | A040765 | 001 | Nov 12, 2008 |
| AB | | MYLAN | 100MG EXTENDED | A040298 | 001 | Dec 28, 1998 |
| AB | | SUN PHARM INDS | 200MG EXTENDED | A040731 | 001 | Jun 30, 2008 |
| AB | | | 300MG EXTENDED | A040731 | 002 | Jun 30, 2008 |
| AB | | SUN PHARM INDS (IN) | 100MG EXTENDED | A040621 | 001 | Dec 11, 2006 |
| AB | | TARO | 100MG EXTENDED | A040684 | 001 | Sep 05, 2006 |

PHENYTEK

| | | | | | | |
|-----------|----------|-------|-----------------------|----------------|------------|--------------|
| AB | | MYLAN | 200MG EXTENDED | A040298 | 002 | Dec 06, 2001 |
| AB | ! | | 300MG EXTENDED | A040298 | 003 | Dec 06, 2001 |

PHENYTOIN SODIUM

| | | | | | | |
|-----------|--|----------------------|-----------------------|----------------|------------|--------------|
| AB | | AUROBINDO PHARMA LTD | 100MG EXTENDED | A204309 | 001 | Jun 10, 2015 |
|-----------|--|----------------------|-----------------------|----------------|------------|--------------|

DILANTIN

! PARKE-DAVIS

| | | | | | | |
|--|--|--|---------------|---------|-----|--|
| | | | 30MG EXTENDED | A084349 | 001 | |
|--|--|--|---------------|---------|-----|--|

INJECTABLE; INJECTION

PHENYTOIN SODIUM

| | | | | | | |
|-----------|----------|----------------------|----------------|----------------|------------|--------------|
| AP | | ACELLA PHARMS LLC | 50MG/ML | A040573 | 001 | Sep 13, 2006 |
| AP | | LUITPOLD | 50MG/ML | A040781 | 001 | Dec 04, 2007 |
| AP | ! | WEST-WARD PHARMS INT | 50MG/ML | A084307 | 001 | |

PHYTONADIONE

INJECTABLE; INJECTION

PHYTONADIONE

| | | | | | | |
|----|----------|-----------------|-----------|---------|-----|--|
| BP | ! | INTL MEDICATION | 1MG/0.5ML | A083722 | 001 | |
|----|----------|-----------------|-----------|---------|-----|--|

VITAMIN K1

| | | | | | | |
|----|----------|---------|-----------|---------|-----|--------------|
| BP | ! | HOSPIRA | 1MG/0.5ML | A087954 | 001 | Jul 25, 1983 |
| | ! | | 10MG/ML | A087955 | 001 | Jul 25, 1983 |

TABLET; ORAL

MEPHYTON

| | | | | | | |
|-----------|------------|----------------|------------|----------------|------------|--|
| AB | + ! | VALEANT PHARMS | 5MG | N010104 | 003 | |
|-----------|------------|----------------|------------|----------------|------------|--|

PHYTONADIONE

| | | | | | | |
|-----------|--|------------------|------------|----------------|------------|--------------|
| AB | | AMNEAL PHARMS CO | 5MG | A209373 | 001 | May 11, 2018 |
|-----------|--|------------------|------------|----------------|------------|--------------|

PILOCARPINE HYDROCHLORIDE

SOLUTION; OPHTHALMIC

ISOPTO CARPINE

| | | | | | | |
|-----------|------------|----------------------|-----------|----------------|------------|--------------|
| AT | + | NOVARTIS PHARMS CORP | 1% | N200890 | 001 | Jun 22, 2010 |
| AT | + | | 2% | N200890 | 002 | Jun 22, 2010 |
| AT | + ! | | 4% | N200890 | 003 | Jun 22, 2010 |

PILOCARPINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|-----------|-----------|----------------|------------|--------------|
| AT | | AKORN INC | 1% | A204398 | 001 | Sep 27, 2017 |
| AT | | | 2% | A204398 | 002 | Sep 27, 2017 |
| AT | | | 4% | A204398 | 003 | Sep 27, 2017 |

TABLET; ORAL

PILOCARPINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|----------------|--------------|----------------|------------|--------------|
| AB | | IMPAX LABS | 5MG | A077248 | 001 | Mar 31, 2006 |
| AB | | | 7.5MG | A077248 | 002 | Mar 31, 2006 |
| AB | | INNOGENIX | 5MG | A076963 | 001 | Dec 22, 2004 |
| AB | | | 7.5MG | A076963 | 002 | Feb 27, 2007 |
| AB | | LANNETT CO INC | 5MG | A077220 | 001 | Oct 14, 2005 |

PRESCRIPTION DRUG PRODUCT LIST

PILOCARPINE HYDROCHLORIDE

TABLET;ORAL

PILOCARPINE HYDROCHLORIDE

| | | | | |
|-----------|---------------------|--------------|--------------------|--------------|
| AB | | 7.5MG | A077220 002 | May 06, 2009 |
| AB | PERRIGO PHARMA INTL | 5MG | A076746 001 | Nov 16, 2004 |

SALAGEN

| | | | | |
|-----------|-------------|--------------|--------------------|--------------|
| AB | + EISAI INC | 5MG | N020237 001 | Mar 22, 1994 |
| AB | +! | 7.5MG | N020237 002 | Apr 18, 2003 |

PIMAVANSERIN TARTRATE

CAPSULE;ORAL

NUPLAZID

| | | | | |
|----|-------------------|--------------|-------------|--------------|
| +! | ACADIA PHARMS INC | EQ 34MG BASE | N210793 001 | Jun 28, 2018 |
|----|-------------------|--------------|-------------|--------------|

TABLET;ORAL

NUPLAZID

| | | | | |
|----|-------------------|--------------|-------------|--------------|
| +! | ACADIA PHARMS INC | EQ 10MG BASE | N207318 002 | Jun 28, 2018 |
| +! | | EQ 17MG BASE | N207318 001 | Apr 29, 2016 |

PIMECROLIMUS

CREAM;TOPICAL

ELIDEL

| | | | | | |
|-----------|----|-----------------|-----------|--------------------|--------------|
| AB | +! | VALEANT BERMUDA | 1% | N021302 001 | Dec 13, 2001 |
|-----------|----|-----------------|-----------|--------------------|--------------|

PIMECROLIMUS

| | | | | | |
|-----------|--|---------------------|-----------|--------------------|--------------|
| AB | | ACTAVIS LABS UT INC | 1% | A209345 001 | Dec 27, 2018 |
|-----------|--|---------------------|-----------|--------------------|--------------|

PIMOZIDE

TABLET;ORAL

ORAP

| | | | | |
|-----------|--------|------------|--------------------|--------------|
| AB | + TEVA | 1MG | N017473 003 | Aug 27, 1997 |
| AB | +! | | N017473 001 | Jul 31, 1984 |

PIMOZIDE

| | | | | | |
|-----------|--|-----------|------------|--------------------|--------------|
| AB | | PAR PHARM | 1MG | A204521 001 | Sep 28, 2015 |
| AB | | | 2MG | A204521 002 | Sep 28, 2015 |

PINDOLOL

TABLET;ORAL

PINDOLOL

| | | | | | |
|-----------|---|----------------------|-------------|--------------------|--------------|
| AB | | ANI PHARMS INC | 5MG | A073609 002 | Mar 29, 1993 |
| AB | | | 10MG | A073609 001 | Mar 29, 1993 |
| AB | | MYLAN PHARMS INC | 5MG | A074019 001 | Sep 03, 1992 |
| AB | ! | | 10MG | A074019 002 | Sep 03, 1992 |
| AB | | NOSTRUM LABS INC | 5MG | A205415 001 | Jan 13, 2016 |
| AB | | | 10MG | A205415 002 | Jan 13, 2016 |
| AB | | SUN PHARM INDUSTRIES | 5MG | A074063 001 | Jan 27, 1994 |
| AB | | | 10MG | A074063 002 | Jan 27, 1994 |
| AB | | ZYDUS PHARMS USA INC | 5MG | A209866 001 | Aug 18, 2017 |
| AB | | | 10MG | A209866 002 | Aug 18, 2017 |

PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

ACTOS

| | | | | |
|-----------|---------------------|---------------------|--------------------|--------------|
| AB | + TAKEDA PHARMS USA | EQ 15MG BASE | N021073 001 | Jul 15, 1999 |
| AB | + | EQ 30MG BASE | N021073 002 | Jul 15, 1999 |
| AB | +! | EQ 45MG BASE | N021073 003 | Jul 15, 1999 |

PIOGLITAZONE HYDROCHLORIDE

| | | | | | |
|-----------|--|----------------------|---------------------|--------------------|--------------|
| AB | | ACCORD HLTHCARE | EQ 15MG BASE | A200044 001 | Feb 13, 2013 |
| AB | | | EQ 30MG BASE | A200044 002 | Feb 13, 2013 |
| AB | | | EQ 45MG BASE | A200044 003 | Feb 13, 2013 |
| AB | | AUROBINDO PHARMA LTD | EQ 15MG BASE | A200268 001 | Feb 13, 2013 |
| AB | | | EQ 30MG BASE | A200268 002 | Feb 13, 2013 |
| AB | | | EQ 45MG BASE | A200268 003 | Feb 13, 2013 |
| AB | | BRECKENRIDGE PHARM | EQ 15MG BASE | A078472 001 | Feb 13, 2013 |
| AB | | | EQ 30MG BASE | A078472 002 | Feb 13, 2013 |
| AB | | | EQ 45MG BASE | A078472 003 | Feb 13, 2013 |
| AB | | CELLTRION | EQ 15MG BASE | A076798 001 | Oct 26, 2012 |
| AB | | | EQ 30MG BASE | A076798 002 | Oct 26, 2012 |
| AB | | | EQ 45MG BASE | A076798 003 | Oct 26, 2012 |
| AB | | LUPIN LTD | EQ 15MG BASE | A204133 001 | Apr 07, 2014 |
| AB | | | EQ 30MG BASE | A204133 002 | Apr 07, 2014 |
| AB | | | EQ 45MG BASE | A204133 003 | Apr 07, 2014 |
| AB | | MACLEODS PHARMS LTD | EQ 15MG BASE | A202467 001 | Feb 06, 2013 |
| AB | | | EQ 30MG BASE | A202467 002 | Feb 06, 2013 |

PRESCRIPTION DRUG PRODUCT LIST

PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

PIOGLITAZONE HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|---------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A202467 003</u> | Feb 06, 2013 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 15MG BASE</u> | <u>A076801 001</u> | Aug 17, 2012 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A076801 002</u> | Aug 17, 2012 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A076801 003</u> | Aug 17, 2012 |
| <u>AB</u> | NEOPHARMA | <u>EQ 15MG BASE</u> | <u>A078383 001</u> | Mar 12, 2013 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A078383 002</u> | Mar 12, 2013 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A078383 003</u> | Mar 12, 2013 |
| <u>AB</u> | PRINSTON INC | <u>EQ 15MG BASE</u> | <u>A207806 001</u> | Apr 17, 2018 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A207806 002</u> | Apr 17, 2018 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A207806 003</u> | Apr 17, 2018 |
| <u>AB</u> | PURACAP PHARM LLC | <u>EQ 15MG BASE</u> | <u>A206738 001</u> | Oct 06, 2017 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A206738 002</u> | Oct 06, 2017 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A206738 003</u> | Oct 06, 2017 |
| <u>AB</u> | SANDOZ | <u>EQ 15MG BASE</u> | <u>A078670 001</u> | Feb 13, 2013 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A078670 002</u> | Feb 13, 2013 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A078670 003</u> | Feb 13, 2013 |
| <u>AB</u> | TEVA PHARMS USA | <u>EQ 15MG BASE</u> | <u>A077210 001</u> | Jan 10, 2014 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A077210 002</u> | Jan 10, 2014 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A077210 003</u> | Jan 10, 2014 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>EQ 15MG BASE</u> | <u>A091298 001</u> | Feb 13, 2013 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A091298 002</u> | Feb 13, 2013 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A091298 003</u> | Feb 13, 2013 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 15MG BASE</u> | <u>A202456 001</u> | Feb 13, 2013 |
| <u>AB</u> | | <u>EQ 30MG BASE</u> | <u>A202456 002</u> | Feb 13, 2013 |
| <u>AB</u> | | <u>EQ 45MG BASE</u> | <u>A202456 003</u> | Feb 13, 2013 |

PIPERACILLIN SODIUM

INJECTABLE; INJECTION

PIPERACILLIN

| | | | | |
|---|-------------------------|-------------------|-------------|--------------|
| ! | ISTITUTO BIO ITA SPA | EQ 2GM BASE/VIAL | A065114 001 | Nov 14, 2003 |
| ! | | EQ 3GM BASE/VIAL | A065114 002 | Nov 14, 2003 |
| ! | | EQ 4GM BASE/VIAL | A065114 003 | Nov 14, 2003 |
| ! | | EQ 40GM BASE/VIAL | A065157 001 | Jul 12, 2004 |

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

PIPERACILLIN AND TAZOBACTAM

| | | | | |
|-----------|-------------------------|---------------------------------------------|--------------------|--------------|
| <u>AP</u> | APOLLO PHARMS INC | <u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u> | <u>A207847 001</u> | Jan 13, 2017 |
| <u>AP</u> | | <u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u> | <u>A207847 002</u> | Jan 13, 2017 |
| <u>AP</u> | | <u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u> | <u>A207848 002</u> | Jan 13, 2017 |
| <u>AP</u> | | <u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A207847 003</u> | Jan 13, 2017 |
| <u>AP</u> | | <u>EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL</u> | <u>A207848 001</u> | May 11, 2018 |
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u> | <u>A065498 001</u> | May 23, 2011 |
| <u>AP</u> | | <u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u> | <u>A065498 002</u> | May 23, 2011 |
| <u>AP</u> | | <u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A065498 003</u> | May 23, 2011 |
| <u>AP</u> | FRESENIUS KABI | <u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u> | <u>A203719 001</u> | May 18, 2018 |
| <u>AP</u> | | <u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u> | <u>A203719 002</u> | May 18, 2018 |
| <u>AP</u> | | <u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A203719 003</u> | May 18, 2018 |
| <u>AP</u> | | <u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u> | <u>A203720 001</u> | May 11, 2018 |
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL</u> | <u>A206204 001</u> | May 11, 2018 |
| <u>AP</u> | HOSPIRA INC | <u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u> | <u>A065386 001</u> | Sep 15, 2009 |
| <u>AP</u> | | <u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u> | <u>A065386 002</u> | Sep 15, 2009 |
| <u>AP</u> | | <u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A065386 003</u> | Sep 15, 2009 |
| <u>AP</u> | | <u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u> | <u>A065446 001</u> | Sep 15, 2009 |
| <u>AP</u> | ISTITUTO BIO ITA SPA | <u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u> | <u>A065523 001</u> | May 31, 2011 |
| <u>AP</u> | | <u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u> | <u>A065523 002</u> | May 31, 2011 |
| <u>AP</u> | | <u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A065523 003</u> | May 31, 2011 |
| <u>AP</u> | | <u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u> | <u>A090498 001</u> | May 31, 2011 |
| <u>AP</u> | MYLAN LABS LTD | <u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u> | <u>A065458 001</u> | Aug 15, 2014 |
| <u>AP</u> | | <u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u> | <u>A065458 002</u> | Aug 15, 2014 |
| <u>AP</u> | | <u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A065458 003</u> | Aug 15, 2014 |
| <u>AP</u> | QILU TIANHE | <u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u> | <u>A204959 001</u> | Aug 10, 2018 |
| <u>AP</u> | | <u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u> | <u>A204959 002</u> | Aug 10, 2018 |
| <u>AP</u> | | <u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A204959 003</u> | Aug 10, 2018 |
| <u>AP</u> | SANDOZ | <u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u> | <u>A065362 001</u> | Oct 21, 2010 |
| <u>AP</u> | | <u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u> | <u>A065362 001</u> | Oct 21, 2010 |
| <u>AP</u> | | <u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u> | <u>A065362 002</u> | Oct 21, 2010 |

PRESCRIPTION DRUG PRODUCT LIST

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

PIPERACILLIN AND TAZOBACTAM

| | | | | |
|-----------|------------------|---------------------------------------------|--------------------|--------------|
| <u>AP</u> | | <u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u> | <u>A065363 002</u> | Oct 21, 2010 |
| <u>AP</u> | | <u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A065362 003</u> | Oct 21, 2010 |
| <u>AP</u> | | <u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A065363 003</u> | Oct 21, 2010 |
| <u>AP</u> | SANDOZ INC | <u>EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL</u> | <u>A203557 001</u> | Oct 29, 2014 |
| <u>AP</u> | WOCKHARDT BIO AG | <u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u> | <u>A206996 001</u> | Mar 22, 2017 |
| <u>AP</u> | | <u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u> | <u>A206996 002</u> | Mar 22, 2017 |
| <u>AP</u> | | <u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>A206996 003</u> | Mar 22, 2017 |
| <u>AP</u> | | <u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u> | <u>A207146 001</u> | Mar 17, 2017 |

ZOSYN

| | | | | | |
|-----------|----|----------------------------|---------------------------------------------|--------------------|--------------|
| <u>AP</u> | +! | WYETH PHARMS | <u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u> | <u>N050684 001</u> | Oct 22, 1993 |
| <u>AP</u> | +! | | <u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u> | <u>N050684 002</u> | Oct 22, 1993 |
| <u>AP</u> | +! | | <u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u> | <u>N050684 003</u> | Oct 22, 1993 |
| <u>AP</u> | +! | | <u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u> | <u>N050684 004</u> | Oct 22, 1993 |
| | | ZOSYN IN PLASTIC CONTAINER | | | |
| | +! | WYETH PHARMS | EQ 40MG BASE/ML;EQ 5MG BASE/ML | N050750 001 | Feb 24, 1998 |
| | +! | | EQ 60MG BASE/ML;EQ 7.5MG BASE/ML | N050750 002 | Feb 24, 1998 |
| | +! | | EQ 4GM BASE/100ML;EQ 500MG BASE/100ML | N050750 003 | Feb 24, 1998 |

PIRFENIDONE

CAPSULE; ORAL

ESBRIET

+! GENENTECH INC 267MG N022535 001 Oct 15, 2014

TABLET; ORAL

ESBRIET

+ GENENTECH INC 267MG N208780 001 Jan 11, 2017

+! 801MG N208780 003 Jan 11, 2017

PIROXICAM

CAPSULE; ORAL

FELDENE

| | | | | | |
|-----------|----|--------|-------------|--------------------|--------------|
| <u>AB</u> | + | PFIZER | <u>10MG</u> | <u>N018147 002</u> | Apr 06, 1982 |
| <u>AB</u> | +! | | <u>20MG</u> | <u>N018147 003</u> | Apr 06, 1982 |

PIROXICAM

| | | | | | |
|-----------|--|----------------------|-------------|--------------------|--------------|
| <u>AB</u> | | BRECKENRIDGE PHARM | <u>10MG</u> | <u>A208991 001</u> | Feb 21, 2018 |
| <u>AB</u> | | | <u>20MG</u> | <u>A208991 002</u> | Feb 21, 2018 |
| <u>AB</u> | | FLAMINGO PHARMS | <u>10MG</u> | <u>A207938 001</u> | Sep 09, 2016 |
| <u>AB</u> | | | <u>20MG</u> | <u>A207938 002</u> | Sep 09, 2016 |
| <u>AB</u> | | HIKMA PHARMS | <u>10MG</u> | <u>A209256 001</u> | Aug 11, 2017 |
| <u>AB</u> | | | <u>20MG</u> | <u>A209256 002</u> | Aug 11, 2017 |
| <u>AB</u> | | MICRO LABS | <u>10MG</u> | <u>A206152 001</u> | Dec 29, 2017 |
| <u>AB</u> | | | <u>20MG</u> | <u>A206152 002</u> | Dec 29, 2017 |
| <u>AB</u> | | MYLAN IRELAND LTD | <u>10MG</u> | <u>A074116 001</u> | Jun 15, 1993 |
| <u>AB</u> | | | <u>20MG</u> | <u>A074118 001</u> | Jun 15, 1993 |
| <u>AB</u> | | PII | <u>10MG</u> | <u>A206136 001</u> | Jun 20, 2017 |
| <u>AB</u> | | | <u>20MG</u> | <u>A206136 002</u> | Jun 20, 2017 |
| <u>AB</u> | | STRIDES PHARMA | <u>10MG</u> | <u>A210347 001</u> | Jan 26, 2018 |
| <u>AB</u> | | | <u>20MG</u> | <u>A210347 002</u> | Jan 26, 2018 |
| <u>AB</u> | | SUN PHARM INDUSTRIES | <u>10MG</u> | <u>A073536 002</u> | Jan 23, 2008 |
| <u>AB</u> | | | <u>20MG</u> | <u>A073536 001</u> | Mar 12, 1993 |
| <u>AB</u> | | TEVA | <u>10MG</u> | <u>A074131 001</u> | Dec 11, 1992 |
| <u>AB</u> | | | <u>20MG</u> | <u>A074131 002</u> | Dec 11, 1992 |
| <u>AB</u> | | UNICHEM LABS LTD | <u>10MG</u> | <u>A208340 001</u> | Apr 13, 2017 |
| <u>AB</u> | | | <u>20MG</u> | <u>A208340 002</u> | Apr 13, 2017 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>10MG</u> | <u>A205585 001</u> | Jul 17, 2018 |
| <u>AB</u> | | | <u>20MG</u> | <u>A205585 002</u> | Jul 17, 2018 |

PITAVASTATIN CALCIUM

TABLET; ORAL

LIVALO

| | | | | | |
|-----------|----|---------|--------------------|--------------------|--------------|
| <u>AB</u> | + | KOWA CO | <u>EQ 1MG BASE</u> | <u>N022363 001</u> | Aug 03, 2009 |
| <u>AB</u> | + | | <u>EQ 2MG BASE</u> | <u>N022363 002</u> | Aug 03, 2009 |
| <u>AB</u> | +! | | <u>EQ 4MG BASE</u> | <u>N022363 003</u> | Aug 03, 2009 |

PITAVASTATIN CALCIUM

| | | | | | |
|-----------|--|----------------------|--------------------|--------------------|--------------|
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>EQ 1MG BASE</u> | <u>A206015 001</u> | Dec 20, 2016 |
| <u>AB</u> | | | <u>EQ 2MG BASE</u> | <u>A206015 002</u> | Dec 20, 2016 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A206015 003</u> | Dec 20, 2016 |
| <u>AB</u> | | ORIENT PHARMA CO LTD | <u>EQ 1MG BASE</u> | <u>A205932 001</u> | Feb 03, 2017 |
| <u>AB</u> | | | <u>EQ 2MG BASE</u> | <u>A205932 002</u> | Feb 03, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

PITAVASTATIN CALCIUM

TABLET; ORAL

PITAVASTATIN CALCIUM

| | | | | |
|-----------|-----------|--------------------|--------------------|--------------|
| AB | | EQ 4MG BASE | A205932 003 | Feb 03, 2017 |
| AB | SAWAI USA | EQ 1MG BASE | A205955 001 | Feb 03, 2017 |
| AB | | EQ 2MG BASE | A205955 002 | Feb 03, 2017 |
| AB | | EQ 4MG BASE | A205955 003 | Feb 03, 2017 |

PITAVASTATIN MAGNESIUM

TABLET; ORAL

ZYPITAMAG

| | | | | |
|---|-------------------------|-------------|-------------|--------------|
| + | ZYDUS PHARMS USA INC | EQ 1MG BASE | N208379 001 | Jul 14, 2017 |
| + | | EQ 2MG BASE | N208379 002 | Jul 14, 2017 |
| + | + | EQ 4MG BASE | N208379 003 | Jul 14, 2017 |

PLAZOMICIN SULFATE

SOLUTION; INTRAVENOUS

ZEMDRI

| | | | | |
|---|--------------|--------------------------------------|-------------|--------------|
| + | ACHAOGEN INC | EQ 500MG BASE/10ML (EQ 50MG BASE/ML) | N210303 001 | Jun 25, 2018 |
|---|--------------|--------------------------------------|-------------|--------------|

PLECANATIDE

TABLET; ORAL

TRULANCE

| | | | | |
|---|----------------|-----|-------------|--------------|
| + | SYNERGY PHARMS | 3MG | N208745 001 | Jan 19, 2017 |
|---|----------------|-----|-------------|--------------|

PLERIXAFOR

SOLUTION; SUBCUTANEOUS

MOZOBIL

| | | | | |
|---|---------|----------------------|-------------|--------------|
| + | GENZYME | 24MG/1.2ML (20MG/ML) | N022311 001 | Dec 15, 2008 |
|---|---------|----------------------|-------------|--------------|

PODOFILOX

GEL; TOPICAL

CONDYLOX

| | | | | |
|---|--------------------|------|-------------|--------------|
| + | ALLERGAN SALES LLC | 0.5% | N020529 001 | Mar 13, 1997 |
|---|--------------------|------|-------------|--------------|

SOLUTION; TOPICAL

CONDYLOX

| | | | | | |
|-----------|---|--------------------|-------------|--------------------|--------------|
| AT | + | ALLERGAN SALES LLC | 0.5% | N019795 001 | Dec 13, 1990 |
|-----------|---|--------------------|-------------|--------------------|--------------|

PODOFILOX

| | | | | | |
|-----------|--|-------------|-------------|--------------------|--------------|
| AT | | PADDOCK LLC | 0.5% | A075600 001 | Jan 29, 2002 |
|-----------|--|-------------|-------------|--------------------|--------------|

POLIDOCANOL

SOLUTION; INTRAVENOUS

ASCLERA

| | | | | |
|---|-------------------------|-------------------|-------------|--------------|
| + | CHEMISCH FBRK KRSSLR | 10MG/2ML (5MG/ML) | N021201 001 | Mar 30, 2010 |
|---|-------------------------|-------------------|-------------|--------------|

| | | | | |
|---|---|--------------------|-------------|--------------|
| + | + | 20MG/2ML (10MG/ML) | N021201 002 | Mar 30, 2010 |
|---|---|--------------------|-------------|--------------|

VARITHENA

| | | | | |
|---|-----------|-------------------------|-------------|--------------|
| + | PROVENSIS | 77.5MG/7.75ML (10MG/ML) | N205098 002 | Dec 21, 2017 |
|---|-----------|-------------------------|-------------|--------------|

| | | | | |
|---|---|----------------------|-------------|--------------|
| + | + | 180MG/18ML (10MG/ML) | N205098 001 | Nov 25, 2013 |
|---|---|----------------------|-------------|--------------|

POLYETHYLENE GLYCOL 3350

FOR SOLUTION; ORAL

GLYCOLAX

| | | | | | |
|-----------|--|----------------|----------------------|--------------------|--------------|
| AA | | LANNETT CO INC | 17GM/SCOOPFUL | A076652 001 | Jul 02, 2004 |
|-----------|--|----------------|----------------------|--------------------|--------------|

POLYETHYLENE GLYCOL 3350

| | | | | | |
|-----------|--|-------------------|----------------------|--------------------|--------------|
| AA | | NEXGEN PHARMA INC | 17GM/SCOOPFUL | A077706 001 | Sep 27, 2006 |
|-----------|--|-------------------|----------------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------------|----------------------|--------------------|--------------|
| AA | | PADDOCK LLC | 17GM/SCOOPFUL | A077893 001 | May 26, 2006 |
|-----------|--|-------------|----------------------|--------------------|--------------|

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION; ORAL

LAX-LYTE WITH FLAVOR PACKS

| | | | | | |
|-----------|--|-------------|------------------------------------------------------|--------------------|--------------|
| AA | | PADDOCK LLC | 420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/BOT | A079232 001 | Feb 25, 2010 |
|-----------|--|-------------|------------------------------------------------------|--------------------|--------------|

NULYTELY

| | | | | | |
|-----------|---|-----------|------------------------------------------------------|--------------------|--------------|
| AA | + | BRAINTREE | 420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/BOT | N019797 001 | Apr 22, 1991 |
|-----------|---|-----------|------------------------------------------------------|--------------------|--------------|

NULYTELY-FLAVORED

| | | | | | |
|-----------|---|-----------|------------------------------------------------------|--------------------|--------------|
| AA | + | BRAINTREE | 420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/BOT | N019797 002 | Nov 18, 1994 |
|-----------|---|-----------|------------------------------------------------------|--------------------|--------------|

PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE

| | | | | | |
|-----------|--|--------------------|------------------------------------------------------|--------------------|--------------|
| AA | | BRECKENRIDGE PHARM | 420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/BOT | A202060 001 | Mar 08, 2017 |
|-----------|--|--------------------|------------------------------------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------|------------------------------------------------------|--------------------|--------------|
| AA | | NOVEL LABS INC | 420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/BOT | A090019 001 | May 27, 2009 |
|-----------|--|----------------|------------------------------------------------------|--------------------|--------------|

| | | | | | |
|-----------|--|----------------|------------------------------------------------------|--------------------|--------------|
| AA | | STRIDES PHARMA | 420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/BOT | A204559 001 | Apr 13, 2015 |
|-----------|--|----------------|------------------------------------------------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION;ORAL

TRILYTE

| | | | | |
|-----------|------------------|---------------------------------------------------|--------------------|--------------|
| AA | MYLAN PHARMS INC | <u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u> | <u>A076491 001</u> | Feb 05, 2004 |
|-----------|------------------|---------------------------------------------------|--------------------|--------------|

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE ANHYDROUS

FOR SOLUTION;ORAL

COLYTE WITH FLAVOR PACKS

| | | | | |
|-----------|---------------------|---------------------------------------------------------------|--------------------|--------------|
| AA | MYLAN SPECIALITY LP | <u>240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT</u> | <u>N018983 012</u> | Oct 08, 1998 |
|-----------|---------------------|---------------------------------------------------------------|--------------------|--------------|

GOLYTELY

| | | | | |
|-----------|--------------|---------------------------------------------------------------|--------------------|--------------|
| AA | +! BRAINTREE | <u>236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT</u> | <u>N019011 001</u> | Jul 13, 1984 |
|-----------|--------------|---------------------------------------------------------------|--------------------|--------------|

PEG 3350 AND ELECTROLYTES

| | | | | |
|-----------|----------------|---------------------------------------------------------------|--------------------|--------------|
| AA | NOVEL LABS INC | <u>236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT</u> | <u>A090231 001</u> | Jun 01, 2009 |
|-----------|----------------|---------------------------------------------------------------|--------------------|--------------|

| | | | | |
|-----------|--|---------------------------------------------------------------|--------------------|--------------|
| AA | | <u>240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT</u> | <u>A090186 001</u> | Jun 01, 2009 |
|-----------|--|---------------------------------------------------------------|--------------------|--------------|

POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES

| | | | | |
|-----------|----------------|---------------------------------------------------------------|--------------------|--------------|
| AA | STRIDES PHARMA | <u>236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT</u> | <u>A204558 001</u> | Dec 21, 2018 |
|-----------|----------------|---------------------------------------------------------------|--------------------|--------------|

GOLYTELY

| | | | | |
|--|--------------|------------------------------------------------------------------------|-------------|--------------|
| | +! BRAINTREE | 227.1GM/PACKET;2.82GM/PACKET;6.36GM/PACKET;5.53GM/PACKET;21.5GM/PACKET | N019011 002 | Jun 02, 1992 |
|--|--------------|------------------------------------------------------------------------|-------------|--------------|

POLYMYXIN B SULFATE

INJECTABLE;INJECTION

POLYMYXIN B SULFATE

| | | | | |
|-----------|----------------------|-----------------------------------|--------------------|--------------|
| AP | AUROBINDO PHARMA LTD | <u>EQ 500,000 UNITS BASE/VIAL</u> | <u>A206589 001</u> | Apr 04, 2016 |
|-----------|----------------------|-----------------------------------|--------------------|--------------|

| | | | | |
|-----------|--------------------|-----------------------------------|--------------------|--------------|
| AP | FRESENIUS KABI USA | <u>EQ 500,000 UNITS BASE/VIAL</u> | <u>A065372 001</u> | Jan 10, 2008 |
|-----------|--------------------|-----------------------------------|--------------------|--------------|

| | | | | |
|-----------|------------------|-----------------------------------|--------------------|--------------|
| AP | GLAND PHARMA LTD | <u>EQ 500,000 UNITS BASE/VIAL</u> | <u>A207322 001</u> | Apr 14, 2016 |
|-----------|------------------|-----------------------------------|--------------------|--------------|

| | | | | |
|-----------|-----------|-----------------------------------|--------------------|--------------|
| AP | MYLAN ASI | <u>EQ 500,000 UNITS BASE/VIAL</u> | <u>A090110 001</u> | Jun 29, 2011 |
|-----------|-----------|-----------------------------------|--------------------|--------------|

| | | | | |
|-----------|------------------------|-----------------------------------|--------------------|--|
| AP | ! WEST-WARD PHARMS INT | <u>EQ 500,000 UNITS BASE/VIAL</u> | <u>A060716 001</u> | |
|-----------|------------------------|-----------------------------------|--------------------|--|

| | | | | |
|-----------|--------------|-----------------------------------|--------------------|--------------|
| AP | X GEN PHARMS | <u>EQ 500,000 UNITS BASE/VIAL</u> | <u>A063000 001</u> | Sep 30, 1994 |
|-----------|--------------|-----------------------------------|--------------------|--------------|

| | | | | |
|-----------|-------------------|-----------------------------------|--------------------|--------------|
| AP | XELLIA PHARMS APS | <u>EQ 500,000 UNITS BASE/VIAL</u> | <u>A202766 001</u> | Jan 15, 2014 |
|-----------|-------------------|-----------------------------------|--------------------|--------------|

POLYMYXIN B SULFATE; TRIMETHOPRIM SULFATE

SOLUTION/DROPS;OPHTHALMIC

POLYTRIM

| | | | | |
|-----------|-------------|---------------------------------------|--------------------|--------------|
| AT | +! ALLERGAN | <u>10,000 UNITS/ML;EQ 1MG BASE/ML</u> | <u>N050567 001</u> | Oct 20, 1988 |
|-----------|-------------|---------------------------------------|--------------------|--------------|

TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE

| | | | | |
|-----------|-----------|---------------------------------------|--------------------|--------------|
| AT | AKORN INC | <u>10,000 UNITS/ML;EQ 1MG BASE/ML</u> | <u>A065006 001</u> | Dec 17, 1998 |
|-----------|-----------|---------------------------------------|--------------------|--------------|

| | | | | |
|-----------|-----------------|---------------------------------------|--------------------|--------------|
| AT | BAUSCH AND LOMB | <u>10,000 UNITS/ML;EQ 1MG BASE/ML</u> | <u>A064120 001</u> | Feb 14, 1997 |
|-----------|-----------------|---------------------------------------|--------------------|--------------|

| | | | | |
|-----------|------------|---------------------------------------|--------------------|--------------|
| AT | SANDOZ INC | <u>10,000 UNITS/ML;EQ 1MG BASE/ML</u> | <u>A064211 001</u> | Apr 13, 1998 |
|-----------|------------|---------------------------------------|--------------------|--------------|

POMALIDOMIDE

CAPSULE;ORAL

POMALYST

| | | | | |
|--|-----------|-----|-------------|--------------|
| | + CELGENE | 1MG | N204026 001 | Feb 08, 2013 |
|--|-----------|-----|-------------|--------------|

| | | | | |
|--|---|-----|-------------|--------------|
| | + | 2MG | N204026 002 | Feb 08, 2013 |
|--|---|-----|-------------|--------------|

| | | | | |
|--|---|-----|-------------|--------------|
| | + | 3MG | N204026 003 | Feb 08, 2013 |
|--|---|-----|-------------|--------------|

| | | | | |
|--|----|-----|-------------|--------------|
| | +! | 4MG | N204026 004 | Feb 08, 2013 |
|--|----|-----|-------------|--------------|

PONATINIB HYDROCHLORIDE

TABLET;ORAL

ICLUSIG

| | | | | |
|--|---------|--------------|-------------|--------------|
| | + ARIAD | EQ 15MG BASE | N203469 001 | Dec 14, 2012 |
|--|---------|--------------|-------------|--------------|

| | | | | |
|--|---|--------------|-------------|--------------|
| | + | EQ 30MG BASE | N203469 003 | Apr 23, 2015 |
|--|---|--------------|-------------|--------------|

| | | | | |
|--|----|--------------|-------------|--------------|
| | +! | EQ 45MG BASE | N203469 002 | Dec 14, 2012 |
|--|----|--------------|-------------|--------------|

PORACTANT ALFA

SUSPENSION;INTRATRACHEAL

CUROSURF

| | | | | |
|--|-------------------|---------|-------------|--------------|
| | +! CHIESI USA INC | 80MG/ML | N020744 001 | Nov 18, 1999 |
|--|-------------------|---------|-------------|--------------|

PORFIMER SODIUM

INJECTABLE;INJECTION

PHOTOFRIN

| | | | | |
|--|--------------------|-----------|-------------|--------------|
| | CONCORDIA LABS INC | 75MG/VIAL | N020451 001 | Dec 27, 1995 |
|--|--------------------|-----------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

POSACONAZOLE

SOLUTION; INTRAVENOUS

NOXAFIL

+! MERCK SHARP DOHME 300MG/16.7ML (18MG/ML) N205596 001 Mar 13, 2014

SUSPENSION; ORAL

NOXAFIL

+! SCHERING 40MG/ML N022003 001 Sep 15, 2006

TABLET, DELAYED RELEASE; ORAL

NOXAFIL

+! MERCK SHARP DOHME 100MG N205053 001 Nov 25, 2013

POTASSIUM ACETATE

INJECTABLE; INJECTION

POTASSIUM ACETATEAP EXELA PHARMA SCS 2MEQ/ML A206203 001 Dec 29, 2015
LLCAP +! HOSPIRA 2MEQ/ML N018896 001 Jul 20, 1984POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

KLOR-CONAB UPSHER SMITH LABS 8MEQ A203106 001 Jul 10, 2015AB 10MEQ A203106 002 Jul 10, 2015MICRO-KAB + NESHER PHARMS 8MEQ N018238 001MICRO-K 10AB + NESHER PHARMS 10MEQ N018238 002 May 14, 1984POTASSIUM CHLORIDEAB ACTAVIS LABS FL INC 8MEQ A077419 001 Jun 02, 2008AB ! 10MEQ A077419 002 Jun 02, 2008AB ADARE PHARMS INC 8MEQ A208864 001 Mar 17, 2017AB 10MEQ A208864 002 Mar 17, 2017AB AMNEAL PHARMS 10MEQ A202128 001 Feb 22, 2013AB ANCHEN PHARMS 8MEQ A202886 001 Dec 26, 2013AB 10MEQ A202886 002 Dec 26, 2013AB GLENMARK PHARMS LTD 10MEQ A202868 001 Jan 19, 2016AB LANNETT CO INC 8MEQ A204210 001 Mar 28, 2016AB 10MEQ A204210 002 Mar 28, 2016AB LUPIN LTD 8MEQ A203002 001 Dec 18, 2015AB 10MEQ A203002 002 Dec 18, 2015AB NOVEL LABS INC 8MEQ A204828 001 Aug 16, 2016AB 10MEQ A204828 002 Aug 16, 2016AB PADDOCK LLC 8MEQ A200185 001 May 18, 2011AB 10MEQ A200185 002 May 18, 2011AB PII 8MEQ A205549 001 Dec 08, 2015AB 10MEQ A205549 002 Dec 08, 2015AB TRIS PHARMA INC 8MEQ A201944 001 Mar 04, 2016AB 10MEQ A201944 002 Mar 04, 2016

FOR SOLUTION; ORAL

KLOR-CONAA UPSHER SMITH LABS 20MEQ A209662 001 Oct 23, 2017POTASSIUM CHLORIDEAA EPIC PHARMA LLC 20MEQ A210200 001 Nov 23, 2018AA NOVEL LABS INC 20MEQ A210241 001 Nov 21, 2018AA +! PHARMA RES SOFTWARE 20MEQ N208019 001 Aug 19, 2015

INJECTABLE; INJECTION

POTASSIUM CHLORIDEAP B BRAUN 2MEQ/ML A085870 001AP FRESENIUS KABI USA 2MEQ/ML A080225 001AP ! HOSPIRA 2MEQ/ML A080205 001POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINERAP +! BAXTER HLTHCARE 14.9MG/ML N019904 001 Dec 26, 1989AP +! 746MG/100ML N019904 005 Dec 17, 1990AP + ICU MEDICAL INC 14.9MG/ML N020161 005 Nov 30, 1992AP + 745MG/100ML N020161 001 Nov 30, 1992POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINERAP +! BAXTER HLTHCARE 29.8MG/ML N019904 002 Dec 26, 1989AP +! 1.49GM/100ML N019904 006 Dec 17, 1990AP +! ICU MEDICAL INC 29.8MG/ML N020161 006 Aug 11, 1998AP + 1.49GM/100ML N020161 002 Nov 30, 1992POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINERAP +! BAXTER HLTHCARE 2.98GM/100ML N019904 004 Dec 26, 1989AP +! ICU MEDICAL INC 2.98GM/100ML N020161 004 Aug 11, 1998

PRESCRIPTION DRUG PRODUCT LIST

POTASSIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE IN PLASTIC CONTAINER

| | | | | |
|-----------|--------------------|----------------|--------------------|--------------|
| <u>AP</u> | FRESENIUS KABI USA | <u>2MEQ/ML</u> | <u>A088901 001</u> | Jan 25, 1985 |
| <u>AP</u> | | <u>2MEQ/ML</u> | <u>A088908 001</u> | Jan 25, 1985 |

POTASSIUM CHLORIDE

| | | | | |
|---|--------------------|---------|---------|-----|
| ! | FRESENIUS KABI USA | 3MEQ/ML | A080225 | 003 |
|---|--------------------|---------|---------|-----|

POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER

| | | | | | |
|---|---|-----------------|--------------|-------------|--------------|
| + | ! | BAXTER HLTHCARE | 2.24GM/100ML | N019904 003 | Dec 26, 1989 |
|---|---|-----------------|--------------|-------------|--------------|

SOLUTION; ORAL

POTASSIUM CHLORIDE

| | | | | | |
|-----------|-------------------|--------------------|--------------------|--------------------|--------------|
| <u>AA</u> | AMNEAL PHARMS LLC | <u>20MEQ/15ML</u> | <u>A210041 001</u> | Jul 19, 2018 | |
| <u>AA</u> | | <u>40MEQ/15ML</u> | <u>A210041 002</u> | Jul 19, 2018 | |
| <u>AA</u> | APOTEX INC | <u>20MEQ/15ML</u> | <u>A211067 001</u> | Aug 08, 2018 | |
| <u>AA</u> | | <u>40MEQ/15ML</u> | <u>A211067 002</u> | Aug 08, 2018 | |
| <u>AA</u> | + | GENUS LIFESCIENCES | <u>20MEQ/15ML</u> | <u>N206814 001</u> | Dec 22, 2014 |
| <u>AA</u> | + | ! | <u>40MEQ/15ML</u> | <u>N206814 002</u> | Dec 22, 2014 |
| <u>AA</u> | NOVEL LABS INC | <u>20MEQ/15ML</u> | <u>A209786 001</u> | Aug 29, 2018 | |
| <u>AA</u> | | <u>40MEQ/15ML</u> | <u>A209786 002</u> | Aug 29, 2018 | |

TABLET, EXTENDED RELEASE; ORAL

KLOR-CON M10

| | | | | |
|------------|-------------------|--------------|--------------------|--------------|
| <u>AB1</u> | UPSHER SMITH LABS | <u>10MEQ</u> | <u>A074726 002</u> | Aug 09, 2000 |
|------------|-------------------|--------------|--------------------|--------------|

KLOR-CON M20

| | | | | | |
|------------|---|-------------------|--------------|--------------------|--------------|
| <u>AB1</u> | ! | UPSHER SMITH LABS | <u>20MEQ</u> | <u>A074726 001</u> | Nov 20, 1998 |
|------------|---|-------------------|--------------|--------------------|--------------|

POTASSIUM CHLORIDE

| | | | | |
|------------|---------------------|--------------|--------------------|--------------|
| <u>AB1</u> | ACTAVIS LABS FL INC | <u>10MEQ</u> | <u>A075604 001</u> | Apr 10, 2002 |
| <u>AB1</u> | | <u>20MEQ</u> | <u>A075604 002</u> | Apr 10, 2002 |
| <u>AB1</u> | ADARE PHARMS INC | <u>20MEQ</u> | <u>A076368 001</u> | Aug 18, 2004 |
| <u>AB1</u> | GLENMARK PHARMS LTD | <u>10MEQ</u> | <u>A203562 001</u> | Jul 26, 2016 |
| <u>AB1</u> | | <u>20MEQ</u> | <u>A203562 002</u> | Jul 26, 2016 |
| <u>AB1</u> | NOVEL LABS INC | <u>10MEQ</u> | <u>A206347 001</u> | Jan 21, 2016 |
| <u>AB1</u> | | <u>20MEQ</u> | <u>A206347 002</u> | Jan 21, 2016 |

K-TAB

| | | | | | | |
|------------|---|---|--------|--------------|--------------------|--------------|
| <u>AB2</u> | + | ! | ABBVIE | <u>20MEQ</u> | <u>N018279 003</u> | Nov 25, 2013 |
|------------|---|---|--------|--------------|--------------------|--------------|

KLOR-CON

| | | | | | | |
|------------|---|-------------------|-------------|--------------------|--------------------|--------------|
| <u>AB2</u> | + | UPSHER SMITH LABS | <u>8MEQ</u> | <u>N019123 001</u> | Apr 17, 1986 | |
| <u>AB2</u> | + | ! | | <u>10MEQ</u> | <u>N019123 002</u> | Apr 17, 1986 |

POTASSIUM CHLORIDE

| | | | | |
|------------|----------------------|--------------|--------------------|--------------|
| <u>AB2</u> | AUROBINDO PHARMA LTD | <u>8MEQ</u> | <u>A210921 001</u> | Dec 19, 2018 |
| <u>AB2</u> | | <u>10MEQ</u> | <u>A210921 002</u> | Dec 19, 2018 |
| <u>AB2</u> | MYLAN PHARMS INC | <u>8MEQ</u> | <u>A204662 001</u> | Aug 21, 2014 |
| <u>AB2</u> | | <u>10MEQ</u> | <u>A204662 002</u> | Aug 21, 2014 |
| <u>AB2</u> | NOVEL LABS INC | <u>8MEQ</u> | <u>A206759 001</u> | Aug 09, 2016 |
| <u>AB2</u> | | <u>10MEQ</u> | <u>A206759 002</u> | Aug 09, 2016 |
| <u>AB2</u> | PADDOCK LLC | <u>8MEQ</u> | <u>A205993 001</u> | Nov 05, 2015 |
| <u>AB2</u> | | <u>10MEQ</u> | <u>A205993 002</u> | Nov 05, 2015 |
| <u>AB2</u> | SIGMAPHARM LABS LLC | <u>8MEQ</u> | <u>A207528 001</u> | Aug 19, 2016 |
| <u>AB2</u> | | <u>10MEQ</u> | <u>A207528 002</u> | Aug 19, 2016 |
| <u>AB2</u> | STRIDES PHARMA | <u>8MEQ</u> | <u>A210733 001</u> | Aug 31, 2018 |
| <u>AB2</u> | | <u>10MEQ</u> | <u>A210733 002</u> | Aug 31, 2018 |
| <u>AB2</u> | VITRUVIAS THERAP | <u>20MEQ</u> | <u>A209688 002</u> | Jan 12, 2018 |
| <u>AB2</u> | YICHANG HUMANWELL | <u>8MEQ</u> | <u>A209314 001</u> | Jun 22, 2018 |
| <u>AB2</u> | | <u>10MEQ</u> | <u>A209314 002</u> | Jun 22, 2018 |

K-TAB

| | | | | |
|------------|---|--------|--------------|--------------------|
| <u>AB3</u> | + | ABBVIE | <u>10MEQ</u> | <u>N018279 001</u> |
|------------|---|--------|--------------|--------------------|

POTASSIUM CHLORIDE

| | | | | |
|------------|------------------|--------------|--------------------|--------------|
| <u>AB3</u> | VITRUVIAS THERAP | <u>10MEQ</u> | <u>A209688 001</u> | Jan 12, 2018 |
|------------|------------------|--------------|--------------------|--------------|

K-TAB

| | | | | | |
|----|---|--------|------|-------------|--------------|
| BC | + | ABBVIE | 8MEQ | N018279 002 | Aug 01, 1988 |
|----|---|--------|------|-------------|--------------|

KLOR-CON M15

| | | | | | |
|--|--|-------------------|-------|-------------|--------------|
| | | UPSHER SMITH LABS | 15MEQ | A074726 003 | Jun 06, 2003 |
|--|--|-------------------|-------|-------------|--------------|

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.149% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

| | | | | |
|-----------|-----------------|---------------------------------|--------------------|--------------|
| <u>AP</u> | ICU MEDICAL INC | <u>149MG/100ML; 450MG/100ML</u> | <u>A078446 001</u> | Sep 10, 2008 |
|-----------|-----------------|---------------------------------|--------------------|--------------|

POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

| | | | | | | |
|-----------|---|---|-----------------|---------------------------------|--------------------|--------------|
| <u>AP</u> | + | ! | BAXTER HLTHCARE | <u>150MG/100ML; 450MG/100ML</u> | <u>N017648 005</u> | Nov 26, 2002 |
|-----------|---|---|-----------------|---------------------------------|--------------------|--------------|

POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | |
|-----------|---------|---------------------------------|---------------------------------|--------------------|
| <u>AP</u> | B BRAUN | <u>150MG/100ML; 900MG/100ML</u> | <u>N019708 004</u> | Sep 29, 1989 |
| <u>AP</u> | + | BAXTER HLTHCARE | <u>150MG/100ML; 900MG/100ML</u> | <u>N017648 001</u> |

PRESCRIPTION DRUG PRODUCT LIST

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | | | |
|------------------------------------------------------------------------------|---|-----------------|--------------------------------|----------------|------------|--------------|
| <u>AP</u> | + | BAXTER HLTHCARE | <u>300MG/100ML;900MG/100ML</u> | <u>N017648</u> | <u>002</u> | |
| <u>POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u> | | | | | | |
| <u>AP</u> | | ICU MEDICAL INC | <u>149MG/100ML;900MG/100ML</u> | <u>N019686</u> | <u>001</u> | Oct 17, 1988 |
| <u>POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u> | | | | | | |
| <u>AP</u> | | ICU MEDICAL INC | <u>298MG/100ML;900MG/100ML</u> | <u>N019686</u> | <u>002</u> | Oct 17, 1988 |

POTASSIUM CITRATE

TABLET, EXTENDED RELEASE; ORAL

POTASSIUM CITRATE

| | | | | | | |
|------------------|---|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | MOUNTAIN | <u>5MEQ</u> | <u>A077440</u> | <u>001</u> | Jun 09, 2006 |
| <u>AB</u> | | | <u>10MEQ</u> | <u>A077440</u> | <u>002</u> | Jun 09, 2006 |
| <u>AB</u> | | STRIDES PHARMA | <u>5MEQ</u> | <u>A206813</u> | <u>001</u> | Sep 11, 2017 |
| <u>AB</u> | | | <u>10MEQ</u> | <u>A206813</u> | <u>002</u> | Sep 11, 2017 |
| <u>AB</u> | | | <u>15MEQ</u> | <u>A206813</u> | <u>003</u> | Sep 11, 2017 |
| <u>AB</u> | | TEVA PHARMS USA INC | <u>5MEQ</u> | <u>A209758</u> | <u>001</u> | Mar 05, 2018 |
| <u>AB</u> | | | <u>10MEQ</u> | <u>A209758</u> | <u>002</u> | Mar 05, 2018 |
| <u>AB</u> | | | <u>15MEQ</u> | <u>A209758</u> | <u>003</u> | Mar 05, 2018 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>5MEQ</u> | <u>A203546</u> | <u>001</u> | Aug 06, 2014 |
| <u>AB</u> | | | <u>10MEQ</u> | <u>A203546</u> | <u>002</u> | Aug 06, 2014 |
| <u>AB</u> | | | <u>15MEQ</u> | <u>A203546</u> | <u>003</u> | Aug 06, 2014 |
| <u>UROCI-T-K</u> | | | | | | |
| <u>AB</u> | + | MISSION PHARMA | <u>5MEQ</u> | <u>N019071</u> | <u>001</u> | Aug 30, 1985 |
| <u>AB</u> | + | | <u>10MEQ</u> | <u>N019071</u> | <u>002</u> | Aug 31, 1992 |
| <u>AB</u> | + | | <u>15MEQ</u> | <u>N019071</u> | <u>003</u> | Dec 30, 2009 |

POVIDONE-IODINE

SOLUTION/DROPS; OPHTHALMIC

BETADINE

| | | | | | | |
|---|---|------------------|----|---------|-----|--------------|
| + | ! | ALCON PHARMS LTD | 5% | N018634 | 001 | Dec 17, 1986 |
|---|---|------------------|----|---------|-----|--------------|

PRALATREXATE

SOLUTION; INTRAVENOUS

FOLOTYN

| | | | | | | |
|---|---|-------|--------------------|---------|-----|--------------|
| + | | ALLOS | 20MG/ML (20MG/ML) | N022468 | 001 | Sep 24, 2009 |
| + | ! | | 40MG/2ML (20MG/ML) | N022468 | 002 | Sep 24, 2009 |

PRALIDOXIME CHLORIDE

INJECTABLE; INJECTION

PRALIDOXIME CHLORIDE

| | | | | | | |
|---|---|-------------------------|----------|---------|-----|--------------|
| + | ! | MERIDIAN MEDCL TECHN | 300MG/ML | N018986 | 001 | Apr 26, 1983 |
|---|---|-------------------------|----------|---------|-----|--------------|

PROTOPAM CHLORIDE

| | | | | | | |
|---|---|-------------------------|----------|---------|-----|--|
| + | ! | BAXTER HLTHCARE CORP | 1GM/VIAL | N014134 | 001 | |
|---|---|-------------------------|----------|---------|-----|--|

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

MIRAPEX

| | | | | | | |
|-----------|---|-------------------------|----------------|----------------|------------|--------------|
| <u>AB</u> | + | BOEHRINGER INGELHEIM | <u>0.125MG</u> | <u>N020667</u> | <u>001</u> | Jul 01, 1997 |
| <u>AB</u> | + | | <u>0.25MG</u> | <u>N020667</u> | <u>002</u> | Jul 01, 1997 |
| <u>AB</u> | + | | <u>0.5MG</u> | <u>N020667</u> | <u>006</u> | Feb 12, 1998 |
| <u>AB</u> | + | | <u>0.75MG</u> | <u>N020667</u> | <u>007</u> | Jul 30, 2007 |
| <u>AB</u> | + | | <u>1MG</u> | <u>N020667</u> | <u>003</u> | Jul 01, 1997 |
| <u>AB</u> | + | | <u>1.5MG</u> | <u>N020667</u> | <u>005</u> | Jul 01, 1997 |

PRAMIPEXOLE DIHYDROCHLORIDE

| | | | | | | |
|-----------|--|-------------------------|----------------|----------------|------------|--------------|
| <u>AB</u> | | ALEMBIC PHARMS LTD | <u>0.125MG</u> | <u>A078894</u> | <u>001</u> | Oct 08, 2010 |
| <u>AB</u> | | | <u>0.25MG</u> | <u>A078894</u> | <u>002</u> | Oct 08, 2010 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A078894</u> | <u>003</u> | Oct 08, 2010 |
| <u>AB</u> | | | <u>1MG</u> | <u>A078894</u> | <u>004</u> | Oct 08, 2010 |
| <u>AB</u> | | | <u>1.5MG</u> | <u>A078894</u> | <u>005</u> | Oct 08, 2010 |
| <u>AB</u> | | APOTEX INC | <u>0.125MG</u> | <u>A090151</u> | <u>001</u> | Apr 30, 2012 |
| <u>AB</u> | | | <u>0.25MG</u> | <u>A090151</u> | <u>002</u> | Apr 30, 2012 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A090151</u> | <u>003</u> | Apr 30, 2012 |
| <u>AB</u> | | | <u>0.75MG</u> | <u>A090151</u> | <u>006</u> | Apr 30, 2012 |
| <u>AB</u> | | | <u>1MG</u> | <u>A090151</u> | <u>004</u> | Apr 30, 2012 |
| <u>AB</u> | | | <u>1.5MG</u> | <u>A090151</u> | <u>005</u> | Apr 30, 2012 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>0.125MG</u> | <u>A202633</u> | <u>001</u> | Oct 26, 2012 |
| <u>AB</u> | | | <u>0.25MG</u> | <u>A202633</u> | <u>002</u> | Oct 26, 2012 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A202633</u> | <u>003</u> | Oct 26, 2012 |
| <u>AB</u> | | | <u>0.75MG</u> | <u>A202633</u> | <u>004</u> | Oct 26, 2012 |

PRESCRIPTION DRUG PRODUCT LIST

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

| | | | | |
|-----------|---------------------|----------------|--------------------|--------------|
| <u>AB</u> | | <u>1MG</u> | <u>A202633 005</u> | Oct 26, 2012 |
| <u>AB</u> | | <u>1.5MG</u> | <u>A202633 006</u> | Oct 26, 2012 |
| <u>AB</u> | BARR | <u>0.125MG</u> | <u>A077724 001</u> | Feb 19, 2008 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A077724 002</u> | Feb 19, 2008 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A077724 003</u> | Feb 19, 2008 |
| <u>AB</u> | | <u>1MG</u> | <u>A077724 004</u> | Feb 19, 2008 |
| <u>AB</u> | | <u>1.5MG</u> | <u>A077724 005</u> | Feb 19, 2008 |
| <u>AB</u> | BRECKENRIDGE PHARM | <u>0.125MG</u> | <u>A091450 001</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A091450 002</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A091450 003</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>1MG</u> | <u>A091450 004</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>1.5MG</u> | <u>A091450 005</u> | Oct 08, 2010 |
| <u>AB</u> | CSPC OUYI PHARM CO | <u>0.25MG</u> | <u>A211088 001</u> | Oct 03, 2018 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A211088 002</u> | Oct 03, 2018 |
| <u>AB</u> | | <u>0.75MG</u> | <u>A211088 003</u> | Oct 03, 2018 |
| <u>AB</u> | | <u>1MG</u> | <u>A211088 004</u> | Oct 03, 2018 |
| <u>AB</u> | | <u>1.5MG</u> | <u>A211088 005</u> | Oct 03, 2018 |
| <u>AB</u> | GLENMARK GENERICS | <u>0.125MG</u> | <u>A090781 001</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A090781 002</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A090781 003</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>0.75MG</u> | <u>A090781 006</u> | Sep 11, 2015 |
| <u>AB</u> | | <u>1MG</u> | <u>A090781 004</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>1.5MG</u> | <u>A090781 005</u> | Oct 08, 2010 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>0.125MG</u> | <u>A202164 001</u> | Sep 20, 2012 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A202164 002</u> | Sep 20, 2012 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A202164 003</u> | Sep 20, 2012 |
| <u>AB</u> | | <u>1MG</u> | <u>A202164 004</u> | Sep 20, 2012 |
| <u>AB</u> | | <u>1.5MG</u> | <u>A202164 005</u> | Sep 20, 2012 |
| <u>AB</u> | MYLAN | <u>0.125MG</u> | <u>A077854 001</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A077854 002</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A077854 003</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>0.75MG</u> | <u>A090764 001</u> | Apr 09, 2010 |
| <u>AB</u> | | <u>1MG</u> | <u>A077854 004</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>1.5MG</u> | <u>A077854 005</u> | Oct 08, 2010 |
| <u>AB</u> | SCIEGEN PHARMS INC | <u>0.125MG</u> | <u>A203855 001</u> | Oct 28, 2014 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A203855 002</u> | Oct 28, 2014 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A203855 003</u> | Oct 28, 2014 |
| <u>AB</u> | | <u>0.75MG</u> | <u>A203855 004</u> | Oct 28, 2014 |
| <u>AB</u> | | <u>1MG</u> | <u>A203855 005</u> | Oct 28, 2014 |
| <u>AB</u> | | <u>1.5MG</u> | <u>A203855 006</u> | Oct 28, 2014 |
| <u>AB</u> | STRIDES PHARMA | <u>0.125MG</u> | <u>A202702 001</u> | Jun 03, 2014 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A202702 002</u> | Jun 03, 2014 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A202702 003</u> | Jun 03, 2014 |
| <u>AB</u> | | <u>0.75MG</u> | <u>A202702 004</u> | Jun 03, 2014 |
| <u>AB</u> | | <u>1MG</u> | <u>A202702 005</u> | Jun 03, 2014 |
| <u>AB</u> | | <u>1.5MG</u> | <u>A202702 006</u> | Jun 03, 2014 |
| <u>AB</u> | SUN PHARM INDS INC | <u>0.125MG</u> | <u>A091683 001</u> | Mar 27, 2013 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A091683 002</u> | Mar 27, 2013 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A091683 003</u> | Mar 27, 2013 |
| <u>AB</u> | | <u>0.75MG</u> | <u>A091683 004</u> | Mar 27, 2013 |
| <u>AB</u> | | <u>1MG</u> | <u>A091683 005</u> | Mar 27, 2013 |
| <u>AB</u> | | <u>1.5MG</u> | <u>A091683 006</u> | Mar 27, 2013 |
| <u>AB</u> | TEVA PHARMS | <u>0.125MG</u> | <u>A090241 001</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A090241 002</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A090241 003</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>0.75MG</u> | <u>A090241 004</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>1MG</u> | <u>A090241 005</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>1.5MG</u> | <u>A090241 006</u> | Oct 08, 2010 |
| <u>AB</u> | TORRENT PHARMS | <u>0.125MG</u> | <u>A090865 001</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A090865 002</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A090865 003</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>0.75MG</u> | <u>A090865 004</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>1MG</u> | <u>A090865 005</u> | Oct 08, 2010 |
| <u>AB</u> | | <u>1.5MG</u> | <u>A090865 006</u> | Oct 08, 2010 |
| <u>AB</u> | UNICHEM LABS LTD | <u>0.125MG</u> | <u>A207011 001</u> | Dec 19, 2018 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A207011 002</u> | Dec 19, 2018 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A207011 003</u> | Dec 19, 2018 |
| <u>AB</u> | | <u>0.75MG</u> | <u>A207011 004</u> | Dec 19, 2018 |
| <u>AB</u> | | <u>1MG</u> | <u>A207011 005</u> | Dec 19, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET;ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

| | | | | |
|-----------|-------------------------|----------------|--------------------|--------------|
| <u>AB</u> | | <u>1.5MG</u> | <u>A207011 006</u> | Dec 19, 2018 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>0.125MG</u> | <u>A078920 001</u> | Jul 06, 2010 |
| <u>AB</u> | | <u>0.25MG</u> | <u>A078920 002</u> | Jul 06, 2010 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A078920 003</u> | Jul 06, 2010 |
| <u>AB</u> | | <u>1MG</u> | <u>A078920 004</u> | Jul 06, 2010 |
| <u>AB</u> | | <u>1.5MG</u> | <u>A078920 005</u> | Jul 06, 2010 |

TABLET, EXTENDED RELEASE;ORAL

MIRAPEX ER

| | | | | | |
|-----------|----|-------------------------|----------------|--------------------|--------------|
| <u>AB</u> | +! | BOEHRINGER INGELHEIM | <u>0.375MG</u> | <u>N022421 001</u> | Feb 19, 2010 |
| <u>AB</u> | + | | <u>0.75MG</u> | <u>N022421 002</u> | Feb 19, 2010 |
| <u>AB</u> | + | | <u>1.5MG</u> | <u>N022421 003</u> | Feb 19, 2010 |
| <u>AB</u> | + | | <u>2.25MG</u> | <u>N022421 006</u> | Jun 17, 2011 |
| <u>AB</u> | + | | <u>3MG</u> | <u>N022421 004</u> | Feb 19, 2010 |
| <u>AB</u> | + | | <u>3.75MG</u> | <u>N022421 007</u> | Jun 17, 2011 |
| <u>AB</u> | + | | <u>4.5MG</u> | <u>N022421 005</u> | Feb 19, 2010 |

PRAMIPEXOLE DIHYDROCHLORIDE

| | | | | | |
|-----------|--|-------------------------|----------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS ELIZABETH | <u>0.375MG</u> | <u>A201963 001</u> | Apr 21, 2016 |
| <u>AB</u> | | | <u>0.75MG</u> | <u>A201963 002</u> | Apr 21, 2016 |
| <u>AB</u> | | | <u>1.5MG</u> | <u>A201963 003</u> | Apr 21, 2016 |
| <u>AB</u> | | | <u>2.25MG</u> | <u>A203615 001</u> | Oct 14, 2016 |
| <u>AB</u> | | | <u>3MG</u> | <u>A201963 004</u> | Apr 21, 2016 |
| <u>AB</u> | | | <u>3.75MG</u> | <u>A203615 002</u> | Jan 03, 2017 |
| <u>AB</u> | | | <u>4.5MG</u> | <u>A201963 005</u> | Apr 21, 2016 |
| <u>AB</u> | | ALEMbic PHARMS LTD | <u>0.375MG</u> | <u>A204518 001</u> | Jan 02, 2019 |
| <u>AB</u> | | | <u>0.75MG</u> | <u>A204518 002</u> | Jan 02, 2019 |
| <u>AB</u> | | | <u>1.5MG</u> | <u>A204518 003</u> | Jan 02, 2019 |
| <u>AB</u> | | | <u>2.25MG</u> | <u>A204518 004</u> | Jan 02, 2019 |
| <u>AB</u> | | | <u>3MG</u> | <u>A204518 005</u> | Jan 02, 2019 |
| <u>AB</u> | | | <u>3.75MG</u> | <u>A204518 006</u> | Jan 02, 2019 |
| <u>AB</u> | | | <u>4.5MG</u> | <u>A204518 007</u> | Jan 02, 2019 |
| <u>AB</u> | | ANCHEN PHARMS | <u>0.375MG</u> | <u>A202206 001</u> | Feb 06, 2014 |
| <u>AB</u> | | | <u>0.75MG</u> | <u>A202206 002</u> | Feb 06, 2014 |
| <u>AB</u> | | | <u>1.5MG</u> | <u>A202206 003</u> | Feb 06, 2014 |
| <u>AB</u> | | | <u>2.25MG</u> | <u>A202206 004</u> | Feb 06, 2014 |
| <u>AB</u> | | | <u>3MG</u> | <u>A202206 005</u> | Feb 06, 2014 |
| <u>AB</u> | | | <u>3.75MG</u> | <u>A202206 006</u> | Feb 06, 2014 |
| <u>AB</u> | | | <u>4.5MG</u> | <u>A202206 007</u> | Feb 06, 2014 |
| <u>AB</u> | | DR REDDYS LABS LTD | <u>0.375MG</u> | <u>A203354 001</u> | Aug 07, 2015 |
| <u>AB</u> | | | <u>0.75MG</u> | <u>A203354 002</u> | Aug 07, 2015 |
| <u>AB</u> | | | <u>1.5MG</u> | <u>A203354 003</u> | Aug 07, 2015 |
| <u>AB</u> | | | <u>3MG</u> | <u>A203354 004</u> | Aug 07, 2015 |
| <u>AB</u> | | | <u>4.5MG</u> | <u>A203354 005</u> | Aug 07, 2015 |
| <u>AB</u> | | MACLEODS PHARMS LTD | <u>0.375MG</u> | <u>A206156 001</u> | Jun 24, 2016 |
| <u>AB</u> | | | <u>0.75MG</u> | <u>A206156 002</u> | Jun 24, 2016 |
| <u>AB</u> | | | <u>1.5MG</u> | <u>A206156 003</u> | Jun 24, 2016 |
| <u>AB</u> | | | <u>2.25MG</u> | <u>A206156 004</u> | Jun 24, 2016 |
| <u>AB</u> | | | <u>3MG</u> | <u>A206156 005</u> | Jun 24, 2016 |
| <u>AB</u> | | | <u>3.75MG</u> | <u>A206156 007</u> | Jan 23, 2017 |
| <u>AB</u> | | | <u>4.5MG</u> | <u>A206156 006</u> | Jun 24, 2016 |
| <u>AB</u> | | SANDOZ INC | <u>0.375MG</u> | <u>A202353 001</u> | Dec 04, 2014 |
| <u>AB</u> | | | <u>0.75MG</u> | <u>A202353 002</u> | Dec 04, 2014 |
| <u>AB</u> | | | <u>1.5MG</u> | <u>A202353 003</u> | Dec 04, 2014 |
| <u>AB</u> | | | <u>3MG</u> | <u>A202353 004</u> | Dec 04, 2014 |
| <u>AB</u> | | | <u>4.5MG</u> | <u>A202353 005</u> | Dec 04, 2014 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>0.375MG</u> | <u>A202891 001</u> | Dec 12, 2017 |
| <u>AB</u> | | | <u>0.75MG</u> | <u>A202891 002</u> | Dec 12, 2017 |
| <u>AB</u> | | | <u>1.5MG</u> | <u>A202891 003</u> | Dec 12, 2017 |
| <u>AB</u> | | | <u>2.25MG</u> | <u>A202891 004</u> | Dec 12, 2017 |
| <u>AB</u> | | | <u>3MG</u> | <u>A202891 005</u> | Dec 12, 2017 |
| <u>AB</u> | | | <u>3.75MG</u> | <u>A202891 006</u> | Dec 12, 2017 |
| <u>AB</u> | | | <u>4.5MG</u> | <u>A202891 007</u> | Dec 12, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

PRAMLINTIDE ACETATE

INJECTABLE; SUBCUTANEOUS

SYMLIN

+ ASTRAZENECA AB

EQ 1.5MG BASE/1.5ML (EQ 1MG BASE/ML)

N021332 002 Sep 25, 2007

+!

EQ 2.7MG BASE/2.7ML (EQ 1MG BASE/ML)

N021332 003 Sep 25, 2007

PRASTERONE

INSERT; VAGINAL

INTRAROSA

+! AMAG PHARMS INC

6.5MG

N208470 001 Nov 16, 2016

PRASUGREL HYDROCHLORIDE

TABLET; ORAL

EFFIENTAB + ELI LILLY AND COEQ 5MG BASEN022307 001 Jul 10, 2009AB +!EQ 10MG BASEN022307 002 Jul 10, 2009PRASUGRELAB ACCORD HLTHCAREEQ 5MG BASEA205987 001 Feb 02, 2018ABEQ 10MG BASEA205987 002 Feb 02, 2018AB

AMNEAL PHARMS

EQ 5MG BASEA205913 001 Jun 19, 2018ABEQ 10MG BASEA205913 002 Jun 19, 2018AB

AUROBINDO PHARMA LTD

EQ 5MG BASEA205888 001 Oct 16, 2017ABEQ 10MG BASEA205888 002 Oct 16, 2017AB

HEC PHARM

EQ 5MG BASEA206021 001 Jan 16, 2019ABEQ 10MG BASEA206021 002 Jan 16, 2019AB

MYLAN PHARMS INC

EQ 5MG BASEA205927 001 Jul 12, 2017ABEQ 10MG BASEA205927 002 Jul 12, 2017AB

PANACEA BIOTEC LTD

EQ 5MG BASEA205897 001 Oct 16, 2017ABEQ 10MG BASEA205897 002 Oct 16, 2017AB

USPHARMA WINDLAS

EQ 5MG BASEA205790 001 Oct 16, 2017ABEQ 10MG BASEA205790 002 Oct 16, 2017PRAVASTATIN SODIUM

TABLET; ORAL

PRAVACHOLAB + BRISTOL MYERS20MGN019898 003 Oct 31, 1991

SQUIBB

AB +40MGN019898 004 Mar 22, 1993AB +!80MGN019898 008 Dec 18, 2001PRAVASTATIN SODIUMAB ACCORD HLTHCARE10MGA207068 001 Nov 17, 2016AB20MGA207068 002 Nov 17, 2016AB40MGA207068 003 Nov 17, 2016AB80MGA207068 004 Nov 17, 2016AB

APOTEX INC

10MGA076341 001 Oct 23, 2006AB20MGA076341 002 Oct 23, 2006AB40MGA076341 003 Oct 23, 2006AB80MGA076341 004 Dec 28, 2007AB

AUROBINDO PHARMA LTD

10MGA203367 001 Feb 02, 2017AB20MGA203367 002 Feb 02, 2017AB40MGA203367 003 Feb 02, 2017AB80MGA203367 004 Feb 02, 2017AB

CHARTWELL RX

10MGA209869 001 Apr 13, 2018AB20MGA209869 002 Apr 13, 2018AB40MGA209869 003 Apr 13, 2018AB80MGA209869 004 Apr 13, 2018AB

CIPLA

10MGA077904 001 Oct 23, 2006AB20MGA077904 002 Oct 23, 2006AB40MGA077904 003 Oct 23, 2006AB80MGA077904 004 Mar 22, 2016AB

DR REDDYS LABS INC

10MGA076714 001 Oct 23, 2006AB20MGA076714 002 Oct 23, 2006AB40MGA076714 003 Oct 23, 2006AB80MGA076714 004 Dec 28, 2007AB

GLENMARK GENERICS

10MGA077987 001 May 11, 2007AB20MGA077987 002 May 11, 2007AB40MGA077987 003 May 11, 2007AB80MGA077987 004 Dec 28, 2007AB

HISUN PHARM

20MGA206061 001 Nov 23, 2018

HANGZHOU

AB40MGA206061 002 Nov 23, 2018AB80MGA206061 003 Nov 23, 2018AB

LUPIN PHARMS

10MGA077917 001 Jan 08, 2008

PRESCRIPTION DRUG PRODUCT LIST

PRAVASTATIN SODIUM

TABLET; ORAL

PRAVASTATIN SODIUM

| | | | | |
|-----------|------------------|-------------|--------------------|--------------|
| <u>AB</u> | | <u>20MG</u> | <u>A077917 002</u> | Jan 08, 2008 |
| <u>AB</u> | | <u>40MG</u> | <u>A077917 003</u> | Jan 08, 2008 |
| <u>AB</u> | | <u>80MG</u> | <u>A077917 004</u> | Jan 08, 2008 |
| <u>AB</u> | MYLAN PHARMS INC | <u>10MG</u> | <u>A079187 001</u> | May 27, 2010 |
| <u>AB</u> | | <u>20MG</u> | <u>A079187 002</u> | May 27, 2010 |
| <u>AB</u> | | <u>40MG</u> | <u>A079187 003</u> | May 27, 2010 |
| <u>AB</u> | | <u>80MG</u> | <u>A079187 004</u> | May 27, 2010 |
| <u>AB</u> | SANDOZ | <u>10MG</u> | <u>A076397 003</u> | Oct 23, 2006 |
| <u>AB</u> | | <u>20MG</u> | <u>A076397 002</u> | Oct 23, 2006 |
| <u>AB</u> | | <u>40MG</u> | <u>A076397 001</u> | Oct 23, 2006 |
| <u>AB</u> | | <u>80MG</u> | <u>A077491 001</u> | Feb 11, 2008 |
| <u>AB</u> | TEVA | <u>10MG</u> | <u>A076056 001</u> | Apr 24, 2006 |
| <u>AB</u> | | <u>20MG</u> | <u>A076056 002</u> | Apr 24, 2006 |
| <u>AB</u> | | <u>40MG</u> | <u>A076056 003</u> | Apr 24, 2006 |
| <u>AB</u> | TEVA PHARMS | <u>80MG</u> | <u>A077793 001</u> | Jan 15, 2008 |
| <u>AB</u> | WATSON LABS | <u>10MG</u> | <u>A076939 004</u> | Oct 23, 2006 |
| <u>AB</u> | | <u>20MG</u> | <u>A076939 003</u> | Oct 23, 2006 |
| <u>AB</u> | | <u>40MG</u> | <u>A076939 002</u> | Oct 23, 2006 |
| <u>AB</u> | | <u>80MG</u> | <u>A076939 001</u> | Dec 28, 2007 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>10MG</u> | <u>A077751 001</u> | Apr 30, 2008 |
| <u>AB</u> | | <u>20MG</u> | <u>A077751 002</u> | Apr 30, 2008 |
| <u>AB</u> | | <u>40MG</u> | <u>A077751 003</u> | Apr 30, 2008 |
| <u>AB</u> | | <u>80MG</u> | <u>A077751 004</u> | Apr 30, 2008 |

PRAZICQUANTEL

TABLET; ORAL

BILTRICIDE

| | | | | | |
|-----------|----|----------------|--------------|--------------------|--------------|
| <u>AB</u> | +! | BAYER HLTHCARE | <u>600MG</u> | <u>N018714 001</u> | Dec 29, 1982 |
| <u>AB</u> | | PAR PHARM INC | <u>600MG</u> | <u>A208820 001</u> | Nov 27, 2017 |

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

MINIPRESS

| | | | | | |
|-----------|----|-----------------|--------------------|--------------------|--------------|
| <u>AB</u> | + | PFIZER | <u>EQ 1MG BASE</u> | <u>N017442 002</u> | |
| <u>AB</u> | +! | | <u>EQ 2MG BASE</u> | <u>N017442 003</u> | |
| <u>AB</u> | + | | <u>EQ 5MG BASE</u> | <u>N017442 001</u> | |
| <u>AB</u> | | MYLAN | <u>EQ 1MG BASE</u> | <u>A072575 003</u> | May 16, 1989 |
| <u>AB</u> | | | <u>EQ 2MG BASE</u> | <u>A072575 002</u> | May 16, 1989 |
| <u>AB</u> | | | <u>EQ 5MG BASE</u> | <u>A072575 001</u> | May 16, 1989 |
| <u>AB</u> | | NOVITIUM PHARMA | <u>EQ 1MG BASE</u> | <u>A210971 001</u> | Oct 03, 2018 |
| <u>AB</u> | | | <u>EQ 2MG BASE</u> | <u>A210971 002</u> | Oct 03, 2018 |
| <u>AB</u> | | | <u>EQ 5MG BASE</u> | <u>A210971 003</u> | Oct 03, 2018 |
| <u>AB</u> | | TEVA PHARMS | <u>EQ 1MG BASE</u> | <u>A071745 002</u> | Sep 12, 1988 |
| <u>AB</u> | | | <u>EQ 2MG BASE</u> | <u>A071745 003</u> | Sep 12, 1988 |
| <u>AB</u> | | | <u>EQ 5MG BASE</u> | <u>A071745 001</u> | Sep 12, 1988 |

PREDNICARBATE

CREAM; TOPICAL

DERMATOP E EMOLLIENT

| | | | | | |
|-----------|----|----------------------|-------------|--------------------|--------------|
| <u>AB</u> | +! | VALEANT BERMUDA | <u>0.1%</u> | <u>N020279 001</u> | Oct 29, 1993 |
| <u>AB</u> | | FOUGERA PHARMS | <u>0.1%</u> | <u>A077287 001</u> | Sep 19, 2006 |
| <u>AB</u> | | VALEANT PHARMS NORTH | <u>0.1%</u> | <u>N019568 001</u> | Sep 23, 1991 |
| <u>AB</u> | | FOUGERA PHARMS | <u>0.1%</u> | <u>A077236 001</u> | Mar 09, 2007 |

PREDNISOLONE

SYRUP; ORAL

PREDNISOLONE

| | | | | | |
|-----------|---|-------------------|-----------------|--------------------|--------------|
| <u>AA</u> | ! | HI TECH PHARMA CO | <u>15MG/5ML</u> | <u>A040401 001</u> | Feb 27, 2003 |
| <u>AA</u> | | LANNETT CO INC | <u>15MG/5ML</u> | <u>A040775 001</u> | Sep 21, 2007 |
| <u>AA</u> | | PHARM ASSOC | <u>15MG/5ML</u> | <u>A040399 001</u> | Mar 05, 2003 |
| <u>AA</u> | | VISTAPHARM | <u>15MG/5ML</u> | <u>A040323 001</u> | May 13, 1999 |
| <u>AA</u> | | WOCKHARDT BIO AG | <u>15MG/5ML</u> | <u>A040313 001</u> | Sep 10, 2003 |
| <u>AA</u> | | TEVA | <u>15MG/5ML</u> | <u>A089081 001</u> | Feb 04, 1986 |

PRESCRIPTION DRUG PRODUCT LIST

PREDNISOLONE

TABLET; ORAL

PREDNISOLONE

! WATSON LABS 5MG A080354 001

PREDNISOLONE ACETATE

SUSPENSION/DROPS; OPHTHALMIC

OMNIPRED**AB** NOVARTIS PHARMS **1%** **N017469 001**
CORPPRED FORTE**AB** +! ALLERGAN **1%** **N017011 001**

PRED MILD

+! ALLERGAN 0.12% N017100 001

PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC

BLEPHAMIDE S.O.P.

! ALLERGAN 0.2%;10% A087748 001 Dec 03, 1986

SUSPENSION; OPHTHALMIC

BLEPHAMIDE

+! ALLERGAN 0.2%;10% N012813 002

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION; ORAL

PEDIAPRED**AA** +! SETON PHARM **EQ 5MG BASE/5ML** **N019157 001** May 28, 1986PREDNISOLONE SODIUM PHOSPHATE**AA** CHARTWELL RX **EQ 5MG BASE/5ML** **A075988 001** May 25, 2004**AA** EDENBRIDGE PHARMS **EQ 10MG BASE/5ML** **A203559 001** Dec 20, 2016**AA** **EQ 20MG BASE/5ML** **A203559 002** Dec 20, 2016**AA** HI TECH PHARMA **EQ 5MG BASE/5ML** **A075183 001** Mar 26, 2003**AA** ! PHARM ASSOC **EQ 10MG BASE/5ML** **A078465 001** Mar 07, 2008**AA** **EQ 15MG BASE/5ML** **A076913 001** Apr 25, 2005**AA** ! **EQ 20MG BASE/5ML** **A078988 001** Jun 09, 2008**AA** VINTAGE **EQ 15MG BASE/5ML** **A079010 001** May 26, 2009**AA** WOCKHARDT BIO AG **EQ 5MG BASE/5ML** **A075099 001** Jun 28, 2002**AA** ! **EQ 15MG BASE/5ML** **A076895 001** Oct 04, 2004

! MISSION PHARMA EQ 25MG BASE/5ML A091396 001 Sep 13, 2010

SOLUTION/DROPS; OPHTHALMIC

PREDNISOLONE SODIUM PHOSPHATE

! BAUSCH AND LOMB EQ 0.9% PHOSPHATE A040070 001 Jul 29, 1994

TABLET, ORALLY DISINTEGRATING; ORAL

ORAPRED ODT**AB** + CONCORDIA PHARMS **EQ 10MG BASE** **N021959 001** Jun 01, 2006
INC**AB** + **EQ 15MG BASE** **N021959 002** Jun 01, 2006**AB** +! **EQ 30MG BASE** **N021959 003** Jun 01, 2006PREDNISOLONE SODIUM PHOSPHATE**AB** MYLAN PHARMS INC **EQ 10MG BASE** **A202179 001** Apr 10, 2013**AB** **EQ 15MG BASE** **A202179 002** Apr 10, 2013**AB** **EQ 30MG BASE** **A202179 003** Apr 10, 2013PREDNISOLONE SODIUM PHOSPHATE; SULFACETAMIDE SODIUM

SOLUTION/DROPS; OPHTHALMIC

SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE**AT** ! BAUSCH AND LOMB **EQ 0.23% PHOSPHATE;10%** **A074449 001** Dec 29, 1995**AT** SANDOZ INC **EQ 0.23% PHOSPHATE;10%** **A073630 001** May 27, 1993PREDNISON

SOLUTION; ORAL

PREDNISON

! WEST-WARD PHARMS 5MG/5ML A088703 001 Nov 08, 1984

INT

PREDNISON INTENSOL

! WEST-WARD PHARMS 5MG/ML A088810 001 Feb 20, 1985

INT

TABLET; ORAL

PREDNISON**AB** GENEYORK PHARMS **1MG** **A211496 001** Dec 28, 2018**AB** **2.5MG** **A211495 001** Dec 07, 2018**AB** **5MG** **A211495 002** Dec 07, 2018**AB** **10MG** **A210525 001** Dec 04, 2018**AB** **20MG** **A210525 002** Dec 04, 2018**AB** **50MG** **A210525 003** Dec 04, 2018**AB** HIKMA PHARMS **50MG** **A088465 001** Jun 01, 1984

PRESCRIPTION DRUG PRODUCT LISTPREDNISONE

TABLET; ORAL

PREDNISONE

| | | | | | |
|-----------|---------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | JUBILANT CADISTA | <u>1MG</u> | <u>A040611</u> | <u>001</u> | Jun 06, 2005 |
| <u>AB</u> | | <u>5MG</u> | <u>A040362</u> | <u>002</u> | Aug 29, 2001 |
| <u>AB</u> | | <u>10MG</u> | <u>A040362</u> | <u>001</u> | Aug 29, 2001 |
| <u>AB</u> | | <u>20MG</u> | <u>A040362</u> | <u>003</u> | Jun 29, 2005 |
| <u>AB</u> | MUTUAL PHARM | <u>5MG</u> | <u>A089245</u> | <u>001</u> | Dec 04, 1985 |
| <u>AB</u> | MYLAN PHARMS INC | <u>5MG</u> | <u>A080292</u> | <u>001</u> | |
| <u>AB</u> | | <u>10MG</u> | <u>A088832</u> | <u>001</u> | Dec 04, 1985 |
| <u>AB</u> | | <u>20MG</u> | <u>A083677</u> | <u>001</u> | |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>10MG</u> | <u>A089246</u> | <u>001</u> | Dec 04, 1985 |
| <u>AB</u> | | <u>20MG</u> | <u>A089247</u> | <u>001</u> | Dec 04, 1985 |
| <u>AB</u> | VINTAGE PHARMS | <u>1MG</u> | <u>A040584</u> | <u>001</u> | Dec 21, 2004 |
| <u>AB</u> | | <u>2.5MG</u> | <u>A040581</u> | <u>001</u> | Dec 21, 2004 |
| <u>AB</u> | | <u>5MG</u> | <u>A040256</u> | <u>001</u> | Jul 12, 2002 |
| <u>AB</u> | | <u>10MG</u> | <u>A040256</u> | <u>002</u> | Jul 12, 2002 |
| <u>AB</u> | | <u>20MG</u> | <u>A040392</u> | <u>001</u> | Feb 12, 2003 |
| <u>AB</u> | WATSON LABS | <u>5MG</u> | <u>A080356</u> | <u>001</u> | |
| <u>AB</u> | | <u>10MG</u> | <u>A085162</u> | <u>001</u> | |
| <u>AB</u> | | <u>20MG</u> | <u>A085161</u> | <u>001</u> | |
| <u>AB</u> | ! WEST-WARD PHARMS INT | <u>1MG</u> | <u>A087800</u> | <u>001</u> | Apr 22, 1982 |
| <u>AB</u> | ! | <u>2.5MG</u> | <u>A087801</u> | <u>001</u> | Apr 22, 1982 |
| <u>AB</u> | ! | <u>5MG</u> | <u>A080352</u> | <u>001</u> | |
| <u>AB</u> | ! | <u>10MG</u> | <u>A084122</u> | <u>001</u> | |
| <u>AB</u> | ! | <u>20MG</u> | <u>A087342</u> | <u>001</u> | |
| <u>AB</u> | ! | <u>50MG</u> | <u>A084283</u> | <u>001</u> | |

TABLET, DELAYED RELEASE; ORAL

PREDNISONE

| | | | | | |
|-----------|----------------------|------------|----------------|------------|--------------|
| <u>AB</u> | ACTAVIS LABS FL INC | <u>1MG</u> | <u>A204867</u> | <u>001</u> | Apr 25, 2017 |
| <u>AB</u> | | <u>2MG</u> | <u>A204867</u> | <u>002</u> | Apr 25, 2017 |
| <u>AB</u> | | <u>5MG</u> | <u>A204867</u> | <u>003</u> | Apr 25, 2017 |
| | <u>RAYOS</u> | | | | |
| <u>AB</u> | + HORIZON PHARMA USA | <u>1MG</u> | <u>N202020</u> | <u>001</u> | Jul 26, 2012 |
| <u>AB</u> | + | <u>2MG</u> | <u>N202020</u> | <u>002</u> | Jul 26, 2012 |
| <u>AB</u> | +! | <u>5MG</u> | <u>N202020</u> | <u>003</u> | Jul 26, 2012 |

PREGABALIN

CAPSULE; ORAL

LYRICA

| | | | | | |
|---|-------------|-------|---------|-----|--------------|
| + | PF PRISM CV | 25MG | N021446 | 001 | Dec 30, 2004 |
| + | | 50MG | N021446 | 002 | Dec 30, 2004 |
| + | | 75MG | N021446 | 003 | Dec 30, 2004 |
| + | | 100MG | N021446 | 004 | Dec 30, 2004 |
| + | | 150MG | N021446 | 005 | Dec 30, 2004 |
| + | | 200MG | N021446 | 006 | Dec 30, 2004 |
| + | | 225MG | N021446 | 007 | Dec 30, 2004 |
| + | ! | 300MG | N021446 | 008 | Dec 30, 2004 |

SOLUTION; ORAL

LYRICA

| | | | | | |
|---|-------------|---------|---------|-----|--------------|
| + | PF PRISM CV | 20MG/ML | N022488 | 001 | Jan 04, 2010 |
|---|-------------|---------|---------|-----|--------------|

TABLET, EXTENDED RELEASE; ORAL

LYRICA CR

| | | | | | |
|---|-------------|--------|---------|-----|--------------|
| + | PF PRISM CV | 82.5MG | N209501 | 001 | Oct 11, 2017 |
| + | | 165MG | N209501 | 002 | Oct 11, 2017 |
| + | ! | 330MG | N209501 | 003 | Oct 11, 2017 |

PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

PRILOCAINE HYDROCHLORIDE

| | | | | | |
|---|---------------|----|---------|-----|--------------|
| ! | SEPTODONT INC | 4% | A079235 | 001 | Sep 29, 2010 |
|---|---------------|----|---------|-----|--------------|

PRIMAQUINE PHOSPHATE

TABLET; ORAL

PRIMAQUINE

| | | | | | |
|-----------|-----------------------------|---------------------|----------------|------------|--------------|
| <u>AB</u> | +! SANOFI AVENTIS US | <u>EQ 15MG BASE</u> | <u>N008316</u> | <u>001</u> | |
| | <u>PRIMAQUINE PHOSPHATE</u> | | | | |
| <u>AB</u> | ALVOGEN INC | <u>EQ 15MG BASE</u> | <u>A203924</u> | <u>001</u> | Feb 03, 2014 |
| <u>AB</u> | BAYSHORE PHARMS LLC | <u>EQ 15MG BASE</u> | <u>A204476</u> | <u>001</u> | Feb 25, 2014 |
| <u>AB</u> | NOVAST LABS | <u>EQ 15MG BASE</u> | <u>A206043</u> | <u>001</u> | Jun 23, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

PRIMIDONE

TABLET; ORAL

MYSOLINE

| | | | | | | |
|-----------|---|---------|--------------|----------------|------------|--|
| <u>AB</u> | + | VALEANT | <u>50MG</u> | <u>N009170</u> | <u>003</u> | |
| <u>AB</u> | + | | <u>250MG</u> | <u>N009170</u> | <u>002</u> | |

PRIMIDONE

| | | | | | | |
|-----------|--|-------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | AMNEAL PHARM | <u>50MG</u> | <u>A040866</u> | <u>001</u> | Apr 23, 2008 |
| <u>AB</u> | | | <u>250MG</u> | <u>A040866</u> | <u>002</u> | Apr 23, 2008 |
| <u>AB</u> | | ANDA REPOSITORY | <u>50MG</u> | <u>A040626</u> | <u>001</u> | Sep 29, 2005 |
| <u>AB</u> | | | <u>250MG</u> | <u>A040626</u> | <u>002</u> | Sep 29, 2005 |
| <u>AB</u> | | HIKMA INTL PHARMS | <u>250MG</u> | <u>A040667</u> | <u>002</u> | Jul 27, 2006 |
| <u>AB</u> | | LANNETT | <u>50MG</u> | <u>A084903</u> | <u>002</u> | May 24, 2001 |
| <u>AB</u> | | | <u>250MG</u> | <u>A084903</u> | <u>001</u> | |
| <u>AB</u> | | OXFORD PHARMS | <u>50MG</u> | <u>A040586</u> | <u>001</u> | Feb 24, 2005 |
| <u>AB</u> | | | <u>250MG</u> | <u>A040586</u> | <u>002</u> | Feb 24, 2005 |
| <u>AB</u> | | WATSON LABS | <u>250MG</u> | <u>A083551</u> | <u>001</u> | |

PROBENECID

TABLET; ORAL

PROBALAN

| | | | | | | |
|-----------|--|---------|--------------|----------------|------------|--|
| <u>AB</u> | | LANNETT | <u>500MG</u> | <u>A080966</u> | <u>001</u> | |
|-----------|--|---------|--------------|----------------|------------|--|

PROBENECID

| | | | | | | |
|-----------|---|------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ! | MYLAN | <u>500MG</u> | <u>A084211</u> | <u>002</u> | |
| <u>AB</u> | | WATSON LABS TEVA | <u>500MG</u> | <u>A084442</u> | <u>004</u> | Mar 29, 1983 |

PROCAINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

| | | | | | | |
|-----------|---|-----------------|-----------------|----------------|------------|--------------|
| <u>AP</u> | ! | HOSPIRA | <u>100MG/ML</u> | <u>A089069</u> | <u>001</u> | Feb 12, 1986 |
| <u>AP</u> | | INTL MEDICATION | <u>100MG/ML</u> | <u>A088636</u> | <u>001</u> | Jul 31, 1984 |
| <u>AP</u> | | NEXUS PHARMS | <u>100MG/ML</u> | <u>A206332</u> | <u>001</u> | Oct 13, 2017 |
| <u>AP</u> | | | <u>500MG/ML</u> | <u>A206332</u> | <u>002</u> | Oct 13, 2017 |
| | ! | HOSPIRA | 500MG/ML | A089070 | 001 | Feb 12, 1986 |

PROCARBAZINE HYDROCHLORIDE

CAPSULE; ORAL

MATULANE

| | | | | | | |
|---|---|---------------------|--------------|---------|-----|--|
| + | ! | LEADIANT BIOSCI INC | EQ 50MG BASE | N016785 | 001 | |
|---|---|---------------------|--------------|---------|-----|--|

PROCHLORPERAZINE

SUPPOSITORY; RECTAL

COMPRO

| | | | | | | |
|-----------|--|-------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | PADDOCK LLC | <u>25MG</u> | <u>A040246</u> | <u>001</u> | Jun 28, 2000 |
|-----------|--|-------------|-------------|----------------|------------|--------------|

PROCHLORPERAZINE

| | | | | | | |
|-----------|---|--------------|-------------|----------------|------------|--------------|
| <u>AB</u> | ! | G AND W LABS | <u>25MG</u> | <u>A040058</u> | <u>001</u> | Nov 24, 1993 |
|-----------|---|--------------|-------------|----------------|------------|--------------|

PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION

PROCHLORPERAZINE EDISYLATE

| | | | | | | |
|-----------|---|-------------------------|-----------------------|----------------|------------|--------------|
| <u>AP</u> | | ATHENEX INC | <u>EQ 5MG BASE/ML</u> | <u>A040540</u> | <u>001</u> | May 28, 2004 |
| <u>AP</u> | ! | EMCURE PHARMS LTD | <u>EQ 5MG BASE/ML</u> | <u>A204147</u> | <u>001</u> | Oct 15, 2013 |
| <u>AP</u> | | MYLAN LABS LTD | <u>EQ 5MG BASE/ML</u> | <u>A210710</u> | <u>001</u> | Oct 25, 2018 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>EQ 5MG BASE/ML</u> | <u>A089903</u> | <u>001</u> | Aug 29, 1989 |

PROCHLORPERAZINE MALEATE

TABLET; ORAL

PROCHLORPERAZINE MALEATE

| | | | | | | |
|-----------|---|-------------|---------------------|----------------|------------|--------------|
| <u>AB</u> | | MYLAN | <u>EQ 5MG BASE</u> | <u>A040185</u> | <u>002</u> | Oct 28, 1996 |
| <u>AB</u> | | | <u>EQ 10MG BASE</u> | <u>A040185</u> | <u>001</u> | Oct 28, 1996 |
| <u>AB</u> | | SANDOZ | <u>EQ 5MG BASE</u> | <u>A040101</u> | <u>001</u> | Jul 19, 1996 |
| <u>AB</u> | ! | | <u>EQ 10MG BASE</u> | <u>A040101</u> | <u>002</u> | Jul 19, 1996 |
| <u>AB</u> | | TEVA PHARMS | <u>EQ 5MG BASE</u> | <u>A040120</u> | <u>001</u> | Jul 11, 1996 |
| <u>AB</u> | | | <u>EQ 10MG BASE</u> | <u>A040120</u> | <u>002</u> | Jul 11, 1996 |

PROCOMP

| | | | | | | |
|-----------|--|------------------|---------------------|----------------|------------|--------------|
| <u>AB</u> | | JUBILANT CADISTA | <u>EQ 5MG BASE</u> | <u>A040268</u> | <u>001</u> | Feb 27, 1998 |
| <u>AB</u> | | | <u>EQ 10MG BASE</u> | <u>A040268</u> | <u>002</u> | Feb 27, 1998 |

PROGESTERONE

CAPSULE; ORAL

PROGESTERONE

| | | | | | | |
|-----------|--|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | AMNEAL PHARMS NY | <u>100MG</u> | <u>A207724</u> | <u>001</u> | Sep 07, 2017 |
| <u>AB</u> | | | <u>200MG</u> | <u>A207724</u> | <u>002</u> | Sep 07, 2017 |
| <u>AB</u> | | BIONPHARMA INC | <u>100MG</u> | <u>A200900</u> | <u>001</u> | Aug 16, 2013 |
| <u>AB</u> | | | <u>200MG</u> | <u>A200900</u> | <u>002</u> | Aug 16, 2013 |
| <u>AB</u> | | DR REDDYS LABS INC | <u>100MG</u> | <u>A208801</u> | <u>001</u> | Feb 28, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

PROGESTERONE

CAPSULE; ORAL

PROGESTERONE

| | | | | | |
|-----------|---------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | <u>200MG</u> | <u>A208801</u> | <u>002</u> | Feb 28, 2017 |
| <u>AB</u> | EUGIA PHARMA | <u>100MG</u> | <u>A211285</u> | <u>001</u> | Oct 26, 2018 |
| <u>AB</u> | | <u>200MG</u> | <u>A211285</u> | <u>002</u> | Oct 26, 2018 |
| <u>AB</u> | SANDOZ INC | <u>100MG</u> | <u>A205229</u> | <u>001</u> | Oct 20, 2017 |
| <u>AB</u> | | <u>200MG</u> | <u>A205229</u> | <u>002</u> | Oct 20, 2017 |
| <u>AB</u> | SOFGEN PHARMS | <u>100MG</u> | <u>A200456</u> | <u>001</u> | Sep 28, 2012 |
| <u>AB</u> | | <u>200MG</u> | <u>A200456</u> | <u>002</u> | Sep 28, 2012 |

PROMETRIUM

| | | | | | | |
|-----------|---|---------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | VIRTUS PHARMS | <u>100MG</u> | <u>N019781</u> | <u>001</u> | May 14, 1998 |
| <u>AB</u> | + | ! | <u>200MG</u> | <u>N019781</u> | <u>002</u> | Oct 15, 1999 |

GEL; VAGINAL

CRINONE

| | | | | | | |
|---|---|--------------------|----|---------|-----|--------------|
| + | ! | ALLERGAN SALES LLC | 4% | N020701 | 001 | Jul 31, 1997 |
| + | ! | | 8% | N020701 | 002 | Jul 31, 1997 |

INJECTABLE; INJECTION

PROGESTERONE

| | | | | | | | |
|-----------|---|---|---------------------|----------------|----------------|------------|--------------|
| <u>AO</u> | + | ! | ACTAVIS LABS UT INC | <u>50MG/ML</u> | <u>N017362</u> | <u>002</u> | |
| <u>AO</u> | | | EUGIA PHARMA | <u>50MG/ML</u> | <u>A210965</u> | <u>001</u> | Dec 06, 2018 |
| <u>AO</u> | | | FRESENIUS KABI USA | <u>50MG/ML</u> | <u>A075906</u> | <u>001</u> | Apr 25, 2001 |
| <u>AO</u> | | | HIKMA FARMACEUTICA | <u>50MG/ML</u> | <u>A091033</u> | <u>001</u> | Oct 28, 2010 |
| <u>AO</u> | | | LUITPOLD | <u>50MG/ML</u> | <u>A090845</u> | <u>001</u> | Jun 22, 2009 |

INSERT; VAGINAL

ENDOMETRIN

| | | | | | | |
|---|---|---------|-------|---------|-----|--------------|
| + | ! | FERRING | 100MG | N022057 | 001 | Jun 21, 2007 |
|---|---|---------|-------|---------|-----|--------------|

PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROMETHAZINE HYDROCHLORIDE

| | | | | | | |
|-----------|---|------------------|----------------|----------------|------------|--------------|
| <u>AP</u> | ! | WEST-WARD PHARMS | <u>25MG/ML</u> | <u>A083312</u> | <u>001</u> | |
| <u>AP</u> | ! | INT | <u>50MG/ML</u> | <u>A083312</u> | <u>002</u> | |
| <u>AP</u> | | X-GEN PHARMS | <u>25MG/ML</u> | <u>A040737</u> | <u>001</u> | Apr 24, 2008 |
| <u>AP</u> | | | <u>50MG/ML</u> | <u>A040737</u> | <u>002</u> | Apr 24, 2008 |

SUPPOSITORY; RECTAL

PROMETHAZINE HYDROCHLORIDE

| | | | | | | |
|-----------|---|------------------|---------------|----------------|------------|--------------|
| <u>AB</u> | | G AND W LABS | <u>12.5MG</u> | <u>A040428</u> | <u>002</u> | Mar 31, 2003 |
| <u>AB</u> | ! | | <u>25MG</u> | <u>A040428</u> | <u>001</u> | Feb 05, 2002 |
| <u>AB</u> | | PERRIGO NEW YORK | <u>12.5MG</u> | <u>A040500</u> | <u>001</u> | Jun 30, 2003 |
| <u>AB</u> | | | <u>25MG</u> | <u>A040500</u> | <u>002</u> | Jun 30, 2003 |
| <u>AB</u> | | TARO | <u>12.5MG</u> | <u>A040603</u> | <u>001</u> | Oct 26, 2006 |
| <u>AB</u> | | | <u>25MG</u> | <u>A040603</u> | <u>002</u> | Oct 26, 2006 |
| <u>AB</u> | | WATSON LABS INC | <u>12.5MG</u> | <u>A040479</u> | <u>001</u> | Jun 24, 2003 |
| <u>AB</u> | | | <u>25MG</u> | <u>A040479</u> | <u>002</u> | Jun 24, 2003 |

PROMETHEGAN

| | | | | | | |
|---|---|--------------|------|---------|-----|--------------|
| ! | ! | G AND W LABS | 50MG | A087165 | 001 | Aug 14, 1987 |
|---|---|--------------|------|---------|-----|--------------|

SYRUP; ORAL

PROMETHAZINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|------------------|-------------------|----------------|------------|--------------|
| <u>AA</u> | | AMNEAL PHARMS | <u>6.25MG/5ML</u> | <u>A040882</u> | <u>001</u> | Dec 30, 2009 |
| <u>AA</u> | | HI TECH PHARMA | <u>6.25MG/5ML</u> | <u>A040026</u> | <u>001</u> | Sep 25, 1998 |
| <u>AA</u> | | NOSTRUM LABS INC | <u>6.25MG/5ML</u> | <u>A040891</u> | <u>001</u> | Mar 13, 2009 |
| <u>AA</u> | | TARO | <u>6.25MG/5ML</u> | <u>A040718</u> | <u>001</u> | Apr 04, 2007 |
| <u>AA</u> | | TRIS PHARMA INC | <u>6.25MG/5ML</u> | <u>A091675</u> | <u>001</u> | Jun 28, 2012 |
| <u>AA</u> | | VINTAGE | <u>6.25MG/5ML</u> | <u>A040643</u> | <u>001</u> | Apr 26, 2006 |

PROMETHAZINE PLAIN

| | | | | | | |
|-----------|---|------------------|-------------------|----------------|------------|--------------|
| <u>AA</u> | ! | WOCKHARDT BIO AG | <u>6.25MG/5ML</u> | <u>A087953</u> | <u>001</u> | Nov 15, 1982 |
|-----------|---|------------------|-------------------|----------------|------------|--------------|

TABLET; ORAL

PROMETHAZINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|------------------|---------------|----------------|------------|--------------|
| <u>AB</u> | | AMNEAL PHARMS NY | <u>12.5MG</u> | <u>A091179</u> | <u>001</u> | Dec 13, 2010 |
| <u>AB</u> | | | <u>25MG</u> | <u>A091179</u> | <u>002</u> | Dec 13, 2010 |
| <u>AB</u> | | | <u>50MG</u> | <u>A091179</u> | <u>003</u> | Dec 13, 2010 |
| <u>AB</u> | | KVK TECH | <u>12.5MG</u> | <u>A040712</u> | <u>002</u> | May 04, 2007 |
| <u>AB</u> | | | <u>25MG</u> | <u>A040712</u> | <u>001</u> | Jul 31, 2006 |
| <u>AB</u> | | | <u>50MG</u> | <u>A040712</u> | <u>003</u> | Jul 31, 2006 |
| <u>AB</u> | | PRINSTON INC | <u>12.5MG</u> | <u>A040622</u> | <u>001</u> | Jul 18, 2006 |
| <u>AB</u> | | | <u>25MG</u> | <u>A040622</u> | <u>002</u> | Jul 18, 2006 |
| <u>AB</u> | | | <u>50MG</u> | <u>A040622</u> | <u>003</u> | Jul 18, 2006 |
| <u>AB</u> | | QUAGEN | <u>12.5MG</u> | <u>A040673</u> | <u>001</u> | Mar 05, 2008 |
| <u>AB</u> | | | <u>25MG</u> | <u>A040673</u> | <u>002</u> | Mar 05, 2008 |
| <u>AB</u> | | | <u>50MG</u> | <u>A040673</u> | <u>003</u> | Mar 05, 2008 |
| <u>AB</u> | | SANDOZ | <u>25MG</u> | <u>A084176</u> | <u>003</u> | |

PRESCRIPTION DRUG PRODUCT LIST

PROMETHAZINE HYDROCHLORIDE

TABLET; ORAL

PROMETHAZINE HYDROCHLORIDE

| | | | | | |
|-----------|--------------------|---------------|----------------|------------|--------------|
| <u>AB</u> | ! | <u>50MG</u> | <u>A084176</u> | <u>001</u> | |
| <u>AB</u> | STRIDES PHARMA | <u>12.5MG</u> | <u>A209177</u> | <u>001</u> | Jun 30, 2017 |
| <u>AB</u> | | <u>25MG</u> | <u>A209177</u> | <u>002</u> | Jun 30, 2017 |
| <u>AB</u> | | <u>50MG</u> | <u>A209177</u> | <u>003</u> | Jun 30, 2017 |
| <u>AB</u> | SUN PHARM INDS INC | <u>12.5MG</u> | <u>A040863</u> | <u>001</u> | Dec 30, 2008 |
| <u>AB</u> | | <u>25MG</u> | <u>A040863</u> | <u>002</u> | Dec 30, 2008 |
| <u>AB</u> | | <u>50MG</u> | <u>A040863</u> | <u>003</u> | Dec 30, 2008 |
| <u>AB</u> | WATSON LABS | <u>25MG</u> | <u>A083426</u> | <u>001</u> | |
| <u>AB</u> | | <u>50MG</u> | <u>A083711</u> | <u>001</u> | |
| <u>AB</u> | ZYDUS PHARMS USA | <u>12.5MG</u> | <u>A040596</u> | <u>001</u> | Nov 18, 2005 |
| <u>AB</u> | | <u>25MG</u> | <u>A040596</u> | <u>002</u> | Nov 18, 2005 |
| <u>AB</u> | | <u>50MG</u> | <u>A040596</u> | <u>003</u> | Nov 18, 2005 |
| | IMPAX LABS | 12.5MG | A040791 | 002 | Feb 12, 2008 |
| | | 25MG | A040791 | 003 | Feb 12, 2008 |

PROPAFENONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PROPAFENONE HYDROCHLORIDE

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | GLENMARK PHARMS LTD | <u>225MG</u> | <u>A205268</u> | <u>001</u> | Sep 08, 2017 |
| <u>AB</u> | | <u>325MG</u> | <u>A205268</u> | <u>002</u> | Sep 08, 2017 |
| <u>AB</u> | | <u>425MG</u> | <u>A205268</u> | <u>003</u> | Sep 08, 2017 |
| <u>AB</u> | MYLAN PHARMS INC | <u>225MG</u> | <u>A203803</u> | <u>001</u> | Apr 29, 2016 |
| <u>AB</u> | | <u>325MG</u> | <u>A203803</u> | <u>002</u> | Apr 29, 2016 |
| <u>AB</u> | | <u>425MG</u> | <u>A203803</u> | <u>003</u> | Apr 29, 2016 |
| <u>AB</u> | PAR PHARM | <u>225MG</u> | <u>A078540</u> | <u>001</u> | Oct 18, 2010 |
| <u>AB</u> | | <u>325MG</u> | <u>A078540</u> | <u>002</u> | Oct 18, 2010 |
| <u>AB</u> | | <u>425MG</u> | <u>A078540</u> | <u>003</u> | Oct 18, 2010 |
| <u>AB</u> | SINOTHERAPEUTICS INC | <u>225MG</u> | <u>A210339</u> | <u>001</u> | Jan 04, 2019 |
| <u>AB</u> | | <u>325MG</u> | <u>A210339</u> | <u>002</u> | Jan 04, 2019 |
| <u>AB</u> | | <u>425MG</u> | <u>A210339</u> | <u>003</u> | Jan 04, 2019 |
| <u>AB</u> | WATSON LABS INC | <u>225MG</u> | <u>A202688</u> | <u>001</u> | Aug 24, 2015 |
| <u>AB</u> | | <u>325MG</u> | <u>A202688</u> | <u>002</u> | Aug 24, 2015 |
| <u>AB</u> | | <u>425MG</u> | <u>A202688</u> | <u>003</u> | Aug 24, 2015 |
| <u>AB</u> | WILSHIRE PHARMS INC | <u>225MG</u> | <u>A205956</u> | <u>001</u> | Jul 02, 2018 |
| <u>AB</u> | | <u>325MG</u> | <u>A205956</u> | <u>002</u> | Jul 02, 2018 |
| <u>AB</u> | | <u>425MG</u> | <u>A205956</u> | <u>003</u> | Jul 02, 2018 |
| | <u>RYTHMOL SR</u> | | | | |
| <u>AB</u> | + GLAXOSMITHKLINE LLC | <u>225MG</u> | <u>N021416</u> | <u>001</u> | Sep 04, 2003 |
| <u>AB</u> | + | <u>325MG</u> | <u>N021416</u> | <u>002</u> | Sep 04, 2003 |
| <u>AB</u> | + | <u>425MG</u> | <u>N021416</u> | <u>003</u> | Sep 04, 2003 |

TABLET; ORAL

PROPAFENONE HYDROCHLORIDE

| | | | | | |
|-----------|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ANI PHARMS INC | <u>150MG</u> | <u>A076550</u> | <u>001</u> | Apr 23, 2004 |
| <u>AB</u> | | <u>225MG</u> | <u>A076550</u> | <u>002</u> | Apr 23, 2004 |
| <u>AB</u> | | <u>300MG</u> | <u>A076550</u> | <u>003</u> | Apr 23, 2004 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>150MG</u> | <u>A202445</u> | <u>001</u> | May 11, 2016 |
| <u>AB</u> | | <u>225MG</u> | <u>A202445</u> | <u>002</u> | May 11, 2016 |
| <u>AB</u> | | <u>300MG</u> | <u>A202445</u> | <u>003</u> | May 11, 2016 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>150MG</u> | <u>A075998</u> | <u>001</u> | Nov 29, 2001 |
| <u>AB</u> | | <u>225MG</u> | <u>A075998</u> | <u>002</u> | Nov 29, 2001 |
| <u>AB</u> | | <u>300MG</u> | <u>A075998</u> | <u>003</u> | Nov 29, 2001 |
| <u>AB</u> | VINTAGE PHARMS | <u>150MG</u> | <u>A075938</u> | <u>001</u> | Oct 17, 2002 |
| <u>AB</u> | | <u>225MG</u> | <u>A075938</u> | <u>002</u> | Oct 17, 2002 |
| <u>AB</u> | ! | <u>300MG</u> | <u>A075938</u> | <u>003</u> | Oct 17, 2002 |
| <u>AB</u> | WATSON LABS | <u>150MG</u> | <u>A075203</u> | <u>001</u> | Oct 24, 2000 |
| <u>AB</u> | | <u>225MG</u> | <u>A075203</u> | <u>002</u> | Oct 24, 2000 |

PROPANTHELINE BROMIDE

TABLET; ORAL

PROPANTHELINE BROMIDE

| | | | | | |
|---|-------------------------|------|---------|-----|--|
| ! | WEST-WARD PHARMS INT | 15MG | A080927 | 002 | |
|---|-------------------------|------|---------|-----|--|

PRESCRIPTION DRUG PRODUCT LIST

PROPARACAINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

ALCAINE

| | | | | |
|------------------------------------------|----------------|-----------------|--------------------|---------------------------------|
| AT | ALCON LABS INC | 0.5% | A080027 001 | |
| <u>PROPARACAINE HYDROCHLORIDE</u> | | | | |
| AT | ! | AKORN INC | 0.5% | A040277 001 Mar 16, 2000 |
| AT | | BAUSCH AND LOMB | 0.5% | A040074 001 Sep 29, 1995 |

PROPOFOL

INJECTABLE; INJECTION

DIPRIVAN

| | | | | |
|-----------|---|--------------------|----------------|---------------------------------|
| AB | + | FRESENIUS KABI USA | 10MG/ML | N019627 002 Jun 11, 1996 |
|-----------|---|--------------------|----------------|---------------------------------|

PROPOFOL

| | | | | |
|-----------|--|--------------------|----------------|---------------------------------|
| AB | | DR REDDYS LABS INC | 10MG/ML | A205067 001 Nov 15, 2018 |
| AB | | HOSPIRA | 10MG/ML | A077908 001 Mar 17, 2006 |
| AB | | SAGENT PHARMS | 10MG/ML | A075102 001 Jan 04, 1999 |
| AB | | WATSON LABS INC | 10MG/ML | A205307 001 Dec 22, 2015 |

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

INDERAL LA

| | | | | |
|-----------|---|----------------|--------------|---------------------------------|
| AB | + | ANI PHARMS INC | 60MG | N018553 004 Mar 18, 1987 |
| AB | + | | 80MG | N018553 002 Apr 19, 1983 |
| AB | + | | 120MG | N018553 003 Apr 19, 1983 |
| AB | + | | 160MG | N018553 001 Apr 19, 1983 |

PROPRANOLOL HYDROCHLORIDE

| | | | | |
|-----------|--|-------------------------|--------------|---------------------------------|
| AB | | ACTAVIS ELIZABETH | 60MG | A078494 001 Aug 10, 2007 |
| AB | | | 80MG | A078494 002 Aug 10, 2007 |
| AB | | | 120MG | A078494 003 Aug 10, 2007 |
| AB | | | 160MG | A078494 004 Aug 10, 2007 |
| AB | | NORTEC DEV ASSOC | 60MG | A078065 001 Jan 26, 2007 |
| AB | | | 80MG | A078065 002 Jan 26, 2007 |
| AB | | | 120MG | A078065 003 Jan 26, 2007 |
| AB | | | 160MG | A078065 004 Jan 26, 2007 |
| AB | | RP SCHERER | 60MG | A078703 001 Jul 15, 2011 |
| AB | | | 80MG | A078703 002 Jul 15, 2011 |
| AB | | | 120MG | A078703 003 Jul 15, 2011 |
| AB | | | 160MG | A078703 004 Jul 15, 2011 |
| AB | | ZYDUS PHARMS USA INC | 60MG | A090321 001 Mar 25, 2011 |
| AB | | | 80MG | A090321 002 Mar 25, 2011 |
| AB | | | 120MG | A090321 003 Mar 25, 2011 |
| AB | | | 160MG | A090321 004 Mar 25, 2011 |

INNOPRAN XL

| | | | | |
|----|--|----------------|-------|--------------------------|
| BX | | ANI PHARMS INC | 80MG | N021438 001 Mar 12, 2003 |
| BX | | | 120MG | N021438 002 Mar 12, 2003 |

INJECTABLE; INJECTION

PROPRANOLOL HYDROCHLORIDE

| | | | | |
|-----------|---|--------------------|---------------|---------------------------------|
| AP | | ATHENEX INC | 1MG/ML | A075792 001 Aug 29, 2000 |
| AP | | FRESENIUS KABI USA | 1MG/ML | A075826 001 Aug 31, 2001 |
| | ! | HIKMA FARMACEUTICA | 1MG/ML | A077760 001 Jan 31, 2008 |

SOLUTION; ORAL

HEMANGEOL

| | | | | |
|--|---|--------------------|-----------|--------------------------|
| | + | PIERRE FABRE DERMA | 4.28MG/ML | N205410 001 Mar 14, 2014 |
|--|---|--------------------|-----------|--------------------------|

PROPRANOLOL HYDROCHLORIDE

| | | | | |
|--|---|-------------------------|----------|--------------------------|
| | ! | WEST-WARD PHARMS INT | 20MG/5ML | A070979 001 May 15, 1987 |
|--|---|-------------------------|----------|--------------------------|

| | | | | |
|--|---|--|----------|--------------------------|
| | ! | | 40MG/5ML | A070690 001 May 15, 1987 |
|--|---|--|----------|--------------------------|

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

| | | | | |
|-----------|---|----------------|-------------|---------------------------------|
| AB | | IMPAX LABS INC | 10MG | A071972 001 Apr 06, 1988 |
| AB | | | 20MG | A071972 002 Apr 06, 1988 |
| AB | | | 40MG | A071972 003 Apr 06, 1988 |
| AB | | | 60MG | A071976 002 May 13, 1986 |
| AB | ! | | 80MG | A071976 001 Apr 06, 1988 |
| AB | | IPCA LABS LTD | 10MG | A078955 001 Jun 02, 2008 |
| AB | | | 20MG | A078955 002 Jun 02, 2008 |
| AB | | | 40MG | A078955 003 Jun 02, 2008 |
| AB | | | 60MG | A078955 004 Jun 02, 2008 |
| AB | | | 80MG | A078955 005 Jun 02, 2008 |
| AB | | MYLAN | 10MG | A070213 002 Nov 19, 1985 |
| AB | | | 20MG | A070213 003 Nov 19, 1985 |
| AB | | | 40MG | A070213 001 Nov 19, 1985 |
| AB | | | 60MG | A070213 005 Apr 08, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

| | | | | | |
|-----------|--------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | <u>80MG</u> | <u>A070213</u> | <u>004</u> | Nov 19, 1985 |
| <u>AB</u> | NORTHSTAR HLTHCARE | <u>10MG</u> | <u>A078213</u> | <u>001</u> | Jan 10, 2008 |
| <u>AB</u> | | <u>20MG</u> | <u>A078213</u> | <u>002</u> | Jan 10, 2008 |
| <u>AB</u> | | <u>40MG</u> | <u>A078213</u> | <u>003</u> | Jan 10, 2008 |
| <u>AB</u> | | <u>60MG</u> | <u>A078213</u> | <u>004</u> | Jan 10, 2008 |
| <u>AB</u> | | <u>80MG</u> | <u>A078213</u> | <u>005</u> | Jan 10, 2008 |
| <u>AB</u> | VINTAGE PHARMS | <u>10MG</u> | <u>A070221</u> | <u>002</u> | Aug 01, 1986 |
| <u>AB</u> | | <u>20MG</u> | <u>A070221</u> | <u>003</u> | Aug 01, 1986 |
| <u>AB</u> | | <u>40MG</u> | <u>A070219</u> | <u>001</u> | Aug 01, 1986 |
| <u>AB</u> | | <u>40MG</u> | <u>A070221</u> | <u>004</u> | Aug 01, 1986 |
| <u>AB</u> | | <u>60MG</u> | <u>A070221</u> | <u>005</u> | Sep 24, 1986 |
| <u>AB</u> | | <u>80MG</u> | <u>A070221</u> | <u>001</u> | Apr 14, 1986 |
| <u>AB</u> | WATSON LABS | <u>10MG</u> | <u>A070175</u> | <u>001</u> | May 13, 1986 |
| <u>AB</u> | | <u>20MG</u> | <u>A070176</u> | <u>001</u> | May 13, 1986 |
| <u>AB</u> | | <u>40MG</u> | <u>A070177</u> | <u>001</u> | May 13, 1986 |
| <u>AB</u> | | <u>60MG</u> | <u>A070178</u> | <u>002</u> | Apr 23, 2018 |
| <u>AB</u> | | <u>80MG</u> | <u>A070178</u> | <u>001</u> | May 13, 1986 |

PROPYLTHIOURACIL

TABLET; ORAL

PROPYLTHIOURACIL

| | | | | | |
|----|-------------------|------|---------|-----|--|
| BD | ACTAVIS ELIZABETH | 50MG | A080172 | 001 | |
| BD | ! DAVA PHARMS INC | 50MG | N006188 | 001 | |

PROTAMINE SULFATE

INJECTABLE; INJECTION

PROTAMINE SULFATE

! FRESENIUS KABI USA 10MG/ML A089454 001 Apr 07, 1987

PROTRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

PROTRIPTYLINE HYDROCHLORIDE

| | | | | | |
|-----------|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | EPIC PHARMA LLC | <u>5MG</u> | <u>A202220</u> | <u>001</u> | Nov 19, 2012 |
| <u>AB</u> | | <u>10MG</u> | <u>A202220</u> | <u>002</u> | Nov 19, 2012 |
| <u>AB</u> | SIGMAPHARM LABS LLC | <u>5MG</u> | <u>A090462</u> | <u>001</u> | May 03, 2010 |
| <u>AB</u> | | <u>10MG</u> | <u>A090462</u> | <u>002</u> | May 03, 2010 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>5MG</u> | <u>A078913</u> | <u>001</u> | Sep 16, 2008 |
| <u>AB</u> | | <u>10MG</u> | <u>A078913</u> | <u>002</u> | Sep 16, 2008 |
| | <u>VIVACTIL</u> | | | | |
| <u>AB</u> | ODYSSEY PHARMS | <u>5MG</u> | <u>A073644</u> | <u>001</u> | Aug 24, 1995 |
| <u>AB</u> | ! | <u>10MG</u> | <u>A073645</u> | <u>001</u> | Aug 24, 1995 |

PRUCALOPRIDE SUCCINATE

TABLET; ORAL

MOTTEGRITY+ SHIRE DEV LLC EQ 1MG BASE N210166 001 Dec 14, 2018
+! EQ 2MG BASE N210166 002 Dec 14, 2018PYRAZINAMIDE

TABLET; ORAL

PYRAZINAMIDE

| | | | | | |
|-----------|-------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | AKORN | <u>500MG</u> | <u>A081319</u> | <u>001</u> | Jun 30, 1992 |
| <u>AB</u> | ! DAVA PHARMS INC | <u>500MG</u> | <u>A080157</u> | <u>001</u> | |

PYRIDOSTIGMINE BROMIDE

INJECTABLE; INJECTION

MESTINONAP +! VALEANT PHARM INTL 5MG/ML N009830 001REGONOLAP SANDOZ INC 5MG/ML N017398 001

SYRUP; ORAL

MESTINON

+! VALEANT PHARMS 60MG/5ML N015193 001

TABLET; ORAL

MESTINONAB +! VALEANT PHARMS LLC 60MG N009829 002PYRIDOSTIGMINE BROMIDE

| | | | | | |
|-----------|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | APNAR PHARMA LP | <u>60MG</u> | <u>A211181</u> | <u>001</u> | Jul 20, 2018 |
| <u>AB</u> | IMPAX LABS | <u>60MG</u> | <u>A040502</u> | <u>001</u> | Apr 24, 2003 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>60MG</u> | <u>A205650</u> | <u>001</u> | Jun 22, 2015 |

PRESCRIPTION DRUG PRODUCT LIST

PYRIDOSTIGMINE BROMIDE

TABLET, EXTENDED RELEASE;ORAL

MESTINON

| | | | | | | |
|-----------|-----------|---------------------------|--------------|-----------------------|-------------------|--|
| AB | +! | VALEANT PHARMS LLC | 180MG | <u>N011665</u> | <u>001</u> | |
|-----------|-----------|---------------------------|--------------|-----------------------|-------------------|--|

PYRIDOSTIGMINE BROMIDE

| | | | | | | |
|-----------|--|-----------------------|--------------|-----------------------|-------------------|---------------------|
| AB | | ALVOGEN MALTA | 180MG | <u>A204737</u> | <u>001</u> | Jun 26, 2015 |
| AB | | IMPAX LABS INC | 180MG | <u>A203184</u> | <u>001</u> | Sep 15, 2015 |
| AB | | RISING PHARMS | 180MG | <u>A205464</u> | <u>001</u> | Aug 15, 2017 |

PYRIDOXINE HYDROCHLORIDE

INJECTABLE; INJECTION

PYRIDOXINE HYDROCHLORIDE

| | | | | | |
|----------|---------------------------|-----------------|-----------------------|-------------------|--|
| ! | FRESENIUS KABI USA | 100MG/ML | <u>A080618</u> | <u>001</u> | |
|----------|---------------------------|-----------------|-----------------------|-------------------|--|

PYRIMETHAMINE

TABLET; ORAL

DARAPRIM

| | | | | | |
|-----------|-------------------------|-------------|-----------------------|-------------------|--|
| +! | VYERA PHARMS LLC | 25MG | <u>N008578</u> | <u>001</u> | |
|-----------|-------------------------|-------------|-----------------------|-------------------|--|

QUAZEPAM

TABLET; ORAL

DORAL

| | | | | | |
|-----------|--------------------|-------------|-----------------------|-------------------|---------------------|
| +! | GALT PHARMS | 15MG | <u>N018708</u> | <u>001</u> | Dec 27, 1985 |
|-----------|--------------------|-------------|-----------------------|-------------------|---------------------|

QUETIAPINE FUMARATE

TABLET; ORAL

QUETIAPINE FUMARATE

| | | | | | | |
|-----------|--|-----------------------------|-----------------------------|-----------------------|-------------------|---------------------|
| AB | | ACCORD HLTHCARE | <u>EQ 25MG BASE</u> | <u>A202152</u> | <u>001</u> | Mar 27, 2012 |
| AB | | | <u>EQ 50MG BASE</u> | <u>A202152</u> | <u>002</u> | Mar 27, 2012 |
| AB | | | <u>EQ 100MG BASE</u> | <u>A202152</u> | <u>003</u> | Mar 27, 2012 |
| AB | | | <u>EQ 200MG BASE</u> | <u>A202152</u> | <u>004</u> | Mar 27, 2012 |
| AB | | | <u>EQ 300MG BASE</u> | <u>A202152</u> | <u>005</u> | Mar 27, 2012 |
| AB | | | <u>EQ 400MG BASE</u> | <u>A202152</u> | <u>006</u> | Mar 27, 2012 |
| AB | | ALEMBIC PHARMS LTD | <u>EQ 25MG BASE</u> | <u>A203390</u> | <u>001</u> | Oct 28, 2014 |
| AB | | | <u>EQ 50MG BASE</u> | <u>A203390</u> | <u>002</u> | Oct 28, 2014 |
| AB | | | <u>EQ 100MG BASE</u> | <u>A203390</u> | <u>003</u> | Oct 28, 2014 |
| AB | | | <u>EQ 200MG BASE</u> | <u>A203390</u> | <u>004</u> | Oct 28, 2014 |
| AB | | | <u>EQ 300MG BASE</u> | <u>A203390</u> | <u>005</u> | Oct 28, 2014 |
| AB | | | <u>EQ 400MG BASE</u> | <u>A203390</u> | <u>006</u> | Oct 28, 2014 |
| AB | | ALKEM LABS LTD | <u>EQ 25MG BASE</u> | <u>A201504</u> | <u>001</u> | Feb 12, 2013 |
| AB | | | <u>EQ 50MG BASE</u> | <u>A201504</u> | <u>002</u> | Feb 12, 2013 |
| AB | | | <u>EQ 100MG BASE</u> | <u>A201504</u> | <u>003</u> | Feb 12, 2013 |
| AB | | | <u>EQ 150MG BASE</u> | <u>A201504</u> | <u>004</u> | Feb 12, 2013 |
| AB | | | <u>EQ 200MG BASE</u> | <u>A201504</u> | <u>005</u> | Feb 12, 2013 |
| AB | | | <u>EQ 300MG BASE</u> | <u>A201504</u> | <u>006</u> | Feb 12, 2013 |
| AB | | | <u>EQ 400MG BASE</u> | <u>A201504</u> | <u>007</u> | Feb 12, 2013 |
| AB | | APOTEX INC | <u>EQ 25MG BASE</u> | <u>A090960</u> | <u>001</u> | Mar 27, 2012 |
| AB | | | <u>EQ 50MG BASE</u> | <u>A090960</u> | <u>002</u> | Mar 27, 2012 |
| AB | | | <u>EQ 100MG BASE</u> | <u>A090960</u> | <u>003</u> | Mar 27, 2012 |
| AB | | | <u>EQ 200MG BASE</u> | <u>A090960</u> | <u>004</u> | Mar 27, 2012 |
| AB | | | <u>EQ 300MG BASE</u> | <u>A090960</u> | <u>005</u> | Mar 27, 2012 |
| AB | | | <u>EQ 400MG BASE</u> | <u>A090960</u> | <u>006</u> | Mar 27, 2012 |
| AB | | AUROBINDO PHARMA LTD | <u>EQ 25MG BASE</u> | <u>A091388</u> | <u>001</u> | Mar 27, 2012 |
| AB | | | <u>EQ 50MG BASE</u> | <u>A091388</u> | <u>002</u> | Mar 27, 2012 |
| AB | | | <u>EQ 100MG BASE</u> | <u>A091388</u> | <u>003</u> | Mar 27, 2012 |
| AB | | | <u>EQ 150MG BASE</u> | <u>A091388</u> | <u>004</u> | Mar 27, 2012 |
| AB | | | <u>EQ 200MG BASE</u> | <u>A091388</u> | <u>005</u> | Mar 27, 2012 |
| AB | | | <u>EQ 300MG BASE</u> | <u>A091388</u> | <u>006</u> | Mar 27, 2012 |
| AB | | | <u>EQ 400MG BASE</u> | <u>A091388</u> | <u>007</u> | Mar 27, 2012 |
| AB | | DR REDDYS LABS LTD | <u>EQ 25MG BASE</u> | <u>A077380</u> | <u>001</u> | Mar 27, 2012 |
| AB | | | <u>EQ 50MG BASE</u> | <u>A077380</u> | <u>002</u> | Mar 27, 2012 |
| AB | | | <u>EQ 100MG BASE</u> | <u>A077380</u> | <u>003</u> | Mar 27, 2012 |
| AB | | | <u>EQ 150MG BASE</u> | <u>A077380</u> | <u>004</u> | Mar 27, 2012 |
| AB | | | <u>EQ 200MG BASE</u> | <u>A077380</u> | <u>005</u> | Mar 27, 2012 |
| AB | | | <u>EQ 300MG BASE</u> | <u>A077380</u> | <u>006</u> | Mar 27, 2012 |
| AB | | | <u>EQ 400MG BASE</u> | <u>A077380</u> | <u>007</u> | Mar 27, 2012 |
| AB | | JUBILANT GENERICS | <u>EQ 25MG BASE</u> | <u>A203150</u> | <u>001</u> | Nov 26, 2013 |
| AB | | LUPIN LTD | <u>EQ 25MG BASE</u> | <u>A201109</u> | <u>001</u> | Mar 27, 2012 |
| AB | | | <u>EQ 50MG BASE</u> | <u>A201109</u> | <u>002</u> | Mar 27, 2012 |
| AB | | | <u>EQ 100MG BASE</u> | <u>A201109</u> | <u>003</u> | Mar 27, 2012 |
| AB | | | <u>EQ 200MG BASE</u> | <u>A201109</u> | <u>004</u> | Mar 27, 2012 |
| AB | | | <u>EQ 300MG BASE</u> | <u>A201109</u> | <u>005</u> | Mar 27, 2012 |
| AB | | | <u>EQ 400MG BASE</u> | <u>A201109</u> | <u>006</u> | Mar 27, 2012 |
| AB | | MACLEODS PHARMS LTD | <u>EQ 25MG BASE</u> | <u>A203359</u> | <u>001</u> | May 17, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

QUETIAPINE FUMARATE

TABLET; ORAL

QUETIAPINE FUMARATE

| | | | | | |
|--------------------------------|-------------------------|-------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A203359 002</u> | May 17, 2016 | |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A203359 003</u> | May 17, 2016 | |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A203359 004</u> | May 17, 2016 | |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A203359 005</u> | May 17, 2016 | |
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A203359 006</u> | May 17, 2016 | |
| <u>AB</u> | SANDOZ | <u>EQ 25MG BASE</u> | <u>A078679 001</u> | Dec 14, 2012 | |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A078679 002</u> | Dec 14, 2012 | |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A078679 003</u> | Dec 14, 2012 | |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A078679 004</u> | Dec 14, 2012 | |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A078679 005</u> | Dec 14, 2012 | |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A078679 006</u> | Dec 14, 2012 | |
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A078679 007</u> | Dec 14, 2012 | |
| <u>AB</u> | SUN PHARMA GLOBAL | <u>EQ 25MG BASE</u> | <u>A201190 001</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A201190 002</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A201190 003</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A201190 004</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A201190 005</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A201190 006</u> | Mar 27, 2012 | |
| <u>AB</u> | TEVA PHARMS | <u>EQ 25MG BASE</u> | <u>A077745 001</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A077745 002</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A077745 003</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A077745 004</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A077745 005</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A077745 006</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A077745 007</u> | Mar 27, 2012 | |
| <u>AB</u> | TORRENT PHARMS LTD | <u>EQ 25MG BASE</u> | <u>A200363 001</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A200363 002</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A200363 003</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A200363 004</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A200363 005</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A200363 006</u> | Mar 27, 2012 | |
| <u>AB</u> | UNICHEM LABS LTD | <u>EQ 25MG BASE</u> | <u>A202674 001</u> | Mar 08, 2016 | |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A202674 002</u> | Mar 08, 2016 | |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A202674 003</u> | Mar 08, 2016 | |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A202674 004</u> | Mar 08, 2016 | |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A202674 005</u> | Mar 08, 2016 | |
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A202674 006</u> | Mar 08, 2016 | |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>EQ 25MG BASE</u> | <u>A090120 001</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A090749 001</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A090749 002</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A090749 003</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A090749 004</u> | Mar 27, 2012 | |
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A090749 005</u> | Mar 27, 2012 | |
| <u>SEROQUEL</u> | | | | | |
| <u>AB</u> | +! | ASTRAZENECA PHARMS | <u>EQ 25MG BASE</u> | <u>N020639 001</u> | Sep 26, 1997 |
| <u>AB</u> | + | | <u>EQ 50MG BASE</u> | <u>N020639 007</u> | Oct 04, 2005 |
| <u>AB</u> | + | | <u>EQ 100MG BASE</u> | <u>N020639 002</u> | Sep 26, 1997 |
| <u>AB</u> | + | | <u>EQ 200MG BASE</u> | <u>N020639 003</u> | Sep 26, 1997 |
| <u>AB</u> | +! | | <u>EQ 300MG BASE</u> | <u>N020639 005</u> | Jul 26, 2000 |
| <u>AB</u> | + | | <u>EQ 400MG BASE</u> | <u>N020639 006</u> | Oct 04, 2005 |
| TABLET, EXTENDED RELEASE; ORAL | | | | | |
| <u>QUETIAPINE FUMARATE</u> | | | | | |
| <u>AB</u> | | ACCORD HLTHCARE | <u>EQ 50MG BASE</u> | <u>A206252 001</u> | Nov 29, 2017 |
| <u>AB</u> | | | <u>EQ 150MG BASE</u> | <u>A090681 001</u> | May 09, 2017 |
| <u>AB</u> | | | <u>EQ 200MG BASE</u> | <u>A090681 002</u> | May 09, 2017 |
| <u>AB</u> | | | <u>EQ 300MG BASE</u> | <u>A090681 003</u> | May 09, 2017 |
| <u>AB</u> | | | <u>EQ 400MG BASE</u> | <u>A090681 004</u> | Nov 01, 2016 |
| <u>AB</u> | | ALIGNSCIENCE PHARMA | <u>EQ 150MG BASE</u> | <u>A209497 001</u> | Sep 28, 2018 |
| <u>AB</u> | | | <u>EQ 200MG BASE</u> | <u>A209497 002</u> | Sep 28, 2018 |
| <u>AB</u> | | AMNEAL PHARMS | <u>EQ 400MG BASE</u> | <u>A211405 001</u> | Oct 26, 2018 |
| <u>AB</u> | | ANCHEN PHARMS | <u>EQ 150MG BASE</u> | <u>A090757 001</u> | Dec 01, 2017 |
| <u>AB</u> | | | <u>EQ 200MG BASE</u> | <u>A090757 002</u> | Dec 01, 2017 |
| <u>AB</u> | | | <u>EQ 300MG BASE</u> | <u>A090757 003</u> | Dec 01, 2017 |
| <u>AB</u> | | | <u>EQ 400MG BASE</u> | <u>A090757 004</u> | Dec 01, 2017 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>EQ 50MG BASE</u> | <u>A207655 001</u> | Nov 29, 2017 |
| <u>AB</u> | | | <u>EQ 150MG BASE</u> | <u>A207655 002</u> | Nov 29, 2017 |
| <u>AB</u> | | | <u>EQ 200MG BASE</u> | <u>A207655 003</u> | Nov 29, 2017 |
| <u>AB</u> | | | <u>EQ 300MG BASE</u> | <u>A207655 004</u> | Nov 29, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

QUETIAPINE FUMARATE

TABLET, EXTENDED RELEASE;ORAL

QUETIAPINE FUMARATE

| | | | | |
|-----------|--------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A207655 005</u> | Nov 29, 2017 |
| <u>AB</u> | INTELLIPHARMACEUTIC S | <u>EQ 50MG BASE</u> | <u>A202939 001</u> | May 09, 2017 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A202939 002</u> | May 09, 2017 |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A202939 003</u> | May 09, 2017 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A202939 004</u> | May 09, 2017 |
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A202939 005</u> | May 09, 2017 |
| <u>AB</u> | LUPIN LTD | <u>EQ 50MG BASE</u> | <u>A204203 001</u> | May 17, 2017 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A204203 002</u> | May 17, 2017 |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A204203 003</u> | May 17, 2017 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A204203 004</u> | May 17, 2017 |
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A204203 005</u> | May 17, 2017 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>EQ 150MG BASE</u> | <u>A204253 001</u> | Nov 29, 2017 |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A204253 002</u> | Nov 29, 2017 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A204253 003</u> | Nov 29, 2017 |
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A204253 004</u> | Nov 29, 2017 |
| <u>AB</u> | NOVAST LABS | <u>EQ 50MG BASE</u> | <u>A208947 001</u> | Nov 29, 2017 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A208947 002</u> | Nov 29, 2017 |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A208947 003</u> | Nov 29, 2017 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A208947 004</u> | Nov 29, 2017 |
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A208947 005</u> | Nov 29, 2017 |
| <u>AB</u> | PAR PHARM | <u>EQ 50MG BASE</u> | <u>A090482 001</u> | May 09, 2017 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A090482 002</u> | May 09, 2017 |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A090482 003</u> | May 09, 2017 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A090482 004</u> | May 09, 2017 |
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A090482 005</u> | May 09, 2017 |
| <u>AB</u> | PHARMADAX INC | <u>EQ 50MG BASE</u> | <u>A206260 001</u> | May 09, 2017 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A206260 002</u> | May 09, 2017 |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A206260 003</u> | May 09, 2017 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A206260 004</u> | May 09, 2017 |
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A206260 005</u> | May 09, 2017 |
| <u>AB</u> | SCIEGEN PHARMS INC | <u>EQ 50MG BASE</u> | <u>A209635 005</u> | Nov 16, 2018 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A209635 001</u> | Nov 29, 2017 |
| <u>AB</u> | | <u>EQ 200MG BASE</u> | <u>A209635 002</u> | Nov 29, 2017 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A209635 003</u> | Nov 29, 2017 |
| <u>AB</u> | | <u>EQ 400MG BASE</u> | <u>A209635 004</u> | Nov 29, 2017 |

SEROQUEL XR

| | | | | | |
|-----------|---|-------------|----------------------|--------------------|--------------|
| <u>AB</u> | + | ASTRAZENECA | <u>EQ 50MG BASE</u> | <u>N022047 001</u> | May 17, 2007 |
| <u>AB</u> | + | | <u>EQ 150MG BASE</u> | <u>N022047 005</u> | Aug 11, 2008 |
| <u>AB</u> | + | ! | <u>EQ 200MG BASE</u> | <u>N022047 002</u> | May 17, 2007 |
| <u>AB</u> | + | | <u>EQ 300MG BASE</u> | <u>N022047 003</u> | May 17, 2007 |
| <u>AB</u> | + | | <u>EQ 400MG BASE</u> | <u>N022047 004</u> | May 17, 2007 |

QUINAPRIL HYDROCHLORIDE

TABLET;ORAL

ACCUPRIL

| | | | | | |
|-----------|---|---------------|---------------------|--------------------|--------------|
| <u>AB</u> | + | PFIZER PHARMS | <u>EQ 5MG BASE</u> | <u>N019885 001</u> | Nov 19, 1991 |
| <u>AB</u> | + | | <u>EQ 10MG BASE</u> | <u>N019885 002</u> | Nov 19, 1991 |
| <u>AB</u> | + | | <u>EQ 20MG BASE</u> | <u>N019885 003</u> | Nov 19, 1991 |
| <u>AB</u> | + | ! | <u>EQ 40MG BASE</u> | <u>N019885 004</u> | Nov 19, 1991 |

QUINAPRIL HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|---------------------|--------------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 5MG BASE</u> | <u>A202725 001</u> | Apr 29, 2013 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A202725 002</u> | Apr 29, 2013 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A202725 003</u> | Apr 29, 2013 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A202725 004</u> | Apr 29, 2013 |
| <u>AB</u> | INVAGEN PHARMS | <u>EQ 5MG BASE</u> | <u>A078457 001</u> | Aug 24, 2007 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A078457 002</u> | Aug 24, 2007 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A078457 003</u> | Aug 24, 2007 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A078457 004</u> | Aug 24, 2007 |
| <u>AB</u> | LUPIN | <u>EQ 5MG BASE</u> | <u>A077690 001</u> | Jun 20, 2006 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A077690 002</u> | Jun 20, 2006 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A077690 003</u> | Jun 20, 2006 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077690 004</u> | Jun 20, 2006 |
| <u>AB</u> | MYLAN | <u>EQ 5MG BASE</u> | <u>A076694 001</u> | Dec 23, 2004 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A076694 002</u> | Dec 23, 2004 |
| <u>AB</u> | | <u>EQ 20MG BASE</u> | <u>A076694 003</u> | Dec 23, 2004 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A076694 004</u> | Dec 23, 2004 |
| <u>AB</u> | PRINSTON INC | <u>EQ 5MG BASE</u> | <u>A205823 001</u> | Sep 15, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A205823 002</u> | Sep 15, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

QUINAPRIL HYDROCHLORIDE

| | | | | |
|-----------|------|---------------------|--------------------|--------------|
| AB | | EQ 20MG BASE | A205823 003 | Sep 15, 2016 |
| AB | | EQ 40MG BASE | A205823 004 | Sep 15, 2016 |
| AB | TEVA | EQ 5MG BASE | A075504 001 | Aug 24, 2007 |
| AB | | EQ 10MG BASE | A075504 002 | Aug 24, 2007 |
| AB | | EQ 20MG BASE | A075504 003 | Aug 24, 2007 |
| AB | | EQ 40MG BASE | A075504 004 | Aug 24, 2007 |

QUINIDINE GLUCONATE

TABLET, EXTENDED RELEASE; ORAL

QUINIDINE GLUCONATE

| | | | | | |
|----|---|----------------------|-------|-------------|--------------|
| BX | ! | SUN PHARM INDUSTRIES | 324MG | A089338 001 | Feb 11, 1987 |
|----|---|----------------------|-------|-------------|--------------|

QUINIDINE SULFATE

TABLET; ORAL

QUINIDINE SULFATE

| | | | | | |
|-----------|---|----------------------|--------------|--------------------|--------------|
| AB | | SANDOZ | 200MG | A088072 002 | |
| AB | | | 300MG | A088072 001 | Sep 26, 1983 |
| AB | | SUN PHARM INDUSTRIES | 200MG | A081030 001 | Apr 14, 1989 |
| AB | | | 300MG | A081031 001 | Apr 14, 1989 |
| AB | ! | WATSON LABS | 200MG | A083288 001 | |
| AB | ! | | 300MG | A085583 001 | |

QUININE SULFATE

CAPSULE; ORAL

QUALAQUIN

| | | | | | | |
|-----------|---|---|----------------------|--------------|--------------------|--------------|
| AB | + | ! | SUN PHARM INDUSTRIES | 324MG | N021799 001 | Aug 12, 2005 |
|-----------|---|---|----------------------|--------------|--------------------|--------------|

QUININE SULFATE

| | | | | | |
|-----------|--|------------------|--------------|--------------------|--------------|
| AB | | AMNEAL PHARMS | 324MG | A203729 001 | Jul 15, 2015 |
| AB | | LUPIN LTD | 324MG | A203112 001 | Apr 24, 2015 |
| AB | | MYLAN PHARMS INC | 324MG | A202581 001 | Dec 14, 2012 |
| AB | | NOVAST LABS | 324MG | A204372 001 | Jul 22, 2015 |
| AB | | TEVA PHARMS | 324MG | A091661 001 | Sep 28, 2012 |

RABEPRAZOLE SODIUM

CAPSULE, DELAYED RELEASE; ORAL

ACIPHEX SPRINKLE

+ CERECOR INC

5MG

N204736 001

Mar 26, 2013

+!

+ CERECOR INC

10MG

N204736 002

Mar 26, 2013

TABLET, DELAYED RELEASE; ORAL

ACIPHEX

| | | | | | | |
|-----------|---|---|-----------|-------------|--------------------|--------------|
| AB | + | ! | EISAI INC | 20MG | N020973 002 | Aug 19, 1999 |
|-----------|---|---|-----------|-------------|--------------------|--------------|

RABEPRAZOLE SODIUM

| | | | | | |
|-----------|--|----------------------|-------------|--------------------|--------------|
| AB | | ALKEM LABS LTD | 20MG | A208644 001 | Apr 24, 2018 |
| AB | | AMNEAL PHARMS | 20MG | A204179 001 | Jul 31, 2015 |
| AB | | AUROBINDO PHARMA LTD | 20MG | A205761 001 | Feb 17, 2017 |
| AB | | BRECKENRIDGE PHARM | 20MG | A204237 001 | Nov 18, 2015 |
| AB | | DR REDDYS LABS LTD | 20MG | A076824 001 | Nov 08, 2013 |
| AB | | LANNETT CO INC | 20MG | A090678 001 | Nov 08, 2013 |
| AB | | LUPIN LTD | 20MG | A078964 001 | Nov 08, 2013 |
| AB | | MYLAN PHARMS INC | 20MG | A076885 001 | Nov 08, 2013 |
| AB | | TEVA PHARMS USA | 20MG | A076822 001 | Nov 08, 2013 |
| AB | | TORRENT PHARMS LTD | 20MG | A202376 001 | Nov 08, 2013 |

RADIUM RA-223 DICHLORIDE

SOLUTION; INTRAVENOUS

XOFIGO

+! BAYER HLTHCARE

162mCi/6ML (27mCi/ML)

N203971 001

May 15, 2013

RALOXIFENE HYDROCHLORIDE

TABLET; ORAL

EVISTA

| | | | | | | |
|-----------|---|---|-------|-------------|--------------------|--------------|
| AB | + | ! | LILLY | 60MG | N020815 001 | Dec 09, 1997 |
|-----------|---|---|-------|-------------|--------------------|--------------|

RALOXIFENE HYDROCHLORIDE

| | | | | | |
|-----------|--|----------------------|-------------|--------------------|--------------|
| AB | | AMNEAL PHARMS | 60MG | A208206 001 | Apr 08, 2016 |
| AB | | AUROBINDO PHARMA LTD | 60MG | A204310 001 | Aug 28, 2015 |
| AB | | GLENMARK PHARMS LTD | 60MG | A204491 001 | Mar 22, 2016 |
| AB | | INVAGEN PHARMS | 60MG | A090842 001 | Sep 24, 2014 |
| AB | | SCIEGEN PHARMS INC | 60MG | A206384 001 | Oct 12, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

RALOXIFENE HYDROCHLORIDE

TABLET; ORAL

RALOXIFENE HYDROCHLORIDE

| | | | | | |
|-----------|-----------------|-------------|----------------|------------|--------------|
| <u>AB</u> | TEVA PHARMS USA | <u>60MG</u> | <u>A078193</u> | <u>001</u> | Mar 04, 2014 |
| <u>AB</u> | WATSON LABS INC | <u>60MG</u> | <u>A200825</u> | <u>001</u> | Jan 21, 2015 |

RALTEGRAVIR POTASSIUM

POWDER; ORAL

ISENTRESS

| | | | | | | |
|---|---|-------------------|----------------------|---------|-----|--------------|
| + | ! | MERCK SHARP DOHME | EQ 100MG BASE/PACKET | N205786 | 001 | Dec 20, 2013 |
|---|---|-------------------|----------------------|---------|-----|--------------|

TABLET; ORAL

ISENTRESS

| | | | | | | |
|---|---|-------------------|---------------|---------|-----|--------------|
| + | ! | MERCK SHARP DOHME | EQ 400MG BASE | N022145 | 001 | Oct 12, 2007 |
|---|---|-------------------|---------------|---------|-----|--------------|

ISENTRESS HD

| | | | | | | |
|---|---|-------------------|---------------|---------|-----|--------------|
| + | ! | MERCK SHARP DOHME | EQ 600MG BASE | N022145 | 002 | May 26, 2017 |
|---|---|-------------------|---------------|---------|-----|--------------|

TABLET, CHEWABLE; ORAL

ISENTRESS

| | | | | | | |
|---|--|-------------------|--------------|---------|-----|--------------|
| + | | MERCK SHARP DOHME | EQ 25MG BASE | N203045 | 001 | Dec 21, 2011 |
|---|--|-------------------|--------------|---------|-----|--------------|

| | | | | | | |
|---|---|--|---------------|---------|-----|--------------|
| + | ! | | EQ 100MG BASE | N203045 | 002 | Dec 21, 2011 |
|---|---|--|---------------|---------|-----|--------------|

RAMELTEON

TABLET; ORAL

RAMELTEON

| | | | | | |
|-----------|---------------------|------------|----------------|------------|--------------|
| <u>AB</u> | ACTAVIS LABS FL INC | <u>8MG</u> | <u>A091610</u> | <u>001</u> | Aug 19, 2015 |
| <u>AB</u> | DR REDDYS LABS SA | <u>8MG</u> | <u>A091693</u> | <u>001</u> | Jul 26, 2013 |

ROZEREM

| | | | | | | | |
|-----------|---|---|-------------------|------------|----------------|------------|--------------|
| <u>AB</u> | + | ! | TAKEDA PHARMS USA | <u>8MG</u> | <u>N021782</u> | <u>001</u> | Jul 22, 2005 |
|-----------|---|---|-------------------|------------|----------------|------------|--------------|

RAMIPRIL

CAPSULE; ORAL

ALTACE

| | | | | | | | |
|-----------|---|---|-----------------|---------------|----------------|------------|--------------|
| <u>AB</u> | + | | KING PHARMS LLC | <u>1.25MG</u> | <u>N019901</u> | <u>001</u> | Jan 28, 1991 |
| <u>AB</u> | + | | | <u>2.5MG</u> | <u>N019901</u> | <u>002</u> | Jan 28, 1991 |
| <u>AB</u> | + | | | <u>5MG</u> | <u>N019901</u> | <u>003</u> | Jan 28, 1991 |
| <u>AB</u> | + | ! | | <u>10MG</u> | <u>N019901</u> | <u>004</u> | Jan 28, 1991 |

RAMIPRIL

| | | | | | | | |
|-----------|--|--|----------------------|---------------|----------------|------------|--------------|
| <u>AB</u> | | | ACCORD HLTHCARE | <u>1.25MG</u> | <u>A202392</u> | <u>001</u> | Apr 15, 2014 |
| <u>AB</u> | | | | <u>2.5MG</u> | <u>A202392</u> | <u>002</u> | Apr 15, 2014 |
| <u>AB</u> | | | | <u>5MG</u> | <u>A202392</u> | <u>003</u> | Apr 15, 2014 |
| <u>AB</u> | | | | <u>10MG</u> | <u>A202392</u> | <u>004</u> | Apr 15, 2014 |
| <u>AB</u> | | | APOTEX | <u>1.25MG</u> | <u>A079116</u> | <u>001</u> | Jun 20, 2008 |
| <u>AB</u> | | | | <u>2.5MG</u> | <u>A079116</u> | <u>002</u> | Jun 20, 2008 |
| <u>AB</u> | | | | <u>5MG</u> | <u>A079116</u> | <u>003</u> | Jun 20, 2008 |
| <u>AB</u> | | | | <u>10MG</u> | <u>A079116</u> | <u>004</u> | Jun 20, 2008 |
| <u>AB</u> | | | AUROBINDO PHARMA LTD | <u>1.25MG</u> | <u>A091604</u> | <u>001</u> | Jun 08, 2011 |
| <u>AB</u> | | | | <u>2.5MG</u> | <u>A091604</u> | <u>002</u> | Jun 08, 2011 |
| <u>AB</u> | | | | <u>5MG</u> | <u>A091604</u> | <u>003</u> | Jun 08, 2011 |
| <u>AB</u> | | | | <u>10MG</u> | <u>A091604</u> | <u>004</u> | Jun 08, 2011 |
| <u>AB</u> | | | CHARTWELL MOLECULAR | <u>1.25MG</u> | <u>A078745</u> | <u>001</u> | Jun 18, 2008 |
| <u>AB</u> | | | | <u>2.5MG</u> | <u>A078745</u> | <u>002</u> | Jun 18, 2008 |
| <u>AB</u> | | | | <u>5MG</u> | <u>A078745</u> | <u>003</u> | Jun 18, 2008 |
| <u>AB</u> | | | | <u>10MG</u> | <u>A078745</u> | <u>004</u> | Jun 18, 2008 |
| <u>AB</u> | | | DR REDDYS LABS LTD | <u>1.25MG</u> | <u>A078191</u> | <u>001</u> | Jun 18, 2008 |
| <u>AB</u> | | | | <u>2.5MG</u> | <u>A078191</u> | <u>002</u> | Jun 18, 2008 |
| <u>AB</u> | | | | <u>5MG</u> | <u>A078191</u> | <u>003</u> | Jun 18, 2008 |
| <u>AB</u> | | | | <u>10MG</u> | <u>A078191</u> | <u>004</u> | Jun 18, 2008 |
| <u>AB</u> | | | LUPIN | <u>1.25MG</u> | <u>A077626</u> | <u>001</u> | Jun 09, 2008 |
| <u>AB</u> | | | | <u>2.5MG</u> | <u>A077626</u> | <u>002</u> | Jun 09, 2008 |
| <u>AB</u> | | | | <u>5MG</u> | <u>A077626</u> | <u>003</u> | Jun 09, 2008 |
| <u>AB</u> | | | | <u>10MG</u> | <u>A077626</u> | <u>004</u> | Jun 09, 2008 |
| <u>AB</u> | | | TEVA PHARMS | <u>1.25MG</u> | <u>A077470</u> | <u>001</u> | Jun 18, 2008 |
| <u>AB</u> | | | | <u>2.5MG</u> | <u>A077470</u> | <u>002</u> | Jun 18, 2008 |
| <u>AB</u> | | | | <u>5MG</u> | <u>A077470</u> | <u>003</u> | Jun 18, 2008 |
| <u>AB</u> | | | | <u>10MG</u> | <u>A077470</u> | <u>004</u> | Jun 18, 2008 |
| <u>AB</u> | | | WATSON LABS | <u>1.25MG</u> | <u>A076549</u> | <u>001</u> | Oct 24, 2005 |
| <u>AB</u> | | | | <u>2.5MG</u> | <u>A076549</u> | <u>002</u> | Oct 24, 2005 |
| <u>AB</u> | | | | <u>10MG</u> | <u>A076549</u> | <u>004</u> | Oct 24, 2005 |
| <u>AB</u> | | | WEST-WARD PHARMS INT | <u>1.25MG</u> | <u>A077900</u> | <u>001</u> | Jun 18, 2008 |
| <u>AB</u> | | | | <u>2.5MG</u> | <u>A077900</u> | <u>002</u> | Jun 18, 2008 |
| <u>AB</u> | | | | <u>5MG</u> | <u>A077900</u> | <u>003</u> | Jun 18, 2008 |
| <u>AB</u> | | | | <u>10MG</u> | <u>A077900</u> | <u>004</u> | Jun 18, 2008 |
| <u>AB</u> | | | ZYDUS PHARMS USA | <u>1.25MG</u> | <u>A078832</u> | <u>001</u> | Sep 02, 2008 |

PRESCRIPTION DRUG PRODUCT LIST

RAMIPRIL

CAPSULE; ORAL

RAMIPRIL

| | | | | | |
|-----------|--|--------------|----------------|------------|--------------|
| <u>AB</u> | | <u>2.5MG</u> | <u>A078832</u> | <u>002</u> | Sep 02, 2008 |
| <u>AB</u> | | <u>5MG</u> | <u>A078832</u> | <u>003</u> | Sep 02, 2008 |
| <u>AB</u> | | <u>10MG</u> | <u>A078832</u> | <u>004</u> | Sep 02, 2008 |

RANITIDINE

CAPSULE; ORAL

RANITIDINE

| | | | | | |
|-----------|-----------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | NOVITIUM PHARMA | <u>EQ 150MG BASE</u> | <u>A210681</u> | <u>001</u> | Nov 23, 2018 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A210681</u> | <u>002</u> | Nov 23, 2018 |

RANITIDINE HYDROCHLORIDE

CAPSULE; ORAL

RANITIDINE HYDROCHLORIDE

| | | | | | |
|-----------|----------------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | AJANTA PHARMA LTD | <u>EQ 150MG BASE</u> | <u>A209859</u> | <u>001</u> | Sep 27, 2018 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A209859</u> | <u>002</u> | Sep 27, 2018 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 150MG BASE</u> | <u>A211058</u> | <u>001</u> | Jul 16, 2018 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A211058</u> | <u>002</u> | Jul 16, 2018 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 150MG BASE</u> | <u>A075742</u> | <u>001</u> | Nov 29, 2000 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A075742</u> | <u>002</u> | Nov 29, 2000 |
| <u>AB</u> | SANDOZ | <u>EQ 150MG BASE</u> | <u>A074655</u> | <u>001</u> | Oct 22, 1997 |
| <u>AB</u> | ! | <u>EQ 300MG BASE</u> | <u>A074655</u> | <u>002</u> | Oct 22, 1997 |

INJECTABLE; INJECTION

RANITIDINE HYDROCHLORIDE

| | | | | | |
|-----------|----------------------|------------------------|----------------|------------|--------------|
| <u>AP</u> | MYLAN LABS LTD | <u>EQ 25MG BASE/ML</u> | <u>A079076</u> | <u>001</u> | Jun 09, 2016 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>EQ 25MG BASE/ML</u> | <u>A074777</u> | <u>001</u> | Mar 02, 2005 |
| <u>AP</u> | | <u>EQ 25MG BASE/ML</u> | <u>A077458</u> | <u>001</u> | Feb 16, 2006 |
| <u>AP</u> | ZYDUS PHARMS USA INC | <u>EQ 25MG BASE/ML</u> | <u>A091534</u> | <u>001</u> | Feb 22, 2013 |

ZANTAC

| | | | | | |
|-----------|-------------|------------------------|----------------|------------|--------------|
| <u>AP</u> | +! TELIGENT | <u>EQ 25MG BASE/ML</u> | <u>N019090</u> | <u>001</u> | Oct 19, 1984 |
|-----------|-------------|------------------------|----------------|------------|--------------|

SYRUP; ORAL

RANITIDINE HYDROCHLORIDE

| | | | | | |
|-----------|----------------------|------------------------|----------------|------------|--------------|
| <u>AA</u> | ACTAVIS MID ATLANTIC | <u>EQ 15MG BASE/ML</u> | <u>A076124</u> | <u>001</u> | Feb 21, 2007 |
| <u>AA</u> | AMNEAL PHARMS | <u>EQ 15MG BASE/ML</u> | <u>A078312</u> | <u>001</u> | Sep 02, 2008 |
| <u>AA</u> | ANDA REPOSITORY | <u>EQ 15MG BASE/ML</u> | <u>A090054</u> | <u>001</u> | Nov 15, 2010 |
| <u>AA</u> | AUROBINDO PHARMA LTD | <u>EQ 15MG BASE/ML</u> | <u>A090623</u> | <u>001</u> | Jul 28, 2010 |
| <u>AA</u> | BIO PHARM INC | <u>EQ 15MG BASE/ML</u> | <u>A090102</u> | <u>001</u> | May 26, 2009 |
| <u>AA</u> | BRECKENRIDGE PHARM | <u>EQ 15MG BASE/ML</u> | <u>A078684</u> | <u>001</u> | Aug 27, 2009 |
| <u>AA</u> | HI TECH PHARMA | <u>EQ 15MG BASE/ML</u> | <u>A091078</u> | <u>001</u> | Mar 22, 2011 |
| <u>AA</u> | LANNETT CO INC | <u>EQ 15MG BASE/ML</u> | <u>A078890</u> | <u>001</u> | Jul 01, 2010 |
| <u>AA</u> | | <u>EQ 15MG BASE/ML</u> | <u>A091288</u> | <u>001</u> | Dec 09, 2010 |
| <u>AA</u> | NOSTRUM LABS INC | <u>EQ 15MG BASE/ML</u> | <u>A091091</u> | <u>001</u> | Sep 20, 2011 |
| <u>AA</u> | ! | <u>EQ 15MG BASE/ML</u> | <u>A077405</u> | <u>001</u> | Sep 21, 2007 |
| <u>AA</u> | TARO | <u>EQ 15MG BASE/ML</u> | <u>A077476</u> | <u>001</u> | Jun 13, 2011 |

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

| | | | | | |
|-----------|----------------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | ACIC PHARMS | <u>EQ 150MG BASE</u> | <u>A203694</u> | <u>001</u> | Nov 30, 2017 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A203694</u> | <u>002</u> | Nov 30, 2017 |
| <u>AB</u> | AMNEAL PHARMS NY | <u>EQ 150MG BASE</u> | <u>A077824</u> | <u>001</u> | Oct 13, 2006 |
| <u>AB</u> | ! | <u>EQ 300MG BASE</u> | <u>A077824</u> | <u>002</u> | Oct 13, 2006 |
| <u>AB</u> | APOTEX | <u>EQ 150MG BASE</u> | <u>A074680</u> | <u>001</u> | Sep 12, 1997 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A074680</u> | <u>002</u> | Sep 12, 1997 |
| <u>AB</u> | DR REDDYS LABS INC | <u>EQ 150MG BASE</u> | <u>A076705</u> | <u>001</u> | Jul 27, 2005 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A076705</u> | <u>002</u> | Jul 27, 2005 |
| <u>AB</u> | GLENMARK PHARMS INC | <u>EQ 150MG BASE</u> | <u>A078542</u> | <u>001</u> | Nov 19, 2008 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A078542</u> | <u>002</u> | Nov 19, 2008 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>EQ 150MG BASE</u> | <u>A075165</u> | <u>001</u> | Sep 30, 1998 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A075165</u> | <u>002</u> | Sep 30, 1998 |
| <u>AB</u> | PAR PHARM | <u>EQ 150MG BASE</u> | <u>A075180</u> | <u>001</u> | Jan 28, 1999 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A075180</u> | <u>002</u> | Jan 28, 1999 |
| <u>AB</u> | SANDOZ | <u>EQ 150MG BASE</u> | <u>A074467</u> | <u>001</u> | Aug 29, 1997 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A074467</u> | <u>002</u> | Aug 29, 1997 |
| <u>AB</u> | STRIDES PHARMA | <u>EQ 150MG BASE</u> | <u>A205512</u> | <u>001</u> | Aug 22, 2016 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A205512</u> | <u>002</u> | Aug 22, 2016 |
| <u>AB</u> | TEVA | <u>EQ 150MG BASE</u> | <u>A074488</u> | <u>001</u> | Jul 31, 1997 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A074488</u> | <u>002</u> | Jul 31, 1997 |
| <u>AB</u> | VIVIMED GLOBAL | <u>EQ 150MG BASE</u> | <u>A210010</u> | <u>001</u> | Aug 01, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

RANITIDINE HYDROCHLORIDE

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

| | | | | |
|-----------|---------------|----------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A210010 002</u> | Aug 01, 2018 |
| <u>AB</u> | WOCKHARDT LTD | <u>EQ 150MG BASE</u> | <u>A075208 001</u> | Dec 17, 1998 |
| <u>AB</u> | | <u>EQ 300MG BASE</u> | <u>A075208 002</u> | Dec 17, 1998 |

RANOLAZINE

TABLET, EXTENDED RELEASE; ORAL

RANEXA

| | | | | |
|---|--------|-------|-------------|--------------|
| + | GILEAD | 500MG | N021526 002 | Jan 27, 2006 |
| + | ! | 1GM | N021526 001 | Feb 12, 2007 |

RASAGILINE MESYLATE

TABLET; ORAL

AZILECT

| | | | | | |
|-----------|---|------|----------------------|--------------------|--------------|
| <u>AB</u> | + | TEVA | <u>EQ 0.5MG BASE</u> | <u>N021641 001</u> | May 16, 2006 |
| <u>AB</u> | + | ! | <u>EQ 1MG BASE</u> | <u>N021641 002</u> | May 16, 2006 |

RASAGILINE MESYLATE

| | | | | | |
|-----------|--|------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | ALKEM LABS LTD | <u>EQ 0.5MG BASE</u> | <u>A201889 001</u> | Oct 30, 2017 |
| <u>AB</u> | | | <u>EQ 1MG BASE</u> | <u>A201889 002</u> | Oct 30, 2017 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>EQ 0.5MG BASE</u> | <u>A201971 001</u> | May 15, 2017 |
| <u>AB</u> | | | <u>EQ 1MG BASE</u> | <u>A201971 002</u> | May 15, 2017 |
| <u>AB</u> | | ORCHID HLTHCARE | <u>EQ 0.5MG BASE</u> | <u>A201970 001</u> | Mar 15, 2016 |
| <u>AB</u> | | | <u>EQ 1MG BASE</u> | <u>A201970 002</u> | Mar 15, 2016 |
| <u>AB</u> | | SANDOZ INC | <u>EQ 0.5MG BASE</u> | <u>A201892 001</u> | Jul 27, 2018 |
| <u>AB</u> | | | <u>EQ 1MG BASE</u> | <u>A201892 002</u> | Jul 27, 2018 |
| <u>AB</u> | | WATSON LABS INC | <u>EQ 0.5MG BASE</u> | <u>A201823 001</u> | Jul 01, 2013 |
| <u>AB</u> | | | <u>EQ 1MG BASE</u> | <u>A201823 002</u> | Jul 01, 2013 |

REGADENOSON

SOLUTION; INTRAVENOUS

LEXISCAN

| | | | | | |
|---|---|----------|-----------------------|-------------|--------------|
| + | ! | ASTELLAS | 0.4MG/5ML (0.08MG/ML) | N022161 001 | Apr 10, 2008 |
|---|---|----------|-----------------------|-------------|--------------|

REGORAFENIB

TABLET; ORAL

STIVARGA

| | | | | | |
|---|---|----------------|------|-------------|--------------|
| + | ! | BAYER HLTHCARE | 40MG | N203085 001 | Sep 27, 2012 |
|---|---|----------------|------|-------------|--------------|

REMIFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

REMIFENTANIL HYDROCHLORIDE

| | | | | | |
|-----------|--|--------------------|-------------------------|--------------------|--------------|
| <u>AP</u> | | FRESENIUS KABI USA | <u>EQ 1MG BASE/VIAL</u> | <u>A206223 001</u> | Jan 16, 2018 |
| <u>AP</u> | | | <u>EQ 2MG BASE/VIAL</u> | <u>A206223 002</u> | Jan 16, 2018 |
| <u>AP</u> | | | <u>EQ 5MG BASE/VIAL</u> | <u>A206223 003</u> | Jan 16, 2018 |

ULTIVA

| | | | | | |
|-----------|---|---------------------|-------------------------|--------------------|--------------|
| <u>AP</u> | + | MYLAN INSTITUTIONAL | <u>EQ 1MG BASE/VIAL</u> | <u>N020630 001</u> | Jul 12, 1996 |
| <u>AP</u> | + | | <u>EQ 2MG BASE/VIAL</u> | <u>N020630 002</u> | Jul 12, 1996 |
| <u>AP</u> | + | ! | <u>EQ 5MG BASE/VIAL</u> | <u>N020630 003</u> | Jul 12, 1996 |

REPAGLINIDE

TABLET; ORAL

PRANDIN

| | | | | | |
|-----------|---|-----------------|--------------|--------------------|--------------|
| <u>AB</u> | + | GEMINI LABS LLC | <u>0.5MG</u> | <u>N020741 001</u> | Dec 22, 1997 |
| <u>AB</u> | + | | <u>1MG</u> | <u>N020741 002</u> | Dec 22, 1997 |
| <u>AB</u> | + | ! | <u>2MG</u> | <u>N020741 003</u> | Dec 22, 1997 |

REPAGLINIDE

| | | | | | |
|-----------|--|----------------------|--------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS TOTOWA | <u>0.5MG</u> | <u>A090008 001</u> | Jan 22, 2014 |
| <u>AB</u> | | | <u>1MG</u> | <u>A090008 002</u> | Jan 22, 2014 |
| <u>AB</u> | | | <u>2MG</u> | <u>A090008 003</u> | Jan 22, 2014 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>0.5MG</u> | <u>A203820 001</u> | Jan 22, 2014 |
| <u>AB</u> | | | <u>1MG</u> | <u>A203820 002</u> | Jan 22, 2014 |
| <u>AB</u> | | | <u>2MG</u> | <u>A203820 003</u> | Jan 22, 2014 |
| <u>AB</u> | | BOSCOGEN | <u>0.5MG</u> | <u>A091517 001</u> | Apr 24, 2015 |
| <u>AB</u> | | | <u>1MG</u> | <u>A091517 002</u> | Apr 24, 2015 |
| <u>AB</u> | | | <u>2MG</u> | <u>A091517 003</u> | Apr 24, 2015 |
| <u>AB</u> | | CASI PHARMS INC | <u>0.5MG</u> | <u>A078555 001</u> | Nov 22, 2013 |
| <u>AB</u> | | | <u>1MG</u> | <u>A078555 002</u> | Jan 22, 2014 |
| <u>AB</u> | | | <u>2MG</u> | <u>A078555 003</u> | Jan 22, 2014 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>0.5MG</u> | <u>A090252 001</u> | Aug 23, 2013 |
| <u>AB</u> | | | <u>1MG</u> | <u>A090252 002</u> | Jan 22, 2014 |
| <u>AB</u> | | | <u>2MG</u> | <u>A090252 003</u> | Jan 22, 2014 |
| <u>AB</u> | | PADDOCK LLC | <u>0.5MG</u> | <u>A201189 001</u> | Jul 17, 2013 |
| <u>AB</u> | | | <u>1MG</u> | <u>A201189 002</u> | Jan 22, 2014 |

PRESCRIPTION DRUG PRODUCT LIST

REPAGLINIDE

TABLET; ORAL

REPAGLINIDE

| | | | | |
|-----------|--------------------|------------|--------------------|--------------|
| AB | | 2MG | A201189 003 | Jan 22, 2014 |
| AB | SUN PHARM INDS INC | 1MG | A077571 002 | Jul 11, 2013 |
| AB | | 2MG | A077571 003 | Jul 11, 2013 |

RETAPAMULIN

OINTMENT; TOPICAL

ALTABAX

| | | | | |
|---|-----------------|----|-------------|--------------|
| + | AQUA PHARMS LLC | 1% | N022055 001 | Apr 12, 2007 |
|---|-----------------|----|-------------|--------------|

REVEFENACIN

SOLUTION; INHALATION

YUPELRI

| | | | | |
|---|-------------------|------------|-------------|--------------|
| + | MYLAN IRELAND LTD | 175MCG/3ML | N210598 001 | Nov 09, 2018 |
|---|-------------------|------------|-------------|--------------|

RIBAVIRIN

CAPSULE; ORAL

REBETOL

| | | | | | |
|-----------|---|-------------------|--------------|--------------------|--------------|
| AB | + | MERCK SHARP DOHME | 200MG | N020903 002 | Jul 25, 2001 |
|-----------|---|-------------------|--------------|--------------------|--------------|

RIBASPHERE

| | | | | | |
|-----------|--|-------------------|--------------|--------------------|--------------|
| AB | | KADMON PHARMS LLC | 200MG | A076203 001 | Apr 06, 2004 |
|-----------|--|-------------------|--------------|--------------------|--------------|

RIBAVIRIN

| | | | | | |
|-----------|--|------------------|--------------|--------------------|--------------|
| AB | | AUROBINDO PHARMA | 200MG | A079117 001 | Sep 17, 2009 |
|-----------|--|------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|-----------------|--------------|--------------------|--------------|
| AB | | CASI PHARMS INC | 200MG | A076192 001 | Apr 06, 2004 |
|-----------|--|-----------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|------|--------------|--------------------|--------------|
| AB | | TEVA | 200MG | A076277 001 | Oct 04, 2004 |
|-----------|--|------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------------|--------------|--------------------|--------------|
| AB | | ZYDUS PHARMS USA | 200MG | A077224 001 | Oct 28, 2005 |
|-----------|--|------------------|--------------|--------------------|--------------|

FOR SOLUTION; INHALATION

RIBAVIRIN

| | | | | | |
|-----------|--|-------------|-----------------|--------------------|--------------|
| AN | | NAVINTA LLC | 6GM/VIAL | A207366 001 | Oct 06, 2016 |
|-----------|--|-------------|-----------------|--------------------|--------------|

VIRAZOLE

| | | | | | |
|-----------|---|--------------------|-----------------|--------------------|--------------|
| AN | + | VALEANT PHARM INTL | 6GM/VIAL | N018859 001 | Dec 31, 1985 |
|-----------|---|--------------------|-----------------|--------------------|--------------|

SOLUTION; ORAL

REBETOL

| | | | | |
|---|----------|---------|-------------|--------------|
| + | SCHERING | 40MG/ML | N021546 001 | Jul 29, 2003 |
|---|----------|---------|-------------|--------------|

TABLET; ORAL

RIBAVIRIN

| | | | | | |
|-----------|--|------------------|--------------|--------------------|--------------|
| AB | | AUROBINDO PHARMA | 200MG | A079111 001 | Sep 17, 2009 |
|-----------|--|------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|-------------------|--------------|--------------------|--------------|
| AB | | KADMON PHARMS LLC | 200MG | A077456 001 | Dec 05, 2005 |
|-----------|--|-------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 400MG | A077456 002 | Dec 05, 2005 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|---|--|--------------|--------------------|--------------|
| AB | ! | | 600MG | A077456 003 | Dec 05, 2005 |
|-----------|---|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--------|--------------|--------------------|--------------|
| AB | | SANDOZ | 200MG | A077743 001 | Oct 03, 2006 |
|-----------|--|--------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------|--------------|--------------------|--------------|
| AB | | SANDOZ INC | 200MG | A202546 001 | Aug 12, 2014 |
|-----------|--|------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 400MG | A202546 002 | Aug 12, 2014 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 500MG | A202546 003 | Aug 12, 2014 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 600MG | A202546 004 | Aug 12, 2014 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|------|--------------|--------------------|--------------|
| AB | | TEVA | 200MG | A077053 001 | Dec 05, 2005 |
|-----------|--|------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|------------------|--------------|--------------------|--------------|
| AB | | ZYDUS PHARMS USA | 200MG | A077094 001 | Dec 05, 2005 |
|-----------|--|------------------|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 400MG | A077094 002 | Mar 16, 2007 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 500MG | A077094 004 | Apr 18, 2008 |
|-----------|--|--|--------------|--------------------|--------------|

| | | | | | |
|-----------|--|--|--------------|--------------------|--------------|
| AB | | | 600MG | A077094 003 | Mar 16, 2007 |
|-----------|--|--|--------------|--------------------|--------------|

RIBOCICLIB SUCCINATE

TABLET; ORAL

KISQALI

| | | | | |
|---|----------------------|---------------|-------------|--------------|
| + | NOVARTIS PHARMS CORP | EQ 200MG BASE | N209092 001 | Mar 13, 2017 |
|---|----------------------|---------------|-------------|--------------|

RIBOFLAVIN 5'-PHOSPHATE SODIUM

SOLUTION/DROPS; OPHTHALMIC

PHOTREXA

| | | | | |
|---|------------|--------|-------------|--------------|
| + | AVEDRO INC | 0.146% | N203324 001 | Apr 15, 2016 |
|---|------------|--------|-------------|--------------|

| | | | | |
|---|-----------------------------|-----|-------------|--------------|
| + | PHOTREXA VISCOUS IN DEXTRAN | 20% | N203324 002 | Apr 15, 2016 |
|---|-----------------------------|-----|-------------|--------------|

| | | | | |
|---|------------|--------|-------------|--------------|
| + | AVEDRO INC | 0.146% | N203324 002 | Apr 15, 2016 |
|---|------------|--------|-------------|--------------|

RIFABUTIN

CAPSULE; ORAL

MYCOBUTIN

| | | | | | |
|-----------|---|----------------------|--------------|--------------------|--------------|
| AB | + | PHARMACIA AND UPJOHN | 150MG | N050689 001 | Dec 23, 1992 |
|-----------|---|----------------------|--------------|--------------------|--------------|

RIFABUTIN

| | | | | | |
|-----------|--|-----------|--------------|--------------------|--------------|
| AB | | LUPIN LTD | 150MG | A090033 001 | Feb 24, 2014 |
|-----------|--|-----------|--------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

RIFAMPIN

CAPSULE; ORAL

RIFADIN

| | | | | | |
|------------------------|----------|-------------------|--------------|--------------------|--------------|
| AB | | SANOFI AVENTIS US | 150MG | A062303 001 | |
| AB | + | | 300MG | N050420 001 | |
| <u>RIFAMPIN</u> | | | | | |
| AB | | AKORN | 150MG | A065028 001 | Mar 14, 2001 |
| AB | | | 300MG | A065028 002 | Mar 14, 2001 |
| AB | | LANNETT CO INC | 150MG | A065390 001 | Mar 28, 2008 |
| AB | | | 300MG | A065390 002 | Mar 28, 2008 |
| AB | | LUPIN PHARMS | 150MG | A090034 001 | Aug 21, 2013 |
| AB | | | 300MG | A090034 002 | Aug 21, 2013 |
| AB | | SANDOZ | 150MG | A064150 002 | Jan 02, 1998 |
| AB | | | 300MG | A064150 001 | May 28, 1997 |

RIMACTANE

| | | | | | |
|-----------|--|---------------|--------------|--------------------|--|
| AB | | OXFORD PHARMS | 300MG | N050429 001 | |
|-----------|--|---------------|--------------|--------------------|--|

INJECTABLE; INJECTION

RIFADIN

| | | | | | |
|-----------|----------|-------------------|-------------------|--------------------|--------------|
| AP | + | SANOFI AVENTIS US | 600MG/VIAL | N050627 001 | May 25, 1989 |
|-----------|----------|-------------------|-------------------|--------------------|--------------|

RIFAMPIN

| | | | | | |
|-----------|--|-------------------------|-------------------|--------------------|--------------|
| AP | | AKORN | 600MG/VIAL | A065502 001 | Sep 21, 2010 |
| AP | | EMCURE PHARMS LTD | 600MG/VIAL | A204101 001 | Aug 18, 2014 |
| AP | | FRESENIUS KABI USA | 600MG/VIAL | A091181 001 | Aug 21, 2014 |
| AP | | HIKMA PHARMS | 600MG/VIAL | A205039 001 | Mar 03, 2016 |
| AP | | MYLAN LABS LTD | 600MG/VIAL | A065421 001 | May 22, 2008 |
| AP | | WATSON PHARMS TEVA | 600MG/VIAL | A206736 001 | Jan 19, 2016 |
| AP | | WEST-WARD PHARMS INT | 600MG/VIAL | A064217 001 | Oct 29, 1999 |

RIFAMYCIN

TABLET, DELAYED RELEASE; ORAL

AEMCOLO

| | | | | | |
|--|----------|--------------------|-------|-------------|--------------|
| | + | COSMO TECHNOLOGIES | 194MG | N210910 001 | Nov 16, 2018 |
|--|----------|--------------------|-------|-------------|--------------|

RIFAPENTINE

TABLET; ORAL

PRIFTIN

| | | | | | |
|--|----------|-------------------|-------|-------------|--------------|
| | + | SANOFI AVENTIS US | 150MG | N021024 001 | Jun 22, 1998 |
|--|----------|-------------------|-------|-------------|--------------|

RIFAXIMIN

TABLET; ORAL

XIFAXAN

| | | | | | |
|--|----------|--------------|-------|-------------|--------------|
| | + | SALIX PHARMS | 200MG | N021361 001 | May 25, 2004 |
| | + | | 550MG | N022554 001 | Mar 24, 2010 |

RILPIVIRINE HYDROCHLORIDE

TABLET; ORAL

EDURANT

| | | | | | |
|--|----------|---------------|--------------|-------------|--------------|
| | + | JANSSEN PRODS | EQ 25MG BASE | N202022 001 | May 20, 2011 |
|--|----------|---------------|--------------|-------------|--------------|

RILUZOLE

SUSPENSION; ORAL

TIGLUTIK KIT

| | | | | | |
|--|----------|-----------------|-----------|-------------|--------------|
| | + | ITALFARMACO SPA | 50MG/10ML | N209080 001 | Sep 05, 2018 |
|--|----------|-----------------|-----------|-------------|--------------|

TABLET; ORAL

RILUTEK

| | | | | | |
|-----------|----------|-----------------|-------------|--------------------|--------------|
| AB | + | COVIS PHARMA BV | 50MG | N020599 001 | Dec 12, 1995 |
|-----------|----------|-----------------|-------------|--------------------|--------------|

RILUZOLE

| | | | | | |
|-----------|--|---------------------|-------------|--------------------|--------------|
| AB | | ALKEM LABS LTD | 50MG | A204048 001 | Mar 30, 2016 |
| AB | | APOTEX CORP | 50MG | A091300 001 | Jun 18, 2013 |
| AB | | DAITO PHARMS CO LTD | 50MG | A204430 001 | Oct 16, 2018 |
| AB | | GLENMARK PHARMS LTD | 50MG | A091394 001 | Jun 18, 2013 |
| AB | | IMPAX LABS | 50MG | A076173 001 | Jan 29, 2003 |
| AB | | MYLAN PHARMS INC | 50MG | A203042 001 | Jul 01, 2013 |
| AB | | SUN PHARM INDS LTD | 50MG | A091417 001 | Jun 18, 2013 |

RIMANTADINE HYDROCHLORIDE

TABLET; ORAL

FLUMADINE

| | | | | | |
|-----------|----------|--------------------|--------------|--------------------|--------------|
| AB | + | SUN PHARM INDS INC | 100MG | N019649 001 | Sep 17, 1993 |
|-----------|----------|--------------------|--------------|--------------------|--------------|

RIMANTADINE HYDROCHLORIDE

| | | | | | |
|-----------|--|------------|--------------|--------------------|--------------|
| AB | | IMPAX LABS | 100MG | A076132 001 | Aug 30, 2002 |
|-----------|--|------------|--------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

RIOCIGUAT

TABLET;ORAL

ADEMPAS

| | | | | |
|---|----------------|-------|-------------|--------------|
| + | BAYER HLTHCARE | 0.5MG | N204819 001 | Oct 08, 2013 |
| + | | 1MG | N204819 002 | Oct 08, 2013 |
| + | | 1.5MG | N204819 003 | Oct 08, 2013 |
| + | | 2MG | N204819 004 | Oct 08, 2013 |
| + | | 2.5MG | N204819 005 | Oct 08, 2013 |

RISEDRONATE SODIUM

TABLET;ORAL

ACTONEL

| | | | | | |
|-----------|---|------|--------------|--------------------|--------------|
| <u>AB</u> | + | APIL | <u>5MG</u> | <u>N020835 002</u> | Apr 14, 2000 |
| <u>AB</u> | + | | <u>30MG</u> | <u>N020835 001</u> | Mar 27, 1998 |
| <u>AB</u> | + | ! | <u>35MG</u> | <u>N020835 003</u> | May 25, 2002 |
| <u>AB</u> | + | ! | <u>150MG</u> | <u>N020835 005</u> | Apr 22, 2008 |

RISEDRONATE SODIUM

| | | | | | |
|-----------|--|----------------------|--------------|--------------------|--------------|
| <u>AB</u> | | APOTEX INC | <u>35MG</u> | <u>A090877 001</u> | Nov 30, 2015 |
| <u>AB</u> | | | <u>75MG</u> | <u>A090877 002</u> | Jun 10, 2014 |
| <u>AB</u> | | | <u>150MG</u> | <u>A090877 003</u> | Jun 10, 2014 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>5MG</u> | <u>A200296 001</u> | Nov 30, 2015 |
| <u>AB</u> | | | <u>30MG</u> | <u>A200296 002</u> | Nov 30, 2015 |
| <u>AB</u> | | | <u>35MG</u> | <u>A200296 003</u> | Nov 30, 2015 |
| <u>AB</u> | | | <u>150MG</u> | <u>A206768 001</u> | Oct 21, 2016 |
| <u>AB</u> | | MACLEODS PHARMS LTD | <u>5MG</u> | <u>A203533 001</u> | Dec 09, 2015 |
| <u>AB</u> | | | <u>30MG</u> | <u>A203533 002</u> | Dec 09, 2015 |
| <u>AB</u> | | | <u>35MG</u> | <u>A203533 003</u> | Nov 29, 2016 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>5MG</u> | <u>A200477 001</u> | Nov 30, 2015 |
| <u>AB</u> | | | <u>30MG</u> | <u>A200477 002</u> | Nov 30, 2015 |
| <u>AB</u> | | | <u>35MG</u> | <u>A200477 003</u> | Nov 30, 2015 |
| <u>AB</u> | | | <u>75MG</u> | <u>A200477 004</u> | Jun 10, 2014 |
| <u>AB</u> | | | <u>150MG</u> | <u>A200477 005</u> | Jun 10, 2014 |
| <u>AB</u> | | SUN PHARMA GLOBAL | <u>5MG</u> | <u>A090886 001</u> | Nov 30, 2015 |
| <u>AB</u> | | | <u>30MG</u> | <u>A090886 002</u> | Nov 30, 2015 |
| <u>AB</u> | | | <u>35MG</u> | <u>A090886 003</u> | Nov 30, 2015 |
| <u>AB</u> | | | <u>75MG</u> | <u>A090886 004</u> | Jun 10, 2014 |
| <u>AB</u> | | | <u>150MG</u> | <u>A090886 005</u> | Jun 10, 2014 |
| <u>AB</u> | | TEVA PHARMS USA | <u>5MG</u> | <u>A077132 001</u> | Oct 05, 2007 |
| <u>AB</u> | | | <u>30MG</u> | <u>A077132 002</u> | Oct 05, 2007 |
| <u>AB</u> | | | <u>35MG</u> | <u>A077132 003</u> | Oct 05, 2007 |
| <u>AB</u> | | | <u>150MG</u> | <u>A079215 001</u> | Jun 13, 2014 |

TABLET, DELAYED RELEASE;ORAL

ATELVIA

| | | | | | |
|-----------|---|------|-------------|--------------------|--------------|
| <u>AB</u> | + | APIL | <u>35MG</u> | <u>N022560 001</u> | Oct 08, 2010 |
|-----------|---|------|-------------|--------------------|--------------|

RISEDRONATE SODIUM

| | | | | | |
|-----------|--|----------------------|-------------|--------------------|--------------|
| <u>AB</u> | | TEVA PHARMS USA | <u>35MG</u> | <u>A203217 001</u> | May 18, 2015 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>35MG</u> | <u>A203822 001</u> | Sep 11, 2018 |

RISPERIDONE

FOR SUSPENSION, EXTENDED RELEASE;SUBCUTANEOUS

PERSERIS KIT

| | | | | |
|---|--------------|-------|-------------|--------------|
| + | INDIVIOR INC | 90MG | N210655 001 | Jul 27, 2018 |
| + | ! | 120MG | N210655 002 | Jul 27, 2018 |

INJECTABLE;INTRAMUSCULAR

RISPERDAL CONSTA

| | | | | |
|---|----------------|-------------|-------------|--------------|
| + | JANSSEN PHARMS | 12.5MG/VIAL | N021346 004 | Apr 12, 2007 |
| + | ! | 25MG/VIAL | N021346 001 | Oct 29, 2003 |
| + | | 37.5MG/VIAL | N021346 002 | Oct 29, 2003 |
| + | | 50MG/VIAL | N021346 003 | Oct 29, 2003 |

SOLUTION;ORAL

RISPERDAL

| | | | | | |
|-----------|---|----------------|---------------|--------------------|--------------|
| <u>AA</u> | + | JANSSEN PHARMS | <u>1MG/ML</u> | <u>N020588 001</u> | Jun 10, 1996 |
|-----------|---|----------------|---------------|--------------------|--------------|

RISPERIDONE

| | | | | | |
|-----------|--|----------------------|---------------|--------------------|--------------|
| <u>AA</u> | | AMNEAL PHARMS | <u>1MG/ML</u> | <u>A091384 001</u> | May 25, 2011 |
| <u>AA</u> | | ANI PHARMS INC | <u>1MG/ML</u> | <u>A076440 001</u> | Jan 30, 2009 |
| <u>AA</u> | | APOTEX INC | <u>1MG/ML</u> | <u>A077719 001</u> | Jul 29, 2009 |
| <u>AA</u> | | AUROBINDO PHARMA LTD | <u>1MG/ML</u> | <u>A078452 001</u> | Sep 04, 2009 |
| <u>AA</u> | | BIO PHARM INC | <u>1MG/ML</u> | <u>A078909 001</u> | Jul 29, 2009 |
| <u>AA</u> | | LANNETT CO INC | <u>1MG/ML</u> | <u>A079158 001</u> | Dec 03, 2010 |
| <u>AA</u> | | TARO | <u>1MG/ML</u> | <u>A090347 001</u> | Feb 07, 2011 |
| <u>AA</u> | | TRIS PHARMA INC | <u>1MG/ML</u> | <u>A079059 001</u> | Dec 12, 2012 |

PRESCRIPTION DRUG PRODUCT LIST

RISPERIDONE

SOLUTION;ORAL

RISPERIDONE

| | | | | | |
|-----------|-------------------------|---------------|----------------|------------|--------------|
| <u>AA</u> | WEST-WARD PHARMS INT | <u>1MG/ML</u> | <u>A076904</u> | <u>001</u> | Jul 29, 2009 |
|-----------|-------------------------|---------------|----------------|------------|--------------|

TABLET;ORAL

RISPERDAL

| | | | | | | |
|-----------|---|----------------|---------------|----------------|------------|--------------|
| <u>AB</u> | + | JANSSEN PHARMS | <u>0.25MG</u> | <u>N020272</u> | <u>008</u> | May 10, 1999 |
| <u>AB</u> | + | | <u>0.5MG</u> | <u>N020272</u> | <u>007</u> | Jan 27, 1999 |
| <u>AB</u> | + | ! | <u>1MG</u> | <u>N020272</u> | <u>001</u> | Dec 29, 1993 |
| <u>AB</u> | + | | <u>2MG</u> | <u>N020272</u> | <u>002</u> | Dec 29, 1993 |
| <u>AB</u> | + | | <u>3MG</u> | <u>N020272</u> | <u>003</u> | Dec 29, 1993 |
| <u>AB</u> | + | | <u>4MG</u> | <u>N020272</u> | <u>004</u> | Dec 29, 1993 |

RISPERIDONE

| | | | | | | |
|-----------|--|---------------------|---------------|----------------|------------|--------------|
| <u>AB</u> | | AJANTA PHARMA LTD | <u>0.25MG</u> | <u>A201003</u> | <u>001</u> | Aug 24, 2011 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A201003</u> | <u>002</u> | Aug 24, 2011 |
| <u>AB</u> | | | <u>1MG</u> | <u>A201003</u> | <u>003</u> | Aug 24, 2011 |
| <u>AB</u> | | | <u>2MG</u> | <u>A201003</u> | <u>004</u> | Aug 24, 2011 |
| <u>AB</u> | | | <u>3MG</u> | <u>A201003</u> | <u>005</u> | Aug 24, 2011 |
| <u>AB</u> | | | <u>4MG</u> | <u>A201003</u> | <u>006</u> | Aug 24, 2011 |
| <u>AB</u> | | APOTEX INC | <u>0.25MG</u> | <u>A077953</u> | <u>001</u> | Sep 15, 2008 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A077953</u> | <u>002</u> | Sep 15, 2008 |
| <u>AB</u> | | | <u>1MG</u> | <u>A077953</u> | <u>003</u> | Sep 15, 2008 |
| <u>AB</u> | | | <u>2MG</u> | <u>A077953</u> | <u>004</u> | Sep 15, 2008 |
| <u>AB</u> | | | <u>3MG</u> | <u>A077953</u> | <u>005</u> | Sep 15, 2008 |
| <u>AB</u> | | | <u>4MG</u> | <u>A077953</u> | <u>006</u> | Sep 15, 2008 |
| <u>AB</u> | | CEYONE | <u>0.25MG</u> | <u>A078269</u> | <u>001</u> | Oct 08, 2008 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A078269</u> | <u>002</u> | Oct 08, 2008 |
| <u>AB</u> | | | <u>1MG</u> | <u>A078269</u> | <u>003</u> | Oct 08, 2008 |
| <u>AB</u> | | | <u>2MG</u> | <u>A078269</u> | <u>004</u> | Oct 08, 2008 |
| <u>AB</u> | | | <u>3MG</u> | <u>A078269</u> | <u>005</u> | Oct 08, 2008 |
| <u>AB</u> | | | <u>4MG</u> | <u>A078269</u> | <u>006</u> | Oct 08, 2008 |
| <u>AB</u> | | CHARTWELL MOLECULAR | <u>0.25MG</u> | <u>A077543</u> | <u>001</u> | May 18, 2011 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A077543</u> | <u>002</u> | May 18, 2011 |
| <u>AB</u> | | | <u>1MG</u> | <u>A077543</u> | <u>003</u> | May 18, 2011 |
| <u>AB</u> | | | <u>2MG</u> | <u>A077543</u> | <u>004</u> | May 18, 2011 |
| <u>AB</u> | | | <u>3MG</u> | <u>A077543</u> | <u>005</u> | May 18, 2011 |
| <u>AB</u> | | | <u>4MG</u> | <u>A077543</u> | <u>006</u> | May 18, 2011 |
| <u>AB</u> | | DR REDDYS LABS LTD | <u>0.25MG</u> | <u>A076879</u> | <u>001</u> | Oct 24, 2008 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A076879</u> | <u>002</u> | Oct 24, 2008 |
| <u>AB</u> | | | <u>1MG</u> | <u>A076879</u> | <u>003</u> | Oct 24, 2008 |
| <u>AB</u> | | | <u>2MG</u> | <u>A076879</u> | <u>004</u> | Oct 24, 2008 |
| <u>AB</u> | | | <u>3MG</u> | <u>A076879</u> | <u>005</u> | Oct 24, 2008 |
| <u>AB</u> | | | <u>4MG</u> | <u>A076879</u> | <u>006</u> | Oct 24, 2008 |
| <u>AB</u> | | MYLAN | <u>0.25MG</u> | <u>A076288</u> | <u>001</u> | Sep 15, 2008 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A076288</u> | <u>002</u> | Sep 15, 2008 |
| <u>AB</u> | | | <u>1MG</u> | <u>A076288</u> | <u>003</u> | Sep 15, 2008 |
| <u>AB</u> | | | <u>2MG</u> | <u>A076288</u> | <u>004</u> | Sep 15, 2008 |
| <u>AB</u> | | | <u>3MG</u> | <u>A076288</u> | <u>005</u> | Sep 15, 2008 |
| <u>AB</u> | | | <u>4MG</u> | <u>A076288</u> | <u>006</u> | Sep 15, 2008 |
| <u>AB</u> | | OXFORD PHARMS | <u>0.25MG</u> | <u>A078071</u> | <u>001</u> | Jun 17, 2009 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A078071</u> | <u>002</u> | Jun 17, 2009 |
| <u>AB</u> | | | <u>1MG</u> | <u>A078071</u> | <u>003</u> | Jun 17, 2009 |
| <u>AB</u> | | | <u>2MG</u> | <u>A078071</u> | <u>004</u> | Jun 17, 2009 |
| <u>AB</u> | | | <u>3MG</u> | <u>A078071</u> | <u>005</u> | Jun 17, 2009 |
| <u>AB</u> | | | <u>4MG</u> | <u>A078071</u> | <u>006</u> | Jun 17, 2009 |
| <u>AB</u> | | PLIVA HRVATSKA DOO | <u>0.25MG</u> | <u>A077769</u> | <u>001</u> | Oct 16, 2008 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A077769</u> | <u>002</u> | Oct 16, 2008 |
| <u>AB</u> | | | <u>1MG</u> | <u>A077769</u> | <u>003</u> | Oct 16, 2008 |
| <u>AB</u> | | | <u>2MG</u> | <u>A077769</u> | <u>004</u> | Oct 16, 2008 |
| <u>AB</u> | | | <u>3MG</u> | <u>A077769</u> | <u>005</u> | Oct 16, 2008 |
| <u>AB</u> | | | <u>4MG</u> | <u>A077769</u> | <u>006</u> | Oct 16, 2008 |
| <u>AB</u> | | PRINSTON INC | <u>0.25MG</u> | <u>A077493</u> | <u>001</u> | Nov 29, 2011 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A077493</u> | <u>002</u> | Nov 29, 2011 |
| <u>AB</u> | | | <u>1MG</u> | <u>A077493</u> | <u>003</u> | Nov 29, 2011 |
| <u>AB</u> | | | <u>2MG</u> | <u>A077493</u> | <u>004</u> | Nov 29, 2011 |
| <u>AB</u> | | | <u>3MG</u> | <u>A077493</u> | <u>005</u> | Nov 29, 2011 |
| <u>AB</u> | | | <u>4MG</u> | <u>A077493</u> | <u>006</u> | Nov 29, 2011 |
| <u>AB</u> | | RENATA | <u>0.25MG</u> | <u>A078707</u> | <u>001</u> | Dec 29, 2008 |
| <u>AB</u> | | | <u>0.5MG</u> | <u>A078707</u> | <u>002</u> | Dec 29, 2008 |
| <u>AB</u> | | | <u>1MG</u> | <u>A078707</u> | <u>003</u> | Dec 29, 2008 |
| <u>AB</u> | | | <u>2MG</u> | <u>A078707</u> | <u>004</u> | Dec 29, 2008 |
| <u>AB</u> | | | <u>3MG</u> | <u>A078707</u> | <u>005</u> | Dec 29, 2008 |

PRESCRIPTION DRUG PRODUCT LIST

RISPERIDONE

TABLET; ORAL

RISPERIDONE

| | | | | |
|-----------|-------------------------|---------------|--------------------|--------------|
| <u>AB</u> | | <u>4MG</u> | <u>A078707 006</u> | Dec 29, 2008 |
| <u>AB</u> | SANDOZ | <u>0.25MG</u> | <u>A078528 001</u> | Oct 16, 2009 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A078528 002</u> | Oct 16, 2009 |
| <u>AB</u> | | <u>1MG</u> | <u>A078528 003</u> | Oct 16, 2009 |
| <u>AB</u> | | <u>2MG</u> | <u>A078528 004</u> | Oct 16, 2009 |
| <u>AB</u> | | <u>3MG</u> | <u>A078528 005</u> | Oct 16, 2009 |
| <u>AB</u> | | <u>4MG</u> | <u>A078528 006</u> | Oct 16, 2009 |
| <u>AB</u> | SUN PHARM INDS INC | <u>0.25MG</u> | <u>A078036 001</u> | Mar 10, 2014 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A078036 002</u> | Mar 10, 2014 |
| <u>AB</u> | | <u>1MG</u> | <u>A078036 003</u> | Mar 10, 2014 |
| <u>AB</u> | | <u>2MG</u> | <u>A078036 004</u> | Mar 10, 2014 |
| <u>AB</u> | | <u>3MG</u> | <u>A078036 005</u> | Mar 10, 2014 |
| <u>AB</u> | | <u>4MG</u> | <u>A078036 006</u> | Mar 10, 2014 |
| <u>AB</u> | TEVA | <u>0.25MG</u> | <u>A076228 001</u> | Jun 30, 2008 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A076228 002</u> | Jun 30, 2008 |
| <u>AB</u> | | <u>1MG</u> | <u>A076228 003</u> | Jun 30, 2008 |
| <u>AB</u> | | <u>2MG</u> | <u>A076228 004</u> | Jun 30, 2008 |
| <u>AB</u> | | <u>3MG</u> | <u>A076228 005</u> | Jun 30, 2008 |
| <u>AB</u> | | <u>4MG</u> | <u>A076228 006</u> | Jun 30, 2008 |
| <u>AB</u> | TORRENT PHARMS | <u>0.25MG</u> | <u>A079088 001</u> | Oct 30, 2008 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A079088 002</u> | Oct 30, 2008 |
| <u>AB</u> | | <u>1MG</u> | <u>A079088 003</u> | Oct 30, 2008 |
| <u>AB</u> | | <u>2MG</u> | <u>A079088 004</u> | Oct 30, 2008 |
| <u>AB</u> | | <u>3MG</u> | <u>A079088 005</u> | Oct 30, 2008 |
| <u>AB</u> | | <u>4MG</u> | <u>A079088 006</u> | Oct 30, 2008 |
| <u>AB</u> | WOCKHARDT | <u>0.25MG</u> | <u>A078871 001</u> | Oct 09, 2008 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A078871 002</u> | Oct 09, 2008 |
| <u>AB</u> | | <u>1MG</u> | <u>A078871 003</u> | Oct 09, 2008 |
| <u>AB</u> | | <u>2MG</u> | <u>A078871 004</u> | Oct 09, 2008 |
| <u>AB</u> | | <u>3MG</u> | <u>A078871 005</u> | Oct 09, 2008 |
| <u>AB</u> | | <u>4MG</u> | <u>A078871 006</u> | Oct 09, 2008 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>0.25MG</u> | <u>A078040 001</u> | Oct 16, 2008 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A078040 002</u> | Oct 16, 2008 |
| <u>AB</u> | | <u>1MG</u> | <u>A078040 003</u> | Oct 16, 2008 |
| <u>AB</u> | | <u>2MG</u> | <u>A078040 004</u> | Oct 16, 2008 |
| <u>AB</u> | | <u>3MG</u> | <u>A078040 005</u> | Oct 16, 2008 |
| <u>AB</u> | | <u>4MG</u> | <u>A078040 006</u> | Oct 16, 2008 |

TABLET, ORALLY DISINTEGRATING; ORAL

RISPERDAL

| | | | | | |
|-----------|---|----------------|--------------|--------------------|--------------|
| <u>AB</u> | + | JANSSEN PHARMS | <u>0.5MG</u> | <u>N021444 001</u> | Apr 02, 2003 |
| <u>AB</u> | + | ! | <u>1MG</u> | <u>N021444 002</u> | Apr 02, 2003 |
| <u>AB</u> | + | | <u>2MG</u> | <u>N021444 003</u> | Apr 02, 2003 |
| <u>AB</u> | + | | <u>3MG</u> | <u>N021444 004</u> | Dec 23, 2004 |
| <u>AB</u> | + | | <u>4MG</u> | <u>N021444 005</u> | Dec 23, 2004 |

RISPERIDONE

| | | | | | |
|-----------|--|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS LABS FL INC | <u>0.5MG</u> | <u>A076996 001</u> | Apr 19, 2011 |
| <u>AB</u> | | | <u>1MG</u> | <u>A076996 002</u> | Apr 19, 2011 |
| <u>AB</u> | | | <u>2MG</u> | <u>A076996 003</u> | Apr 19, 2011 |
| <u>AB</u> | | | <u>3MG</u> | <u>A076996 004</u> | Apr 19, 2011 |
| <u>AB</u> | | | <u>4MG</u> | <u>A076996 005</u> | Apr 19, 2011 |
| <u>AB</u> | | DR REDDYS LABS LTD | <u>0.5MG</u> | <u>A077328 001</u> | Feb 24, 2009 |
| <u>AB</u> | | | <u>1MG</u> | <u>A077328 002</u> | Oct 05, 2009 |
| <u>AB</u> | | | <u>2MG</u> | <u>A077328 003</u> | Feb 24, 2009 |
| <u>AB</u> | | | <u>3MG</u> | <u>A077328 004</u> | Nov 30, 2009 |
| <u>AB</u> | | | <u>4MG</u> | <u>A077328 005</u> | Nov 30, 2009 |
| <u>AB</u> | | JUBILANT GENERICS | <u>0.5MG</u> | <u>A090839 001</u> | Nov 04, 2011 |
| <u>AB</u> | | | <u>1MG</u> | <u>A090839 002</u> | Nov 04, 2011 |
| <u>AB</u> | | | <u>2MG</u> | <u>A090839 003</u> | Nov 04, 2011 |
| <u>AB</u> | | | <u>3MG</u> | <u>A090839 004</u> | Nov 04, 2011 |
| <u>AB</u> | | | <u>4MG</u> | <u>A090839 005</u> | Nov 04, 2011 |
| <u>AB</u> | | PAR PHARM | <u>0.5MG</u> | <u>A077494 002</u> | Apr 30, 2009 |
| <u>AB</u> | | | <u>1MG</u> | <u>A077494 003</u> | Oct 26, 2009 |
| <u>AB</u> | | | <u>2MG</u> | <u>A077494 004</u> | Apr 30, 2009 |
| <u>AB</u> | | | <u>3MG</u> | <u>A077494 005</u> | Apr 30, 2009 |
| <u>AB</u> | | | <u>4MG</u> | <u>A077494 006</u> | Apr 30, 2009 |
| <u>AB</u> | | SANDOZ | <u>0.5MG</u> | <u>A078116 001</u> | Dec 22, 2009 |
| <u>AB</u> | | | <u>1MG</u> | <u>A078116 002</u> | Dec 22, 2009 |
| <u>AB</u> | | | <u>2MG</u> | <u>A078116 003</u> | Dec 22, 2009 |
| <u>AB</u> | | | <u>3MG</u> | <u>A078116 004</u> | Dec 22, 2009 |

PRESCRIPTION DRUG PRODUCT LIST

RISPERIDONE

TABLET, ORALLY DISINTEGRATING;ORAL

RISPERIDONE

| | | | | | |
|-----------|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | <u>4MG</u> | <u>A078116</u> | <u>005</u> | Dec 22, 2009 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>0.5MG</u> | <u>A077542</u> | <u>001</u> | Aug 06, 2010 |
| <u>AB</u> | | <u>0.5MG</u> | <u>A078464</u> | <u>001</u> | Apr 08, 2013 |
| <u>AB</u> | | <u>1MG</u> | <u>A077542</u> | <u>002</u> | Aug 06, 2010 |
| <u>AB</u> | | <u>1MG</u> | <u>A078464</u> | <u>002</u> | Apr 08, 2013 |
| <u>AB</u> | | <u>2MG</u> | <u>A077542</u> | <u>003</u> | Aug 06, 2010 |
| <u>AB</u> | | <u>2MG</u> | <u>A078464</u> | <u>003</u> | Apr 08, 2013 |
| <u>AB</u> | | <u>3MG</u> | <u>A078464</u> | <u>004</u> | Apr 08, 2013 |
| <u>AB</u> | | <u>3MG</u> | <u>A078474</u> | <u>001</u> | Aug 06, 2010 |
| <u>AB</u> | | <u>4MG</u> | <u>A078464</u> | <u>005</u> | Apr 08, 2013 |
| <u>AB</u> | | <u>4MG</u> | <u>A078474</u> | <u>002</u> | Aug 06, 2010 |
| <u>AB</u> | TEVA | <u>0.5MG</u> | <u>A076908</u> | <u>001</u> | Mar 12, 2012 |
| <u>AB</u> | | <u>1MG</u> | <u>A076908</u> | <u>002</u> | Mar 12, 2012 |
| <u>AB</u> | | <u>2MG</u> | <u>A076908</u> | <u>003</u> | Mar 12, 2012 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>0.5MG</u> | <u>A078516</u> | <u>001</u> | May 01, 2009 |
| <u>AB</u> | | <u>2MG</u> | <u>A078516</u> | <u>003</u> | May 01, 2009 |
| | PAR PHARM | 0.25MG | A077494 | 001 | Apr 30, 2009 |

RITONAVIR

POWDER;ORAL

NORVIR

+! ABBVIE INC

100MG/PACKET

N209512 001 Jun 07, 2017

SOLUTION;ORAL

NORVIR

+! ABBVIE

80MG/ML

N020659 001 Mar 01, 1996

TABLET;ORAL

NORVIRAB +! ABBVIE100MGN022417 001 Feb 10, 2010RITONAVIRAB AMNEAL PHARMS LLC100MGA208890 001 Sep 17, 2018AB AUROBINDO PHARMA LTD100MGA206614 001 Sep 17, 2018AB HETERO LABS LTD III100MGA204587 001 Sep 17, 2018AB WEST-WARD PHARMS INT100MGA202573 001 Jan 15, 2015RIVAROXABAN

TABLET;ORAL

XARELTO

+ JANSSEN PHARMS

2.5MG

N022406 004 Oct 11, 2018

+

10MG

N022406 001 Jul 01, 2011

+

15MG

N022406 002 Nov 04, 2011

+!

20MG

N022406 003 Nov 04, 2011

RIVASTIGMINE

FILM, EXTENDED RELEASE;TRANSDERMAL

EXELONAB + NOVARTIS4.6MG/24HRN022083 001 Jul 06, 2007AB +!9.5MG/24HRN022083 002 Jul 06, 2007AB +13.3MG/24HRN022083 005 Aug 31, 2012RIVASTIGMINEAB ALVOGEN MALTA4.6MG/24HRA204403 001 Sep 03, 2015AB9.5MG/24HRA204403 002 Sep 03, 2015AB13.3MG/24HRA204403 003 Aug 31, 2015AB AMNEAL PHARMS4.6MG/24HRA207308 001 Jan 08, 2019AB9.5MG/24HRA207308 002 Jan 08, 2019AB13.3MG/24HRA207308 003 Jan 08, 2019AB MYLAN TECHNOLOGIES4.6MG/24HRA205622 001 Jun 20, 2018AB9.5MG/24HRA205622 002 Jun 20, 2018AB13.3MG/24HRA205622 003 Jun 20, 2018RIVASTIGMINE TARTRATE

CAPSULE;ORAL

EXELONAB +! NOVARTISEQ 1.5MG BASEN020823 003 Apr 21, 2000AB +EQ 3MG BASEN020823 004 Apr 21, 2000AB +EQ 4.5MG BASEN020823 005 Apr 21, 2000AB +!EQ 6MG BASEN020823 006 Apr 21, 2000RIVASTIGMINE TARTRATEAB ALEMBIC PHARMS LTDEQ 1.5MG BASEA091689 001 Jun 12, 2012ABEQ 3MG BASEA091689 002 Jun 12, 2012ABEQ 4.5MG BASEA091689 003 Jun 12, 2012

PRESCRIPTION DRUG PRODUCT LIST

RIVASTIGMINE TARTRATE

CAPSULE; ORAL

RIVASTIGMINE TARTRATE

| | | | | |
|-----------|----------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A091689 004</u> | Jun 12, 2012 |
| <u>AB</u> | APOTEX INC | <u>EQ 1.5MG BASE</u> | <u>A091072 001</u> | May 16, 2013 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A091072 002</u> | May 16, 2013 |
| <u>AB</u> | | <u>EQ 4.5MG BASE</u> | <u>A091072 003</u> | May 16, 2013 |
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A091072 004</u> | May 16, 2013 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 1.5MG BASE</u> | <u>A204572 001</u> | Mar 25, 2016 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A204572 002</u> | Mar 25, 2016 |
| <u>AB</u> | | <u>EQ 4.5MG BASE</u> | <u>A204572 003</u> | Mar 25, 2016 |
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A204572 004</u> | Mar 25, 2016 |
| <u>AB</u> | CADILA PHARMS LTD | <u>EQ 1.5MG BASE</u> | <u>A203844 001</u> | Feb 13, 2017 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A203844 002</u> | Feb 13, 2017 |
| <u>AB</u> | | <u>EQ 4.5MG BASE</u> | <u>A203844 003</u> | Feb 13, 2017 |
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A203844 004</u> | Feb 13, 2017 |
| <u>AB</u> | CHARTWELL RX | <u>EQ 1.5MG BASE</u> | <u>A207797 001</u> | Sep 28, 2017 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A207797 002</u> | Sep 28, 2017 |
| <u>AB</u> | | <u>EQ 4.5MG BASE</u> | <u>A207797 003</u> | Sep 28, 2017 |
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A207797 004</u> | Sep 28, 2017 |
| <u>AB</u> | DR REDDYS LABS INC | <u>EQ 1.5MG BASE</u> | <u>A077130 001</u> | Oct 31, 2007 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A077130 002</u> | Oct 31, 2007 |
| <u>AB</u> | | <u>EQ 4.5MG BASE</u> | <u>A077130 003</u> | Oct 31, 2007 |
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A077130 004</u> | Oct 31, 2007 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>EQ 1.5MG BASE</u> | <u>A203148 001</u> | Aug 22, 2014 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A203148 002</u> | Aug 22, 2014 |
| <u>AB</u> | | <u>EQ 4.5MG BASE</u> | <u>A203148 003</u> | Aug 22, 2014 |
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A203148 004</u> | Aug 22, 2014 |
| <u>AB</u> | ORCHID HLTHCARE | <u>EQ 1.5MG BASE</u> | <u>A090879 001</u> | Jun 10, 2015 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A090879 002</u> | Jun 10, 2015 |
| <u>AB</u> | | <u>EQ 4.5MG BASE</u> | <u>A090879 003</u> | Jun 10, 2015 |
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A090879 004</u> | Jun 10, 2015 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>EQ 1.5MG BASE</u> | <u>A077131 001</u> | Oct 22, 2007 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A077131 002</u> | Oct 22, 2007 |
| <u>AB</u> | | <u>EQ 4.5MG BASE</u> | <u>A077131 003</u> | Oct 22, 2007 |
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A077131 004</u> | Oct 22, 2007 |
| <u>AB</u> | WATSON LABS | <u>EQ 1.5MG BASE</u> | <u>A077129 001</u> | Jan 08, 2008 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A077129 002</u> | Jan 08, 2008 |
| <u>AB</u> | | <u>EQ 4.5MG BASE</u> | <u>A077129 003</u> | Jan 08, 2008 |
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A077129 004</u> | Jan 08, 2008 |

RIZATRIPTAN BENZOATE

TABLET; ORAL

MAXALT

| | | | | |
|-----------|----------|---------------------|--------------------|--------------|
| <u>AB</u> | +! MERCK | <u>EQ 10MG BASE</u> | <u>N020864 002</u> | Jun 29, 1998 |
|-----------|----------|---------------------|--------------------|--------------|

RIZATRIPTAN BENZOATE

| | | | | |
|-----------|----------------------|---------------------|--------------------|--------------|
| <u>AB</u> | ALKEM LABS LTD | <u>EQ 5MG BASE</u> | <u>A203269 001</u> | Feb 18, 2016 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A203269 002</u> | Feb 18, 2016 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 5MG BASE</u> | <u>A202490 001</u> | Dec 31, 2012 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A202490 002</u> | Dec 31, 2012 |
| <u>AB</u> | CELLTRION | <u>EQ 5MG BASE</u> | <u>A077526 001</u> | Mar 26, 2013 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A077526 002</u> | Mar 26, 2013 |
| <u>AB</u> | ECI PHARMS LLC | <u>EQ 5MG BASE</u> | <u>A202047 001</u> | Dec 31, 2012 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A202047 002</u> | Dec 31, 2012 |
| <u>AB</u> | EMCURE PHARMS LTD | <u>EQ 5MG BASE</u> | <u>A204090 001</u> | Nov 26, 2013 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A204090 002</u> | Nov 26, 2013 |
| <u>AB</u> | GLENMARK GENERICS | <u>EQ 5MG BASE</u> | <u>A201967 001</u> | Dec 31, 2012 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A201967 002</u> | Dec 31, 2012 |
| <u>AB</u> | INVAGEN PHARMS | <u>EQ 5MG BASE</u> | <u>A204339 001</u> | Jul 01, 2013 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A204339 002</u> | Jul 01, 2013 |
| <u>AB</u> | JUBILANT GENERICS | <u>EQ 5MG BASE</u> | <u>A203252 001</u> | Dec 31, 2014 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A203252 002</u> | Dec 31, 2014 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>EQ 5MG BASE</u> | <u>A203147 001</u> | Feb 11, 2014 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A203147 002</u> | Feb 11, 2014 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 5MG BASE</u> | <u>A201993 001</u> | Dec 31, 2012 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A201993 002</u> | Dec 31, 2012 |
| <u>AB</u> | NATCO PHARMA LTD | <u>EQ 5MG BASE</u> | <u>A200482 001</u> | Dec 31, 2012 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A200482 002</u> | Dec 31, 2012 |
| <u>AB</u> | SANDOZ | <u>EQ 5MG BASE</u> | <u>A079230 001</u> | Dec 31, 2012 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A079230 002</u> | Dec 31, 2012 |
| <u>AB</u> | TEVA PHARMS | <u>EQ 5MG BASE</u> | <u>A077263 001</u> | Dec 31, 2012 |

PRESCRIPTION DRUG PRODUCT LIST

RIZATRIPTAN BENZOATE

TABLET; ORAL

RIZATRIPTAN BENZOATE

| | | | | |
|-----------|------------------|---------------------|--------------------|--------------|
| AB | | <u>EQ 10MG BASE</u> | <u>A077263 002</u> | Dec 31, 2012 |
| AB | UNICHEM LABS LTD | <u>EQ 5MG BASE</u> | <u>A207836 001</u> | Mar 07, 2017 |
| AB | | <u>EQ 10MG BASE</u> | <u>A207836 002</u> | Mar 07, 2017 |

TABLET, ORALLY DISINTEGRATING; ORAL

MAXALT-MLT

| | | | | |
|-----------|----------|---------------------|--------------------|--------------|
| AB | +! MERCK | <u>EQ 10MG BASE</u> | <u>N020865 002</u> | Jun 29, 1998 |
|-----------|----------|---------------------|--------------------|--------------|

RIZATRIPTAN BENZOATE

| | | | | |
|-----------|----------------------|---------------------|--------------------|--------------|
| AB | AUROBINDO PHARMA LTD | <u>EQ 5MG BASE</u> | <u>A203062 001</u> | Jul 01, 2013 |
| AB | | <u>EQ 10MG BASE</u> | <u>A203062 002</u> | Jul 01, 2013 |
| AB | GLENMARK PHARMS LTD | <u>EQ 5MG BASE</u> | <u>A201914 001</u> | Jul 01, 2013 |
| AB | | <u>EQ 10MG BASE</u> | <u>A201914 002</u> | Jul 01, 2013 |
| AB | JUBILANT GENERICS | <u>EQ 5MG BASE</u> | <u>A203334 001</u> | Oct 16, 2015 |
| AB | | <u>EQ 10MG BASE</u> | <u>A203334 002</u> | Oct 16, 2015 |
| AB | MACLEODS PHARMS LTD | <u>EQ 5MG BASE</u> | <u>A203146 001</u> | Sep 19, 2014 |
| AB | | <u>EQ 10MG BASE</u> | <u>A203146 002</u> | Sep 19, 2014 |
| AB | MYLAN PHARMS INC | <u>EQ 5MG BASE</u> | <u>A078173 001</u> | Dec 31, 2012 |
| AB | | <u>EQ 10MG BASE</u> | <u>A078173 002</u> | Dec 31, 2012 |
| AB | NATCO PHARMA LTD | <u>EQ 5MG BASE</u> | <u>A203478 001</u> | Jul 01, 2013 |
| AB | | <u>EQ 10MG BASE</u> | <u>A203478 002</u> | Jul 01, 2013 |
| AB | PANACEA BIOTEC LTD | <u>EQ 5MG BASE</u> | <u>A204722 001</u> | Jan 11, 2017 |
| AB | | <u>EQ 10MG BASE</u> | <u>A204722 002</u> | Jan 11, 2017 |
| AB | SANDOZ | <u>EQ 5MG BASE</u> | <u>A078739 001</u> | Jul 01, 2013 |
| AB | | <u>EQ 10MG BASE</u> | <u>A078739 002</u> | Jul 01, 2013 |
| AB | UNICHEM LABS LTD | <u>EQ 5MG BASE</u> | <u>A207835 001</u> | Mar 07, 2017 |
| AB | | <u>EQ 10MG BASE</u> | <u>A207835 002</u> | Mar 07, 2017 |

ROCURONIUM BROMIDE

INJECTABLE; INJECTION

ROCURONIUM BROMIDE

| | | | | |
|-----------|----------------------|-----------------------------|--------------------|--------------|
| AP | AUROBINDO PHARMA LTD | <u>50MG/5ML (10MG/ML)</u> | <u>A206206 001</u> | Apr 12, 2017 |
| AP | | <u>100MG/10ML (10MG/ML)</u> | <u>A206206 002</u> | Apr 12, 2017 |
| AP | FRESENIUS KABI USA | <u>50MG/5ML (10MG/ML)</u> | <u>A078651 001</u> | Dec 29, 2008 |
| AP | | <u>100MG/10ML (10MG/ML)</u> | <u>A078651 002</u> | Dec 29, 2008 |
| AP | GLAND PHARMA LTD | <u>50MG/5ML (10MG/ML)</u> | <u>A205656 001</u> | Apr 04, 2018 |
| AP | | <u>100MG/10ML (10MG/ML)</u> | <u>A205656 002</u> | Apr 04, 2018 |
| AP | HOSPIRA | <u>50MG/5ML (10MG/ML)</u> | <u>A078519 001</u> | Nov 26, 2008 |
| AP | | <u>100MG/10ML (10MG/ML)</u> | <u>A078519 002</u> | Nov 26, 2008 |
| AP | MYLAN INSTITUTIONAL | <u>50MG/5ML (10MG/ML)</u> | <u>A079199 001</u> | Nov 26, 2008 |
| AP | | <u>100MG/10ML (10MG/ML)</u> | <u>A079199 002</u> | Nov 26, 2008 |
| AP | SAGENT PHARMS | <u>50MG/5ML (10MG/ML)</u> | <u>A091458 001</u> | Jul 28, 2010 |
| AP | | <u>100MG/10ML (10MG/ML)</u> | <u>A091458 002</u> | Jul 28, 2010 |
| AP | ! SANDOZ INC | <u>50MG/5ML (10MG/ML)</u> | <u>A079195 001</u> | Dec 05, 2008 |
| AP | ! | <u>100MG/10ML (10MG/ML)</u> | <u>A079195 002</u> | Dec 05, 2008 |
| AP | TAMARANG | <u>50MG/5ML (10MG/ML)</u> | <u>A091115 001</u> | Aug 27, 2012 |
| AP | | <u>100MG/10ML (10MG/ML)</u> | <u>A091115 002</u> | Aug 27, 2012 |
| AP | TEVA PHARMS | <u>50MG/5ML (10MG/ML)</u> | <u>A078717 001</u> | Nov 26, 2008 |
| AP | | <u>100MG/10ML (10MG/ML)</u> | <u>A078717 002</u> | Nov 26, 2008 |
| AP | WEST WARD PHARM CORP | <u>50MG/5ML (10MG/ML)</u> | <u>A204679 001</u> | Feb 28, 2017 |
| AP | | <u>100MG/10ML (10MG/ML)</u> | <u>A204679 002</u> | Feb 28, 2017 |

ROFLUMILAST

TABLET; ORAL

DALIRESP

| | | | | |
|-----------|-----------------------|---------------|--------------------|--------------|
| AB | +! ASTRAZENECA PHARMS | <u>500MCG</u> | <u>N022522 001</u> | Feb 28, 2011 |
|-----------|-----------------------|---------------|--------------------|--------------|

ROFLUMILAST

| | | | | |
|-----------|---------------------|---------------|--------------------|--------------|
| AB | BRECKENRIDGE PHARM | <u>500MCG</u> | <u>A208236 001</u> | Oct 03, 2018 |
| AB | HETERO LABS LTD III | <u>500MCG</u> | <u>A208213 001</u> | Nov 23, 2018 |
| AB | TORRENT PHARMS LTD | <u>500MCG</u> | <u>A208272 001</u> | Aug 06, 2018 |

DALIRESP

| | | | | |
|---|--------------------|--------|-------------|--------------|
| + | ASTRAZENECA PHARMS | 250MCG | N022522 002 | Jan 23, 2018 |
|---|--------------------|--------|-------------|--------------|

ROLAPITANT HYDROCHLORIDE

TABLET; ORAL

VARUBI

| | | | | |
|---|---------------------|--------------|-------------|--------------|
| + | TERSERA THERAPS LLC | EQ 90MG BASE | N206500 001 | Sep 01, 2015 |
|---|---------------------|--------------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

ROMIDEPSIN

POWDER; INTRAVENOUS

ISTODAX

+! CELGENE

10MG/VIAL

N022393 001 Nov 05, 2009

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

REQUIP

| | | | | | |
|-----------|----|---------------------|-----------------------|--------------------|--------------|
| <u>AB</u> | +! | GLAXOSMITHKLINE LLC | <u>EQ 0.25MG BASE</u> | <u>N020658 001</u> | Sep 19, 1997 |
| <u>AB</u> | + | | <u>EQ 0.5MG BASE</u> | <u>N020658 002</u> | Sep 19, 1997 |
| <u>AB</u> | + | | <u>EQ 1MG BASE</u> | <u>N020658 003</u> | Sep 19, 1997 |
| <u>AB</u> | + | | <u>EQ 2MG BASE</u> | <u>N020658 004</u> | Sep 19, 1997 |
| <u>AB</u> | + | | <u>EQ 3MG BASE</u> | <u>N020658 006</u> | Jan 27, 1999 |
| <u>AB</u> | + | | <u>EQ 4MG BASE</u> | <u>N020658 007</u> | Jan 27, 1999 |
| <u>AB</u> | + | | <u>EQ 5MG BASE</u> | <u>N020658 005</u> | Sep 19, 1997 |

ROPINIROLE HYDROCHLORIDE

| | | | | | |
|-----------|--|-------------------------|-----------------------|--------------------|--------------|
| <u>AB</u> | | ACCORD HLTHCARE | <u>EQ 0.25MG BASE</u> | <u>A204022 001</u> | Feb 28, 2017 |
| <u>AB</u> | | | <u>EQ 0.5MG BASE</u> | <u>A204022 002</u> | Feb 28, 2017 |
| <u>AB</u> | | | <u>EQ 1MG BASE</u> | <u>A204022 003</u> | Feb 28, 2017 |
| <u>AB</u> | | | <u>EQ 2MG BASE</u> | <u>A204022 004</u> | Feb 28, 2017 |
| <u>AB</u> | | | <u>EQ 3MG BASE</u> | <u>A204022 005</u> | Feb 28, 2017 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A204022 006</u> | Feb 28, 2017 |
| <u>AB</u> | | | <u>EQ 5MG BASE</u> | <u>A204022 007</u> | Feb 28, 2017 |
| <u>AB</u> | | ALEMBIC LTD | <u>EQ 0.25MG BASE</u> | <u>A090429 001</u> | Mar 24, 2010 |
| <u>AB</u> | | | <u>EQ 0.5MG BASE</u> | <u>A090429 002</u> | Mar 24, 2010 |
| <u>AB</u> | | | <u>EQ 1MG BASE</u> | <u>A090429 003</u> | Mar 24, 2010 |
| <u>AB</u> | | | <u>EQ 2MG BASE</u> | <u>A090429 004</u> | Mar 24, 2010 |
| <u>AB</u> | | | <u>EQ 3MG BASE</u> | <u>A090429 005</u> | Mar 24, 2010 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A090429 006</u> | Mar 24, 2010 |
| <u>AB</u> | | | <u>EQ 5MG BASE</u> | <u>A090429 007</u> | Mar 24, 2010 |
| <u>AB</u> | | APOTEX | <u>EQ 0.25MG BASE</u> | <u>A079165 001</u> | Feb 07, 2012 |
| <u>AB</u> | | | <u>EQ 0.5MG BASE</u> | <u>A079165 002</u> | Feb 07, 2012 |
| <u>AB</u> | | | <u>EQ 1MG BASE</u> | <u>A079165 003</u> | Feb 07, 2012 |
| <u>AB</u> | | | <u>EQ 2MG BASE</u> | <u>A079165 004</u> | Feb 07, 2012 |
| <u>AB</u> | | | <u>EQ 3MG BASE</u> | <u>A079165 005</u> | Feb 07, 2012 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A079165 006</u> | Feb 07, 2012 |
| <u>AB</u> | | | <u>EQ 5MG BASE</u> | <u>A079165 007</u> | Feb 07, 2012 |
| <u>AB</u> | | GLENMARK GENERICS | <u>EQ 0.25MG BASE</u> | <u>A090135 001</u> | Feb 25, 2010 |
| <u>AB</u> | | | <u>EQ 0.5MG BASE</u> | <u>A090135 002</u> | Feb 25, 2010 |
| <u>AB</u> | | | <u>EQ 1MG BASE</u> | <u>A090135 003</u> | Feb 25, 2010 |
| <u>AB</u> | | | <u>EQ 2MG BASE</u> | <u>A090135 004</u> | Feb 25, 2010 |
| <u>AB</u> | | | <u>EQ 3MG BASE</u> | <u>A090135 005</u> | Feb 25, 2010 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A090135 006</u> | Feb 25, 2010 |
| <u>AB</u> | | | <u>EQ 5MG BASE</u> | <u>A090135 007</u> | Feb 25, 2010 |
| <u>AB</u> | | MYLAN | <u>EQ 0.25MG BASE</u> | <u>A078881 001</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 0.5MG BASE</u> | <u>A078881 002</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 1MG BASE</u> | <u>A078881 003</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 2MG BASE</u> | <u>A078881 004</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 3MG BASE</u> | <u>A078881 005</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A078881 006</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 5MG BASE</u> | <u>A078881 007</u> | May 19, 2008 |
| <u>AB</u> | | ORCHID HLTHCARE | <u>EQ 0.25MG BASE</u> | <u>A079229 001</u> | Nov 28, 2012 |
| <u>AB</u> | | | <u>EQ 0.5MG BASE</u> | <u>A079229 002</u> | Nov 28, 2012 |
| <u>AB</u> | | | <u>EQ 1MG BASE</u> | <u>A079229 003</u> | Nov 28, 2012 |
| <u>AB</u> | | | <u>EQ 2MG BASE</u> | <u>A079229 004</u> | Nov 28, 2012 |
| <u>AB</u> | | | <u>EQ 3MG BASE</u> | <u>A079229 005</u> | Nov 28, 2012 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A079229 006</u> | Nov 28, 2012 |
| <u>AB</u> | | | <u>EQ 5MG BASE</u> | <u>A079229 007</u> | Nov 28, 2012 |
| <u>AB</u> | | PRINSTON INC | <u>EQ 0.25MG BASE</u> | <u>A078110 001</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 0.5MG BASE</u> | <u>A078110 002</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 1MG BASE</u> | <u>A078110 003</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 2MG BASE</u> | <u>A078110 004</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 3MG BASE</u> | <u>A078110 005</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A078110 006</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 5MG BASE</u> | <u>A078110 007</u> | Jul 11, 2008 |
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>EQ 0.25MG BASE</u> | <u>A077852 001</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 0.5MG BASE</u> | <u>A077852 002</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 1MG BASE</u> | <u>A077852 003</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 2MG BASE</u> | <u>A077852 004</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 3MG BASE</u> | <u>A077852 005</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A077852 006</u> | May 05, 2008 |
| <u>AB</u> | | | <u>EQ 5MG BASE</u> | <u>A077852 007</u> | May 19, 2008 |

PRESCRIPTION DRUG PRODUCT LIST

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

ROPINIROLE HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|-----------------------|--------------------|--------------|
| <u>AB</u> | WOCKHARDT | <u>EQ 0.25MG BASE</u> | <u>A079050 001</u> | May 29, 2008 |
| <u>AB</u> | | <u>EQ 0.5MG BASE</u> | <u>A079050 002</u> | May 29, 2008 |
| <u>AB</u> | | <u>EQ 1MG BASE</u> | <u>A079050 003</u> | May 29, 2008 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A079050 004</u> | May 29, 2008 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A079050 005</u> | May 29, 2008 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A079050 006</u> | May 29, 2008 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A079050 007</u> | May 29, 2008 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 0.25MG BASE</u> | <u>A090411 001</u> | Jun 01, 2009 |
| <u>AB</u> | | <u>EQ 0.5MG BASE</u> | <u>A090411 002</u> | Jun 01, 2009 |
| <u>AB</u> | | <u>EQ 1MG BASE</u> | <u>A090411 003</u> | Jun 01, 2009 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A090411 004</u> | Jun 01, 2009 |
| <u>AB</u> | | <u>EQ 3MG BASE</u> | <u>A090411 005</u> | Jun 01, 2009 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A090411 006</u> | Jun 01, 2009 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A090411 007</u> | Jun 01, 2009 |

TABLET, EXTENDED RELEASE; ORAL

REQUIP XL

| | | | | | |
|-----------|----|---------------------|---------------------|--------------------|--------------|
| <u>AB</u> | +! | GLAXOSMITHKLINE LLC | <u>EQ 2MG BASE</u> | <u>N022008 001</u> | Jun 13, 2008 |
| <u>AB</u> | + | | <u>EQ 4MG BASE</u> | <u>N022008 003</u> | Jun 13, 2008 |
| <u>AB</u> | + | | <u>EQ 6MG BASE</u> | <u>N022008 006</u> | Apr 10, 2009 |
| <u>AB</u> | + | | <u>EQ 8MG BASE</u> | <u>N022008 004</u> | Jun 13, 2008 |
| <u>AB</u> | + | | <u>EQ 12MG BASE</u> | <u>N022008 005</u> | Oct 31, 2008 |

ROPINIROLE HYDROCHLORIDE

| | | | | | |
|-----------|--|--------------------|---------------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS ELIZABETH | <u>EQ 2MG BASE</u> | <u>A090869 001</u> | May 17, 2012 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A090869 002</u> | May 17, 2012 |
| <u>AB</u> | | | <u>EQ 6MG BASE</u> | <u>A090869 003</u> | May 17, 2012 |
| <u>AB</u> | | | <u>EQ 8MG BASE</u> | <u>A090869 004</u> | May 17, 2012 |
| <u>AB</u> | | | <u>EQ 12MG BASE</u> | <u>A090869 005</u> | May 17, 2012 |
| <u>AB</u> | | ALEMBIC PHARMS LTD | <u>EQ 2MG BASE</u> | <u>A202786 001</u> | Apr 22, 2013 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A202786 002</u> | Apr 22, 2013 |
| <u>AB</u> | | | <u>EQ 6MG BASE</u> | <u>A202786 003</u> | Apr 22, 2013 |
| <u>AB</u> | | | <u>EQ 8MG BASE</u> | <u>A202786 004</u> | Apr 22, 2013 |
| <u>AB</u> | | | <u>EQ 12MG BASE</u> | <u>A202786 005</u> | Apr 22, 2013 |
| <u>AB</u> | | DR REDDYS LABS LTD | <u>EQ 2MG BASE</u> | <u>A201576 001</u> | Jun 06, 2012 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A201576 002</u> | Jun 06, 2012 |
| <u>AB</u> | | | <u>EQ 6MG BASE</u> | <u>A201576 003</u> | Jun 06, 2012 |
| <u>AB</u> | | | <u>EQ 8MG BASE</u> | <u>A201576 004</u> | Jun 06, 2012 |
| <u>AB</u> | | | <u>EQ 12MG BASE</u> | <u>A201576 005</u> | Jun 06, 2012 |
| <u>AB</u> | | SANDOZ INC | <u>EQ 2MG BASE</u> | <u>A201047 001</u> | Jun 06, 2012 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A201047 003</u> | Jun 06, 2012 |
| <u>AB</u> | | | <u>EQ 6MG BASE</u> | <u>A201047 004</u> | Jun 06, 2012 |
| <u>AB</u> | | | <u>EQ 8MG BASE</u> | <u>A201047 005</u> | Jun 06, 2012 |
| <u>AB</u> | | | <u>EQ 12MG BASE</u> | <u>A201047 006</u> | Jun 06, 2012 |
| <u>AB</u> | | WATSON LABS INC | <u>EQ 2MG BASE</u> | <u>A200431 001</u> | Jun 06, 2012 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A200431 002</u> | Jun 06, 2012 |
| <u>AB</u> | | | <u>EQ 6MG BASE</u> | <u>A200431 003</u> | Jun 06, 2012 |
| <u>AB</u> | | | <u>EQ 8MG BASE</u> | <u>A200431 004</u> | Jun 06, 2012 |
| <u>AB</u> | | | <u>EQ 12MG BASE</u> | <u>A200431 005</u> | Jun 06, 2012 |
| <u>AB</u> | | WOCKHARDT LTD | <u>EQ 2MG BASE</u> | <u>A091395 001</u> | Aug 27, 2012 |
| <u>AB</u> | | | <u>EQ 4MG BASE</u> | <u>A091395 002</u> | Aug 27, 2012 |
| <u>AB</u> | | | <u>EQ 6MG BASE</u> | <u>A091395 003</u> | Aug 27, 2012 |
| <u>AB</u> | | | <u>EQ 8MG BASE</u> | <u>A091395 004</u> | Aug 27, 2012 |
| <u>AB</u> | | | <u>EQ 12MG BASE</u> | <u>A091395 005</u> | Aug 27, 2012 |

ROPIVACAINE HYDROCHLORIDE

SOLUTION; INJECTION

NAROPIN

| | | | | | |
|-----------|---|--------------------|------------------------------|--------------------|--------------|
| <u>AP</u> | + | FRESENIUS KABI USA | <u>20MG/10ML (2MG/ML)</u> | <u>N020533 001</u> | May 01, 1998 |
| <u>AP</u> | + | | <u>40MG/20ML (2MG/ML)</u> | <u>N020533 002</u> | Sep 24, 1996 |
| <u>AP</u> | + | | <u>100MG/20ML (5MG/ML)</u> | <u>N020533 003</u> | May 01, 1998 |
| <u>AP</u> | + | | <u>100MG/10ML (10MG/ML)</u> | <u>N020533 005</u> | Sep 24, 1996 |
| <u>AP</u> | + | | <u>150MG/20ML (7.5MG/ML)</u> | <u>N020533 004</u> | Sep 24, 1996 |
| <u>AP</u> | + | | <u>150MG/30ML (5MG/ML)</u> | <u>N020533 008</u> | Sep 24, 1996 |
| <u>AP</u> | + | | <u>200MG/100ML (2MG/ML)</u> | <u>N020533 006</u> | Sep 24, 1996 |
| <u>AP</u> | + | | <u>200MG/20ML (10MG/ML)</u> | <u>N020533 011</u> | Sep 24, 1996 |
| <u>AP</u> | + | | <u>400MG/200ML (2MG/ML)</u> | <u>N020533 007</u> | Sep 24, 1996 |
| <u>AP</u> | + | | <u>500MG/100ML (5MG/ML)</u> | <u>N020533 009</u> | Jan 04, 2011 |
| <u>AP</u> | + | | <u>1GM/200ML (5MG/ML)</u> | <u>N020533 010</u> | Jan 04, 2011 |

ROPIVACAINE HYDROCHLORIDE

| | | | | | |
|-----------|--|-------|-----------------------------|--------------------|--------------|
| <u>AP</u> | | AKORN | <u>200MG/100ML (2MG/ML)</u> | <u>A204636 001</u> | Mar 16, 2018 |
|-----------|--|-------|-----------------------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

ROPIVACAINE HYDROCHLORIDE

SOLUTION; INJECTION

ROPIVACAINE HYDROCHLORIDE

| | | | | | |
|-----------|----------------------|------------------------------|----------------|------------|--------------|
| <u>AP</u> | | <u>400MG/200ML (2MG/ML)</u> | <u>A204636</u> | <u>002</u> | Mar 16, 2018 |
| <u>AP</u> | AKORN INC | <u>150MG/30ML (5MG/ML)</u> | <u>A203955</u> | <u>001</u> | Apr 11, 2016 |
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>40MG/20ML (2MG/ML)</u> | <u>A205612</u> | <u>001</u> | Jul 13, 2016 |
| <u>AP</u> | | <u>100MG/20ML (5MG/ML)</u> | <u>A205612</u> | <u>003</u> | Jul 13, 2016 |
| <u>AP</u> | | <u>100MG/10ML (10MG/ML)</u> | <u>A205612</u> | <u>006</u> | Jul 13, 2016 |
| <u>AP</u> | | <u>150MG/30ML (5MG/ML)</u> | <u>A205612</u> | <u>004</u> | Jul 13, 2016 |
| <u>AP</u> | | <u>150MG/20ML (7.5MG/ML)</u> | <u>A205612</u> | <u>005</u> | Jul 13, 2016 |
| <u>AP</u> | | <u>200MG/100ML (2MG/ML)</u> | <u>A205612</u> | <u>002</u> | Jul 13, 2016 |
| <u>AP</u> | | <u>200MG/20ML (10MG/ML)</u> | <u>A205612</u> | <u>007</u> | Jul 13, 2016 |
| <u>AP</u> | HOSPIRA | <u>20MG/10ML (2MG/ML)</u> | <u>A090194</u> | <u>001</u> | Sep 23, 2014 |
| <u>AP</u> | | <u>40MG/20ML (2MG/ML)</u> | <u>A090194</u> | <u>005</u> | Sep 23, 2014 |
| <u>AP</u> | | <u>100MG/10ML (10MG/ML)</u> | <u>A090194</u> | <u>004</u> | Sep 23, 2014 |
| <u>AP</u> | | <u>150MG/30ML (5MG/ML)</u> | <u>A090194</u> | <u>002</u> | Sep 23, 2014 |
| <u>AP</u> | | <u>150MG/20ML (7.5MG/ML)</u> | <u>A090194</u> | <u>003</u> | Sep 23, 2014 |
| <u>AP</u> | | <u>200MG/20ML (10MG/ML)</u> | <u>A090194</u> | <u>006</u> | Sep 23, 2014 |
| <u>AP</u> | INFORLIFE | <u>200MG/100ML (2MG/ML)</u> | <u>A206166</u> | <u>001</u> | Jun 11, 2018 |
| <u>AP</u> | | <u>400MG/200ML (2MG/ML)</u> | <u>A206166</u> | <u>002</u> | Jun 11, 2018 |
| <u>AP</u> | | <u>500MG/100ML (5MG/ML)</u> | <u>A206166</u> | <u>003</u> | Jun 11, 2018 |
| <u>AP</u> | | <u>1GM/200ML (5MG/ML)</u> | <u>A206166</u> | <u>004</u> | Jun 11, 2018 |
| <u>AP</u> | MYLAN ASI | <u>40MG/20ML (2MG/ML)</u> | <u>A090318</u> | <u>001</u> | Sep 23, 2014 |
| <u>AP</u> | | <u>150MG/30ML (5MG/ML)</u> | <u>A090318</u> | <u>002</u> | Sep 23, 2014 |
| <u>AP</u> | | <u>150MG/20ML (7.5MG/ML)</u> | <u>A090318</u> | <u>003</u> | Sep 23, 2014 |
| <u>AP</u> | | <u>200MG/20ML (10MG/ML)</u> | <u>A090318</u> | <u>004</u> | Sep 23, 2014 |
| <u>AP</u> | NAVINTA LLC | <u>150MG/30ML (5MG/ML)</u> | <u>A078601</u> | <u>002</u> | Jul 17, 2014 |
| <u>AP</u> | | <u>200MG/20ML (10MG/ML)</u> | <u>A078601</u> | <u>003</u> | Jul 17, 2014 |
| <u>AP</u> | SOMERSET THERAPS LLC | <u>20MG/10ML (2MG/ML)</u> | <u>A207636</u> | <u>001</u> | Jun 15, 2018 |
| <u>AP</u> | | <u>40MG/20ML (2MG/ML)</u> | <u>A207636</u> | <u>002</u> | Jun 15, 2018 |
| <u>AP</u> | | <u>100MG/20ML (5MG/ML)</u> | <u>A207636</u> | <u>003</u> | Jun 15, 2018 |
| <u>AP</u> | | <u>100MG/10ML (10MG/ML)</u> | <u>A207636</u> | <u>006</u> | Jun 15, 2018 |
| <u>AP</u> | | <u>150MG/30ML (5MG/ML)</u> | <u>A207636</u> | <u>004</u> | Jun 15, 2018 |
| <u>AP</u> | | <u>150MG/20ML (7.5MG/ML)</u> | <u>A207636</u> | <u>005</u> | Jun 15, 2018 |
| <u>AP</u> | | <u>200MG/20ML (10MG/ML)</u> | <u>A207636</u> | <u>007</u> | Jun 15, 2018 |

ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDIA

+ SB PHARMCO
+

EQ 2MG BASE
EQ 4MG BASE

N021071 002 May 25, 1999
N021071 003 May 25, 1999

ROSUVASTATIN CALCIUM

CAPSULE; ORAL

EZALLOR

+ SUN PHARMA GLOBAL
+
+
+!

EQ 5MG BASE
EQ 10MG BASE
EQ 20MG BASE
EQ 40MG BASE

N208647 001 Dec 18, 2018
N208647 002 Dec 18, 2018
N208647 003 Dec 18, 2018
N208647 004 Dec 18, 2018

TABLET; ORAL

CRESTOR

| | | | | | |
|-----------|-------|-------------|----------------|------------|--------------|
| <u>AB</u> | + IPR | <u>5MG</u> | <u>N021366</u> | <u>002</u> | Aug 12, 2003 |
| <u>AB</u> | + | <u>10MG</u> | <u>N021366</u> | <u>003</u> | Aug 12, 2003 |
| <u>AB</u> | + | <u>20MG</u> | <u>N021366</u> | <u>004</u> | Aug 12, 2003 |
| <u>AB</u> | +! | <u>40MG</u> | <u>N021366</u> | <u>005</u> | Aug 12, 2003 |

ROSUVASTATIN CALCIUM

| | | | | | |
|-----------|------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>5MG</u> | <u>A206434</u> | <u>001</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A206434</u> | <u>002</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A206434</u> | <u>003</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A206434</u> | <u>004</u> | Oct 31, 2016 |
| <u>AB</u> | ALKEM LABS LTD | <u>5MG</u> | <u>A206465</u> | <u>001</u> | Mar 21, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A206465</u> | <u>002</u> | Mar 21, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A206465</u> | <u>003</u> | Mar 21, 2017 |
| <u>AB</u> | | <u>40MG</u> | <u>A206465</u> | <u>004</u> | Mar 21, 2017 |
| <u>AB</u> | ALLIED | <u>5MG</u> | <u>A079168</u> | <u>001</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A079168</u> | <u>002</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A079168</u> | <u>003</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A079168</u> | <u>004</u> | Jul 19, 2016 |
| <u>AB</u> | AMNEAL PHARMS CO | <u>5MG</u> | <u>A208850</u> | <u>001</u> | Oct 16, 2018 |
| <u>AB</u> | | <u>10MG</u> | <u>A208850</u> | <u>002</u> | Oct 16, 2018 |
| <u>AB</u> | | <u>20MG</u> | <u>A208850</u> | <u>003</u> | Oct 16, 2018 |
| <u>AB</u> | | <u>40MG</u> | <u>A208850</u> | <u>004</u> | Oct 16, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

ROSUVASTATIN CALCIUM

TABLET; ORAL

ROSUVASTATIN CALCIUM

| | | | | |
|-----------|----------------------|-------------|--------------------|--------------|
| <u>AB</u> | APOTEX INC | <u>5MG</u> | <u>A079145 001</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A079145 002</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A079145 003</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A079145 004</u> | Jul 19, 2016 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>5MG</u> | <u>A079170 001</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A079170 002</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A079170 003</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A079170 004</u> | Jul 19, 2016 |
| <u>AB</u> | BIOCON LTD | <u>5MG</u> | <u>A207752 001</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A207752 002</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A207752 003</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A207752 004</u> | Oct 31, 2016 |
| <u>AB</u> | CADILA PHARMS LTD | <u>5MG</u> | <u>A207453 001</u> | Nov 23, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A207453 002</u> | Nov 23, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A207453 003</u> | Nov 23, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A207453 004</u> | Nov 23, 2016 |
| <u>AB</u> | CHANGZHOU PHARM | <u>5MG</u> | <u>A207408 001</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A207408 002</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A207408 003</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A207408 004</u> | Oct 31, 2016 |
| <u>AB</u> | GLENMARK PHARMS | <u>5MG</u> | <u>A079172 001</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A079172 002</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A079172 003</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A079172 004</u> | Jul 19, 2016 |
| <u>AB</u> | HETERO LABS LTD V | <u>5MG</u> | <u>A207616 001</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A207616 002</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A207616 003</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A207616 004</u> | Oct 31, 2016 |
| <u>AB</u> | JUBILANT GENERICS | <u>5MG</u> | <u>A207062 001</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A207062 002</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A207062 003</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A207062 004</u> | Oct 31, 2016 |
| <u>AB</u> | LUPIN LTD | <u>5MG</u> | <u>A205587 001</u> | Jul 31, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A205587 002</u> | Jul 31, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A205587 003</u> | Jul 31, 2017 |
| <u>AB</u> | | <u>40MG</u> | <u>A205587 004</u> | Jul 31, 2017 |
| <u>AB</u> | MSN LABS PVT LTD | <u>5MG</u> | <u>A208898 001</u> | Nov 22, 2017 |
| <u>AB</u> | | <u>10MG</u> | <u>A208898 002</u> | Nov 22, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A208898 003</u> | Nov 22, 2017 |
| <u>AB</u> | | <u>40MG</u> | <u>A208898 004</u> | Nov 22, 2017 |
| <u>AB</u> | MYLAN PHARMS INC | <u>5MG</u> | <u>A079161 001</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A079161 002</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A079161 003</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A079161 004</u> | Jul 19, 2016 |
| <u>AB</u> | SANDOZ INC | <u>5MG</u> | <u>A079171 001</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A079171 002</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A079171 003</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A079171 004</u> | Jul 19, 2016 |
| <u>AB</u> | SUN PHARMA GLOBAL | <u>5MG</u> | <u>A079169 001</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A079169 002</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A079169 003</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A079169 004</u> | Jul 19, 2016 |
| <u>AB</u> | TEVA PHARMS USA | <u>5MG</u> | <u>A079166 001</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A079166 002</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A079166 003</u> | Jul 19, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A079166 004</u> | Jul 19, 2016 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>5MG</u> | <u>A201619 001</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A201619 002</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A201619 003</u> | Oct 31, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A201619 004</u> | Oct 31, 2016 |
| <u>AB</u> | WATSON LABS INC | <u>5MG</u> | <u>A079167 001</u> | Apr 29, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A079167 002</u> | Apr 29, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A079167 003</u> | Apr 29, 2016 |
| <u>AB</u> | | <u>40MG</u> | <u>A079167 004</u> | Apr 29, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

ROTIGOTINE

FILM, EXTENDED RELEASE;TRANSDERMAL

NEUPRO

| | | | | | |
|---|---------|----------|---------|-----|--------------|
| + | UCB INC | 1MG/24HR | N021829 | 004 | Apr 02, 2012 |
| + | ! | 2MG/24HR | N021829 | 001 | May 09, 2007 |
| + | | 3MG/24HR | N021829 | 005 | Apr 02, 2012 |
| + | | 4MG/24HR | N021829 | 002 | May 09, 2007 |
| + | | 6MG/24HR | N021829 | 003 | May 09, 2007 |
| + | | 8MG/24HR | N021829 | 006 | Apr 02, 2012 |

RUBIDIUM CHLORIDE RB-82

INJECTABLE; INJECTION

CARDIOGEN-82

BRACCO

N/A

N019414 001 Dec 29, 1989

SOLUTION; INTRAVENOUS

RUBY-FILL

JUBILANT DRAXIMAGE

N/A

N202153 001 Sep 30, 2016

RUCAPARIB CAMSYLATE

TABLET; ORAL

RUBRACA

| | | | | | |
|---|---------------------|---------------|---------|-----|--------------|
| + | CLOVIS ONCOLOGY INC | EQ 200MG BASE | N209115 | 001 | Dec 19, 2016 |
| + | | EQ 250MG BASE | N209115 | 003 | May 01, 2017 |
| + | ! | EQ 300MG BASE | N209115 | 002 | Dec 19, 2016 |

RUFINAMIDE

SUSPENSION; ORAL

BANZEL

| | | | | | |
|---|-----------|---------|---------|-----|--------------|
| + | EISAI INC | 40MG/ML | N201367 | 001 | Mar 03, 2011 |
|---|-----------|---------|---------|-----|--------------|

TABLET; ORAL

BANZEL

| | | | | | |
|-----------|---|-----------|--------------|--------------------|--------------|
| AB | + | EISAI INC | 200MG | N021911 002 | Nov 14, 2008 |
| AB | + | ! | 400MG | N021911 003 | Nov 14, 2008 |

RUFINAMIDE

| | | | | | |
|-----------|--|-------------------------|--------------|--------------------|--------------|
| AB | | GLENMARK PHARMS LTD | 200MG | A205075 001 | May 16, 2016 |
| AB | | | 400MG | A205075 002 | May 16, 2016 |
| AB | | MYLAN PHARMS INC | 200MG | A205095 001 | May 16, 2016 |
| AB | | | 400MG | A205095 002 | May 16, 2016 |
| AB | | WEST-WARD PHARMS INT | 200MG | A204988 001 | May 16, 2016 |
| AB | | | 400MG | A204988 002 | May 16, 2016 |

RUXOLITINIB PHOSPHATE

TABLET; ORAL

JAKAFI

| | | | | | |
|---|-------------|--------------|---------|-----|--------------|
| + | INCYTE CORP | EQ 5MG BASE | N202192 | 001 | Nov 16, 2011 |
| + | | EQ 10MG BASE | N202192 | 002 | Nov 16, 2011 |
| + | | EQ 15MG BASE | N202192 | 003 | Nov 16, 2011 |
| + | | EQ 20MG BASE | N202192 | 004 | Nov 16, 2011 |
| + | ! | EQ 25MG BASE | N202192 | 005 | Nov 16, 2011 |

SACROSIDASE

SOLUTION; ORAL

SUCRAID

| | | | | | |
|---|-----------|-------------|---------|-----|--------------|
| + | QOL MEDCL | 8,500 IU/ML | N020772 | 001 | Apr 09, 1998 |
|---|-----------|-------------|---------|-----|--------------|

SACUBITRIL; VALSARTAN

TABLET; ORAL

ENTRESTO

| | | | | | |
|---|-------------------------|------------|---------|-----|--------------|
| + | NOVARTIS PHARMS CORP | 24MG;26MG | N207620 | 001 | Jul 07, 2015 |
| + | | 49MG;51MG | N207620 | 002 | Jul 07, 2015 |
| + | ! | 97MG;103MG | N207620 | 003 | Jul 07, 2015 |

SAFINAMIDE MESYLATE

TABLET; ORAL

XADAGO

| | | | | | |
|---|------------------|-------|---------|-----|--------------|
| + | US WORLDMEDS LLC | 50MG | N207145 | 001 | Mar 21, 2017 |
| + | ! | 100MG | N207145 | 002 | Mar 21, 2017 |

SALMETEROL XINAFOATE

POWDER; INHALATION

SEREVENT

| | | | | | |
|---|-----------------|--------------------|---------|-----|--------------|
| + | GLAXOSMITHKLINE | EQ 0.05MG BASE/INH | N020692 | 001 | Sep 19, 1997 |
|---|-----------------|--------------------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

SAMARIUM SM-153 LEXIDRONAM PENTASODIUM

INJECTABLE; INJECTION

QUADRAMET

+! LANTHEUS MEDICAL 50mCi/ML N020570 001 Mar 28, 1997

SAPROPTERIN DIHYDROCHLORIDE

POWDER; ORAL

KUVAN

+! BIOMARIN PHARM 100MG/PACKET N205065 001 Dec 19, 2013

+ 500MG/PACKET N205065 002 Oct 27, 2015

TABLET; ORAL

KUVAN

+! BIOMARIN PHARM 100MG N022181 001 Dec 13, 2007

SAQUINAVIR MESYLATE

TABLET; ORAL

INVIRASE

+! HOFFMANN-LA ROCHE EQ 500MG BASE N021785 001 Dec 17, 2004

SARECYCLINE HYDROCHLORIDE

TABLET; ORAL

SEYSARA

+ ALMIRALL EQ 60MG BASE N209521 001 Oct 01, 2018

+ EQ 100MG BASE N209521 002 Oct 01, 2018

+! EQ 150MG BASE N209521 003 Oct 01, 2018

SAXAGLIPTIN HYDROCHLORIDE

TABLET; ORAL

ONGLYZA

+ ASTRAZENECA AB EQ 2.5MG BASE N022350 001 Jul 31, 2009

+! EQ 5MG BASE N022350 002 Jul 31, 2009

SCOPOLAMINE

FILM, EXTENDED RELEASE; TRANSDERMAL

SCOPOLAMINE**AB** PERRIGO PHARMS CO **1MG/72HR** **A078830 001** Jan 30, 2015TRANSDERM SCOP**AB** +! GLAXOSMITHKLINE CON **1MG/72HR** **N017874 001**SECNIDAZOLE

GRANULE; ORAL

SOLOSEC

+! LUPIN 2GM/PACKET N209363 001 Sep 15, 2017

SECOBARBITAL SODIUM

CAPSULE; ORAL

SECONAL SODIUM

! VALEANT PHARMS 50MG A086101 001 Oct 03, 1983

! NORTH 100MG A086101 002 Oct 03, 1983

SECRETIN SYNTHETIC HUMAN

FOR SOLUTION; INTRAVENOUS

CHIRHOSTIM

+! CHIRHOCLIN 16MCG/VIAL N021256 001 Apr 09, 2004

+ 40MCG/VIAL N021256 002 Jun 21, 2007

SELEGILINE

FILM, EXTENDED RELEASE; TRANSDERMAL

EMSAM

+! SOMERSET 6MG/24HR N021336 001 Feb 27, 2006

+ 9MG/24HR N021336 002 Feb 27, 2006

+ 12MG/24HR N021336 003 Feb 27, 2006

SELEGILINE HYDROCHLORIDE

CAPSULE; ORAL

SELEGILINE HYDROCHLORIDE**AB** ! APOTEX **5MG** **A075321 001** Dec 04, 1998**AB** DAVA PHARMS INC **5MG** **A075352 001** Nov 30, 1998

TABLET; ORAL

SELEGILINE HYDROCHLORIDE**AB** ! APOTEX INC **5MG** **A074871 001** Jun 06, 1997**AB** BOSCOGEN **5MG** **A074912 001** Apr 30, 1998**AB** MYLAN **5MG** **A074866 001** Nov 26, 1997

TABLET, ORALLY DISINTEGRATING; ORAL

ZELAPAR

+! VALEANT PHARM INTL 1.25MG N021479 001 Jun 14, 2006

PRESCRIPTION DRUG PRODUCT LIST

SELENIUM SULFIDE

LOTION;SHAMPOO;TOPICAL

SELENIUM SULFIDE

| | | | | | | |
|-----------|---|------------------|-------------|----------------|------------|--------------|
| <u>AT</u> | ! | PERRIGO NEW YORK | <u>2.5%</u> | <u>A089996</u> | <u>001</u> | Jan 10, 1991 |
| <u>AT</u> | | WOCKHARDT BIO AG | <u>2.5%</u> | <u>A088228</u> | <u>001</u> | Sep 01, 1983 |

SELEXIPAG

TABLET;ORAL

UPTRAVI

| | | | | | |
|---|---------------------|-------|---------|-----|--------------|
| + | ACTELION PHARMS LTD | 0.2MG | N207947 | 001 | Dec 21, 2015 |
| + | | 0.4MG | N207947 | 002 | Dec 21, 2015 |
| + | | 0.6MG | N207947 | 003 | Dec 21, 2015 |
| + | | 0.8MG | N207947 | 004 | Dec 21, 2015 |
| + | | 1MG | N207947 | 005 | Dec 21, 2015 |
| + | | 1.2MG | N207947 | 006 | Dec 21, 2015 |
| + | | 1.4MG | N207947 | 007 | Dec 21, 2015 |
| + | ! | 1.6MG | N207947 | 008 | Dec 21, 2015 |

SEMAGLUTIDE

SOLUTION;SUBCUTANEOUS

OZEMPIC

| | | | | | | |
|---|---|------|-----------------------|---------|-----|--------------|
| + | ! | NOVO | 2MG/1.5ML (1.34MG/ML) | N209637 | 001 | Dec 05, 2017 |
|---|---|------|-----------------------|---------|-----|--------------|

SERTACONAZOLE NITRATE

CREAM;TOPICAL

ERTACZO

| | | | | | | |
|---|---|--------------------|----|---------|-----|--------------|
| + | ! | VALEANT LUXEMBOURG | 2% | N021385 | 001 | Dec 10, 2003 |
|---|---|--------------------|----|---------|-----|--------------|

SERTRALINE HYDROCHLORIDE

CONCENTRATE;ORAL

SERTRALINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|------------------|------------------------|----------------|------------|--------------|
| <u>AA</u> | | AUROBINDO PHARMA | <u>EQ 20MG BASE/ML</u> | <u>A078861</u> | <u>001</u> | Oct 31, 2008 |
|-----------|--|------------------|------------------------|----------------|------------|--------------|

ZOLOFT

| | | | | | | | |
|-----------|---|---|--------|------------------------|----------------|------------|--------------|
| <u>AA</u> | + | ! | PFIZER | <u>EQ 20MG BASE/ML</u> | <u>N020990</u> | <u>001</u> | Dec 07, 1999 |
|-----------|---|---|--------|------------------------|----------------|------------|--------------|

TABLET;ORAL

SERTRALINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|--------------------|----------------------|----------------|------------|--------------|
| <u>AB</u> | | ACCORD HLTHCARE | <u>EQ 25MG BASE</u> | <u>A202825</u> | <u>001</u> | Nov 07, 2014 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A202825</u> | <u>002</u> | Nov 07, 2014 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A202825</u> | <u>003</u> | Nov 07, 2014 |
| <u>AB</u> | | APOTEX INC | <u>EQ 25MG BASE</u> | <u>A076882</u> | <u>001</u> | Feb 06, 2007 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A076882</u> | <u>002</u> | Feb 06, 2007 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A076882</u> | <u>003</u> | Feb 06, 2007 |
| <u>AB</u> | | AUROBINDO PHARMA | <u>EQ 25MG BASE</u> | <u>A077206</u> | <u>001</u> | Feb 06, 2007 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A077206</u> | <u>002</u> | Feb 06, 2007 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A077206</u> | <u>003</u> | Feb 06, 2007 |
| <u>AB</u> | | AUSTARPHARMA LLC | <u>EQ 25MG BASE</u> | <u>A078677</u> | <u>001</u> | Mar 04, 2009 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A078677</u> | <u>002</u> | Mar 04, 2009 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A078677</u> | <u>003</u> | Mar 04, 2009 |
| <u>AB</u> | | INVAGEN PHARMS | <u>EQ 25MG BASE</u> | <u>A077397</u> | <u>001</u> | Feb 06, 2007 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A077397</u> | <u>002</u> | Feb 06, 2007 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A077397</u> | <u>003</u> | Feb 06, 2007 |
| <u>AB</u> | | LUPIN | <u>EQ 25MG BASE</u> | <u>A077670</u> | <u>001</u> | Feb 06, 2007 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A077670</u> | <u>002</u> | Feb 06, 2007 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A077670</u> | <u>003</u> | Feb 06, 2007 |
| <u>AB</u> | | MYLAN PHARMS INC | <u>EQ 25MG BASE</u> | <u>A078626</u> | <u>001</u> | Jan 31, 2008 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A078626</u> | <u>002</u> | Jan 31, 2008 |
| <u>AB</u> | | OXFORD PHARMS | <u>EQ 25MG BASE</u> | <u>A078175</u> | <u>001</u> | Jul 21, 2010 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A078175</u> | <u>002</u> | Jul 21, 2010 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A078175</u> | <u>003</u> | Jul 21, 2010 |
| <u>AB</u> | | SUN PHARM INDS LTD | <u>EQ 25MG BASE</u> | <u>A077977</u> | <u>001</u> | Feb 06, 2007 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A077977</u> | <u>002</u> | Feb 06, 2007 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A077977</u> | <u>003</u> | Feb 06, 2007 |
| <u>AB</u> | | TEVA | <u>EQ 25MG BASE</u> | <u>A076465</u> | <u>001</u> | Aug 11, 2006 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A076465</u> | <u>002</u> | Aug 11, 2006 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A076465</u> | <u>003</u> | Aug 11, 2006 |
| <u>AB</u> | | TORRENT PHARMS | <u>EQ 25MG BASE</u> | <u>A077765</u> | <u>001</u> | Feb 06, 2007 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A077765</u> | <u>002</u> | Feb 06, 2007 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A077765</u> | <u>003</u> | Feb 06, 2007 |
| <u>AB</u> | | WOCKHARDT | <u>EQ 25MG BASE</u> | <u>A078403</u> | <u>001</u> | Jan 08, 2008 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A078403</u> | <u>002</u> | Jan 08, 2008 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A078403</u> | <u>003</u> | Jan 08, 2008 |
| <u>AB</u> | | ZYDUS PHARMS USA | <u>EQ 25MG BASE</u> | <u>A077106</u> | <u>001</u> | Feb 06, 2007 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A077106</u> | <u>002</u> | Feb 06, 2007 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A077106</u> | <u>003</u> | Feb 06, 2007 |

PRESCRIPTION DRUG PRODUCT LIST

SERTRALINE HYDROCHLORIDE

TABLET; ORAL

ZOLOFT

| | | | | | |
|-----------|---|--------|----------------------|--------------------|--------------|
| <u>AB</u> | + | PFIZER | <u>EQ 25MG BASE</u> | <u>N019839 005</u> | Mar 06, 1996 |
| <u>AB</u> | + | | <u>EQ 50MG BASE</u> | <u>N019839 001</u> | Dec 30, 1991 |
| <u>AB</u> | + | ! | <u>EQ 100MG BASE</u> | <u>N019839 002</u> | Dec 30, 1991 |

SERTRALINE HYDROCHLORIDE

SUN PHARM INDS LTD

EQ 150MG BASE

A077977 004 Feb 06, 2007

EQ 200MG BASE

A077977 005 Feb 06, 2007

SEVELAMER CARBONATE

FOR SUSPENSION; ORAL

RENVELA

| | | | | | |
|-----------|---|---------|---------------------|--------------------|--------------|
| <u>AB</u> | + | GENZYME | <u>800MG/PACKET</u> | <u>N022318 001</u> | Aug 12, 2009 |
| <u>AB</u> | + | ! | <u>2.4GM/PACKET</u> | <u>N022318 002</u> | Feb 18, 2009 |

SEVELAMER CARBONATE

| | | | | | |
|-----------|--|----------------------|---------------------|--------------------|--------------|
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>800MG/PACKET</u> | <u>A207624 001</u> | Jun 13, 2017 |
| <u>AB</u> | | | <u>2.4GM/PACKET</u> | <u>A207624 002</u> | Jun 13, 2017 |
| <u>AB</u> | | DR REDDYS LABS LTD | <u>800MG/PACKET</u> | <u>A210464 001</u> | Oct 25, 2018 |
| <u>AB</u> | | | <u>2.4GM/PACKET</u> | <u>A210464 002</u> | Oct 25, 2018 |

TABLET; ORAL

RENVELA

| | | | | | |
|-----------|---|--------|--------------|--------------------|--------------|
| <u>AB</u> | + | SANOFI | <u>800MG</u> | <u>N022127 001</u> | Oct 19, 2007 |
|-----------|---|--------|--------------|--------------------|--------------|

SEVELAMER CARBONATE

| | | | | | |
|-----------|--|----------------------|--------------|--------------------|--------------|
| <u>AB</u> | | AMNEAL PHARMS CO | <u>800MG</u> | <u>A207288 001</u> | Nov 28, 2017 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>800MG</u> | <u>A207179 001</u> | Jul 17, 2017 |
| <u>AB</u> | | DR REDDYS LABS LTD | <u>800MG</u> | <u>A206094 001</u> | Sep 29, 2017 |
| <u>AB</u> | | IMPAX LABS INC | <u>800MG</u> | <u>A090975 001</u> | Oct 23, 2017 |
| <u>AB</u> | | INVAGEN PHARMS | <u>800MG</u> | <u>A203860 001</u> | Oct 26, 2017 |
| <u>AB</u> | | TWI PHARMS | <u>800MG</u> | <u>A200959 001</u> | Mar 20, 2018 |
| <u>AB</u> | | WILSHIRE PHARMS INC | <u>800MG</u> | <u>A204451 001</u> | Nov 29, 2018 |

SEVELAMER HYDROCHLORIDE

TABLET; ORAL

RENAGEL

| | | | | | |
|--|---|---------|-------|-------------|--------------|
| | + | GENZYME | 400MG | N021179 001 | Jul 12, 2000 |
| | + | ! | 800MG | N021179 002 | Jul 12, 2000 |

SEVOFLURANE

LIQUID; INHALATION

SEVOFLURANE

| | | | | | |
|-----------|--|------------------|-------------|--------------------|--------------|
| <u>AN</u> | | BAXTER HLTHCARE | <u>100%</u> | <u>A075895 001</u> | Jul 02, 2002 |
| <u>AN</u> | | HALOCARBON PRODS | <u>100%</u> | <u>A078650 001</u> | Nov 19, 2007 |
| <u>AN</u> | | SHANGHAI HENGRUI | <u>100%</u> | <u>A203793 001</u> | Nov 03, 2015 |

SOJOURN

| | | | | | |
|-----------|--|------------------|-------------|--------------------|--------------|
| <u>AN</u> | | PIRAMAL CRITICAL | <u>100%</u> | <u>A077867 001</u> | May 02, 2007 |
|-----------|--|------------------|-------------|--------------------|--------------|

ULTANE

| | | | | | | |
|-----------|---|---|--------|-------------|--------------------|--------------|
| <u>AN</u> | + | ! | ABEVIE | <u>100%</u> | <u>N020478 001</u> | Jun 07, 1995 |
|-----------|---|---|--------|-------------|--------------------|--------------|

SILDENAFIL CITRATE

FOR SUSPENSION; ORAL

REVATIO

| | | | | | |
|--|---|--------|-----------------|-------------|--------------|
| | + | PFIZER | EQ 10MG BASE/ML | N203109 001 | Aug 30, 2012 |
|--|---|--------|-----------------|-------------|--------------|

SOLUTION; INTRAVENOUS

REVATIO

| | | | | | | |
|-----------|---|---|--------|-----------------------------------------------|--------------------|--------------|
| <u>AP</u> | + | ! | PFIZER | <u>EQ 10MG BASE/12.5ML (EQ 0.8MG BASE/ML)</u> | <u>N022473 001</u> | Nov 18, 2009 |
|-----------|---|---|--------|-----------------------------------------------|--------------------|--------------|

SILDENAFIL CITRATE

| | | | | | |
|-----------|--|----------------------|-----------------------------------------------|--------------------|--------------|
| <u>AP</u> | | AUROBINDO PHARMA LTD | <u>EQ 10MG BASE/12.5ML (EQ 0.8MG BASE/ML)</u> | <u>A203988 001</u> | Apr 01, 2015 |
|-----------|--|----------------------|-----------------------------------------------|--------------------|--------------|

TABLET; ORAL

REVATIO

| | | | | | | |
|-----------|---|---|--------|---------------------|--------------------|--------------|
| <u>AB</u> | + | ! | PFIZER | <u>EQ 20MG BASE</u> | <u>N021845 001</u> | Jun 03, 2005 |
|-----------|---|---|--------|---------------------|--------------------|--------------|

SILDENAFIL CITRATE

| | | | | | |
|-----------|--|----------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | AJANTA PHARMA LTD | <u>EQ 20MG BASE</u> | <u>A210394 001</u> | May 04, 2018 |
| <u>AB</u> | | | <u>EQ 25MG BASE</u> | <u>A206401 001</u> | Oct 12, 2018 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A206401 002</u> | Oct 12, 2018 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A206401 003</u> | Oct 12, 2018 |
| <u>AB</u> | | AMNEAL PHARMS | <u>EQ 20MG BASE</u> | <u>A202025 001</u> | Feb 28, 2013 |
| <u>AB</u> | | AMNEAL PHARMS NY | <u>EQ 25MG BASE</u> | <u>A202023 001</u> | Jun 27, 2018 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A202023 002</u> | Jun 27, 2018 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A202023 003</u> | Jun 27, 2018 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>EQ 20MG BASE</u> | <u>A203963 001</u> | Nov 18, 2015 |

PRESCRIPTION DRUG PRODUCT LIST

SILDENAFIL CITRATE

TABLET;ORAL

SILDENAFIL CITRATE

| | | | | |
|---------------|---------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 25MG BASE</u> | <u>A203962 001</u> | Jun 11, 2018 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A203962 002</u> | Jun 11, 2018 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A203962 003</u> | Jun 11, 2018 |
| <u>AB</u> | HEBEI CHANGSHAN | <u>EQ 20MG BASE</u> | <u>A202598 001</u> | Nov 06, 2012 |
| <u>AB</u> | HETERO LABS LTD V | <u>EQ 20MG BASE</u> | <u>A203623 001</u> | Nov 26, 2014 |
| <u>AB</u> | | <u>EQ 25MG BASE</u> | <u>A202659 001</u> | Jun 11, 2018 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A202659 002</u> | Jun 11, 2018 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A202659 003</u> | Jun 11, 2018 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>EQ 20MG BASE</u> | <u>A203814 001</u> | Dec 17, 2013 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 20MG BASE</u> | <u>A201150 001</u> | Nov 09, 2012 |
| <u>AB</u> | RUBICON RES PVT LTD | <u>EQ 20MG BASE</u> | <u>A204883 001</u> | Jun 20, 2016 |
| <u>AB</u> | | <u>EQ 25MG BASE</u> | <u>A204882 001</u> | Jun 11, 2018 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A204882 002</u> | Jun 11, 2018 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A204882 003</u> | Jun 11, 2018 |
| <u>AB</u> | TEVA | <u>EQ 25MG BASE</u> | <u>A077342 001</u> | Mar 09, 2016 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A077342 002</u> | Mar 09, 2016 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A077342 003</u> | Mar 09, 2016 |
| <u>AB</u> | TEVA PHARMS | <u>EQ 20MG BASE</u> | <u>A078380 001</u> | Jan 07, 2013 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>EQ 20MG BASE</u> | <u>A091479 001</u> | Nov 06, 2012 |
| <u>AB</u> | | <u>EQ 25MG BASE</u> | <u>A091448 001</u> | Jun 11, 2018 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A091448 002</u> | Jun 11, 2018 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A091448 003</u> | Jun 11, 2018 |
| <u>AB</u> | WATSON LABS INC | <u>EQ 20MG BASE</u> | <u>A202503 001</u> | Nov 06, 2012 |
| <u>VIAGRA</u> | | | | |
| <u>AB</u> | + | <u>EQ 25MG BASE</u> | <u>N020895 001</u> | Mar 27, 1998 |
| <u>AB</u> | + | <u>EQ 50MG BASE</u> | <u>N020895 002</u> | Mar 27, 1998 |
| <u>AB</u> | + | <u>EQ 100MG BASE</u> | <u>N020895 003</u> | Mar 27, 1998 |

SILODOSIN

CAPSULE;ORAL

RAPAFLO

| | | | | | |
|-----------|---|--------------------|------------|--------------------|--------------|
| <u>AB</u> | + | ALLERGAN SALES LLC | <u>4MG</u> | <u>N022206 001</u> | Oct 08, 2008 |
| <u>AB</u> | + | | <u>8MG</u> | <u>N022206 002</u> | Oct 08, 2008 |

SILODOSIN

| | | | | | |
|-----------|--|----------------------|------------|--------------------|--------------|
| <u>AB</u> | | AJANTA PHARMA LTD | <u>4MG</u> | <u>A211060 001</u> | Dec 03, 2018 |
| <u>AB</u> | | | <u>8MG</u> | <u>A211060 002</u> | Dec 03, 2018 |
| <u>AB</u> | | AMNEAL PHARMS CO | <u>4MG</u> | <u>A209745 001</u> | Dec 03, 2018 |
| <u>AB</u> | | | <u>8MG</u> | <u>A209745 002</u> | Dec 03, 2018 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>4MG</u> | <u>A210626 001</u> | Dec 10, 2018 |
| <u>AB</u> | | | <u>8MG</u> | <u>A210626 002</u> | Dec 10, 2018 |
| <u>AB</u> | | LUPIN LTD | <u>4MG</u> | <u>A206541 001</u> | Dec 03, 2018 |
| <u>AB</u> | | | <u>8MG</u> | <u>A206541 002</u> | Dec 03, 2018 |
| <u>AB</u> | | MACLEODS PHARMS LTD | <u>4MG</u> | <u>A211166 001</u> | Dec 03, 2018 |
| <u>AB</u> | | | <u>8MG</u> | <u>A211166 002</u> | Dec 03, 2018 |
| <u>AB</u> | | MSN LABS PVT LTD | <u>4MG</u> | <u>A210687 001</u> | Dec 03, 2018 |
| <u>AB</u> | | | <u>8MG</u> | <u>A210687 002</u> | Dec 03, 2018 |
| <u>AB</u> | | SANDOZ INC | <u>4MG</u> | <u>A204726 001</u> | Mar 31, 2017 |
| <u>AB</u> | | | <u>8MG</u> | <u>A204726 002</u> | Mar 31, 2017 |

SILVER SULFADIAZINE

CREAM;TOPICAL

SILVADENE

| | | | | | |
|-----------|---|-----------------|-----------|--------------------|--|
| <u>AB</u> | + | KING PHARMS LLC | <u>1%</u> | <u>N017381 001</u> | |
|-----------|---|-----------------|-----------|--------------------|--|

SSD

| | | | | | |
|-----------|--|--------------|-----------|--------------------|--------------|
| <u>AB</u> | | DR REDDYS LA | <u>1%</u> | <u>N018578 001</u> | Feb 25, 1982 |
|-----------|--|--------------|-----------|--------------------|--------------|

THERMAZENE

| | | | | | |
|-----------|--|----------------------|-----------|--------------------|--------------|
| <u>AB</u> | | THEPHARMANETWORK LLC | <u>1%</u> | <u>N018810 001</u> | Dec 23, 1985 |
|-----------|--|----------------------|-----------|--------------------|--------------|

SIMVASTATIN

SUSPENSION;ORAL

FLOLIPID

| | | | | |
|---|-------------------|----------|-------------|--------------|
| + | TCG FLUENT PHARMA | 20MG/5ML | N206679 001 | Apr 21, 2016 |
| + | | 40MG/5ML | N206679 002 | Apr 21, 2016 |

TABLET;ORAL

SIMVASTATIN

| | | | | |
|-----------|-----------------|-------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>5MG</u> | <u>A078155 005</u> | Apr 05, 2013 |
| <u>AB</u> | | <u>10MG</u> | <u>A078155 002</u> | Feb 26, 2008 |
| <u>AB</u> | | <u>20MG</u> | <u>A078155 003</u> | Feb 26, 2008 |
| <u>AB</u> | | <u>40MG</u> | <u>A078155 004</u> | Feb 26, 2008 |

PRESCRIPTION DRUG PRODUCT LIST

SIMVASTATIN

TABLET; ORAL

SIMVASTATIN

| | | | | |
|-----------|---------------------|-------------|--------------------|--------------|
| <u>AB</u> | | <u>80MG</u> | <u>A078155 001</u> | Feb 26, 2008 |
| <u>AB</u> | AUROBINDO PHARMA | <u>5MG</u> | <u>A077691 001</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>10MG</u> | <u>A077691 002</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>20MG</u> | <u>A077691 003</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>40MG</u> | <u>A077691 004</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>80MG</u> | <u>A077691 005</u> | Dec 20, 2006 |
| <u>AB</u> | BIOCON LIMITED | <u>5MG</u> | <u>A078034 001</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>10MG</u> | <u>A078034 002</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>20MG</u> | <u>A078034 003</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>40MG</u> | <u>A078034 004</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>80MG</u> | <u>A078034 005</u> | Dec 20, 2006 |
| <u>AB</u> | DR REDDYS LABS INC | <u>5MG</u> | <u>A077752 005</u> | Jan 23, 2008 |
| <u>AB</u> | | <u>10MG</u> | <u>A077752 001</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>20MG</u> | <u>A077752 002</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>40MG</u> | <u>A077752 003</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>80MG</u> | <u>A077752 004</u> | Dec 20, 2006 |
| <u>AB</u> | HETERO LABS LTD III | <u>5MG</u> | <u>A200895 001</u> | Nov 25, 2014 |
| <u>AB</u> | | <u>10MG</u> | <u>A200895 002</u> | Nov 25, 2014 |
| <u>AB</u> | | <u>20MG</u> | <u>A200895 003</u> | Nov 25, 2014 |
| <u>AB</u> | | <u>40MG</u> | <u>A200895 004</u> | Nov 25, 2014 |
| <u>AB</u> | | <u>80MG</u> | <u>A200895 005</u> | Nov 25, 2014 |
| <u>AB</u> | LUPIN | <u>5MG</u> | <u>A078103 005</u> | Apr 14, 2009 |
| <u>AB</u> | | <u>10MG</u> | <u>A078103 001</u> | May 11, 2007 |
| <u>AB</u> | | <u>20MG</u> | <u>A078103 002</u> | May 11, 2007 |
| <u>AB</u> | | <u>40MG</u> | <u>A078103 003</u> | May 11, 2007 |
| <u>AB</u> | | <u>80MG</u> | <u>A078103 004</u> | May 11, 2007 |
| <u>AB</u> | OXFORD PHARMS | <u>5MG</u> | <u>A078735 001</u> | Aug 30, 2010 |
| <u>AB</u> | | <u>10MG</u> | <u>A078735 002</u> | Aug 30, 2010 |
| <u>AB</u> | | <u>20MG</u> | <u>A078735 003</u> | Aug 30, 2010 |
| <u>AB</u> | | <u>40MG</u> | <u>A078735 004</u> | Aug 30, 2010 |
| <u>AB</u> | | <u>80MG</u> | <u>A078735 005</u> | Aug 30, 2010 |
| <u>AB</u> | VIVA HLTHCARE | <u>5MG</u> | <u>A090383 001</u> | Sep 16, 2011 |
| <u>AB</u> | | <u>10MG</u> | <u>A090383 002</u> | Sep 16, 2011 |
| <u>AB</u> | | <u>20MG</u> | <u>A090383 003</u> | Sep 16, 2011 |
| <u>AB</u> | | <u>40MG</u> | <u>A090383 004</u> | Sep 16, 2011 |
| <u>AB</u> | | <u>80MG</u> | <u>A090383 005</u> | Sep 16, 2011 |
| <u>AB</u> | WATSON LABS TEVA | <u>5MG</u> | <u>A076685 001</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>10MG</u> | <u>A076685 002</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>20MG</u> | <u>A076685 003</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>40MG</u> | <u>A076685 004</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>80MG</u> | <u>A076685 005</u> | Dec 20, 2006 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>5MG</u> | <u>A077837 001</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>10MG</u> | <u>A077837 002</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>20MG</u> | <u>A077837 003</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>40MG</u> | <u>A077837 004</u> | Dec 20, 2006 |
| <u>AB</u> | | <u>80MG</u> | <u>A077837 005</u> | Dec 20, 2006 |

ZOCOR

| | | | | | |
|-----------|---|--------------|-------------|--------------------|--------------|
| <u>AB</u> | + | MSD MERCK CO | <u>5MG</u> | <u>N019766 001</u> | Dec 23, 1991 |
| <u>AB</u> | + | | <u>10MG</u> | <u>N019766 002</u> | Dec 23, 1991 |
| <u>AB</u> | + | | <u>20MG</u> | <u>N019766 003</u> | Dec 23, 1991 |
| <u>AB</u> | + | | <u>40MG</u> | <u>N019766 004</u> | Dec 23, 1991 |
| <u>AB</u> | + | | <u>80MG</u> | <u>N019766 005</u> | Jul 10, 1998 |

SINCALIDE

INJECTABLE; INJECTION

KINEVAC

+! BRACCO

0.005MG/VIAL

N017697 001

SINECATECHINS

OINTMENT; TOPICAL

VEREGEN

+! FOUGERA PHARMS INC

15%

N021902 001 Oct 31, 2006

SIROLIMUS

SOLUTION; ORAL

RAPAMUNE

+! PF PRISM CV

1MG/ML

N021083 001 Sep 15, 1999

TABLET; ORAL

RAPAMUNE

| | | | | | |
|-----------|---|-------------|--------------|--------------------|--------------|
| <u>AB</u> | + | PF PRISM CV | <u>0.5MG</u> | <u>N021110 004</u> | Jan 25, 2010 |
| <u>AB</u> | + | | <u>1MG</u> | <u>N021110 001</u> | Aug 25, 2000 |

PRESCRIPTION DRUG PRODUCT LIST

SIROLIMUS

TABLET; ORAL

RAPAMUNE

| | | | | | | |
|-----------|------------|--|------------|----------------|------------|--------------|
| AB | + ! | | 2MG | N021110 | 002 | Aug 22, 2002 |
|-----------|------------|--|------------|----------------|------------|--------------|

SIROLIMUS

| | | | | | | |
|-----------|--|--------------------|------------|----------------|------------|--------------|
| AB | | DR REDDYS LABS LTD | 1MG | A201578 | 001 | Oct 27, 2014 |
|-----------|--|--------------------|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|------------|----------------|------------|--------------|
| AB | | | 2MG | A201578 | 002 | Oct 27, 2014 |
|-----------|--|--|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-------------------------|--------------|----------------|------------|--------------|
| AB | | ZYDUS PHARMS USA INC | 0.5MG | A201676 | 003 | Jan 08, 2014 |
|-----------|--|-------------------------|--------------|----------------|------------|--------------|

SITAGLIPTIN PHOSPHATE

TABLET; ORAL

JANUVIA

| | | | | | | |
|---|--|-------------------|--------------|---------|-----|--------------|
| + | | MERCK SHARP DOHME | EQ 25MG BASE | N021995 | 001 | Oct 16, 2006 |
|---|--|-------------------|--------------|---------|-----|--------------|

| | | | | | | |
|---|--|--|--------------|---------|-----|--------------|
| + | | | EQ 50MG BASE | N021995 | 002 | Oct 16, 2006 |
|---|--|--|--------------|---------|-----|--------------|

| | | | | | | |
|---|---|--|---------------|---------|-----|--------------|
| + | ! | | EQ 100MG BASE | N021995 | 003 | Oct 16, 2006 |
|---|---|--|---------------|---------|-----|--------------|

SODIUM ACETATE

INJECTABLE; INJECTION

SODIUM ACETATE

| | | | | | | |
|--|--|--------------------|---------|---------|-----|--------------|
| | | FRESENIUS KABI USA | 4MEQ/ML | A206687 | 001 | Oct 30, 2017 |
|--|--|--------------------|---------|---------|-----|--------------|

| | | | | | | |
|---|---|---------|---------|---------|-----|--------------|
| + | ! | HOSPIRA | 2MEQ/ML | N018893 | 001 | May 04, 1983 |
|---|---|---------|---------|---------|-----|--------------|

SODIUM BENZOATE; SODIUM PHENYLACETATE

SOLUTION; INTRAVENOUS

AMMONUL

| | | | | | | |
|-----------|------------|---------|------------------------------------|----------------|------------|--------------|
| AP | + ! | MEDICIS | 10%;10% (5GM/50ML;5GM/50ML) | N020645 | 001 | Feb 17, 2005 |
|-----------|------------|---------|------------------------------------|----------------|------------|--------------|

SODIUM PHENYLACETATE AND SODIUM BENZOATE

| | | | | | | |
|-----------|--|------------------|------------------------------------|----------------|------------|--------------|
| AP | | AILEX PHARMS LLC | 10%;10% (5GM/50ML;5GM/50ML) | A207096 | 001 | Feb 24, 2016 |
|-----------|--|------------------|------------------------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-----------------|------------------------------------|----------------|------------|--------------|
| AP | | MAIA PHARMS INC | 10%;10% (5GM/50ML;5GM/50ML) | A208521 | 001 | May 08, 2017 |
|-----------|--|-----------------|------------------------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-------------|------------------------------------|----------------|------------|--------------|
| AP | | NAVINTA LLC | 10%;10% (5GM/50ML;5GM/50ML) | A205880 | 001 | Aug 04, 2016 |
|-----------|--|-------------|------------------------------------|----------------|------------|--------------|

SODIUM BICARBONATE

INJECTABLE; INJECTION

SODIUM BICARBONATE

| | | | | | | |
|-----------|----------|---------|------------------|----------------|------------|--------------|
| AP | ! | HOSPIRA | 0.9MEQ/ML | A077394 | 001 | Nov 09, 2005 |
|-----------|----------|---------|------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|----------|--|----------------|----------------|------------|--------------|
| AP | ! | | 1MEQ/ML | A077394 | 002 | Nov 09, 2005 |
|-----------|----------|--|----------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-------------|------------------|----------------|------------|--------------|
| AP | | HOSPIRA INC | 0.9MEQ/ML | A202494 | 001 | Mar 06, 2017 |
|-----------|--|-------------|------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|----------------|----------------|------------|--------------|
| AP | | | 1MEQ/ML | A202432 | 001 | Sep 26, 2017 |
|-----------|--|--|----------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|----------------|----------------|------------|--------------|
| AP | | | 1MEQ/ML | A202494 | 002 | Mar 06, 2017 |
|-----------|--|--|----------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|---------------------|----------------|----------------|------------|--------------|
| AP | | INTL MEDICATION SYS | 1MEQ/ML | A203449 | 001 | Sep 19, 2017 |
|-----------|--|---------------------|----------------|----------------|------------|--------------|

| | | | | | | |
|--|--|-------------|----------------|---------|-----|--------------|
| | | HOSPIRA INC | 1MEQ/ML | A202495 | 001 | Mar 06, 2017 |
|--|--|-------------|----------------|---------|-----|--------------|

SODIUM CHLORIDE

INJECTABLE; INJECTION

BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | | | |
|-----------|--|--------------------|---------------|----------------|------------|--------------|
| AP | | FRESENIUS KABI USA | 9MG/ML | A088911 | 001 | Feb 07, 1985 |
|-----------|--|--------------------|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|------------|---------|---------------|----------------|------------|--------------|
| AP | + ! | HOSPIRA | 9MG/ML | N018800 | 001 | Oct 29, 1982 |
|-----------|------------|---------|---------------|----------------|------------|--------------|

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

| | | | | | | |
|-----------|----------|---------|--------------------|----------------|------------|--------------|
| AP | + | B BRAUN | 450MG/100ML | N019635 | 001 | Mar 09, 1988 |
|-----------|----------|---------|--------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-----------------|--------------------|----------------|------------|--|
| AP | | BAXTER HLTHCARE | 450MG/100ML | N018016 | 001 | |
|-----------|--|-----------------|--------------------|----------------|------------|--|

| | | | | | | |
|-----------|--|--------------------|--------------------|----------------|------------|--------------|
| AP | | FRESENIUS KABI USA | 450MG/100ML | A208122 | 001 | Jul 23, 2018 |
|-----------|--|--------------------|--------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|---------|--------------------|----------------|------------|--------------|
| AP | | HOSPIRA | 450MG/100ML | N019759 | 001 | Jun 08, 1988 |
|-----------|--|---------|--------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|------------|-----------------|--------------------|----------------|------------|--|
| AP | + ! | ICU MEDICAL INC | 450MG/100ML | N018090 | 001 | |
|-----------|------------|-----------------|--------------------|----------------|------------|--|

SODIUM CHLORIDE 0.9%

| | | | | | | |
|-----------|--|-------------------------|---------------|----------------|------------|--------------|
| AP | | SPECTRA MDCL DEVICES | 9MG/ML | A206171 | 001 | Jul 21, 2017 |
|-----------|--|-------------------------|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-------------------------|---------------|----------------|------------|--------------|
| AP | | WEST-WARD PHARMS INT | 9MG/ML | A201850 | 001 | Jan 20, 2012 |
|-----------|--|-------------------------|---------------|----------------|------------|--------------|

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | | | |
|-----------|------------|---------|--------------------|----------------|------------|--|
| AP | + ! | B BRAUN | 900MG/100ML | N017464 | 001 | |
|-----------|------------|---------|--------------------|----------------|------------|--|

| | | | | | | |
|-----------|------------|--|--------------------|----------------|------------|--------------|
| AP | + ! | | 900MG/100ML | N019635 | 002 | Mar 09, 1988 |
|-----------|------------|--|--------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|------------|-----------------|---------------|----------------|------------|--------------|
| AP | + ! | BAXTER HLTHCARE | 9MG/ML | N016677 | 004 | Oct 30, 1985 |
|-----------|------------|-----------------|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|----------|--|---------------|----------------|------------|--------------|
| AP | + | | 9MG/ML | N020178 | 002 | Dec 07, 1992 |
|-----------|----------|--|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|------------|--|--------------------|----------------|------------|--|
| AP | + ! | | 900MG/100ML | N016677 | 001 | |
|-----------|------------|--|--------------------|----------------|------------|--|

| | | | | | | |
|-----------|------------|--|--------------------|----------------|------------|--------------|
| AP | + ! | | 900MG/100ML | N020178 | 001 | Dec 07, 1992 |
|-----------|------------|--|--------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|----------|--------------------|---------------|----------------|------------|--------------|
| AP | ! | FRESENIUS KABI USA | 9MG/ML | A088912 | 001 | Jan 10, 1985 |
|-----------|----------|--------------------|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|--------------------|----------------|------------|--------------|
| AP | | | 900MG/100ML | A207310 | 001 | Sep 19, 2017 |
|-----------|--|--|--------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-----------------|--------------------|----------------|------------|--------------|
| AP | | FRESENIUS MEDCL | 900MG/100ML | A078177 | 001 | Apr 12, 2007 |
|-----------|--|-----------------|--------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-------------|--------------------|----------------|------------|--------------|
| AP | | HAEMONETICS | 900MG/100ML | A076316 | 001 | Oct 27, 2004 |
|-----------|--|-------------|--------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|------------|---------|---------------|----------------|------------|--------------|
| AP | + ! | HOSPIRA | 9MG/ML | N018803 | 001 | Oct 29, 1982 |
|-----------|------------|---------|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|----------|--|---------------|----------------|------------|--------------|
| AP | + | | 9MG/ML | N019465 | 002 | Jul 15, 1985 |
|-----------|----------|--|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|------------|--|--------------------|----------------|------------|--------------|
| AP | + ! | | 900MG/100ML | N019465 | 001 | Jul 15, 1985 |
|-----------|------------|--|--------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|------------|--|--------------------|----------------|------------|--------------|
| AP | + ! | | 900MG/100ML | N019480 | 001 | Sep 17, 1985 |
|-----------|------------|--|--------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|------------|-----------------|--------------------|----------------|------------|--|
| AP | + ! | ICU MEDICAL INC | 900MG/100ML | N016366 | 001 | |
|-----------|------------|-----------------|--------------------|----------------|------------|--|

PRESCRIPTION DRUG PRODUCT LIST

SODIUM CHLORIDE

INJECTABLE; INJECTION

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | | |
|-----------|-------------------------|--------------------|----------------|------------|--------------|
| <u>AP</u> | JUBILANT HOLLISTRSTR | <u>9MG/ML</u> | <u>A203352</u> | <u>001</u> | May 18, 2016 |
| <u>AP</u> | LABORATORIOS GRIFOLS | <u>900MG/100ML</u> | <u>A207956</u> | <u>001</u> | May 25, 2017 |
| <u>AP</u> | ! TARO | <u>9MG/ML</u> | <u>A077407</u> | <u>001</u> | Aug 11, 2006 |

SODIUM CHLORIDE 3% IN PLASTIC CONTAINER

| | | | | | |
|-----------|--------------------|------------------|----------------|------------|--------------|
| <u>AP</u> | B BRAUN | <u>3GM/100ML</u> | <u>N019635</u> | <u>003</u> | Mar 09, 1988 |
| <u>AP</u> | +! BAXTER HLTHCARE | <u>3GM/100ML</u> | <u>N019022</u> | <u>001</u> | Nov 01, 1983 |

SODIUM CHLORIDE 5% IN PLASTIC CONTAINER

| | | | | | |
|-----------|-------------------|------------------|----------------|------------|--------------|
| <u>AP</u> | B BRAUN | <u>5GM/100ML</u> | <u>N019635</u> | <u>004</u> | Mar 09, 1988 |
| <u>AP</u> | + BAXTER HLTHCARE | <u>5GM/100ML</u> | <u>N019022</u> | <u>002</u> | Nov 01, 1983 |

SODIUM CHLORIDE 0.9%

| | | | | | |
|---|------------------|--------------------|---------|-----|--------------|
| + | B BRAUN | 900MG/10ML | N019635 | 005 | Aug 11, 2016 |
| + | MEDEFIL INC | 90MG/10ML (9MG/ML) | N202832 | 006 | Jan 06, 2012 |
| | WEST-WARD PHARMS | 9MG/ML | A201833 | 001 | Sep 24, 2013 |

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | | |
|---|--------------------|-------------------------|---------|-----|--------------|
| + | ! LIEBEL-FLARSHEIM | 405MG/50ML (9MG/ML) | N021569 | 001 | Jul 27, 2006 |
| + | | 1012.5MG/125ML (9MG/ML) | N021569 | 002 | Jul 27, 2006 |

SODIUM CHLORIDE IN PLASTIC CONTAINER

| | | | | | |
|---|---------|-----------|---------|-----|--------------|
| + | HOSPIRA | 2.5MEQ/ML | N018897 | 001 | Jul 20, 1984 |
|---|---------|-----------|---------|-----|--------------|

SOLUTION; IRRIGATION

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | | |
|-----------|-----------------|--------------------|----------------|------------|--|
| <u>AT</u> | ! B BRAUN | <u>900MG/100ML</u> | <u>N016733</u> | <u>001</u> | |
| <u>AT</u> | BAXTER HLTHCARE | <u>900MG/100ML</u> | <u>N017427</u> | <u>001</u> | |
| <u>AT</u> | | <u>900MG/100ML</u> | <u>N017867</u> | <u>001</u> | |
| <u>AT</u> | ICU MEDICAL INC | <u>900MG/100ML</u> | <u>N017514</u> | <u>001</u> | |
| <u>AT</u> | | <u>900MG/100ML</u> | <u>N018314</u> | <u>001</u> | |

SOLUTION FOR SLUSH; IRRIGATION

SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER

| | | | | | |
|--|-----------------|-------------|---------|-----|--------------|
| | BAXTER HLTHCARE | 900MG/100ML | N019319 | 002 | May 17, 1985 |
|--|-----------------|-------------|---------|-----|--------------|

SODIUM FERRIC GLUCONATE COMPLEX

INJECTABLE; INJECTION

FERRLECIT

| | | | | | |
|-----------|---------------------|-------------------|----------------|------------|--------------|
| <u>AB</u> | ! SANOFI AVENTIS US | <u>62.5MG/5ML</u> | <u>N020955</u> | <u>001</u> | Feb 18, 1999 |
|-----------|---------------------|-------------------|----------------|------------|--------------|

SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE

| | | | | | |
|-----------|------------------|-------------------|----------------|------------|--------------|
| <u>AB</u> | WEST-WARD PHARMS | <u>62.5MG/5ML</u> | <u>A078215</u> | <u>001</u> | Mar 31, 2011 |
| | INT | | | | |

SODIUM FLUORIDE F-18

INJECTABLE; INTRAVENOUS

SODIUM FLUORIDE F-18

| | | | | | |
|-----------|---------------------|---------------------|----------------|------------|--------------|
| <u>AP</u> | 3D IMAGING DRUG | <u>10-200mCi/ML</u> | <u>A203777</u> | <u>001</u> | Oct 19, 2015 |
| <u>AP</u> | BIOMEDCL RES FDN | <u>10-200mCi/ML</u> | <u>A204351</u> | <u>001</u> | Jan 09, 2015 |
| <u>AP</u> | CARDINAL HEALTH 414 | <u>10-200mCi/ML</u> | <u>A203780</u> | <u>001</u> | Jul 30, 2015 |
| <u>AP</u> | ESSENTIAL ISOTOPEs | <u>10-200mCi/ML</u> | <u>A204541</u> | <u>001</u> | Oct 29, 2014 |
| <u>AP</u> | GLOBAL ISOTOPEs LLC | <u>10-200mCi/ML</u> | <u>A204464</u> | <u>001</u> | Oct 21, 2014 |
| <u>AP</u> | HOT SHOTS NM LLC | <u>10-200mCi/ML</u> | <u>A204530</u> | <u>001</u> | Jul 29, 2015 |
| <u>AP</u> | ! HOUSTON CYCLOTRON | <u>10-200mCi/ML</u> | <u>A203544</u> | <u>001</u> | Dec 26, 2012 |
| <u>AP</u> | JUBILANT DRAXIMAGE | <u>10-200mCi/ML</u> | <u>A203968</u> | <u>001</u> | Oct 23, 2015 |
| <u>AP</u> | KREITCHMAN PET CTR | <u>10-200mCi/ML</u> | <u>A203936</u> | <u>001</u> | May 19, 2016 |
| <u>AP</u> | MIDWEST MEDCL | <u>10-200mCi/ML</u> | <u>A204440</u> | <u>001</u> | Nov 17, 2015 |
| <u>AP</u> | MIPS CRF | <u>10-200mCi/ML</u> | <u>A204517</u> | <u>001</u> | Jul 21, 2015 |
| <u>AP</u> | NCM USA BRONX LLC | <u>10-200mCi/ML</u> | <u>A204513</u> | <u>001</u> | Nov 28, 2014 |
| <u>AP</u> | PETNET | <u>10-200mCi/ML</u> | <u>A203890</u> | <u>001</u> | Sep 28, 2015 |
| <u>AP</u> | PRECISION NUCLEAR | <u>10-200mCi/ML</u> | <u>A204542</u> | <u>001</u> | Feb 27, 2015 |
| <u>AP</u> | SHERTECH LABS LLC | <u>10-200mCi/ML</u> | <u>A204315</u> | <u>001</u> | Sep 22, 2014 |
| <u>AP</u> | SOFIE | <u>10-200mCi/ML</u> | <u>A203592</u> | <u>001</u> | Aug 18, 2015 |
| <u>AP</u> | SPECTRON MRC LLC | <u>10-200mCi/ML</u> | <u>A203912</u> | <u>001</u> | Apr 22, 2015 |
| <u>AP</u> | UCSF RODIOPHARM | <u>10-200mCi/ML</u> | <u>A204437</u> | <u>001</u> | Mar 13, 2014 |
| <u>AP</u> | UNIV UTAH CYCLOTRON | <u>10-200mCi/ML</u> | <u>A204497</u> | <u>001</u> | Apr 20, 2015 |
| | ! MCPRF | 10-91.5mCi/ML | A203605 | 001 | Jun 28, 2013 |
| | THE FEINSTEIN INST | 20-600mCi/ML | A204328 | 001 | Nov 19, 2014 |

SODIUM IODIDE I-123

CAPSULE; ORAL

SODIUM IODIDE I 123

| | | | | | |
|-----------|-----------------------|---------------|----------------|------------|--------------|
| <u>AA</u> | ! CARDINAL HEALTH 418 | <u>100uCi</u> | <u>N018671</u> | <u>001</u> | May 27, 1982 |
| <u>AA</u> | ! MALLINKRODT NUCLEAR | <u>200uCi</u> | <u>N018671</u> | <u>002</u> | May 27, 1982 |
| <u>AA</u> | | <u>100uCi</u> | <u>A071909</u> | <u>001</u> | Feb 28, 1989 |
| <u>AA</u> | | <u>200uCi</u> | <u>A071910</u> | <u>001</u> | Feb 28, 1989 |

PRESCRIPTION DRUG PRODUCT LIST

SODIUM IODIDE I-131

CAPSULE; ORAL

SODIUM IODIDE I 131

+ JUBILANT DRAXIMAGE 0.009-0.1mCi

N021305 006 May 19, 2005

SOLUTION; ORAL

HICON

+! JUBILANT DRAXIMAGE 250-1000mCi

N021305 007 Dec 05, 2011

SODIUM LACTATE

INJECTABLE; INJECTION

SODIUM LACTATE IN PLASTIC CONTAINER

+! HOSPIRA 5MEQ/ML

N018947 001 Sep 05, 1984

SODIUM NITRITE

SOLUTION; INTRAVENOUS

SODIUM NITRITE

+! HOPE PHARMS 300MG/10ML (30MG/ML)

N203922 001 Feb 14, 2012

SODIUM NITRITE; SODIUM THIOSULFATE

SOLUTION, SOLUTION; INTRAVENOUS, INTRAVENOUS

NITHIODOTE

+! HOPE PHARMS 300MG/10ML (30MG/ML), N/A; N/A, 12.5GM/50ML (250MG/ML)

N201444 001 Jan 14, 2011

SODIUM NITROPRUSSIDE

INJECTABLE; INJECTION

NITROPRESSAP +! HOSPIRA25MG/MLA071961 001 Aug 01, 1988SODIUM NITROPRUSSIDEAP AKORN25MG/MLA208635 001 May 04, 2017AP AMNEAL PHARMS CO25MG/MLA209493 001 Nov 07, 2017AP AMPHASTAR PHARMS25MG/MLA209832 001 Dec 18, 2017

INC

AP CIPLA25MG/MLA210855 001 Jul 16, 2018AP MEDICURE25MG/MLA209584 001 Aug 10, 2018AP MICRO LABS25MG/MLA209352 001 Dec 08, 2017AP MYLAN LABS LTD25MG/MLA210763 001 Apr 17, 2018AP NAMIGEN LLC25MG/MLA207426 001 Dec 08, 2016AP NEXUS PHARMS25MG/MLA207499 001 May 25, 2017AP RENAISSANCE SSA LLC25MG/MLA209834 001 Jun 26, 2018AP SOMERSET THERAPS25MG/MLA210882 001 Aug 17, 2018

LLC

AP SUN PHARM INDS LTD25MG/MLA210467 001 Nov 26, 2018

SOLUTION; INTRAVENOUS

NIPRIDE RTU IN SODIUM CHLORIDE 0.9%

+! EXELA PHARMA SCS 10MG/50ML (0.2MG/ML)

N209387 002 Dec 07, 2017

LLC

+! 20MG/100ML (0.2MG/ML)

N209387 003 Jul 13, 2018

+! 50MG/100ML (0.5MG/ML)

N209387 001 Mar 08, 2017

SODIUM OXYBATE

SOLUTION; ORAL

SODIUM OXYBATEAA WEST-WARD PHARMS500MG/MLA202090 001 Jan 17, 2017

INT

XYREMAA +! JAZZ PHARMS500MG/MLN021196 001 Jul 17, 2002SODIUM PHENYLBUTYRATE

POWDER; ORAL

BUPHENYLAB +! HORIZON PHARMA INC3GM/TEASPOONFULN020573 001 Apr 30, 1996SODIUM PHENYLBUTYRATEAB PAR PHARM3GM/TEASPOONFULA203918 001 Jun 15, 2016AB SIGMAPHARM LABS LLC3GM/TEASPOONFULA202819 001 Mar 22, 2013

TABLET; ORAL

BUPHENYLAB +! HORIZON PHARMA INC500MGN020572 001 May 13, 1996SODIUM PHENYLBUTYRATEAB ALVOGEN MALTA500MGA090910 001 Nov 18, 2011AB PAR PHARM500MGA204395 001 Apr 15, 2016

PRESCRIPTION DRUG PRODUCT LIST

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL

OSMOPREP

+! SALIX PHARMS 0.398GM;1.102GM N021892 001 Mar 16, 2006

SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE; SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS

INJECTABLE; INJECTION

SODIUM PHOSPHATES IN PLASTIC CONTAINER

+! HOSPIRA 142MG/ML;276MG/ML N018892 001 May 10, 1983

SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL

KALEXATE**AA** ! KVK TECH **454GM/BOT** **A040905 001** Mar 30, 2009**KIONEX****AA** PADDOCK LLC **454GM/BOT** **A040029 001** Feb 06, 1998**SODIUM POLYSTYRENE SULFONATE****AA** APNAR PHARMA LP **454GM/BOT** **A206815 001** Feb 18, 2016**AA** BELCHER PHARMS LLC **454GM/BOT** **A205727 001** Feb 23, 2016**AA** CMP PHARMA INC **454GM/BOT** **A089910 001** Jan 19, 1989**AA** ECI PHARMS LLC **453.6GM/BOT** **A090313 001** Dec 21, 2011**AA** EPIC PHARMA LLC **453.6GM/BOT** **A202333 001** Mar 19, 2014**AA** NUVO PHARMS INC **454GM/BOT** **A204071 001** Nov 28, 2014

KALEXATE

KVK TECH 15GM/BOT

A040905 002 Apr 03, 2015

SODIUM POLYSTYRENE SULFONATE

NUVO PHARMS INC 15GM/BOT

A204071 002 Nov 28, 2014

SUSPENSION; ORAL, RECTAL

KIONEX**AA** PADDOCK LLC **15GM/60ML** **A040028 001** Sep 17, 2007**SODIUM POLYSTYRENE SULFONATE****AA** PADDOCK LLC **15GM/60ML** **A090590 001** May 13, 2011**AA** WEST-WARD PHARMS **15GM/60ML** **A089049 001** Nov 17, 1986

INT

SPS**AA** ! CMP PHARMA INC **15GM/60ML** **A087859 001** Dec 08, 1982SODIUM TETRADECYL SULFATE

INJECTABLE; INJECTION

SOTRADECOL

! MYLAN INSTITUTIONAL 20MG/2ML (10MG/ML) A040541 001 Nov 12, 2004

! 60MG/2ML (30MG/ML) A040541 002 Nov 12, 2004

SODIUM THIOSULFATE

SOLUTION; INTRAVENOUS

SODIUM THIOSULFATE

+! HOPE PHARMS 12.5GM/50ML (250MG/ML) N203923 001 Feb 14, 2012

SODIUM ZIRCONIUM CYCLOSILICATE

FOR SUSPENSION; ORAL

LOKELMA

+ ASTRAZENECA PHARMS 5GM/PACKET N207078 001 May 18, 2018

+! 10GM/PACKET N207078 002 May 18, 2018

SOFOBUVIR

TABLET; ORAL

SOVALDI

+! GILEAD SCIENCES INC 400MG N204671 001 Dec 06, 2013

SOFOBUVIR; VELPATASVIR

TABLET; ORAL

EPCLUSA

+! GILEAD SCIENCES INC 400MG;100MG N208341 001 Jun 28, 2016

SOFOBUVIR; VELPATASVIR; VOXILAPREVIR

TABLET; ORAL

VOSEVI

+! GILEAD SCIENCES INC 400MG;100MG;100MG N209195 001 Jul 18, 2017

SOLIFENACIN SUCCINATE

TABLET; ORAL

SOLIFENACIN SUCCINATE**AB** TEVA PHARMS USA **5MG** **A091464 001** Apr 02, 2014**AB** **10MG** **A091464 002** Apr 02, 2014**VESICARE****AB** + ASTELLAS **5MG** **N021518 001** Nov 19, 2004**AB** +! **10MG** **N021518 002** Nov 19, 2004

PRESCRIPTION DRUG PRODUCT LIST

SOMATROPIN

INJECTABLE; INJECTION

ZOMACTON

| | | | | | | |
|----|---|---------|-----------|---------|-----|--------------|
| BX | + | FERRING | 5MG/VIAL | N019774 | 002 | Jan 04, 2002 |
| | + | | 10MG/VIAL | N019774 | 003 | Mar 07, 2012 |

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION

GENOTROPIN

| | | | | | | |
|----|---|-------------------------|------------|---------|-----|--------------|
| BX | + | PHARMACIA AND UPJOHN | 5.8MG/VIAL | N020280 | 006 | Aug 24, 1995 |
|----|---|-------------------------|------------|---------|-----|--------------|

HUMATROPE

| | | | | | | |
|----|---|-------|----------|---------|-----|--------------|
| BX | + | LILLY | 5MG/VIAL | N019640 | 004 | Mar 08, 1987 |
| BX | + | | 6MG/VIAL | N019640 | 005 | Feb 04, 1999 |

NORDITROPIN FLEXPRO

| | | | | | | |
|----|--|------------------|------------|---------|-----|--------------|
| BX | | NOVO NORDISK INC | 5MG/1.5ML | N021148 | 008 | Mar 01, 2010 |
| BX | | | 10MG/1.5ML | N021148 | 009 | Mar 01, 2010 |

OMNITROPE

| | | | | | | |
|----|--|--------|------------|---------|-----|--------------|
| BX | | SANDOZ | 1.5MG/VIAL | N021426 | 002 | May 30, 2006 |
| BX | | | 5MG/1.5ML | N021426 | 003 | Jan 16, 2008 |
| BX | | | 5.8MG/VIAL | N021426 | 001 | May 30, 2006 |
| BX | | | 10MG/1.5ML | N021426 | 004 | Aug 25, 2008 |

SAIZEN

| | | | | | | |
|----|---|------------|----------|---------|-----|--------------|
| BX | + | EMD SERONO | 5MG/VIAL | N019764 | 002 | Oct 08, 1996 |
|----|---|------------|----------|---------|-----|--------------|

SEROSTIM

| | | | | | | |
|----|--|------------|----------|---------|-----|--------------|
| BX | | EMD SERONO | 5MG/VIAL | N020604 | 002 | Aug 23, 1996 |
| BX | | | 6MG/VIAL | N020604 | 001 | Aug 23, 1996 |

GENOTROPIN

| | | | | | | |
|--|---|-------------------------|-------------|---------|-----|--------------|
| | + | PHARMACIA AND UPJOHN | 13.8MG/VIAL | N020280 | 007 | Oct 23, 1996 |
|--|---|-------------------------|-------------|---------|-----|--------------|

GENOTROPIN PRESERVATIVE FREE

| | | | | | | |
|--|---|-------------------------|------------|---------|-----|--------------|
| | + | PHARMACIA AND UPJOHN | 0.2MG/VIAL | N020280 | 001 | Jan 27, 1998 |
| | + | | 0.4MG/VIAL | N020280 | 002 | Jan 27, 1998 |
| | + | | 0.6MG/VIAL | N020280 | 003 | Jan 27, 1998 |
| | + | | 0.8MG/VIAL | N020280 | 005 | Jan 27, 1998 |
| | + | | 1MG/VIAL | N020280 | 008 | Jan 27, 1998 |
| | + | | 1.2MG/VIAL | N020280 | 009 | Jan 27, 1998 |
| | + | | 1.4MG/VIAL | N020280 | 010 | Jan 27, 1998 |
| | + | | 1.6MG/VIAL | N020280 | 011 | Jan 27, 1998 |
| | + | | 1.8MG/VIAL | N020280 | 012 | Jan 27, 1998 |
| | + | | 2MG/VIAL | N020280 | 013 | Jan 27, 1998 |

HUMATROPE

| | | | | | | |
|--|---|-------|-----------|---------|-----|--------------|
| | + | LILLY | 12MG/VIAL | N019640 | 006 | Feb 04, 1999 |
| | + | | 24MG/VIAL | N019640 | 007 | Feb 04, 1999 |

NORDITROPIN FLEXPRO

| | | | | | | |
|--|--|------------------|------------|---------|-----|--------------|
| | | NOVO NORDISK INC | 15MG/1.5ML | N021148 | 010 | Mar 01, 2010 |
| | | | 30MG/3ML | N021148 | 011 | Jan 23, 2015 |

NUTROPIN AQ NUSPIN

| | | | | | | |
|--|---|-----------|--------------------|---------|-----|--------------|
| | + | GENENTECH | 5MG/2ML (2.5MG/ML) | N020522 | 003 | Jan 03, 2008 |
| | + | | 10MG/2ML (5MG/ML) | N020522 | 005 | Jan 03, 2008 |
| | + | | 20MG/2ML (10MG/ML) | N020522 | 004 | Jan 03, 2008 |

SAIZEN

| | | | | | | |
|--|---|------------|------------|---------|-----|--------------|
| | + | EMD SERONO | 8.8MG/VIAL | N019764 | 003 | Aug 29, 2000 |
|--|---|------------|------------|---------|-----|--------------|

SEROSTIM

| | | | | | | |
|--|--|------------|----------|---------|-----|--------------|
| | | EMD SERONO | 4MG/VIAL | N020604 | 003 | Jul 25, 1997 |
|--|--|------------|----------|---------|-----|--------------|

ZORBATIVE

| | | | | | | |
|--|---|------------|------------|---------|-----|--------------|
| | + | EMD SERONO | 8.8MG/VIAL | N021597 | 004 | Dec 01, 2003 |
|--|---|------------|------------|---------|-----|--------------|

SONIDEGIB PHOSPHATE

CAPSULE; ORAL

ODOMZO

| | | | | | | |
|--|---|-------------------|---------------|---------|-----|--------------|
| | + | SUN PHARMA GLOBAL | EQ 200MG BASE | N205266 | 001 | Jul 24, 2015 |
|--|---|-------------------|---------------|---------|-----|--------------|

SORAFENIB TOSYLATE

TABLET; ORAL

NEXAVAR

| | | | | | | |
|--|---|----------------|---------------|---------|-----|--------------|
| | + | BAYER HLTHCARE | EQ 200MG BASE | N021923 | 001 | Dec 20, 2005 |
|--|---|----------------|---------------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

SORBITOL

SOLUTION;IRRIGATION

SORBITOL 3% IN PLASTIC CONTAINER

BAXTER HLTHCARE 3GM/100ML

N017863 001

SORBITOL 3.3% IN PLASTIC CONTAINER

B BRAUN 3.3GM/100ML

N016741 001

SOTALOL HYDROCHLORIDE

SOLUTION;INTRAVENOUS

SOTALOL HYDROCHLORIDE

+! ALTATHERA PHARMS 150MG/10ML (15MG/ML)

N022306 001 Jul 02, 2009

LLC

SOLUTION;ORAL

SOTYLIZE

+! ARBOR PHARMS LLC 5MG/ML (5MG/ML)

N205108 001 Oct 22, 2014

TABLET;ORAL

BETAPACEAB1 + COVIS PHARMA BV80MGN019865 001 Oct 30, 1992AB1 +120MGN019865 005 Apr 20, 1994AB1 +!160MGN019865 002 Oct 30, 1992AB1 +240MGN019865 003 Oct 30, 1992SORINEAB1 UPSHER SMITH LABS80MGA075500 001 Apr 27, 2001AB1120MGA075500 004 Apr 27, 2001AB1160MGA075500 002 Apr 27, 2001AB1240MGA075500 003 Apr 27, 2001SOTALOL HYDROCHLORIDEAB1 APOTEX INC80MGA076140 001 Sep 26, 2002AB1120MGA076140 002 Sep 26, 2002AB1160MGA076140 003 Sep 26, 2002AB1240MGA076140 004 Sep 26, 2002AB1 BEXIMCO PHARMS USA80MGA207428 001 Oct 21, 2016AB1120MGA207428 002 Oct 21, 2016AB1160MGA207428 003 Oct 21, 2016AB1 OXFORD PHARMS80MGA075563 001 Nov 07, 2003AB1120MGA075563 002 Nov 07, 2003AB1160MGA075563 003 Nov 07, 2003AB1240MGA075563 004 Nov 07, 2003AB1 TEVA80MGA075429 001 May 01, 2000AB1120MGA075429 002 May 01, 2000AB1160MGA075429 003 May 01, 2000AB1240MGA075429 004 May 01, 2000AB1 UPSHER SMITH LABS80MGA075366 001 May 01, 2000AB1120MGA075366 002 May 01, 2000AB1160MGA075366 003 May 01, 2000AB1240MGA075366 004 May 01, 2000BETAPACE AFAB2 + COVIS PHARMA BV80MGN021151 001 Feb 22, 2000AB2 +120MGN021151 002 Feb 22, 2000AB2 +!160MGN021151 003 Feb 22, 2000SOTALOL HYDROCHLORIDEAB2 APOTEX80MGA076214 001 Aug 27, 2003AB2120MGA076214 002 Aug 27, 2003AB2160MGA076214 003 Aug 27, 2003AB2 BEXIMCO PHARMS USA80MGA207429 001 Nov 02, 2018AB2120MGA207429 002 Nov 02, 2018AB2160MGA207429 003 Nov 02, 2018AB2 EPIC PHARMA INC80MGA077070 001 Nov 04, 2005AB2120MGA077070 002 Nov 04, 2005AB2160MGA077070 003 Nov 04, 2005AB2 MYLAN80MGA077616 001 Feb 07, 2007AB2120MGA077616 002 Feb 07, 2007AB2160MGA077616 003 Feb 07, 2007SOYBEAN OIL

INJECTABLE;INJECTION

INTRALIPID 10%AP +! FRESENIUS10%N017643 001INTRALIPID 20%AP +! FRESENIUS20%N018449 001AP +!20%N020248 001 Aug 07, 1996NUTRILIPID 10%AP +! B BRAUN10%N019531 001 May 28, 1993

PRESCRIPTION DRUG PRODUCT LIST

SOYBEAN OIL

INJECTABLE; INJECTION

NUTRILIPID 20%

| | | | | | | |
|-----------|------------|----------------|------------|----------------|------------|--------------|
| AP | + ! | B BRAUN | 20% | N019531 | 002 | May 28, 1993 |
| | | INTRALIPID 30% | | | | |
| | + ! | FRESENIUS | 30% | N019942 | 001 | Dec 30, 1993 |

SPINOSAD

SUSPENSION; TOPICAL

NATROBA

| | | | | | |
|------------|-------------|-------------|---------|-----|--------------|
| + ! | PARAPRO LLC | 0.9% | N022408 | 001 | Jan 18, 2011 |
|------------|-------------|-------------|---------|-----|--------------|

SPIRONOLACTONE

SUSPENSION; ORAL

CAROSPIR

| | | | | | |
|------------|-------------|-----------------|---------|-----|--------------|
| + ! | CMP DEV LLC | 25MG/5ML | N209478 | 001 | Aug 04, 2017 |
|------------|-------------|-----------------|---------|-----|--------------|

TABLET; ORAL

ALDACTONE

| | | | | | | |
|-----------|------------|---------------|--------------|----------------|------------|--------------|
| AB | + | GD SEARLE LLC | 25MG | N012151 | 009 | Dec 30, 1983 |
| AB | + | | 50MG | N012151 | 008 | Dec 30, 1982 |
| AB | + ! | | 100MG | N012151 | 010 | Dec 30, 1983 |

SPIRONOLACTONE

| | | | | | | |
|-----------|--|----------------------|--------------|----------------|------------|--------------|
| AB | | ACCORD HLTHCARE | 25MG | A203512 | 001 | Sep 19, 2016 |
| AB | | | 50MG | A203512 | 002 | Sep 19, 2016 |
| AB | | | 100MG | A203512 | 003 | Sep 19, 2016 |
| AB | | ACTAVIS ELIZABETH | 25MG | A040353 | 003 | Mar 15, 2006 |
| AB | | | 50MG | A040353 | 001 | Jul 29, 1999 |
| AB | | | 100MG | A040353 | 002 | Jul 29, 1999 |
| AB | | AMNEAL PHARMS | 25MG | A091426 | 001 | Jul 02, 2010 |
| AB | | | 50MG | A091426 | 002 | Jul 02, 2010 |
| AB | | | 100MG | A091426 | 003 | Jul 02, 2010 |
| AB | | CASI PHARMS INC | 25MG | A086809 | 001 | |
| AB | | JUBILANT GENERICS | 25MG | A203253 | 001 | Apr 23, 2014 |
| AB | | | 50MG | A203253 | 002 | Apr 23, 2014 |
| AB | | | 100MG | A203253 | 003 | Apr 23, 2014 |
| AB | | MYLAN | 25MG | A040424 | 001 | Aug 20, 2001 |
| AB | | | 50MG | A040424 | 002 | Aug 20, 2001 |
| AB | | | 100MG | A040424 | 003 | Aug 20, 2001 |
| AB | | OXFORD PHARMS | 25MG | A040750 | 001 | Aug 29, 2006 |
| AB | | | 50MG | A040750 | 002 | Aug 29, 2006 |
| AB | | | 100MG | A040750 | 003 | Aug 29, 2006 |
| AB | | SUN PHARM INDUSTRIES | 25MG | A089424 | 001 | Jul 23, 1986 |
| AB | | | 50MG | A089424 | 002 | Aug 11, 1999 |
| AB | | | 100MG | A089424 | 003 | Aug 11, 1999 |
| AB | | ZYDUS PHARMS USA INC | 25MG | A205936 | 001 | Jul 18, 2018 |
| AB | | | 50MG | A205936 | 002 | Jul 18, 2018 |
| AB | | | 100MG | A205936 | 003 | Jul 18, 2018 |

STAVUDINE

CAPSULE; ORAL

STAVUDINE

| | | | | | | |
|-----------|--|---------------------|-------------|----------------|------------|--------------|
| AB | | AUROBINDO PHARMA | 15MG | A077672 | 003 | Dec 29, 2008 |
| AB | | | 20MG | A077672 | 004 | Dec 29, 2008 |
| AB | | | 30MG | A077672 | 001 | Dec 29, 2008 |
| AB | | | 40MG | A077672 | 002 | Dec 29, 2008 |
| AB | | HETERO LABS LTD III | 15MG | A078957 | 001 | Dec 29, 2008 |
| AB | | | 20MG | A078957 | 002 | Dec 29, 2008 |
| AB | | | 30MG | A078957 | 003 | Dec 29, 2008 |
| AB | | | 40MG | A078957 | 004 | Dec 29, 2008 |
| AB | | MYLAN | 15MG | A079069 | 001 | Dec 29, 2008 |
| AB | | | 20MG | A079069 | 002 | Dec 29, 2008 |
| AB | | | 30MG | A079069 | 003 | Dec 29, 2008 |
| AB | | | 40MG | A079069 | 004 | Dec 29, 2008 |

ZERIT

| | | | | | | |
|-----------|------------|----------------------|-------------|----------------|------------|--------------|
| AB | + | BRISTOL MYERS SQUIBB | 15MG | N020412 | 002 | Jun 24, 1994 |
| AB | + | | 20MG | N020412 | 003 | Jun 24, 1994 |
| AB | + | | 30MG | N020412 | 004 | Jun 24, 1994 |
| AB | + ! | | 40MG | N020412 | 005 | Jun 24, 1994 |

FOR SOLUTION; ORAL

ZERIT

| | | | | | |
|------------|----------------------|---------------|---------|-----|--------------|
| + ! | BRISTOL-MYERS SQUIBB | 1MG/ML | N020413 | 001 | Sep 06, 1996 |
|------------|----------------------|---------------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

STERILE WATER FOR INJECTION

LIQUID;N/A

BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER

| | | | | | | |
|-----------|------------|----------------|-------------|----------------|------------|--------------|
| <u>AP</u> | <u>+</u> ! | <u>HOSPIRA</u> | <u>100%</u> | <u>N018802</u> | <u>001</u> | Oct 27, 1982 |
|-----------|------------|----------------|-------------|----------------|------------|--------------|

STERILE WATER FOR INJECTION

| | | | | | | |
|-----------|--|---------------------------|-------------|----------------|------------|--------------|
| <u>AP</u> | | <u>FRESENIUS KABI USA</u> | <u>100%</u> | <u>A209689</u> | <u>001</u> | Nov 24, 2017 |
| <u>AP</u> | | <u>WEST-WARD PHARMS</u> | <u>100%</u> | <u>A206369</u> | <u>001</u> | Sep 02, 2015 |

INT

STERILE WATER FOR INJECTION IN PLASTIC CONTAINER

| | | | | | | |
|-----------|------------|---------------------------|-------------|----------------|------------|--------------|
| <u>AP</u> | <u>+</u> ! | <u>B BRAUN</u> | <u>100%</u> | <u>N019633</u> | <u>001</u> | Feb 29, 1988 |
| <u>AP</u> | <u>+</u> ! | <u>BAXTER HLTHCARE</u> | <u>100%</u> | <u>N018632</u> | <u>001</u> | Jun 30, 1982 |
| <u>AP</u> | <u>+</u> ! | | <u>100%</u> | <u>N018632</u> | <u>002</u> | Apr 19, 1988 |
| <u>AP</u> | | <u>FRESENIUS KABI USA</u> | <u>100%</u> | <u>A088400</u> | <u>001</u> | Jan 16, 1984 |
| <u>AP</u> | <u>+</u> ! | <u>HOSPIRA</u> | <u>100%</u> | <u>N018801</u> | <u>001</u> | Oct 27, 1982 |
| <u>AP</u> | <u>+</u> ! | <u>ICU MEDICAL INC</u> | <u>100%</u> | <u>N018233</u> | <u>001</u> | |
| <u>AP</u> | <u>+</u> ! | | <u>100%</u> | <u>N019869</u> | <u>001</u> | Dec 26, 1989 |
| <u>AP</u> | | <u>TARO</u> | <u>100%</u> | <u>A077393</u> | <u>001</u> | Aug 11, 2006 |

STERILE WATER FOR IRRIGATION

LIQUID;IRRIGATION

STERILE WATER

| | | | | | | |
|-----------|--|------------------------|-------------|----------------|------------|--|
| <u>AT</u> | | <u>BAXTER HLTHCARE</u> | <u>100%</u> | <u>N017428</u> | <u>001</u> | |
|-----------|--|------------------------|-------------|----------------|------------|--|

STERILE WATER IN PLASTIC CONTAINER

| | | | | | | |
|-----------|----------|------------------------|-------------|----------------|------------|--|
| <u>AT</u> | <u>+</u> | <u>B BRAUN</u> | <u>100%</u> | <u>N016734</u> | <u>001</u> | |
| <u>AT</u> | | <u>BAXTER HLTHCARE</u> | <u>100%</u> | <u>N017866</u> | <u>001</u> | |
| <u>AT</u> | | <u>ICU MEDICAL INC</u> | <u>100%</u> | <u>N017513</u> | <u>001</u> | |
| <u>AT</u> | | | <u>100%</u> | <u>N018313</u> | <u>001</u> | |

STIRIPENTOL

CAPSULE;ORAL

DIACOMIT

| | | | | | |
|------------|--------------------|--------------|----------------|------------|--------------|
| <u>+</u> | <u>BIOCODEX SA</u> | <u>250MG</u> | <u>N206709</u> | <u>001</u> | Aug 20, 2018 |
| <u>+</u> ! | | <u>500MG</u> | <u>N206709</u> | <u>002</u> | Aug 20, 2018 |

FOR SUSPENSION;ORAL

DIACOMIT

| | | | | | |
|------------|--------------------|---------------------|----------------|------------|--------------|
| <u>+</u> | <u>BIOCODEX SA</u> | <u>250MG/PACKET</u> | <u>N207223</u> | <u>001</u> | Aug 20, 2018 |
| <u>+</u> ! | | <u>500MG/PACKET</u> | <u>N207223</u> | <u>002</u> | Aug 20, 2018 |

STREPTOMYCIN SULFATE

INJECTABLE;INJECTION

STREPTOMYCIN SULFATE

| | | | | | |
|----------|---------------------|-------------------------|----------------|------------|--------------|
| <u>!</u> | <u>X GEN PHARMS</u> | <u>EQ 1GM BASE/VIAL</u> | <u>A064210</u> | <u>001</u> | Jun 30, 1998 |
|----------|---------------------|-------------------------|----------------|------------|--------------|

STREPTOZOCIN

INJECTABLE;INJECTION

ZANOSAR

| | | | | | |
|------------|------------------------|-----------------|----------------|------------|--------------|
| <u>+</u> ! | <u>TEVA PHARMS USA</u> | <u>1GM/VIAL</u> | <u>N050577</u> | <u>001</u> | May 07, 1982 |
|------------|------------------------|-----------------|----------------|------------|--------------|

STRONTIUM CHLORIDE SR-89

INJECTABLE;INJECTION

METASTRON

| | | | | | | |
|-----------|------------|----------------------|----------------|----------------|------------|--------------|
| <u>AP</u> | <u>+</u> ! | <u>GE HEALTHCARE</u> | <u>1mCi/ML</u> | <u>N020134</u> | <u>001</u> | Jun 18, 1993 |
|-----------|------------|----------------------|----------------|----------------|------------|--------------|

STRONTIUM CHLORIDE SR-89

| | | | | | | |
|-----------|--|-----------------------|----------------|----------------|------------|--------------|
| <u>AP</u> | | <u>BIO NUCLEONICS</u> | <u>1mCi/ML</u> | <u>A075941</u> | <u>001</u> | Jan 06, 2003 |
|-----------|--|-----------------------|----------------|----------------|------------|--------------|

SUCCIMER

CAPSULE;ORAL

CHEMET

| | | | | | |
|------------|-----------------------|--------------|----------------|------------|--------------|
| <u>+</u> ! | <u>RECORDATI RARE</u> | <u>100MG</u> | <u>N019998</u> | <u>002</u> | Jan 30, 1991 |
|------------|-----------------------|--------------|----------------|------------|--------------|

SUCCINYLCHOLINE CHLORIDE

INJECTABLE;INJECTION

ANECTINE

| | | | | | | |
|-----------|------------|-------------------|----------------|----------------|------------|--|
| <u>AP</u> | <u>+</u> ! | <u>SANDOZ INC</u> | <u>20MG/ML</u> | <u>N008453</u> | <u>002</u> | |
|-----------|------------|-------------------|----------------|----------------|------------|--|

QUELICIN

| | | | | | | |
|-----------|------------|----------------|----------------|----------------|------------|--|
| <u>AP</u> | <u>+</u> ! | <u>HOSPIRA</u> | <u>20MG/ML</u> | <u>N008845</u> | <u>006</u> | |
|-----------|------------|----------------|----------------|----------------|------------|--|

SUCCINYLCHOLINE CHLORIDE

| | | | | | | |
|-----------|----------|----------------------------|----------------|----------------|------------|--------------|
| <u>AP</u> | | <u>AMNEAL PHARMS CO</u> | <u>20MG/ML</u> | <u>A211432</u> | <u>001</u> | Nov 16, 2018 |
| <u>AP</u> | | <u>RENAISSANCE SSA LLC</u> | <u>20MG/ML</u> | <u>A210231</u> | <u>001</u> | Jun 04, 2018 |
| <u>AP</u> | <u>!</u> | <u>ZYDUS PHARMS USA</u> | <u>20MG/ML</u> | <u>A209467</u> | <u>001</u> | May 04, 2018 |

INC

PRESCRIPTION DRUG PRODUCT LIST

SUCRALFATE

SUSPENSION; ORAL

CARAFATE

+! ALLERGAN SALES LLC 1GM/10ML

N019183 001 Dec 16, 1993

TABLET; ORAL

CARAFATE**AB** +! ALLERGAN SALES LLC **1GM****N018333 001**SUCRALFATE**AB** TEVA **1GM****A070848 001** Mar 29, 1996SUCROFERRIC OXYHYDROXIDE

TABLET, CHEWABLE; ORAL

VELPHORO

+! VIFOR FRESENIUS 500MG

N205109 001 Nov 27, 2013

SUFENTANIL CITRATE

INJECTABLE; INJECTION

SUFENTA PRESERVATIVE FREE**AP** +! AKORN **EQ 0.05MG BASE/ML****N019050 001** May 04, 1984SUFENTANIL CITRATE**AP** HOSPIRA **EQ 0.05MG BASE/ML****A074534 001** Dec 11, 1996**AP** WEST-WARD PHARMS **EQ 0.05MG BASE/ML****A074413 001** Dec 15, 1995

INT

TABLET; SUBLINGUAL

DSUVIA

+! ACELRX PHARMS EQ 0.03MG BASE

N209128 001 Nov 02, 2018

SUGAMMADEX SODIUM

SOLUTION; INTRAVENOUS

BRIDION

+ ORGANON SUB MERCK EQ 200MG BASE/2ML (EQ 100MG BASE/ML)

N022225 002 Dec 15, 2015

+! EQ 500MG BASE/5ML (EQ 100MG BASE/ML)

N022225 001 Dec 15, 2015

SULCONAZOLE NITRATE

CREAM; TOPICAL

EXELDERM

+! JOURNEY 1%

N018737 001 Feb 28, 1989

SOLUTION; TOPICAL

EXELDERM

+! JOURNEY 1%

N018738 001 Aug 30, 1985

SULFACETAMIDE SODIUM

LOTION; TOPICAL

KLARON**AB** +! VALEANT PHARMS **10%****N019931 001** Dec 23, 1996

NORTH

SULFACETAMIDE SODIUM**AB** FOUGERA PHARMS **10%****A077015 001** Nov 17, 2006**AB** PERRIGO CO **10%****A078649 001** Mar 23, 2009

TENNESSEE

AB TARO **10%****A078668 001** May 20, 2009

OINTMENT; OPHTHALMIC

SULFACETAMIDE SODIUM

! PERRIGO CO 10%

A080029 001

TENNESSEE

SOLUTION/DROPS; OPHTHALMIC

BLEPH-10**AT** ! ALLERGAN **10%****A080028 001**SULFACETAMIDE SODIUM**AT** AKORN **10%****A040215 001** May 25, 1999**AT** BAUSCH AND LOMB **10%****A040066 001** Dec 28, 1994**AT** SANDOZ INC **10%****A089560 001** Oct 18, 1988SULFADIAZINE

TABLET; ORAL

SULFADIAZINE

! SANDOZ 500MG

A040091 001 Jul 29, 1994

SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE; INJECTION

SULFAMETHOXAZOLE AND TRIMETHOPRIM**AP** MYLAN LABS LTD **80MG/ML; 16MG/ML****A206607 001** Aug 30, 2017**AP** ! TEVA PHARMS USA **80MG/ML; 16MG/ML****A073303 001** Oct 31, 1991

SUSPENSION; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM**AB** AUROBINDO PHARMA **200MG/5ML; 40MG/5ML****A091348 001** Jun 08, 2010**AB** ! HI TECH PHARMA **200MG/5ML; 40MG/5ML****A074650 001** Dec 29, 1997

PRESCRIPTION DRUG PRODUCT LIST

SULFAMETHOXAZOLE; TRIMETHOPRIM

SUSPENSION; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM

| | | | | |
|-----------|----------------|---------------------------|--------------------|--------------|
| AB | LANNETT CO INC | 200MG;5ML;40MG/5ML | A077785 001 | Jan 24, 2007 |
|-----------|----------------|---------------------------|--------------------|--------------|

SULFATRIM PEDIATRIC

| | | | | |
|-----------|-------------|---------------------------|--------------------|--------------|
| AB | PHARM ASSOC | 200MG/5ML;40MG/5ML | N018615 001 | Jan 07, 1983 |
|-----------|-------------|---------------------------|--------------------|--------------|

TABLET; ORAL

BACTRIM

| | | | | |
|-----------|------------------------|-------------------|--------------------|--|
| AB | + SUN PHARM INDUSTRIES | 400MG;80MG | N017377 001 | |
|-----------|------------------------|-------------------|--------------------|--|

BACTRIM DS

| | | | | |
|-----------|-------------------------|--------------------|--------------------|--|
| AB | +! SUN PHARM INDUSTRIES | 800MG;160MG | N017377 002 | |
|-----------|-------------------------|--------------------|--------------------|--|

SEPTRA

| | | | | |
|-----------|----------------|-------------------|--------------------|--|
| AB | MONARCH PHARMS | 400MG;80MG | N017376 001 | |
|-----------|----------------|-------------------|--------------------|--|

SEPTRA DS

| | | | | |
|-----------|----------------|--------------------|--------------------|--|
| AB | MONARCH PHARMS | 800MG;160MG | N017376 002 | |
|-----------|----------------|--------------------|--------------------|--|

SULFAMETHOXAZOLE AND TRIMETHOPRIM

| | | | | |
|-----------|------------------|-------------------|--------------------|--------------|
| AB | AMNEAL PHARMS NY | 400MG;80MG | A076899 001 | Jan 27, 2005 |
|-----------|------------------|-------------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------------|--------------------|--------------|
| AB | | 800MG;160MG | A076899 002 | Jan 27, 2005 |
|-----------|--|--------------------|--------------------|--------------|

| | | | | |
|-----------|------------------|-------------------|--------------------|--------------|
| AB | AUROBINDO PHARMA | 400MG;80MG | A090624 001 | Feb 16, 2010 |
|-----------|------------------|-------------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------------|--------------------|--------------|
| AB | | 800MG;160MG | A090624 002 | Feb 16, 2010 |
|-----------|--|--------------------|--------------------|--------------|

| | | | | |
|-----------|---------------------|-------------------|--------------------|--------------|
| AB | CHARTWELL MOLECULES | 400MG;80MG | A078060 002 | Jan 25, 2007 |
|-----------|---------------------|-------------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------------|--------------------|--------------|
| AB | | 800MG;160MG | A078060 001 | Jan 25, 2007 |
|-----------|--|--------------------|--------------------|--------------|

| | | | | |
|-----------|-------------------|-------------------|--------------------|--------------|
| AB | GLENMARK GENERICS | 400MG;80MG | A090828 002 | Dec 22, 2010 |
|-----------|-------------------|-------------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------------|--------------------|--------------|
| AB | | 800MG;160MG | A090828 001 | Dec 22, 2010 |
|-----------|--|--------------------|--------------------|--------------|

| | | | | |
|-----------|----------------------|-------------------|--------------------|--------------|
| AB | SUN PHARM INDUSTRIES | 400MG;80MG | A071017 002 | Aug 25, 1986 |
|-----------|----------------------|-------------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------------|--------------------|--------------|
| AB | | 800MG;160MG | A071017 001 | Aug 25, 1986 |
|-----------|--|--------------------|--------------------|--------------|

| | | | | |
|-----------|--------------|-------------------|--------------------|--------------|
| AB | VISTA PHARMS | 400MG;80MG | A076817 001 | Oct 07, 2005 |
|-----------|--------------|-------------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------------|--------------------|--------------|
| AB | | 800MG;160MG | A076817 002 | Oct 07, 2005 |
|-----------|--|--------------------|--------------------|--------------|

SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH

| | | | | |
|-----------|------|--------------------|--------------------|--------------|
| AB | TEVA | 800MG;160MG | A070037 001 | Jun 02, 1987 |
|-----------|------|--------------------|--------------------|--------------|

SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH

| | | | | |
|-----------|-------------|-------------------|--------------------|--------------|
| AB | TEVA PHARMS | 400MG;80MG | A070030 001 | Jun 02, 1987 |
|-----------|-------------|-------------------|--------------------|--------------|

| | | | | |
|-----------|--|-------------------|--------------------|--------------|
| AB | | 400MG;80MG | A070030 001 | Jun 02, 1987 |
|-----------|--|-------------------|--------------------|--------------|

SULFANILAMIDE

CREAM; VAGINAL

AVC

| | | | | |
|-----------|------------------------|-----|--------------------|--------------|
| AB | +! MYLAN SPECIALITY LP | 15% | N006530 003 | Jan 27, 1987 |
|-----------|------------------------|-----|--------------------|--------------|

SULFASALAZINE

TABLET; ORAL

AZULFIDINE

| | | | | |
|-----------|-------------------------|--------------|--------------------|--|
| AB | +! PHARMACIA AND UPJOHN | 500MG | N007073 001 | |
|-----------|-------------------------|--------------|--------------------|--|

SULFASALAZINE

| | | | | |
|-----------|----------------|--------------|--------------------|--------------|
| AB | VINTAGE PHARMS | 500MG | A040349 001 | Jan 11, 2002 |
|-----------|----------------|--------------|--------------------|--------------|

| | | | | |
|-----------|-------------|--------------|--------------------|--|
| AB | WATSON LABS | 500MG | A085828 001 | |
|-----------|-------------|--------------|--------------------|--|

TABLET, DELAYED RELEASE; ORAL

AZULFIDINE EN-TABS

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| AB | +! PHARMACIA AND UPJOHN | 500MG | N007073 002 | Apr 06, 1983 |
|-----------|-------------------------|--------------|--------------------|--------------|

SULFASALAZINE

| | | | | |
|-----------|----------------|--------------|--------------------|--------------|
| AB | VINTAGE PHARMS | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|----------------|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

| | | | | |
|-----------|--|--------------|--------------------|--------------|
| AB | | 500MG | A075339 001 | Jan 11, 2002 |
|-----------|--|--------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

SUMATRIPTAN

SPRAY;NASAL

IMITREX

| | | | | | |
|-----------|-----------|------------------------|-------------------|--------------------|--------------|
| <u>AB</u> | <u>+!</u> | <u>GLAXOSMITHKLINE</u> | <u>5MG/SPRAY</u> | <u>N020626 001</u> | Aug 26, 1997 |
| <u>AB</u> | <u>+!</u> | | <u>20MG/SPRAY</u> | <u>N020626 003</u> | Aug 26, 1997 |

SUMATRIPTAN

| | | | | | |
|-----------|--|-----------------------|-------------------|--------------------|--------------|
| <u>AB</u> | | <u>LANNETT CO INC</u> | <u>5MG/SPRAY</u> | <u>A204841 001</u> | Feb 19, 2016 |
| <u>AB</u> | | | <u>20MG/SPRAY</u> | <u>A204841 002</u> | Feb 19, 2016 |

SUMATRIPTAN SUCCINATE

INJECTABLE;SUBCUTANEOUS

IMITREX STATDOSE

| | | | | | |
|-----------|-----------|------------------------|--------------------------------------------|--------------------|--------------|
| <u>AB</u> | <u>+!</u> | <u>GLAXOSMITHKLINE</u> | <u>EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)</u> | <u>N020080 002</u> | Feb 01, 2006 |
| <u>AB</u> | <u>+!</u> | | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>N020080 003</u> | Dec 23, 1996 |

SUMATRIPTAN SUCCINATE

| | | | | | |
|-----------|--|---------------------------|--------------------------------------------|--------------------|--------------|
| <u>AB</u> | | <u>ANTARES PHARMA INC</u> | <u>EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)</u> | <u>A078319 001</u> | Dec 10, 2015 |
| <u>AB</u> | | | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>A078319 002</u> | Dec 10, 2015 |
| <u>AB</u> | | <u>DR REDDYS LABS INC</u> | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>A090495 001</u> | Jan 29, 2014 |
| <u>AB</u> | | <u>SUN PHARMA GLOBAL</u> | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>A090358 001</u> | Jun 21, 2011 |

IMITREX

| | | | | | |
|-----------|-----------|------------------------|--------------------------------------------|--------------------|--------------|
| <u>AP</u> | <u>+!</u> | <u>GLAXOSMITHKLINE</u> | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>N020080 001</u> | Dec 28, 1992 |
|-----------|-----------|------------------------|--------------------------------------------|--------------------|--------------|

SUMATRIPTAN SUCCINATE

| | | | | | |
|-----------|--|-----------------------------|--------------------------------------------|--------------------|--------------|
| <u>AP</u> | | <u>AUROBINDO PHARMA LTD</u> | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>A202758 001</u> | Apr 23, 2013 |
| <u>AP</u> | | <u>FRESENIUS KABI USA</u> | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>A079242 001</u> | Mar 02, 2009 |
| <u>AP</u> | | <u>HIKMA FARMACEUTICA</u> | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>A200183 001</u> | Sep 16, 2013 |
| <u>AP</u> | | <u>MYLAN ASI</u> | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>A090314 001</u> | Jun 10, 2010 |
| <u>AP</u> | | | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>A090641 001</u> | Jul 28, 2010 |
| <u>AP</u> | | <u>MYLAN LABS LTD</u> | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>A203322 001</u> | Apr 14, 2014 |
| <u>AP</u> | | <u>PAR PHARM</u> | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>A077332 001</u> | Oct 09, 2009 |
| <u>AP</u> | | <u>PAR STERILE PRODUCTS</u> | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>A077871 001</u> | Jul 09, 2009 |
| <u>AP</u> | | <u>TEVA PHARMS USA</u> | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>A077907 001</u> | Feb 06, 2009 |
| <u>AP</u> | | <u>WEST-WARD PHARMS INT</u> | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>A079123 001</u> | Feb 06, 2009 |
| <u>AP</u> | | <u>WOCKHARDT</u> | <u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u> | <u>A078593 001</u> | Feb 06, 2009 |

POWDER;INHALATION

ONZETRA XSAIL

| | | | | |
|-----------|----------------------|---------------------|--------------------|--------------|
| <u>+!</u> | <u>AVANIR PHARMS</u> | <u>EQ 11MG BASE</u> | <u>N206099 001</u> | Jan 27, 2016 |
|-----------|----------------------|---------------------|--------------------|--------------|

SOLUTION;SUBCUTANEOUS

ZEMBRACE SYMTOUCH

| | | | | |
|----------|---------------------------|----------------------------------------------|--------------------|--------------|
| <u>+</u> | <u>DR REDDYS LABS LTD</u> | <u>EQ 3MG BASE/0.5ML (EQ 3MG BASE/0.5ML)</u> | <u>N208223 001</u> | Jan 28, 2016 |
|----------|---------------------------|----------------------------------------------|--------------------|--------------|

TABLET;ORAL

IMITREX

| | | | | | |
|-----------|-----------|------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | <u>+</u> | <u>GLAXOSMITHKLINE</u> | <u>EQ 25MG BASE</u> | <u>N020132 002</u> | Jun 01, 1995 |
| <u>AB</u> | <u>+</u> | | <u>EQ 50MG BASE</u> | <u>N020132 003</u> | Jun 01, 1995 |
| <u>AB</u> | <u>+!</u> | | <u>EQ 100MG BASE</u> | <u>N020132 001</u> | Jun 01, 1995 |

SUMATRIPTAN SUCCINATE

| | | | | | |
|-----------|--|---------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | <u>APOTEX INC</u> | <u>EQ 25MG BASE</u> | <u>A200263 001</u> | Jun 19, 2012 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A200263 002</u> | Jun 19, 2012 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A200263 003</u> | Jun 19, 2012 |
| <u>AB</u> | | <u>AUROBINDO PHARMA</u> | <u>EQ 25MG BASE</u> | <u>A078327 001</u> | Aug 10, 2009 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A078327 002</u> | Aug 10, 2009 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A078327 003</u> | Aug 10, 2009 |
| <u>AB</u> | | <u>DR REDDYS LABS INC</u> | <u>EQ 25MG BASE</u> | <u>A076847 001</u> | Aug 10, 2009 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A076847 002</u> | Aug 10, 2009 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A076847 003</u> | Aug 10, 2009 |
| <u>AB</u> | | <u>MYLAN</u> | <u>EQ 25MG BASE</u> | <u>A077744 001</u> | Aug 10, 2009 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A077744 002</u> | Aug 10, 2009 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A077744 003</u> | Aug 10, 2009 |
| <u>AB</u> | | <u>ORCHID HLTHCARE</u> | <u>EQ 25MG BASE</u> | <u>A078284 001</u> | Aug 10, 2009 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A078284 002</u> | Aug 10, 2009 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A078284 003</u> | Aug 10, 2009 |
| <u>AB</u> | | <u>SUN PHARM INDS</u> | <u>EQ 25MG BASE</u> | <u>A078295 001</u> | Aug 10, 2009 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A078295 002</u> | Aug 10, 2009 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A078295 003</u> | Aug 10, 2009 |
| <u>AB</u> | | <u>SUN PHARM INDS LTD</u> | <u>EQ 25MG BASE</u> | <u>A076554 001</u> | Aug 10, 2009 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A076554 002</u> | Aug 10, 2009 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A076572 001</u> | Feb 09, 2009 |
| <u>AB</u> | | <u>WATSON LABS</u> | <u>EQ 25MG BASE</u> | <u>A076933 001</u> | Aug 10, 2009 |
| <u>AB</u> | | | <u>EQ 50MG BASE</u> | <u>A076933 002</u> | Aug 10, 2009 |
| <u>AB</u> | | | <u>EQ 100MG BASE</u> | <u>A076933 003</u> | Aug 10, 2009 |

PRESCRIPTION DRUG PRODUCT LIST

SUNITINIB MALATE

CAPSULE; ORAL

SUTENT

| | | | | |
|---|---------|----------------|-------------|--------------|
| + | CPPI CV | EQ 12.5MG BASE | N021938 001 | Jan 26, 2006 |
| + | | EQ 25MG BASE | N021938 002 | Jan 26, 2006 |
| + | | EQ 37.5MG BASE | N021938 004 | Mar 31, 2009 |
| + | ! | EQ 50MG BASE | N021938 003 | Jan 26, 2006 |

SUVOREXANT

TABLET; ORAL

BELSOMRA

| | | | | |
|---|-------------------|------|-------------|--------------|
| + | MERCK SHARP DOHME | 5MG | N204569 001 | Aug 13, 2014 |
| + | | 10MG | N204569 002 | Aug 13, 2014 |
| + | | 15MG | N204569 003 | Aug 13, 2014 |
| + | ! | 20MG | N204569 004 | Aug 13, 2014 |

TACROLIMUS

CAPSULE; ORAL

PROGRAF

| | | | | | |
|-----------|---|----------|-----------------------------|---------------------------|--------------|
| AB | + | ASTELLAS | <u>EQ 0.5MG BASE</u> | <u>N050708 003</u> | Aug 24, 1998 |
| AB | + | | <u>EQ 1MG BASE</u> | <u>N050708 001</u> | Apr 08, 1994 |
| AB | + | ! | <u>EQ 5MG BASE</u> | <u>N050708 002</u> | Apr 08, 1994 |

TACROLIMUS

| | | | | | |
|-----------|--|--------------------|-----------------------------|---------------------------|--------------|
| AB | | ACCORD HLTHCARE | <u>EQ 0.5MG BASE</u> | <u>A091195 001</u> | Aug 31, 2011 |
| AB | | | <u>EQ 1MG BASE</u> | <u>A091195 002</u> | Aug 31, 2011 |
| AB | | | <u>EQ 5MG BASE</u> | <u>A091195 003</u> | Aug 31, 2011 |
| AB | | BELCHER PHARMS LLC | <u>EQ 0.5MG BASE</u> | <u>A206651 001</u> | Nov 30, 2017 |
| AB | | | <u>EQ 1MG BASE</u> | <u>A206651 002</u> | Nov 30, 2017 |
| AB | | | <u>EQ 5MG BASE</u> | <u>A206651 003</u> | Nov 30, 2017 |
| AB | | DR REDDYS LABS LTD | <u>EQ 0.5MG BASE</u> | <u>A090509 001</u> | May 12, 2010 |
| AB | | | <u>EQ 1MG BASE</u> | <u>A090509 002</u> | May 12, 2010 |
| AB | | | <u>EQ 5MG BASE</u> | <u>A090509 003</u> | May 12, 2010 |
| AB | | MYLAN | <u>EQ 0.5MG BASE</u> | <u>A090596 001</u> | Sep 17, 2010 |
| AB | | | <u>EQ 1MG BASE</u> | <u>A090596 002</u> | Sep 17, 2010 |
| AB | | | <u>EQ 5MG BASE</u> | <u>A090596 003</u> | Sep 17, 2010 |
| AB | | PANACEA BIOTEC LTD | <u>EQ 0.5MG BASE</u> | <u>A090802 001</u> | Sep 28, 2012 |
| AB | | | <u>EQ 1MG BASE</u> | <u>A090802 002</u> | Sep 28, 2012 |
| AB | | | <u>EQ 5MG BASE</u> | <u>A090802 003</u> | Sep 28, 2012 |
| AB | | SANDOZ | <u>EQ 0.5MG BASE</u> | <u>A065461 001</u> | Aug 10, 2009 |
| AB | | | <u>EQ 1MG BASE</u> | <u>A065461 002</u> | Aug 10, 2009 |
| AB | | | <u>EQ 5MG BASE</u> | <u>A065461 003</u> | Aug 10, 2009 |
| AB | | STRIDES PHARMA | <u>EQ 0.5MG BASE</u> | <u>A090687 001</u> | Jul 22, 2014 |
| AB | | | <u>EQ 1MG BASE</u> | <u>A090687 002</u> | Jul 22, 2014 |
| AB | | | <u>EQ 5MG BASE</u> | <u>A090687 003</u> | Jul 22, 2014 |

CAPSULE, EXTENDED RELEASE; ORAL

ASTAGRAF XL

| | | | | |
|---|----------|---------------|-------------|--------------|
| + | ASTELLAS | EQ 0.5MG BASE | N204096 001 | Jul 19, 2013 |
| + | | EQ 1MG BASE | N204096 002 | Jul 19, 2013 |
| + | ! | EQ 5MG BASE | N204096 003 | Jul 19, 2013 |

FOR SUSPENSION; ORAL

PROGRAF

| | | | | |
|---|----------|----------------------|-------------|--------------|
| + | ASTELLAS | EQ 0.2MG BASE/PACKET | N210115 001 | May 24, 2018 |
| + | ! | EQ 1MG BASE/PACKET | N210115 002 | May 24, 2018 |

INJECTABLE; INJECTION

PROGRAF

| | | | | | | |
|-----------|---|---|----------|------------------------------|---------------------------|--------------|
| AP | + | ! | ASTELLAS | <u>EQ 5MG BASE/ML</u> | <u>N050709 001</u> | Apr 08, 1994 |
|-----------|---|---|----------|------------------------------|---------------------------|--------------|

TACROLIMUS

| | | | | | | |
|-----------|--|--|-------------|------------------------------|---------------------------|--------------|
| AP | | | HOSPIRA INC | <u>EQ 5MG BASE/ML</u> | <u>A203900 001</u> | Aug 25, 2017 |
|-----------|--|--|-------------|------------------------------|---------------------------|--------------|

OINTMENT; TOPICAL

PROTOPIC

| | | | | | | |
|-----------|---|---|---------------|---------------------|---------------------------|--------------|
| AB | + | ! | LEO PHARMA AS | <u>0.03%</u> | <u>N050777 001</u> | Dec 08, 2000 |
| AB | + | ! | | <u>0.1%</u> | <u>N050777 002</u> | Dec 08, 2000 |

TACROLIMUS

| | | | | | | |
|-----------|--|--|---------------------|---------------------|---------------------------|--------------|
| AB | | | FOUGERA PHARMS INC | <u>0.03%</u> | <u>A200744 001</u> | Sep 09, 2014 |
| AB | | | | <u>0.1%</u> | <u>A200744 002</u> | Sep 09, 2014 |
| AB | | | GLENMARK PHARMS LTD | <u>0.1%</u> | <u>A210393 001</u> | Apr 16, 2018 |

TABLET, EXTENDED RELEASE; ORAL

ENVARUS XR

| | | | | |
|---|--------------------|----------------|-------------|--------------|
| + | VELOXIS PHARMS INC | EQ 0.75MG BASE | N206406 001 | Jul 10, 2015 |
| + | | EQ 1MG BASE | N206406 002 | Jul 10, 2015 |
| + | ! | EQ 4MG BASE | N206406 003 | Jul 10, 2015 |

PRESCRIPTION DRUG PRODUCT LIST

TADALAFIL

TABLET; ORAL

CIALIS

| | | | | | |
|-----------|---|-------|--------------|--------------------|--------------|
| AB | + | LILLY | 2.5MG | N021368 004 | Jan 07, 2008 |
| AB | + | | 5MG | N021368 001 | Nov 21, 2003 |
| AB | + | | 10MG | N021368 002 | Nov 21, 2003 |

TADALAFIL

| | | | | | |
|-----------|--|-----------------|--------------|--------------------|--------------|
| AB | | TEVA PHARMS USA | 2.5MG | A090141 001 | May 22, 2018 |
| AB | | | 5MG | A090141 002 | May 22, 2018 |
| AB | | | 10MG | A090141 003 | May 22, 2018 |

CIALIS

| | | | | | | |
|------------|---|---|-------|-------------|--------------------|--------------|
| AB1 | + | ! | LILLY | 20MG | N021368 003 | Nov 21, 2003 |
|------------|---|---|-------|-------------|--------------------|--------------|

TADALAFIL

| | | | | | |
|------------|--|-----------------|-------------|--------------------|--------------|
| AB1 | | TEVA PHARMS USA | 20MG | A090141 004 | May 22, 2018 |
|------------|--|-----------------|-------------|--------------------|--------------|

ADCIRCA

| | | | | | | |
|------------|---|---|--------------|-------------|--------------------|--------------|
| AB2 | + | ! | ELI LILLY CO | 20MG | N022332 001 | May 22, 2009 |
|------------|---|---|--------------|-------------|--------------------|--------------|

TADALAFIL

| | | | | | |
|------------|--|------------------|-------------|--------------------|--------------|
| AB2 | | MYLAN PHARMS INC | 20MG | A200630 001 | Aug 03, 2018 |
|------------|--|------------------|-------------|--------------------|--------------|

TAFENOQUINE SUCCINATE

TABLET; ORAL

ARAKODA

| | | | | | | |
|--|---|---|-------------------|---------------|-------------|--------------|
| | + | ! | 60 DEGREES PHARMS | EQ 100MG BASE | N210607 001 | Aug 08, 2018 |
|--|---|---|-------------------|---------------|-------------|--------------|

KRINTAFEL

| | | | | | | |
|--|---|---|-----------------|---------------|-------------|--------------|
| | + | ! | GLAXOSMITHKLINE | EQ 150MG BASE | N210795 001 | Jul 20, 2018 |
|--|---|---|-----------------|---------------|-------------|--------------|

TAFLUPROST

SOLUTION/DROPS; OPHTHALMIC

ZIOPTAN

| | | | | | | |
|--|---|---|----------------|---------|-------------|--------------|
| | + | ! | OAK PHARMS INC | 0.0015% | N202514 001 | Feb 10, 2012 |
|--|---|---|----------------|---------|-------------|--------------|

TALAZOPARIB TOSYLATE

CAPSULE; ORAL

TALZENNA

| | | | | | | |
|--|---|--|------------|----------------|-------------|--------------|
| | + | | PFIZER INC | EQ 0.25MG BASE | N211651 001 | Oct 16, 2018 |
|--|---|--|------------|----------------|-------------|--------------|

| | | | | | | |
|--|---|---|--|-------------|-------------|--------------|
| | + | ! | | EQ 1MG BASE | N211651 002 | Oct 16, 2018 |
|--|---|---|--|-------------|-------------|--------------|

TALC

AEROSOL; INTRAPLEURAL

SCLEROSOL

| | | | | | | |
|--|---|---|-------------|-----------|-------------|--------------|
| | + | ! | LYMOL MEDCL | 4GM/SPRAY | N020587 001 | Dec 24, 1997 |
|--|---|---|-------------|-----------|-------------|--------------|

POWDER; INTRAPLEURAL

STERITALC

| | | | | | | |
|--|---|--|-------------|----------|-------------|--------------|
| | + | | NOVATECH SA | 2GM/VIAL | N205555 001 | May 01, 2017 |
|--|---|--|-------------|----------|-------------|--------------|

| | | | | | | |
|--|---|--|--|----------|-------------|--------------|
| | + | | | 3GM/VIAL | N205555 002 | May 01, 2017 |
|--|---|--|--|----------|-------------|--------------|

| | | | | | | |
|--|---|---|--|----------|-------------|--------------|
| | + | ! | | 4GM/VIAL | N205555 003 | May 01, 2017 |
|--|---|---|--|----------|-------------|--------------|

TALC

| | | | | | | |
|--|---|---|-------------|---------|-------------|--------------|
| | + | ! | LYMOL MEDCL | 5GM/BOT | N021388 001 | Dec 15, 2003 |
|--|---|---|-------------|---------|-------------|--------------|

TALIGLUCERASE ALFA

POWDER; INTRAVENOUS

ELELYSO

| | | | | | | |
|--|---|---|--------|----------------|-------------|--------------|
| | + | ! | PFIZER | 200 UNITS/VIAL | N022458 001 | May 01, 2012 |
|--|---|---|--------|----------------|-------------|--------------|

TAMOXIFEN CITRATE

SOLUTION; ORAL

SOLTAMOX

| | | | | | | |
|--|--|--|--------------------|-------------------|-------------|--------------|
| | | | MIDATECH PHARMA US | EQ 20MG BASE/10ML | N021807 001 | Oct 29, 2005 |
|--|--|--|--------------------|-------------------|-------------|--------------|

TABLET; ORAL

TAMOXIFEN CITRATE

| | | | | | | |
|-----------|--|--|---------------------|---------------------|--------------------|--------------|
| AB | | | ACTAVIS LABS FL INC | EQ 10MG BASE | A070929 001 | Feb 20, 2003 |
|-----------|--|--|---------------------|---------------------|--------------------|--------------|

| | | | | | | |
|-----------|--|--|--|---------------------|--------------------|--------------|
| AB | | | | EQ 20MG BASE | A070929 002 | Feb 20, 2003 |
|-----------|--|--|--|---------------------|--------------------|--------------|

| | | | | | | |
|-----------|--|--|--------|---------------------|--------------------|--------------|
| AB | | | APOTEX | EQ 10MG BASE | A090878 001 | Sep 23, 2011 |
|-----------|--|--|--------|---------------------|--------------------|--------------|

| | | | | | | |
|-----------|--|--|--|---------------------|--------------------|--------------|
| AB | | | | EQ 20MG BASE | A090878 002 | Sep 23, 2011 |
|-----------|--|--|--|---------------------|--------------------|--------------|

| | | | | | | |
|-----------|--|--|--------------|---------------------|--------------------|--------------|
| AB | | | MAYNE PHARMA | EQ 10MG BASE | A075797 001 | Feb 20, 2003 |
|-----------|--|--|--------------|---------------------|--------------------|--------------|

| | | | | | | |
|-----------|---|--|--|---------------------|--------------------|--------------|
| AB | ! | | | EQ 20MG BASE | A075797 002 | Feb 20, 2003 |
|-----------|---|--|--|---------------------|--------------------|--------------|

| | | | | | | |
|-----------|--|--|-------|---------------------|--------------------|--------------|
| AB | | | MYLAN | EQ 10MG BASE | A074732 002 | Feb 20, 2003 |
|-----------|--|--|-------|---------------------|--------------------|--------------|

| | | | | | | |
|-----------|--|--|--|---------------------|--------------------|--------------|
| AB | | | | EQ 20MG BASE | A074732 001 | Feb 20, 2003 |
|-----------|--|--|--|---------------------|--------------------|--------------|

| | | | | | | |
|-----------|--|--|------------------|---------------------|--------------------|--------------|
| AB | | | ZYDUS PHARMS USA | EQ 10MG BASE | A206694 001 | Oct 27, 2017 |
|-----------|--|--|------------------|---------------------|--------------------|--------------|

| | | | | | | |
|-----------|--|--|-----|--|--|--|
| AB | | | INC | | | |
|-----------|--|--|-----|--|--|--|

| | | | | | | |
|-----------|--|--|--|---------------------|--------------------|--------------|
| AB | | | | EQ 20MG BASE | A206694 002 | Oct 27, 2017 |
|-----------|--|--|--|---------------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

TAMSULOSIN HYDROCHLORIDE

CAPSULE; ORAL

FLOMAX

| | | | | | | |
|-----------|------------|-------------------|--------------|----------------|------------|--------------|
| AB | + ! | SANOFI AVENTIS US | 0.4MG | N020579 | 001 | Apr 15, 1997 |
|-----------|------------|-------------------|--------------|----------------|------------|--------------|

TAMSULOSIN HYDROCHLORIDE

| | | | | | | |
|-----------|--|----------------------|--------------|----------------|------------|--------------|
| AB | | ALKEM LABS LTD | 0.4MG | A207405 | 001 | Aug 11, 2017 |
| AB | | ANCHEN PHARMS | 0.4MG | A202010 | 001 | Jan 04, 2013 |
| AB | | AUROBINDO PHARMA LTD | 0.4MG | A202433 | 001 | Apr 30, 2013 |
| AB | | IMPAX LABS | 0.4MG | A090377 | 001 | Mar 02, 2010 |
| AB | | MACLEODS PHARMS LTD | 0.4MG | A204645 | 001 | Jan 20, 2017 |
| AB | | MYLAN | 0.4MG | A090408 | 001 | Apr 27, 2010 |
| AB | | SANDOZ | 0.4MG | A078015 | 001 | Apr 27, 2010 |
| AB | | SUN PHARM INDS LTD | 0.4MG | A090931 | 001 | Jul 15, 2010 |
| AB | | SYNTHON PHARMS | 0.4MG | A078801 | 001 | Apr 27, 2010 |
| AB | | TEVA PHARMS | 0.4MG | A077630 | 001 | Apr 27, 2010 |
| AB | | WOCKHARDT | 0.4MG | A078938 | 001 | Apr 27, 2010 |
| AB | | ZYDUS PHARMS USA INC | 0.4MG | A078225 | 001 | Apr 27, 2010 |

TAPENTADOL HYDROCHLORIDE

TABLET; ORAL

NUCYNTA

| | | | | | |
|---|---------|---------------|---------|-----|--------------|
| + | DEPO NF | EQ 50MG BASE | N022304 | 001 | Nov 20, 2008 |
| + | | EQ 75MG BASE | N022304 | 002 | Nov 20, 2008 |
| + | ! | EQ 100MG BASE | N022304 | 003 | Nov 20, 2008 |

TABLET, EXTENDED RELEASE; ORAL

NUCYNTA ER

| | | | | | |
|---|---------|---------------|---------|-----|--------------|
| + | DEPO NF | EQ 50MG BASE | N200533 | 001 | Aug 25, 2011 |
| + | | EQ 100MG BASE | N200533 | 002 | Aug 25, 2011 |
| + | | EQ 150MG BASE | N200533 | 003 | Aug 25, 2011 |
| + | | EQ 200MG BASE | N200533 | 004 | Aug 25, 2011 |
| + | ! | EQ 250MG BASE | N200533 | 005 | Aug 25, 2011 |

TASIMELTEON

CAPSULE; ORAL

HETLIOZ

| | | | | | | |
|---|---|------------------|------|---------|-----|--------------|
| + | ! | VANDA PHARMS INC | 20MG | N205677 | 001 | Jan 31, 2014 |
|---|---|------------------|------|---------|-----|--------------|

TAVABOROLE

SOLUTION; TOPICAL

KERYDIN

| | | | | | | |
|---|---|-------------------|----|---------|-----|--------------|
| + | ! | ANACOR PHARMS INC | 5% | N204427 | 001 | Jul 07, 2014 |
|---|---|-------------------|----|---------|-----|--------------|

TAZAROTENE

AEROSOL, FOAM; TOPICAL

FABIOR

| | | | | | | |
|---|---|--------------|------|---------|-----|--------------|
| + | ! | MAYNE PHARMA | 0.1% | N202428 | 001 | May 11, 2012 |
|---|---|--------------|------|---------|-----|--------------|

CREAM; TOPICAL

AVAGE

| | | | | | | |
|-----------|------------|----------|-------------|----------------|------------|--------------|
| AB | + ! | ALLERGAN | 0.1% | N021184 | 003 | Sep 30, 2002 |
|-----------|------------|----------|-------------|----------------|------------|--------------|

TAZAROTENE

| | | | | | | |
|-----------|--|------------------|-------------|----------------|------------|--------------|
| AB | | G AND W LABS INC | 0.1% | A208662 | 001 | Dec 22, 2017 |
|-----------|--|------------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-------------|-------------|----------------|------------|--------------|
| AB | | TARO PHARMS | 0.1% | A208258 | 001 | Apr 03, 2017 |
|-----------|--|-------------|-------------|----------------|------------|--------------|

TAZORAC

| | | | | | | |
|-----------|------------|----------|-------------|----------------|------------|--------------|
| AB | + ! | ALLERGAN | 0.1% | N021184 | 002 | Sep 29, 2000 |
|-----------|------------|----------|-------------|----------------|------------|--------------|

| | | | | | | |
|---|---|--|-------|---------|-----|--------------|
| + | ! | | 0.05% | N021184 | 001 | Sep 29, 2000 |
|---|---|--|-------|---------|-----|--------------|

GEL; TOPICAL

TAZORAC

| | | | | | | |
|---|---|----------|-------|---------|-----|--------------|
| + | ! | ALLERGAN | 0.05% | N020600 | 001 | Jun 13, 1997 |
|---|---|----------|-------|---------|-----|--------------|

| | | | | | | |
|---|---|--|------|---------|-----|--------------|
| + | ! | | 0.1% | N020600 | 002 | Jun 13, 1997 |
|---|---|--|------|---------|-----|--------------|

TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

INJECTABLE; INJECTION

TECHNETIUM TC 99M ALBUMIN AGGREGATED KIT

| | | | | | | | |
|----|---|---|-----------|-----|---------|-----|--------------|
| BS | + | ! | DRAXIMAGE | N/A | N017881 | 001 | Dec 30, 1987 |
|----|---|---|-----------|-----|---------|-----|--------------|

TECHNETIUM TC-99M BICISATE KIT

INJECTABLE; INJECTION

NEUROLITE

| | | | | | | |
|---|---|----------------|-----|---------|-----|--------------|
| + | ! | LANTHEUS MEDCL | N/A | N020256 | 001 | Nov 23, 1994 |
|---|---|----------------|-----|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

TECHNETIUM TC-99M DISOFENIN KIT

INJECTABLE; INJECTION

HEPATOLITE

PHARMALUCENCE N/A

N018467 001 Mar 16, 1982

TECHNETIUM TC-99M EXAMETAZIME KIT

INJECTABLE; INJECTION

CERETEC

+! GE HEALTHCARE N/A

N019829 001 Dec 30, 1988

POWDER; INTRAVENOUS

DRAX EXAMETAZIME

JUBILANT DRAXIMAGE N/A

N208870 001 Aug 17, 2017

TECHNETIUM TC-99M MEBROFENIN KIT

INJECTABLE; INJECTION

CHOLETEC**AP +! BRACCO N/A****N018963 001** Jan 21, 1987**TECHNETIUM TC-99M MEBROFENIN****AP PHARMALUCENCE N/A****A078242 001** Jan 29, 2008TECHNETIUM TC-99M MEDRONATE

INJECTABLE; INJECTION

DRAXIMAGE MDP-25

+! JUBILANT DRAXIMAGE N/A

N018035 002 Feb 27, 2004

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION

CIS-MDP

PHARMALUCENCE N/A

N018124 001

TECHNETIUM TC-99M MERTIATIDE KIT

INJECTABLE; INJECTION

TECHNESCAN MAG3

+! MALLINKRODT NUCLEAR N/A

N019882 001 Jun 15, 1990

TECHNETIUM TC-99M OXIDRONATE KIT

INJECTABLE; INJECTION

TECHNESCAN

+! MALLINKRODT NUCLEAR N/A

N018321 001

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION

DTPA

+! JUBILANT DRAXIMAGE N/A

N018511 001 Dec 29, 1989

TECHNETIUM TC-99M PYROPHOSPHATE KIT

INJECTABLE; INJECTION

CIS-PYRO**AP PHARMALUCENCE N/A****N019039 001** Jun 30, 1987**TECHNESCAN PYP KIT****AP MALLINKRODT NUCLEAR N/A****N017538 001**TECHNETIUM TC-99M RED BLOOD CELL KIT

INJECTABLE; INJECTION

ULTRATAG

+! MALLINKRODT NUCLEAR N/A

N019981 001 Jun 10, 1991

TECHNETIUM TC-99M SESTAMIBI KIT

INJECTABLE; INJECTION

CARDIOLITE**AP +! LANTHEUS MEDCL N/A****N019785 001** Dec 21, 1990**TECHNETIUM TC 99M SESTAMIBI****AP CARDINAL HEALTH 414 N/A****A078809 001** Apr 28, 2009**AP JUBILANT DRAXIMAGE N/A****A078806 001** Apr 29, 2009**AP MALLINKRODT NUCLEAR N/A****A078098 001** Sep 22, 2008**AP PHARMALUCENCE 10-30mCi****A079157 001** Jul 10, 2009TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION; INTRAVENOUS

TECHNELITE

+! LANTHEUS MEDCL 1-20 CI/GENERATOR

N017771 002 Feb 12, 2014

ULTRA-TECHNEKOW FM

+! MALLINKRODT NUCLEAR 1-19 CI/GENERATOR

N017243 003 Feb 18, 2014

SOLUTION; INTRAVENOUS, INTRAVESICULAR, OPHTHALMIC

RADIOGENIX SYSTEM

+ NORTHSTAR MEDICAL 30-1153mCi/GENERATOR

N202158 001 Feb 08, 2018

PRESCRIPTION DRUG PRODUCT LIST

TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION;INTRAVENOUS, ORAL

TECHNETIUM TC 99M GENERATOR

+! GE HEALTHCARE 68-2703mCi/GENERATOR

N017693 002 Dec 13, 2013

TECHNETIUM TC-99M SULFUR COLLOID KIT

SOLUTION;INJECTION, ORAL

AN-SULFUR COLLOID

+! PHARMALUCENCE N/A

N017858 001

TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE;INJECTION

MYOVUEW 30ML

+! GE HEALTHCARE N/A

N020372 002 Jul 07, 2005

TECHNETIUM TC-99M TILMANOCEPT

INJECTABLE;INJECTION

LYMPHOSEEK KIT

+! CARDINAL HEALTH 414 N/A

N202207 001 Mar 13, 2013

TECOVIRIMAT

CAPSULE;ORAL

TPOXX

+! SIGA TECHNOLOGIES 200MG

N208627 001 Jul 13, 2018

TEDIZOLID PHOSPHATE

POWDER;INTRAVENOUS

SIVEXTRO

+! CUBIST PHARMS LLC 200MG/VIAL

N205436 001 Jun 20, 2014

TABLET;ORAL

SIVEXTRO

+! CUBIST PHARMS LLC 200MG

N205435 001 Jun 20, 2014

TEDUGLUTIDE RECOMBINANT

POWDER;SUBCUTANEOUS

GATTEX KIT

+! NPS PHARMS INC 5MG/VIAL

N203441 001 Dec 21, 2012

TELAVANCIN HYDROCHLORIDE

POWDER;INTRAVENOUS

VIBATIV

+! CUMBERLAND PHARMS EQ 750MG BASE/VIAL

N022110 002 Sep 11, 2009

TELMISARTAN

TABLET;ORAL

MICARDIS**AB** + BOEHRINGER
INGELHEIM**20MG****N020850 003** Apr 04, 2000**AB** +**40MG****N020850 001** Nov 10, 1998**AB** +!**80MG****N020850 002** Nov 10, 1998**TELMISARTAN****AB** ALEMbic PHARMS LTD**20MG****A202130 001** Jul 07, 2014**AB****40MG****A202130 002** Jul 07, 2014**AB****80MG****A202130 003** Jul 07, 2014**AB** AMNEAL PHARMS**20MG****A204415 001** Sep 08, 2015**AB****40MG****A204415 002** Sep 08, 2015**AB****80MG****A204415 003** Sep 08, 2015**AB** AUROBINDO PHARMA
LTD**20MG****A206511 001** Sep 03, 2015**AB****40MG****A206511 002** Sep 03, 2015**AB****80MG****A206511 003** Sep 03, 2015**AB** CADILA PHARMS LTD**20MG****A208605 001** Jul 25, 2017**AB****40MG****A208605 002** Jul 25, 2017**AB****80MG****A208605 003** Jul 25, 2017**AB** GLENMARK PHARMS LTD**20MG****A090032 001** Jul 07, 2014**AB****40MG****A090032 002** Jul 07, 2014**AB****80MG****A090032 003** Jul 07, 2014**AB** HETERO LABS LTD V**20MG****A205901 001** Apr 22, 2016**AB****40MG****A205901 002** Apr 22, 2016**AB****80MG****A205901 003** Apr 22, 2016**AB** INVENTIA HLTHCARE**20MG****A205150 001** Oct 30, 2015**AB****40MG****A205150 002** Oct 30, 2015**AB****80MG****A205150 003** Oct 30, 2015**AB** JUBILANT GENERICS**20MG****A204164 001** Aug 22, 2016**AB****40MG****A204164 002** Aug 22, 2016**AB****80MG****A204164 003** Aug 22, 2016**AB** MICRO LABS**20MG****A207016 001** Oct 03, 2017

PRESCRIPTION DRUG PRODUCT LIST

TELMISARTAN

TABLET; ORAL

TELMISARTAN

| | | | | | |
|-----------|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | <u>40MG</u> | <u>A207016</u> | <u>002</u> | Oct 03, 2017 |
| <u>AB</u> | | <u>80MG</u> | <u>A207016</u> | <u>003</u> | Oct 03, 2017 |
| <u>AB</u> | MYLAN PHARMS INC | <u>20MG</u> | <u>A202397</u> | <u>001</u> | Jul 07, 2014 |
| <u>AB</u> | | <u>40MG</u> | <u>A202397</u> | <u>002</u> | Jul 07, 2014 |
| <u>AB</u> | | <u>80MG</u> | <u>A202397</u> | <u>003</u> | Jul 07, 2014 |
| <u>AB</u> | PRINSTON INC | <u>20MG</u> | <u>A207882</u> | <u>001</u> | May 03, 2017 |
| <u>AB</u> | | <u>40MG</u> | <u>A207882</u> | <u>002</u> | May 03, 2017 |
| <u>AB</u> | | <u>80MG</u> | <u>A207882</u> | <u>003</u> | May 03, 2017 |
| <u>AB</u> | SANDOZ INC | <u>20MG</u> | <u>A203867</u> | <u>001</u> | Nov 03, 2014 |
| <u>AB</u> | | <u>40MG</u> | <u>A203867</u> | <u>002</u> | Nov 03, 2014 |
| <u>AB</u> | | <u>80MG</u> | <u>A203867</u> | <u>003</u> | Nov 03, 2014 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>20MG</u> | <u>A203171</u> | <u>001</u> | Jul 07, 2014 |
| <u>AB</u> | | <u>40MG</u> | <u>A203171</u> | <u>002</u> | Jul 07, 2014 |
| <u>AB</u> | | <u>80MG</u> | <u>A203171</u> | <u>003</u> | Jul 07, 2014 |
| <u>AB</u> | WATSON LABS | <u>20MG</u> | <u>A078710</u> | <u>001</u> | Jan 08, 2014 |
| <u>AB</u> | | <u>80MG</u> | <u>A078710</u> | <u>003</u> | Jan 08, 2014 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>20MG</u> | <u>A203325</u> | <u>001</u> | Aug 26, 2014 |
| <u>AB</u> | | <u>40MG</u> | <u>A203325</u> | <u>002</u> | Aug 26, 2014 |
| <u>AB</u> | | <u>80MG</u> | <u>A203325</u> | <u>003</u> | Aug 26, 2014 |

TELOTRISTAT ETIPRATE

TABLET; ORAL

XERMELO

+! LEXICON PHARMS INC EQ 250MG BASE N208794 001 Feb 28, 2017

TEMAZEPAM

CAPSULE; ORAL

RESTORIL

| | | | | | | |
|-----------|---|------------|---------------|----------------|------------|--------------|
| <u>AB</u> | + | SPECGX LLC | <u>7.5MG</u> | <u>N018163</u> | <u>003</u> | Oct 25, 1991 |
| <u>AB</u> | + | | <u>15MG</u> | <u>N018163</u> | <u>001</u> | |
| <u>AB</u> | + | | <u>22.5MG</u> | <u>N018163</u> | <u>004</u> | Nov 02, 2004 |
| <u>AB</u> | + | | <u>30MG</u> | <u>N018163</u> | <u>002</u> | |

TEMAZEPAM

| | | | | | | |
|-----------|--|-------------------------|---------------|----------------|------------|--------------|
| <u>AB</u> | | ACTAVIS ELIZABETH | <u>15MG</u> | <u>A071620</u> | <u>002</u> | Aug 07, 1987 |
| <u>AB</u> | | | <u>30MG</u> | <u>A071620</u> | <u>001</u> | Aug 07, 1987 |
| <u>AB</u> | | ALEMBIC PHARMS LTD | <u>7.5MG</u> | <u>A211542</u> | <u>001</u> | Nov 23, 2018 |
| <u>AB</u> | | | <u>15MG</u> | <u>A211542</u> | <u>002</u> | Nov 23, 2018 |
| <u>AB</u> | | | <u>22.5MG</u> | <u>A211542</u> | <u>003</u> | Nov 23, 2018 |
| <u>AB</u> | | | <u>30MG</u> | <u>A211542</u> | <u>004</u> | Nov 23, 2018 |
| <u>AB</u> | | AMNEAL PHARMS | <u>7.5MG</u> | <u>A203482</u> | <u>001</u> | May 23, 2016 |
| <u>AB</u> | | | <u>15MG</u> | <u>A203482</u> | <u>002</u> | May 23, 2016 |
| <u>AB</u> | | | <u>22.5MG</u> | <u>A203482</u> | <u>003</u> | May 23, 2016 |
| <u>AB</u> | | | <u>30MG</u> | <u>A203482</u> | <u>004</u> | May 23, 2016 |
| <u>AB</u> | | MYLAN | <u>7.5MG</u> | <u>A070920</u> | <u>002</u> | May 21, 2010 |
| <u>AB</u> | | | <u>15MG</u> | <u>A070920</u> | <u>004</u> | Jul 07, 1986 |
| <u>AB</u> | | | <u>22.5MG</u> | <u>A070920</u> | <u>003</u> | Jun 12, 2009 |
| <u>AB</u> | | | <u>30MG</u> | <u>A070920</u> | <u>001</u> | Jul 10, 1986 |
| <u>AB</u> | | NOVEL LABS INC | <u>7.5MG</u> | <u>A071457</u> | <u>002</u> | Jun 22, 2012 |
| <u>AB</u> | | | <u>15MG</u> | <u>A071456</u> | <u>001</u> | Apr 21, 1987 |
| <u>AB</u> | | | <u>22.5MG</u> | <u>A071457</u> | <u>003</u> | Jun 22, 2012 |
| <u>AB</u> | | | <u>30MG</u> | <u>A071457</u> | <u>001</u> | Apr 21, 1987 |
| <u>AB</u> | | PRINSTON INC | <u>7.5MG</u> | <u>A201781</u> | <u>001</u> | Jun 04, 2015 |
| <u>AB</u> | | | <u>15MG</u> | <u>A201781</u> | <u>002</u> | Jun 04, 2015 |
| <u>AB</u> | | | <u>22.5MG</u> | <u>A201781</u> | <u>003</u> | Jun 04, 2015 |
| <u>AB</u> | | | <u>30MG</u> | <u>A201781</u> | <u>004</u> | Jun 04, 2015 |
| <u>AB</u> | | SANDOZ | <u>15MG</u> | <u>A071427</u> | <u>001</u> | Jan 12, 1988 |
| <u>AB</u> | | | <u>30MG</u> | <u>A071428</u> | <u>001</u> | Jan 12, 1988 |
| <u>AB</u> | | SUN PHARM INDUSTRIES | <u>7.5MG</u> | <u>A078581</u> | <u>001</u> | Sep 08, 2009 |
| <u>AB</u> | | | <u>22.5MG</u> | <u>A071175</u> | <u>002</u> | Sep 14, 2009 |

TEMOZOLOMIDE

CAPSULE; ORAL

TEMODAR

| | | | | | | |
|-----------|---|-------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | MERCK SHARP DOHME | <u>5MG</u> | <u>N021029</u> | <u>001</u> | Aug 11, 1999 |
| <u>AB</u> | + | | <u>20MG</u> | <u>N021029</u> | <u>002</u> | Aug 11, 1999 |
| <u>AB</u> | + | | <u>100MG</u> | <u>N021029</u> | <u>003</u> | Aug 11, 1999 |
| <u>AB</u> | + | | <u>140MG</u> | <u>N021029</u> | <u>005</u> | Oct 19, 2006 |
| <u>AB</u> | + | | <u>180MG</u> | <u>N021029</u> | <u>006</u> | Oct 19, 2006 |
| <u>AB</u> | + | | <u>250MG</u> | <u>N021029</u> | <u>004</u> | Aug 11, 1999 |

PRESCRIPTION DRUG PRODUCT LIST

TEMOZOLOMIDE

CAPSULE; ORAL

TEMOZOLOMIDE

| | | | | |
|-----------|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>5MG</u> | <u>A201528 001</u> | Feb 27, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A201528 002</u> | Feb 27, 2017 |
| <u>AB</u> | | <u>100MG</u> | <u>A201528 003</u> | Feb 27, 2017 |
| <u>AB</u> | | <u>140MG</u> | <u>A201528 004</u> | Feb 27, 2017 |
| <u>AB</u> | | <u>180MG</u> | <u>A201528 005</u> | Feb 27, 2017 |
| <u>AB</u> | | <u>250MG</u> | <u>A201528 006</u> | Feb 27, 2017 |
| <u>AB</u> | AMERIGEN PHARMS LTD | <u>5MG</u> | <u>A203490 001</u> | Jul 13, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A203490 002</u> | Jul 13, 2016 |
| <u>AB</u> | | <u>100MG</u> | <u>A203490 003</u> | Jul 13, 2016 |
| <u>AB</u> | | <u>140MG</u> | <u>A203490 004</u> | Jul 13, 2016 |
| <u>AB</u> | | <u>180MG</u> | <u>A203490 005</u> | Jul 13, 2016 |
| <u>AB</u> | | <u>250MG</u> | <u>A203490 006</u> | Jul 13, 2016 |
| <u>AB</u> | AMNEAL PHARMS | <u>5MG</u> | <u>A203691 001</u> | May 08, 2015 |
| <u>AB</u> | | <u>20MG</u> | <u>A203691 002</u> | May 08, 2015 |
| <u>AB</u> | | <u>100MG</u> | <u>A203691 003</u> | May 08, 2015 |
| <u>AB</u> | | <u>140MG</u> | <u>A203691 004</u> | May 08, 2015 |
| <u>AB</u> | | <u>180MG</u> | <u>A203691 005</u> | May 08, 2015 |
| <u>AB</u> | | <u>250MG</u> | <u>A203691 006</u> | May 08, 2015 |
| <u>AB</u> | APOTEX INC | <u>5MG</u> | <u>A204159 001</u> | Jul 05, 2018 |
| <u>AB</u> | | <u>20MG</u> | <u>A204159 002</u> | Jul 05, 2018 |
| <u>AB</u> | | <u>100MG</u> | <u>A204159 003</u> | Jul 05, 2018 |
| <u>AB</u> | | <u>140MG</u> | <u>A204159 004</u> | Jul 05, 2018 |
| <u>AB</u> | | <u>180MG</u> | <u>A204159 005</u> | Jul 05, 2018 |
| <u>AB</u> | | <u>250MG</u> | <u>A204159 006</u> | Jul 05, 2018 |
| <u>AB</u> | BARR | <u>5MG</u> | <u>A078879 001</u> | Mar 01, 2010 |
| <u>AB</u> | | <u>20MG</u> | <u>A078879 002</u> | Mar 01, 2010 |
| <u>AB</u> | | <u>100MG</u> | <u>A078879 003</u> | Mar 01, 2010 |
| <u>AB</u> | | <u>140MG</u> | <u>A078879 005</u> | Mar 01, 2010 |
| <u>AB</u> | | <u>180MG</u> | <u>A078879 006</u> | Mar 01, 2010 |
| <u>AB</u> | | <u>250MG</u> | <u>A078879 004</u> | Mar 01, 2010 |
| <u>AB</u> | CHEMI SPA | <u>5MG</u> | <u>A204639 001</u> | Nov 23, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A204639 002</u> | Nov 23, 2016 |
| <u>AB</u> | | <u>100MG</u> | <u>A204639 003</u> | Nov 23, 2016 |
| <u>AB</u> | | <u>140MG</u> | <u>A204639 004</u> | Nov 23, 2016 |
| <u>AB</u> | | <u>180MG</u> | <u>A204639 005</u> | Nov 23, 2016 |
| <u>AB</u> | | <u>250MG</u> | <u>A204639 006</u> | Nov 23, 2016 |
| <u>AB</u> | DEVA HOLDING AS | <u>5MG</u> | <u>A207658 001</u> | Apr 26, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A207658 002</u> | Apr 26, 2017 |
| <u>AB</u> | | <u>100MG</u> | <u>A207658 003</u> | Apr 26, 2017 |
| <u>AB</u> | | <u>140MG</u> | <u>A207658 004</u> | Apr 26, 2017 |
| <u>AB</u> | | <u>180MG</u> | <u>A207658 005</u> | Apr 26, 2017 |
| <u>AB</u> | | <u>250MG</u> | <u>A207658 006</u> | Apr 26, 2017 |
| <u>AB</u> | IDT AUSTRALIA LTD | <u>5MG</u> | <u>A206413 001</u> | Apr 12, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A206413 002</u> | Apr 12, 2016 |
| <u>AB</u> | | <u>100MG</u> | <u>A206413 003</u> | Apr 12, 2016 |
| <u>AB</u> | | <u>140MG</u> | <u>A206413 004</u> | Apr 12, 2016 |
| <u>AB</u> | | <u>180MG</u> | <u>A206413 005</u> | Apr 12, 2016 |
| <u>AB</u> | | <u>250MG</u> | <u>A206413 006</u> | Apr 12, 2016 |
| <u>AB</u> | LANNETT CO INC | <u>5MG</u> | <u>A203898 001</u> | Feb 10, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A203898 002</u> | Feb 10, 2016 |
| <u>AB</u> | | <u>100MG</u> | <u>A203898 003</u> | Feb 10, 2016 |
| <u>AB</u> | | <u>140MG</u> | <u>A203898 004</u> | Feb 10, 2016 |
| <u>AB</u> | | <u>180MG</u> | <u>A203898 005</u> | Feb 10, 2016 |
| <u>AB</u> | | <u>250MG</u> | <u>A203898 006</u> | Feb 10, 2016 |
| <u>AB</u> | RISING PHARMS | <u>5MG</u> | <u>A206309 001</u> | Apr 27, 2016 |
| <u>AB</u> | | <u>20MG</u> | <u>A206309 002</u> | Apr 27, 2016 |
| <u>AB</u> | | <u>100MG</u> | <u>A206309 003</u> | Apr 27, 2016 |
| <u>AB</u> | | <u>140MG</u> | <u>A206309 004</u> | Apr 27, 2016 |
| <u>AB</u> | | <u>180MG</u> | <u>A206309 005</u> | Apr 27, 2016 |
| <u>AB</u> | | <u>250MG</u> | <u>A206309 006</u> | Apr 27, 2016 |
| <u>AB</u> | SUN PHARMA GLOBAL | <u>5MG</u> | <u>A201742 001</u> | Feb 12, 2014 |
| <u>AB</u> | | <u>20MG</u> | <u>A201742 002</u> | Feb 12, 2014 |
| <u>AB</u> | | <u>100MG</u> | <u>A201742 003</u> | Feb 12, 2014 |
| <u>AB</u> | | <u>140MG</u> | <u>A201742 004</u> | Feb 12, 2014 |
| <u>AB</u> | | <u>180MG</u> | <u>A201742 005</u> | Feb 12, 2014 |
| <u>AB</u> | | <u>250MG</u> | <u>A201742 006</u> | Feb 12, 2014 |
| <u>AB</u> | WATSON LABS TEVA | <u>5MG</u> | <u>A203959 001</u> | Apr 18, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A203959 002</u> | Apr 18, 2017 |
| <u>AB</u> | | <u>100MG</u> | <u>A203959 003</u> | Apr 18, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

TEMOZOLOMIDE

CAPSULE; ORAL

TEMOZOLOMIDE

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | | <u>140MG</u> | <u>A203959 004</u> | Apr 18, 2017 |
| <u>AB</u> | | <u>250MG</u> | <u>A203959 005</u> | Apr 18, 2017 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>5MG</u> | <u>A206750 001</u> | Jul 31, 2017 |
| <u>AB</u> | | <u>20MG</u> | <u>A206750 002</u> | Jul 31, 2017 |
| <u>AB</u> | | <u>100MG</u> | <u>A206750 003</u> | Jul 31, 2017 |
| <u>AB</u> | | <u>140MG</u> | <u>A206750 004</u> | Jul 31, 2017 |
| <u>AB</u> | | <u>180MG</u> | <u>A206750 005</u> | Jul 31, 2017 |
| <u>AB</u> | | <u>250MG</u> | <u>A206750 006</u> | Jul 31, 2017 |
| | POWDER; INTRAVENOUS | | | |
| | TEMODAR | | | |
| | +! MERCK SHARP DOHME | 100MG/VIAL | N022277 001 | Feb 27, 2009 |

TEMSIROLIMUS

SOLUTION; INTRAVENOUS

TEMSIROLIMUS

| | | | | |
|-----------|-----------------|--------------------------|--------------------|--------------|
| <u>AP</u> | ACCORD HLTHCARE | <u>25MG/ML (25MG/ML)</u> | <u>A203153 001</u> | Jul 30, 2018 |
| <u>AP</u> | +! PF PRISM CV | <u>25MG/ML (25MG/ML)</u> | <u>N022088 001</u> | May 30, 2007 |

TENOFOVIR ALAFENAMIDE FUMARATE

TABLET; ORAL

VEMLIDY

| | | | | |
|----|---------------------|--------------|-------------|--------------|
| +! | GILEAD SCIENCES INC | EQ 25MG BASE | N208464 001 | Nov 10, 2016 |
|----|---------------------|--------------|-------------|--------------|

TENOFOVIR DISOPROXIL FUMARATE

POWDER; ORAL

VIREAD

| | | | | |
|----|---------------------|---------------|-------------|--------------|
| +! | GILEAD SCIENCES INC | 40MG/SCOOPFUL | N022577 001 | Jan 18, 2012 |
|----|---------------------|---------------|-------------|--------------|

TABLET; ORAL

TENOFOVIR DISOPROXIL FUMARATE

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | APOTEX INC | <u>300MG</u> | <u>A206481 001</u> | Jul 26, 2018 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>150MG</u> | <u>A090647 001</u> | Jan 26, 2018 |
| <u>AB</u> | | <u>200MG</u> | <u>A090647 002</u> | Jan 26, 2018 |
| <u>AB</u> | | <u>250MG</u> | <u>A090647 003</u> | Jan 26, 2018 |
| <u>AB</u> | | <u>300MG</u> | <u>A090647 004</u> | Jan 26, 2018 |
| <u>AB</u> | CASI PHARMS INC | <u>300MG</u> | <u>A209550 001</u> | Feb 26, 2018 |
| <u>AB</u> | CIPLA | <u>300MG</u> | <u>A078800 001</u> | Jan 26, 2018 |
| <u>AB</u> | HETERO LABS LTD III | <u>300MG</u> | <u>A090636 001</u> | Jan 26, 2018 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>300MG</u> | <u>A203232 001</u> | Jan 26, 2018 |
| <u>AB</u> | MYLAN PHARMS INC | <u>150MG</u> | <u>A206569 001</u> | Nov 27, 2018 |
| <u>AB</u> | | <u>200MG</u> | <u>A206569 002</u> | Nov 27, 2018 |
| <u>AB</u> | | <u>250MG</u> | <u>A206569 003</u> | Nov 27, 2018 |
| <u>AB</u> | | <u>300MG</u> | <u>A206569 004</u> | Nov 27, 2018 |
| <u>AB</u> | QILU PHARM CO LTD | <u>200MG</u> | <u>A209498 001</u> | Mar 02, 2018 |
| <u>AB</u> | | <u>250MG</u> | <u>A209498 002</u> | Mar 02, 2018 |
| <u>AB</u> | | <u>300MG</u> | <u>A209498 003</u> | Mar 02, 2018 |
| <u>AB</u> | STRIDES PHARMA | <u>300MG</u> | <u>A090742 001</u> | Jan 26, 2018 |
| <u>AB</u> | TEVA PHARMS USA | <u>150MG</u> | <u>A091612 002</u> | Jan 26, 2018 |
| <u>AB</u> | | <u>200MG</u> | <u>A091612 003</u> | Jan 26, 2018 |
| <u>AB</u> | | <u>250MG</u> | <u>A091612 004</u> | Jan 26, 2018 |
| <u>AB</u> | | <u>300MG</u> | <u>A091612 001</u> | Mar 18, 2015 |
| | <u>VIREAD</u> | | | |
| <u>AB</u> | + GILEAD SCIENCES INC | <u>150MG</u> | <u>N021356 002</u> | Jan 18, 2012 |
| <u>AB</u> | + | <u>200MG</u> | <u>N021356 003</u> | Jan 18, 2012 |
| <u>AB</u> | + | <u>250MG</u> | <u>N021356 004</u> | Jan 18, 2012 |
| <u>AB</u> | +! | <u>300MG</u> | <u>N021356 001</u> | Oct 26, 2001 |

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

TERAZOSIN HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|---------------------|--------------------|--------------|
| <u>AB</u> | APOTEX | <u>EQ 1MG BASE</u> | <u>A075498 001</u> | Apr 12, 2001 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A075498 002</u> | Apr 12, 2001 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A075498 003</u> | Apr 12, 2001 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A075498 004</u> | Apr 12, 2001 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>EQ 1MG BASE</u> | <u>A075614 002</u> | Jan 30, 2001 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A075614 001</u> | Jan 30, 2001 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A075614 003</u> | Jan 30, 2001 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A075614 004</u> | Jan 30, 2001 |
| <u>AB</u> | JUBILANT CADISTA | <u>EQ 1MG BASE</u> | <u>A075317 001</u> | Dec 20, 2004 |

PRESCRIPTION DRUG PRODUCT LIST

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

TERAZOSIN HYDROCHLORIDE

| | | | | |
|-----------|------------------|---------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A075317 002</u> | Dec 20, 2004 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A075317 003</u> | Dec 20, 2004 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A075317 004</u> | Dec 20, 2004 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 1MG BASE</u> | <u>A075140 002</u> | Feb 11, 2000 |
| <u>AB</u> | | <u>EQ 2MG BASE</u> | <u>A075140 003</u> | Feb 11, 2000 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A075140 001</u> | Feb 11, 2000 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A075140 004</u> | Feb 11, 2000 |
| <u>AB</u> | SANDOZ | <u>EQ 1MG BASE</u> | <u>A074823 001</u> | Mar 30, 1998 |
| <u>AB</u> | ! | <u>EQ 2MG BASE</u> | <u>A074823 002</u> | Mar 30, 1998 |
| <u>AB</u> | | <u>EQ 5MG BASE</u> | <u>A074823 003</u> | Mar 30, 1998 |
| <u>AB</u> | | <u>EQ 10MG BASE</u> | <u>A074823 004</u> | Mar 30, 1998 |

TERBINAFFINE HYDROCHLORIDE

TABLET; ORAL

LAMISIL

| | | | | | |
|-----------|----|----------|----------------------|--------------------|--------------|
| <u>AB</u> | +! | NOVARTIS | <u>EQ 250MG BASE</u> | <u>N020539 001</u> | May 10, 1996 |
|-----------|----|----------|----------------------|--------------------|--------------|

TERBINAFFINE HYDROCHLORIDE

| | | | | | |
|-----------|--|--------------------|----------------------|--------------------|--------------|
| <u>AB</u> | | AUROBINDO PHARMA | <u>EQ 250MG BASE</u> | <u>A078297 001</u> | Jul 02, 2007 |
| <u>AB</u> | | BRECKENRIDGE PHARM | <u>EQ 250MG BASE</u> | <u>A077714 001</u> | Jun 04, 2010 |
| <u>AB</u> | | CIPLA | <u>EQ 250MG BASE</u> | <u>A077137 001</u> | Jul 02, 2007 |
| <u>AB</u> | | DR REDDYS LABS INC | <u>EQ 250MG BASE</u> | <u>A076390 001</u> | Jul 02, 2007 |
| <u>AB</u> | | GLENMARK GENERICS | <u>EQ 250MG BASE</u> | <u>A078157 001</u> | Jul 02, 2007 |
| <u>AB</u> | | HARRIS PHARM | <u>EQ 250MG BASE</u> | <u>A077919 001</u> | Jul 02, 2007 |
| <u>AB</u> | | INVAGEN PHARMS | <u>EQ 250MG BASE</u> | <u>A077533 001</u> | Jul 02, 2007 |
| <u>AB</u> | | ORCHID HLTHCARE | <u>EQ 250MG BASE</u> | <u>A078163 001</u> | Jul 02, 2007 |
| <u>AB</u> | | TEVA | <u>EQ 250MG BASE</u> | <u>A076377 001</u> | Jul 02, 2007 |

TERBUTALINE SULFATE

INJECTABLE; INJECTION

TERBUTALINE SULFATE

| | | | | | |
|-----------|---|--------------------|---------------|--------------------|--------------|
| <u>AP</u> | | AKORN | <u>1MG/ML</u> | <u>A078151 001</u> | Jan 07, 2008 |
| <u>AP</u> | ! | ATHENEX INC | <u>1MG/ML</u> | <u>A076770 001</u> | Apr 23, 2004 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>1MG/ML</u> | <u>A076887 001</u> | May 26, 2004 |
| <u>AP</u> | | HIKMA FARMACEUTICA | <u>1MG/ML</u> | <u>A078630 001</u> | May 20, 2009 |
| <u>AP</u> | | UNITED BIOMEDCL | <u>1MG/ML</u> | <u>A200122 001</u> | Nov 08, 2013 |

TABLET; ORAL

BRETHINE

| | | | | | |
|-----------|---|----------------|--------------|--------------------|--|
| <u>AB</u> | + | ANI PHARMS INC | <u>2.5MG</u> | <u>N017849 001</u> | |
| <u>AB</u> | + | | <u>5MG</u> | <u>N017849 002</u> | |

TERBUTALINE SULFATE

| | | | | | |
|-----------|---|----------------|--------------|--------------------|--------------|
| <u>AB</u> | | IMPAX LABS | <u>2.5MG</u> | <u>A075877 001</u> | Jun 26, 2001 |
| <u>AB</u> | | | <u>5MG</u> | <u>A075877 002</u> | Jun 26, 2001 |
| <u>AB</u> | | LANNETT CO INC | <u>2.5MG</u> | <u>A077152 001</u> | Mar 25, 2005 |
| <u>AB</u> | ! | | <u>5MG</u> | <u>A077152 002</u> | Mar 25, 2005 |

TERCONAZOLE

CREAM; VAGINAL

TERCONAZOLE

| | | | | | |
|-----------|----|----------------|-------------|--------------------|--------------|
| <u>AB</u> | | FOUGERA PHARMS | <u>0.4%</u> | <u>A076712 001</u> | Feb 18, 2005 |
| <u>AB</u> | ! | TARO | <u>0.4%</u> | <u>A076043 001</u> | Jan 19, 2005 |
| BX | +! | NYCOMED US | 0.8% | N021735 001 | Oct 01, 2004 |
| BX | ! | TARO | 0.8% | A075953 001 | Apr 06, 2004 |

SUPPOSITORY; VAGINAL

TERCONAZOLE

| | | | | | |
|-----------|---|------------------|-------------|--------------------|--------------|
| <u>AB</u> | ! | PERRIGO NEW YORK | <u>80MG</u> | <u>A077149 001</u> | Mar 17, 2006 |
| <u>AB</u> | | TARO | <u>80MG</u> | <u>A077553 001</u> | Mar 09, 2007 |

TERIFLUNOMIDE

TABLET; ORAL

AUBAGIO

| | | | | | |
|-----------|----|-------------------|-------------|--------------------|--------------|
| <u>AB</u> | + | SANOFI AVENTIS US | <u>7MG</u> | <u>N202992 001</u> | Sep 12, 2012 |
| <u>AB</u> | +! | | <u>14MG</u> | <u>N202992 002</u> | Sep 12, 2012 |

TERIFLUNOMIDE

| | | | | | |
|-----------|--|----------------------|-------------|--------------------|--------------|
| <u>AB</u> | | ACCORD HLTHCARE | <u>7MG</u> | <u>A209690 001</u> | Jan 07, 2019 |
| <u>AB</u> | | | <u>14MG</u> | <u>A209690 002</u> | Jan 07, 2019 |
| <u>AB</u> | | AMNEAL PHARMS CO | <u>7MG</u> | <u>A209613 001</u> | Sep 28, 2018 |
| <u>AB</u> | | | <u>14MG</u> | <u>A209613 002</u> | Sep 28, 2018 |
| <u>AB</u> | | APOTEX INC | <u>7MG</u> | <u>A209601 001</u> | Nov 02, 2018 |
| <u>AB</u> | | | <u>14MG</u> | <u>A209601 002</u> | Nov 02, 2018 |
| <u>AB</u> | | AUROBINDO PHARMA LTD | <u>7MG</u> | <u>A209638 001</u> | Oct 26, 2018 |
| <u>AB</u> | | | <u>14MG</u> | <u>A209638 002</u> | Oct 26, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

TERIFLUNOMIDE

TABLET; ORAL

TERIFLUNOMIDE

| | | | | |
|-----------|-------------------------|-------------|--------------------|--------------|
| <u>AB</u> | GLENMARK PHARMS | <u>7MG</u> | <u>A209663 001</u> | Nov 15, 2018 |
| <u>AB</u> | | <u>14MG</u> | <u>A209663 002</u> | Nov 15, 2018 |
| <u>AB</u> | SANDOZ INC | <u>7MG</u> | <u>A209710 001</u> | Jan 03, 2019 |
| <u>AB</u> | | <u>14MG</u> | <u>A209710 002</u> | Jan 03, 2019 |
| <u>AB</u> | TEVA PHARMS USA | <u>7MG</u> | <u>A209700 001</u> | Sep 04, 2018 |
| <u>AB</u> | | <u>14MG</u> | <u>A209700 002</u> | Sep 04, 2018 |
| <u>AB</u> | WATSON LABS TEVA | <u>7MG</u> | <u>A209549 001</u> | Jul 27, 2018 |
| <u>AB</u> | | <u>14MG</u> | <u>A209549 002</u> | Jul 27, 2018 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>7MG</u> | <u>A209668 001</u> | Nov 30, 2018 |
| <u>AB</u> | | <u>14MG</u> | <u>A209668 002</u> | Nov 30, 2018 |

TERIPARATIDE RECOMBINANT HUMAN

INJECTABLE; SUBCUTANEOUS

FORTEO

+! LILLY 0.6MG/2.4ML (0.25MG/ML) N021318 002 Jun 25, 2008

TESAMORELIN ACETATE

POWDER; SUBCUTANEOUS

EGRIFTA

+! THERATECHNOLOGIES EQ 1MG BASE/VIAL N022505 001 Nov 10, 2010

+! EQ 2MG BASE/VIAL N022505 002 Nov 29, 2011

TESTOSTERONE

FILM, EXTENDED RELEASE; TRANSDERMAL

ANDRODERM

+! ALLERGAN SALES LLC 2MG/24HR N020489 003 Oct 20, 2011

+! 4MG/24HR N020489 004 Oct 20, 2011

GEL; TRANSDERMAL

ANDROGEL

| | | | | |
|------------|----------|--------------------------|--------------------|--------------|
| <u>AB1</u> | + ABBVIE | <u>25MG/2.5GM PACKET</u> | <u>N021015 001</u> | Feb 28, 2000 |
| <u>AB1</u> | +! | <u>50MG/5GM PACKET</u> | <u>N021015 002</u> | Feb 28, 2000 |

TESTOSTERONE

| | | | | |
|------------|---------------------|--------------------------|--------------------|--------------|
| <u>AB1</u> | ACTAVIS LABS UT INC | <u>25MG/2.5GM PACKET</u> | <u>A076737 001</u> | Jan 27, 2006 |
| <u>AB1</u> | | <u>50MG/5GM PACKET</u> | <u>A076737 002</u> | Jan 27, 2006 |
| <u>AB1</u> | PAR PHARM | <u>25MG/2.5GM PACKET</u> | <u>A076744 001</u> | May 23, 2007 |
| <u>AB1</u> | | <u>50MG/5GM PACKET</u> | <u>A076744 002</u> | May 23, 2007 |

ANDROGEL

| | | | | |
|------------|----------|--------------------------------------|--------------------|--------------|
| <u>AB2</u> | + ABBVIE | <u>1.62% (20.25MG/1.25GM PACKET)</u> | <u>N022309 002</u> | Sep 07, 2012 |
| <u>AB2</u> | +! | <u>1.62% (40.5MG/2.5GM PACKET)</u> | <u>N022309 003</u> | Sep 07, 2012 |

TESTIM

| | | | | |
|------------|------------------------|------------------------|--------------------|--------------|
| <u>AB2</u> | +! AUXILIUM PHARMS LLC | <u>50MG/5GM PACKET</u> | <u>N021454 001</u> | Oct 31, 2002 |
|------------|------------------------|------------------------|--------------------|--------------|

TESTOSTERONE

| | | | | |
|------------|---------------------|--------------------------------------|--------------------|--------------|
| <u>AB2</u> | ACTAVIS LABS UT INC | <u>50MG/5GM PACKET</u> | <u>A091073 001</u> | Sep 18, 2017 |
| <u>AB2</u> | PERRIGO UK FINCO | <u>1.62% (20.25MG/1.25GM PACKET)</u> | <u>A205781 001</u> | Jul 12, 2017 |
| <u>AB2</u> | | <u>1.62% (40.5MG/2.5GM PACKET)</u> | <u>A205781 002</u> | Jul 12, 2017 |

VOGELXO

| | | | | |
|------------|-------------------|------------------------|--------------------|--------------|
| <u>AB2</u> | UPSHER SMITH LABS | <u>50MG/5GM PACKET</u> | <u>N204399 002</u> | Jun 04, 2014 |
|------------|-------------------|------------------------|--------------------|--------------|

GEL, METERED; NASAL

NATESTO

AYTU 5.5MG/0.122GM ACTUATION N205488 001 May 28, 2014

GEL, METERED; TRANSDERMAL

ANDROGEL

| | | | | |
|-----------|-----------|-----------------------------------------|--------------------|--------------|
| <u>AB</u> | +! ABBVIE | <u>1.62% (20.25MG/1.25GM ACTUATION)</u> | <u>N022309 001</u> | Apr 29, 2011 |
| <u>AB</u> | +! | <u>12.5MG/1.25GM ACTUATION</u> | <u>N021015 003</u> | Sep 26, 2003 |

FORTESTA

| | | | | |
|-----------|----------------|-----------------------------|--------------------|--------------|
| <u>AB</u> | +! ENDO PHARMS | <u>10MG/0.5GM ACTUATION</u> | <u>N021463 001</u> | Dec 29, 2010 |
|-----------|----------------|-----------------------------|--------------------|--------------|

TESTOSTERONE

| | | | | |
|-----------|---------------------|-----------------------------------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS LABS UT INC | <u>10MG/0.5GM ACTUATION</u> | <u>A204571 001</u> | Aug 05, 2015 |
| <u>AB</u> | | <u>12.5MG/1.25GM ACTUATION</u> | <u>A076737 003</u> | Mar 09, 2015 |
| <u>AB</u> | PERRIGO ISRAEL | <u>1.62% (20.25MG/1.25GM ACTUATION)</u> | <u>A204268 001</u> | Aug 04, 2015 |

VOGELXO

| | | | | |
|----|-------------------|-------------------------|-------------|--------------|
| BX | UPSHER SMITH LABS | 12.5MG/1.25GM ACTUATION | N204399 003 | Jun 04, 2014 |
|----|-------------------|-------------------------|-------------|--------------|

PELLET; IMPLANTATION

TESTOPEL

! AUXILIUM PHARMS INC 75MG A080911 001

SOLUTION, METERED; TRANSDERMAL

TESTOSTERONE

| | | | | |
|-----------|---------------------|-----------------------------|--------------------|--------------|
| <u>AT</u> | ACTAVIS LABS UT INC | <u>30MG/1.5ML ACTUATION</u> | <u>A205328 001</u> | Aug 07, 2017 |
| <u>AT</u> | CIPLA | <u>30MG/1.5ML ACTUATION</u> | <u>A209533 001</u> | Jan 29, 2018 |
| <u>AT</u> | LUPIN LTD | <u>30MG/1.5ML ACTUATION</u> | <u>A208061 001</u> | Oct 23, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

TESTOSTERONE

SOLUTION, METERED; TRANSDERMAL

TESTOSTERONE

| | | | | | | |
|----------------------------------|---|---------------------|-----------------------------|----------------|------------|--------------|
| AT | ! | PERRIGO ISRAEL | 30MG/1.5ML ACTUATION | A204255 | 001 | Feb 28, 2017 |
| TABLET, EXTENDED RELEASE; BUCCAL | | | | | | |
| STRIANT | | | | | | |
| | + | AUXILIUM PHARMS LLC | 30MG | N021543 | 001 | Jun 19, 2003 |

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

DEPO-TESTOSTERONE

| | | | | | | |
|-------------------------------|---|-------------------------|-----------------|----------------|------------|--------------|
| AO | ! | PHARMACIA AND UPJOHN | 100MG/ML | A085635 | 002 | |
| AO | ! | | 200MG/ML | A085635 | 003 | |
| <u>TESTOSTERONE CYPIONATE</u> | | | | | | |
| AO | | CIPLA | 100MG/ML | A210362 | 001 | Jun 19, 2018 |
| AO | | | 200MG/ML | A210362 | 002 | Jun 19, 2018 |
| AO | | HIKMA FARMACEUTICA | 200MG/ML | A091244 | 001 | May 01, 2012 |
| AO | | LUITPOLD | 200MG/ML | A207742 | 001 | Jun 16, 2017 |
| AO | | MYLAN INSTITUTIONAL | 200MG/ML | A040652 | 001 | Dec 11, 2006 |
| AO | | PADDOCK LLC | 200MG/ML | A040530 | 001 | Jan 31, 2005 |
| AO | | SANDOZ INC | 100MG/ML | A040615 | 001 | Aug 10, 2006 |
| AO | | | 200MG/ML | A040615 | 002 | Aug 10, 2006 |
| AO | | SUN PHARM INDS LTD | 100MG/ML | A201720 | 001 | Jun 03, 2013 |
| AO | | | 200MG/ML | A201720 | 002 | Jun 03, 2013 |
| AO | | WATSON PHARMS INC | 200MG/ML | A086030 | 001 | |
| AO | | WEST-WARD PHARMS INT | 100MG/ML | A090387 | 001 | Jul 15, 2010 |
| AO | | | 200MG/ML | A090387 | 002 | Jul 15, 2010 |

TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

TESTOSTERONE ENANTHATE

| | | | | | | |
|------------------------|---|--------------------|---------------------------|----------------|------------|--------------|
| AO | | HIKMA FARMACEUTICA | 200MG/ML | A091120 | 001 | Sep 18, 2012 |
| AO | | NEXUS PHARMS | 200MG/ML | A040575 | 001 | Jun 14, 2006 |
| AO | ! | WATSON PHARMS INC | 200MG/ML | A085598 | 001 | |
| SOLUTION; SUBCUTANEOUS | | | | | | |
| XYOSTED (AUTOINJECTOR) | | | | | | |
| | + | ANTARES PHARMA INC | 50MG/0.5ML (50MG/0.5ML) | N209863 | 001 | Sep 28, 2018 |
| | + | | 75MG/0.5ML (75MG/0.5ML) | N209863 | 002 | Sep 28, 2018 |
| | + | | 100MG/0.5ML (100MG/0.5ML) | N209863 | 003 | Sep 28, 2018 |

TESTOSTERONE UNDECANOATE

INJECTABLE; INTRAMUSCULAR

AVEED

| | | | | | | |
|--|---|-----------------|----------------------|---------|-----|--------------|
| | + | ENDO PHARMS INC | 750MG/3ML (250MG/ML) | N022219 | 001 | Mar 05, 2014 |
|--|---|-----------------|----------------------|---------|-----|--------------|

TETRABENAZINE

TABLET; ORAL

TETRABENAZINE

| | | | | | | |
|-----------|--|---------------------|---------------|----------------|------------|--------------|
| AB | | ACTAVIS LABS FL INC | 25MG | A206686 | 001 | Jul 07, 2017 |
| AB | | APICORE US | 12.5MG | A207682 | 001 | Jan 31, 2017 |
| AB | | | 25MG | A207682 | 002 | Jan 31, 2017 |
| AB | | BIONPHARMA INC | 12.5MG | A208826 | 001 | Dec 18, 2017 |
| AB | | | 25MG | A208826 | 002 | Dec 18, 2017 |
| AB | | DR REDDYS LABS LTD | 12.5MG | A209284 | 001 | Jan 08, 2018 |
| AB | | | 25MG | A209284 | 002 | Jan 08, 2018 |
| AB | | HETERO LABS LTD V | 12.5MG | A204574 | 001 | Feb 03, 2016 |
| AB | | | 25MG | A204574 | 002 | Feb 03, 2016 |
| AB | | LUPIN LTD | 12.5MG | A210544 | 001 | Apr 20, 2018 |
| AB | | | 25MG | A210544 | 002 | Apr 20, 2018 |
| AB | | SUN PHARMA GLOBAL | 12.5MG | A206129 | 001 | Aug 17, 2015 |
| AB | | | 25MG | A206129 | 002 | Aug 17, 2015 |

XENAZINE

| | | | | | | |
|-----------|---|-------------------------|---------------|----------------|------------|--------------|
| AB | + | VALEANT PHARMS NORTH | 12.5MG | N021894 | 001 | Aug 15, 2008 |
| AB | + | ! | 25MG | N021894 | 002 | Aug 15, 2008 |

TETRACAINE HYDROCHLORIDE

SOLUTION; OPHTHALMIC

TETRACAINE HYDROCHLORIDE

| | | | | | | |
|--|---|------------|------|---------|-----|--------------|
| | + | ALCON LABS | 0.5% | N208135 | 001 | Feb 29, 2016 |
|--|---|------------|------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

TETRACYCLINE HYDROCHLORIDE

CAPSULE;ORAL

ACHROMYCIN V

| | | | | | | |
|-----------|---|---------------------|--------------|----------------|------------|--|
| <u>AB</u> | + | HERITAGE PHARMS INC | <u>250MG</u> | <u>N050278</u> | <u>003</u> | |
| <u>AB</u> | + | ! | <u>500MG</u> | <u>N050278</u> | <u>001</u> | |

TETRACYCLINE HYDROCHLORIDE

| | | | | | | |
|-----------|--|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | AMNEAL PHARMS NY | <u>250MG</u> | <u>A210674</u> | <u>001</u> | Sep 18, 2018 |
| <u>AB</u> | | | <u>500MG</u> | <u>A210674</u> | <u>002</u> | Sep 18, 2018 |
| <u>AB</u> | | BRECKENRIDGE PHARM | <u>250MG</u> | <u>A210662</u> | <u>001</u> | Nov 07, 2018 |
| <u>AB</u> | | | <u>500MG</u> | <u>A210662</u> | <u>002</u> | Nov 07, 2018 |
| <u>AB</u> | | CHARTWELL TETRA | <u>250MG</u> | <u>A062752</u> | <u>001</u> | Aug 12, 1988 |
| <u>AB</u> | | | <u>500MG</u> | <u>A062752</u> | <u>002</u> | Aug 12, 1988 |
| <u>AB</u> | | WATSON LABS | <u>250MG</u> | <u>A061837</u> | <u>001</u> | |
| <u>AB</u> | | | <u>500MG</u> | <u>A061837</u> | <u>002</u> | |

TETRAHYDROZOLINE HYDROCHLORIDE

SOLUTION;NASAL

TYZINE

| | | | | | |
|---|----------------|-------|---------|-----|--|
| ! | FOUGERA PHARMS | 0.05% | A086576 | 002 | |
| | | 0.1% | A086576 | 001 | |

SPRAY;NASAL

TYZINE

| | | | | | |
|---|----------------|------|---------|-----|--|
| ! | FOUGERA PHARMS | 0.1% | A086576 | 003 | |
|---|----------------|------|---------|-----|--|

THALIDOMIDE

CAPSULE;ORAL

THALOMID

| | | | | | |
|---|---------|-------|---------|-----|--------------|
| + | CELGENE | 50MG | N020785 | 001 | Jul 16, 1998 |
| + | | 100MG | N020785 | 002 | Jan 17, 2003 |
| + | | 150MG | N020785 | 004 | Jan 10, 2007 |
| + | ! | 200MG | N020785 | 003 | Jan 17, 2003 |

THALLOUS CHLORIDE TL-201

INJECTABLE;INJECTION

THALLOUS CHLORIDE TL 201

| | | | | | | | |
|-----------|---|---|---------------------|----------------|----------------|------------|--------------|
| <u>AP</u> | + | ! | GE HEALTHCARE | <u>1mCi/ML</u> | <u>N018110</u> | <u>002</u> | Feb 27, 1996 |
| <u>AP</u> | + | ! | LANTHEUS MEDCL | <u>1mCi/ML</u> | <u>N017806</u> | <u>001</u> | |
| <u>AP</u> | + | ! | MALLINKRODT NUCLEAR | <u>1mCi/ML</u> | <u>N018150</u> | <u>001</u> | |

THEOPHYLLINE

CAPSULE, EXTENDED RELEASE;ORAL

THEO-24

| | | | | | | |
|---|--|---------------------|-------|---------|-----|--------------|
| | | ACTIENT PHARMS | 100MG | A087942 | 001 | Aug 22, 1983 |
| ! | | AUXILIUM PHARMS INC | 400MG | A081034 | 001 | Feb 28, 1992 |
| | | AUXILIUM PHARMS LLC | 200MG | A087943 | 001 | Aug 22, 1983 |
| | | | 300MG | A087944 | 001 | Aug 22, 1983 |

INJECTABLE;INJECTION

THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | | |
|---|---|---------|------------|---------|-----|--------------|
| + | ! | B BRAUN | 40MG/100ML | N019826 | 001 | Aug 14, 1992 |
|---|---|---------|------------|---------|-----|--------------|

THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | | |
|---|---|---------|------------|---------|-----|--------------|
| + | ! | B BRAUN | 80MG/100ML | N019826 | 002 | Aug 14, 1992 |
|---|---|---------|------------|---------|-----|--------------|

THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | | |
|---|---|---------|-------------|---------|-----|--------------|
| + | ! | B BRAUN | 160MG/100ML | N019826 | 003 | Aug 14, 1992 |
|---|---|---------|-------------|---------|-----|--------------|

THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | | |
|---|---|---------|-------------|---------|-----|--------------|
| + | ! | B BRAUN | 320MG/100ML | N019826 | 006 | Aug 14, 1992 |
|---|---|---------|-------------|---------|-----|--------------|

SOLUTION;ORAL

THEOPHYLLINE

| | | | | | | |
|-----------|---|-----------------|------------------|----------------|------------|--------------|
| <u>AA</u> | ! | LANNETT CO INC | <u>80MG/15ML</u> | <u>A091156</u> | <u>001</u> | Apr 13, 2011 |
| <u>AA</u> | | TRIS PHARMA INC | <u>80MG/15ML</u> | <u>A091586</u> | <u>001</u> | Jun 15, 2012 |

SOLUTION, ELIXIR;ORAL

ELIXOPHYLLIN

| | | | | | | |
|-----------|---|------------------|------------------|----------------|------------|--|
| <u>AA</u> | ! | NOSTRUM LABS INC | <u>80MG/15ML</u> | <u>A085186</u> | <u>001</u> | |
|-----------|---|------------------|------------------|----------------|------------|--|

THEOPHYLLINE

| | | | | | | |
|-----------|--|-------------|------------------|----------------|------------|--------------|
| <u>AA</u> | | PHARM ASSOC | <u>80MG/15ML</u> | <u>A206344</u> | <u>001</u> | Dec 16, 2016 |
|-----------|--|-------------|------------------|----------------|------------|--------------|

TABLET, EXTENDED RELEASE;ORAL

THEOCHRON

| | | | | | | |
|-----------|--|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | NOSTRUM PHARMS LLC | <u>100MG</u> | <u>A087400</u> | <u>003</u> | Feb 21, 1985 |
| <u>AB</u> | | | <u>200MG</u> | <u>A087400</u> | <u>004</u> | Feb 21, 1985 |

THEOPHYLLINE

| | | | | | | |
|-----------|---|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | ALEMBIC PHARMS LTD | <u>300MG</u> | <u>A090430</u> | <u>001</u> | Oct 27, 2010 |
| <u>AB</u> | | GLENMARK GENERICS | <u>400MG</u> | <u>A090355</u> | <u>001</u> | Jul 13, 2010 |
| <u>AB</u> | | | <u>600MG</u> | <u>A090355</u> | <u>002</u> | Jul 13, 2010 |
| <u>AB</u> | | MYLAN IRELAND LTD | <u>400MG</u> | <u>A040560</u> | <u>003</u> | Apr 21, 2006 |
| <u>AB</u> | ! | | <u>600MG</u> | <u>A040560</u> | <u>002</u> | Apr 21, 2006 |

PRESCRIPTION DRUG PRODUCT LIST

THEOPHYLLINE

TABLET, EXTENDED RELEASE;ORAL

THEOPHYLLINE

| | | | | | |
|-----------|---|--------------------|--------------|--------------------|--------------|
| AB | ! | PLIVA | 100MG | A089807 001 | Apr 30, 1990 |
| AB | ! | | 200MG | A089808 001 | Apr 30, 1990 |
| AB | | | 300MG | A089763 001 | Apr 30, 1990 |
| AB | | RHODES PHARMS | 400MG | A087571 001 | Sep 01, 1982 |
| AB | | | 600MG | A040086 001 | Apr 15, 1996 |
| | ! | ALEMBIC PHARMS LTD | 450MG | A090430 002 | Oct 27, 2010 |

THIAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

THIAMINE HYDROCHLORIDE

| | | | | | |
|-----------|---|---------------------|-----------------|--------------------|--------------|
| AP | ! | FRESENIUS KABI USA | 100MG/ML | A080556 001 | |
| AP | | MYLAN INSTITUTIONAL | 100MG/ML | A091623 001 | Jun 25, 2012 |
| AP | | SAGENT PHARMS | 100MG/ML | A206106 001 | Dec 01, 2017 |

THIOGUANINE

TABLET; ORAL

THIOGUANINE

| | | | | | |
|--|---|---|------------------|------|-------------|
| | + | ! | ASPEN GLOBAL INC | 40MG | N012429 001 |
|--|---|---|------------------|------|-------------|

THIORIDAZINE HYDROCHLORIDE

TABLET; ORAL

THIORIDAZINE HYDROCHLORIDE

| | | | | | |
|-----------|---|----------------------|--------------|--------------------|--------------|
| AB | | MYLAN | 10MG | A088004 002 | Mar 15, 1983 |
| AB | | | 25MG | A088004 003 | Mar 15, 1983 |
| AB | | | 50MG | A088004 004 | Mar 15, 1983 |
| AB | ! | | 100MG | A088004 001 | Nov 18, 1983 |
| AB | | SUN PHARM INDUSTRIES | 10MG | A089953 004 | Aug 01, 1986 |
| AB | | | 25MG | A089953 003 | Aug 01, 1986 |
| AB | | | 50MG | A089953 002 | Aug 01, 1986 |
| AB | | | 100MG | A089953 001 | Oct 07, 1988 |

THIOTEPA

INJECTABLE; INJECTION

THIOTEPA

| | | | | | |
|-----------|---|----------------------|------------------|--------------------|--------------------------|
| AP | | DR REDDYS LABS LTD | 15MG/VIAL | A210337 001 | May 04, 2018 |
| AP | | JIANGSU HENGRUI MED | 15MG/VIAL | A209150 001 | May 04, 2018 |
| AP | ! | WEST-WARD PHARMS INT | 15MG/VIAL | A075547 001 | Apr 02, 2001 |
| | | POWDER; INTRAVENOUS | | | |
| | | TEPADINA | | | |
| | + | ! | ADIENNE SA | 15MG/VIAL | N208264 001 Jan 26, 2017 |
| | + | ! | | 100MG/VIAL | N208264 002 Jan 26, 2017 |

THIOTHIXENE

CAPSULE; ORAL

THIOTHIXENE

| | | | | | |
|--|---|-------|------|-------------|--------------|
| | | MYLAN | 1MG | A071093 002 | Jun 23, 1987 |
| | | | 2MG | A071093 003 | Jun 23, 1987 |
| | ! | | 5MG | A071093 004 | Jun 23, 1987 |
| | | | 10MG | A071093 001 | Jun 23, 1987 |

THYROTROPIN ALFA

INJECTABLE; INJECTION

THYROGEN

| | | | | | |
|--|---|---|---------|------------|--------------------------|
| | + | ! | GENZYME | 1.1MG/VIAL | N020898 001 Nov 30, 1998 |
|--|---|---|---------|------------|--------------------------|

TIAGABINE HYDROCHLORIDE

TABLET; ORAL

GABITRIL

| | | | | | |
|-----------|---|----------|-------------|--------------------|--------------|
| AB | + | CEPHALON | 2MG | N020646 005 | Apr 16, 1999 |
| AB | + | ! | 4MG | N020646 001 | Sep 30, 1997 |
| AB | + | | 12MG | N020646 002 | Sep 30, 1997 |
| AB | + | | 16MG | N020646 003 | Sep 30, 1997 |

TIAGABINE HYDROCHLORIDE

| | | | | | |
|-----------|--|---------------------|-------------|--------------------|--------------|
| AB | | AMNEAL PHARMS CO | 2MG | A208181 001 | Dec 08, 2017 |
| AB | | | 4MG | A208181 002 | Dec 08, 2017 |
| AB | | | 12MG | A208181 003 | Dec 08, 2017 |
| AB | | | 16MG | A208181 004 | Dec 08, 2017 |
| AB | | SUN PHARM INDS | 2MG | A077555 001 | Nov 04, 2011 |
| AB | | | 4MG | A077555 002 | Nov 04, 2011 |
| AB | | WILSHIRE PHARMS INC | 2MG | A206857 001 | Oct 13, 2017 |
| AB | | | 4MG | A206857 002 | Oct 13, 2017 |

PRESCRIPTION DRUG PRODUCT LIST

TIAGABINE HYDROCHLORIDE

TABLET; ORAL

TIAGABINE HYDROCHLORIDE

| | | | | |
|-----------|--|-------------|--------------------|--------------|
| <u>AB</u> | | <u>12MG</u> | <u>A206857 003</u> | Oct 13, 2017 |
| <u>AB</u> | | <u>16MG</u> | <u>A206857 004</u> | Oct 13, 2017 |

TICAGRELOR

TABLET; ORAL

BRILINTA

| | | | | | |
|-----------|---|--------------------|-------------|--------------------|--------------|
| <u>AB</u> | + | ASTRAZENECA PHARMS | <u>60MG</u> | <u>N022433 002</u> | Sep 03, 2015 |
| <u>AB</u> | + | ! | <u>90MG</u> | <u>N022433 001</u> | Jul 20, 2011 |

TICAGRELOR

| | | | | | |
|-----------|--|-----------------|-------------|--------------------|--------------|
| <u>AB</u> | | WATSON LABS INC | <u>60MG</u> | <u>A208390 001</u> | Sep 04, 2018 |
| <u>AB</u> | | | <u>90MG</u> | <u>A208390 002</u> | Sep 04, 2018 |

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLOPIDINE HYDROCHLORIDE

| | | | | | |
|-----------|---|--------------------|--------------|--------------------|--------------|
| <u>AB</u> | | APOTEX | <u>250MG</u> | <u>A075089 001</u> | Jul 01, 1999 |
| <u>AB</u> | | SUN PHARM INDS INC | <u>250MG</u> | <u>A075526 001</u> | Sep 26, 2002 |
| <u>AB</u> | ! | TEVA | <u>250MG</u> | <u>A075149 001</u> | Aug 20, 1999 |

TIGECYCLINE

POWDER; INTRAVENOUS

TIGECYCLINE

| | | | | | |
|-----------|--|--------------------|------------------|--------------------|--------------|
| <u>AP</u> | | AMNEAL PHARMS LLC | <u>50MG/VIAL</u> | <u>N211158 001</u> | Aug 02, 2018 |
| <u>AP</u> | | APOTEX INC | <u>50MG/VIAL</u> | <u>A204439 001</u> | Dec 21, 2018 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>50MG/VIAL</u> | <u>N205645 001</u> | Dec 01, 2016 |
| <u>AP</u> | | SANDOZ INC | <u>50MG/VIAL</u> | <u>A091620 001</u> | May 27, 2015 |

TYGACIL

| | | | | | | |
|-----------|---|---|-------------|------------------|--------------------|--------------|
| <u>AP</u> | + | ! | PF PRISM CV | <u>50MG/VIAL</u> | <u>N021821 001</u> | Jun 15, 2005 |
|-----------|---|---|-------------|------------------|--------------------|--------------|

TIGECYCLINE

| | | | | | |
|--|--|---------------------|-----------|-------------|--------------|
| | | ACCORD HLTHCARE INC | 50MG/VIAL | N208744 001 | Jan 18, 2018 |
|--|--|---------------------|-----------|-------------|--------------|

TIMOLOL

SOLUTION/DROPS; OPHTHALMIC

BETIMOL

| | | | | | | |
|-----------|---|---|----------------|----------------------|--------------------|--------------|
| <u>AT</u> | + | ! | OAK PHARMS INC | <u>EQ 0.25% BASE</u> | <u>N020439 001</u> | Mar 31, 1995 |
| <u>AT</u> | + | ! | | <u>EQ 0.5% BASE</u> | <u>N020439 002</u> | Mar 31, 1995 |

TIMOLOL

| | | | | | |
|-----------|--|-------|----------------------|--------------------|--------------|
| <u>AT</u> | | AKORN | <u>EQ 0.25% BASE</u> | <u>A205309 001</u> | Sep 30, 2016 |
| <u>AT</u> | | | <u>EQ 0.5% BASE</u> | <u>A205309 002</u> | Sep 30, 2016 |

TIMOLOL MALEATE

SOLUTION, GEL FORMING/DROPS; OPHTHALMIC

TIMOLOL MALEATE

| | | | | | |
|-----------|--|------------|----------------------|--------------------|--------------|
| <u>AB</u> | | SANDOZ INC | <u>EQ 0.25% BASE</u> | <u>N020963 001</u> | Oct 21, 1998 |
| <u>AB</u> | | | <u>EQ 0.5% BASE</u> | <u>N020963 002</u> | Oct 21, 1998 |

TIMOPTIC-XE

| | | | | | | |
|-----------|---|---|--------------------|----------------------|--------------------|--------------|
| <u>AB</u> | + | ! | VALEANT PHARMS LLC | <u>EQ 0.25% BASE</u> | <u>N020330 001</u> | Nov 04, 1993 |
| <u>AB</u> | + | ! | | <u>EQ 0.5% BASE</u> | <u>N020330 002</u> | Nov 04, 1993 |

SOLUTION/DROPS; OPHTHALMIC

TIMOLOL MALEATE

| | | | | | |
|-----------|---|-----------------|----------------------|--------------------|--------------|
| <u>AT</u> | | BAUSCH AND LOMB | <u>EQ 0.25% BASE</u> | <u>A074778 001</u> | Mar 25, 1997 |
| <u>AT</u> | | FDC LTD | <u>EQ 0.25% BASE</u> | <u>A077259 001</u> | Apr 30, 2008 |
| <u>AT</u> | | PACIFIC PHARMA | <u>EQ 0.25% BASE</u> | <u>A074746 001</u> | Mar 25, 1997 |
| <u>AT</u> | ! | SANDOZ INC | <u>EQ 0.25% BASE</u> | <u>A074261 001</u> | Apr 28, 1995 |
| <u>AT</u> | | WOCKHARDT | <u>EQ 0.25% BASE</u> | <u>A078771 001</u> | Sep 28, 2009 |

TIMOPTIC

| | | | | | |
|-----------|---|------|----------------------|--------------------|--|
| <u>AT</u> | + | ATON | <u>EQ 0.25% BASE</u> | <u>N018086 001</u> | |
|-----------|---|------|----------------------|--------------------|--|

TIMOLOL MALEATE

| | | | | | |
|------------|---|-----------------|---------------------|--------------------|--------------|
| <u>AT1</u> | | AKORN | <u>EQ 0.5% BASE</u> | <u>A074466 001</u> | Mar 25, 1997 |
| <u>AT1</u> | | | <u>EQ 0.5% BASE</u> | <u>A074516 001</u> | Mar 25, 1997 |
| <u>AT1</u> | | BAUSCH AND LOMB | <u>EQ 0.5% BASE</u> | <u>A074776 001</u> | Mar 25, 1997 |
| <u>AT1</u> | | FDC LTD | <u>EQ 0.5% BASE</u> | <u>A077259 002</u> | Apr 30, 2008 |
| <u>AT1</u> | | HI TECH PHARMA | <u>EQ 0.5% BASE</u> | <u>A075163 001</u> | Sep 10, 2002 |
| <u>AT1</u> | | PACIFIC PHARMA | <u>EQ 0.5% BASE</u> | <u>A074747 001</u> | Mar 25, 1997 |
| <u>AT1</u> | ! | SANDOZ INC | <u>EQ 0.5% BASE</u> | <u>A074262 001</u> | Apr 28, 1995 |
| <u>AT1</u> | | WOCKHARDT | <u>EQ 0.5% BASE</u> | <u>A078771 002</u> | Sep 28, 2009 |

TIMOPTIC

| | | | | | |
|------------|---|------|---------------------|--------------------|--|
| <u>AT1</u> | + | ATON | <u>EQ 0.5% BASE</u> | <u>N018086 002</u> | |
|------------|---|------|---------------------|--------------------|--|

ISTALOL

| | | | | | | |
|------------|---|---|-----------------|---------------------|--------------------|--------------|
| <u>AT2</u> | + | ! | BAUSCH AND LOMB | <u>EQ 0.5% BASE</u> | <u>N021516 001</u> | Jun 04, 2004 |
|------------|---|---|-----------------|---------------------|--------------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

TIMOLOL MALEATE

SOLUTION/DROPS;OPHTHALMIC

TIMOLOL MALEATE

| | | | | |
|------------|---------------------|---------------------|--------------------|--------------|
| AT2 | APOTEX INC | EQ 0.5% BASE | A204936 001 | Apr 17, 2015 |
| | TIMOPTIC IN OCUDOSE | | | |
| | +! ATON | EQ 0.25% BASE | N019463 001 | Nov 05, 1986 |
| | +! | EQ 0.5% BASE | N019463 002 | Nov 05, 1986 |
| | TABLET;ORAL | | | |
| | TIMOLOL MALEATE | | | |
| | MYLAN | 5MG | A072668 002 | Jun 08, 1990 |
| | | 10MG | A072668 003 | Jun 08, 1990 |
| | ! | 20MG | A072668 001 | Jun 08, 1990 |

TINIDAZOLE

TABLET;ORAL

TINDAMAX

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| AB | + MISSION PHARMA | 250MG | N021618 001 | May 17, 2004 |
| AB | +! | 500MG | N021618 002 | May 17, 2004 |
| | <u>TINIDAZOLE</u> | | | |
| AB | EDENBRIDGE PHARMS | 250MG | A203808 001 | Aug 04, 2015 |
| AB | | 500MG | A203808 002 | Aug 04, 2015 |
| AB | NOVEL LABS INC | 250MG | A202044 001 | Apr 30, 2012 |
| AB | | 500MG | A202044 002 | Apr 30, 2012 |
| AB | UNIQUE PHARM LABS | 250MG | A202489 001 | Oct 09, 2013 |
| AB | | 500MG | A202489 002 | Oct 09, 2013 |
| AB | WEST-WARD PHARMS INT | 250MG | A201172 001 | Apr 30, 2012 |
| AB | | 500MG | A201172 002 | Apr 30, 2012 |

TIOPRONIN

TABLET;ORAL

THIOLA

| | | | | |
|----|----------------|-------|-------------|--------------|
| +! | MISSION PHARMA | 100MG | N019569 001 | Aug 11, 1988 |
|----|----------------|-------|-------------|--------------|

TIOTROPIUM BROMIDE

POWDER; INHALATION

SPIRIVA

| | | | | |
|----|-------------------------|---------------------|-------------|--------------|
| +! | BOEHRINGER INGELHEIM | EQ 0.018MG BASE/INH | N021395 001 | Jan 30, 2004 |
|----|-------------------------|---------------------|-------------|--------------|

SPRAY, METERED; INHALATION

SPIRIVA RESPIMAT

| | | | | |
|---|-------------------------|-----------------------|-------------|--------------|
| + | BOEHRINGER INGELHEIM | EQ 0.00125MG BASE/INH | N021936 002 | Sep 15, 2015 |
|---|-------------------------|-----------------------|-------------|--------------|

| | | | | |
|----|--|----------------------|-------------|--------------|
| +! | | EQ 0.0025MG BASE/INH | N021936 001 | Sep 24, 2014 |
|----|--|----------------------|-------------|--------------|

TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE

TABLET;ORAL

LONSURF

| | | | | |
|---|----------------|---------------------|-------------|--------------|
| + | TAIHO ONCOLOGY | EQ 6.14MG BASE;15MG | N207981 001 | Sep 22, 2015 |
|---|----------------|---------------------|-------------|--------------|

| | | | | |
|----|--|---------------------|-------------|--------------|
| +! | | EQ 8.19MG BASE;20MG | N207981 002 | Sep 22, 2015 |
|----|--|---------------------|-------------|--------------|

TIPRANAVIR

CAPSULE;ORAL

APTIVUS

| | | | | |
|----|-------------------------|-------|-------------|--------------|
| +! | BOEHRINGER INGELHEIM | 250MG | N021814 001 | Jun 22, 2005 |
|----|-------------------------|-------|-------------|--------------|

SOLUTION;ORAL

APTIVUS

| | | | | |
|----|-------------------------|----------|-------------|--------------|
| +! | BOEHRINGER INGELHEIM | 100MG/ML | N022292 001 | Jun 23, 2008 |
|----|-------------------------|----------|-------------|--------------|

TIROFIBAN HYDROCHLORIDE

INJECTABLE; INJECTION

AGGRASTAT

| | | | | |
|---|----------|---------------------------------------|-------------|--------------|
| + | MEDICURE | EQ 5MG BASE/100ML (EQ 0.05MG BASE/ML) | N020913 002 | May 17, 2002 |
|---|----------|---------------------------------------|-------------|--------------|

| | | | | |
|----|--|---------------------------------------------|-------------|--------------|
| +! | | EQ 12.5MG BASE/250ML (EQ 0.05MG BASE/ML) | N020913 003 | Apr 20, 2000 |
|----|--|---------------------------------------------|-------------|--------------|

SOLUTION; INJECTION

AGGRASTAT

| | | | | |
|----|----------|-----------------------------------------|-------------|--------------|
| +! | MEDICURE | EQ 3.75MG BASE/15ML (EQ 0.25MG BASE/ML) | N020912 002 | Aug 31, 2016 |
|----|----------|-----------------------------------------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

TIZANIDINE HYDROCHLORIDE

CAPSULE; ORAL

TIZANIDINE HYDROCHLORIDE

| | | | | |
|-----------------|-------------------------|--------------------|--------------------|--------------|
| <u>AB</u> | APOTEX INC | <u>EQ 2MG BASE</u> | <u>A078868 001</u> | Feb 03, 2012 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A078868 002</u> | Feb 03, 2012 |
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A078868 003</u> | Feb 03, 2012 |
| <u>AB</u> | JUBILANT GENERICS | <u>EQ 2MG BASE</u> | <u>A209605 001</u> | Aug 04, 2017 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A209605 002</u> | Aug 04, 2017 |
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A209605 003</u> | Aug 04, 2017 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 2MG BASE</u> | <u>A091502 001</u> | Nov 09, 2012 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A091502 002</u> | Nov 09, 2012 |
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A091502 003</u> | Nov 09, 2012 |
| <u>AB</u> | PAR PHARM INC | <u>EQ 2MG BASE</u> | <u>A207199 001</u> | Mar 14, 2017 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A207199 002</u> | Mar 14, 2017 |
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A207199 003</u> | Mar 14, 2017 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 2MG BASE</u> | <u>A208622 001</u> | Mar 03, 2017 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A208622 002</u> | Mar 03, 2017 |
| <u>AB</u> | | <u>EQ 6MG BASE</u> | <u>A208622 003</u> | Mar 03, 2017 |
| <u>ZANAFLEX</u> | | | | |
| <u>AB</u> | + COVIS PHARMA BV | <u>EQ 2MG BASE</u> | <u>N021447 001</u> | Aug 29, 2002 |
| <u>AB</u> | + | <u>EQ 4MG BASE</u> | <u>N021447 002</u> | Aug 29, 2002 |
| <u>AB</u> | +! | <u>EQ 6MG BASE</u> | <u>N021447 003</u> | Aug 29, 2002 |

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE

| | | | | |
|-----------------|-------------------------|--------------------|--------------------|--------------|
| <u>AB</u> | APOTEX | <u>EQ 2MG BASE</u> | <u>A076533 001</u> | Jan 16, 2004 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A076533 002</u> | Jan 16, 2004 |
| <u>AB</u> | CASI PHARMS INC | <u>EQ 2MG BASE</u> | <u>A076280 001</u> | Nov 26, 2002 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A076280 002</u> | Jun 27, 2002 |
| <u>AB</u> | DR REDDYS LABS INC | <u>EQ 2MG BASE</u> | <u>A076286 001</u> | Jul 03, 2002 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A076286 002</u> | Jul 03, 2002 |
| <u>AB</u> | EPIC PHARMA LLC | <u>EQ 2MG BASE</u> | <u>A076347 001</u> | Oct 11, 2002 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A076347 002</u> | Oct 11, 2002 |
| <u>AB</u> | MYLAN | <u>EQ 2MG BASE</u> | <u>A076354 001</u> | Mar 28, 2003 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A076354 002</u> | Mar 28, 2003 |
| <u>AB</u> | OXFORD PHARMS | <u>EQ 2MG BASE</u> | <u>A076281 001</u> | Oct 20, 2003 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A076281 002</u> | Oct 20, 2003 |
| <u>AB</u> | PAR PHARM INC | <u>EQ 2MG BASE</u> | <u>A207170 001</u> | Jan 26, 2017 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A207170 002</u> | Jan 26, 2017 |
| <u>AB</u> | SUN PHARM INDS INC | <u>EQ 2MG BASE</u> | <u>A076416 001</u> | Sep 29, 2003 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A076416 002</u> | Sep 29, 2003 |
| <u>AB</u> | TEVA | <u>EQ 2MG BASE</u> | <u>A076284 001</u> | Jul 03, 2002 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A076284 002</u> | Jul 03, 2002 |
| <u>AB</u> | UNICHEM LABS LTD | <u>EQ 2MG BASE</u> | <u>A091283 001</u> | Nov 28, 2012 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A091283 002</u> | Nov 28, 2012 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 2MG BASE</u> | <u>A208187 001</u> | Mar 09, 2018 |
| <u>AB</u> | | <u>EQ 4MG BASE</u> | <u>A208187 002</u> | Mar 09, 2018 |
| <u>ZANAFLEX</u> | | | | |
| <u>AB</u> | +! COVIS PHARMA BV | <u>EQ 4MG BASE</u> | <u>N020397 001</u> | Nov 27, 1996 |

TOBRAMYCIN

OINTMENT; OPHTHALMIC

TOBREX

| | | | | |
|----|-------------------------|------|-------------|--|
| +! | NOVARTIS PHARMS CORP | 0.3% | N050555 001 | |
|----|-------------------------|------|-------------|--|

POWDER; INHALATION

TOBI PODHALER

| | | | | |
|----|---------------------|------|-------------|--------------|
| +! | MYLAN SPECIALITY LP | 28MG | N201688 001 | Mar 22, 2013 |
|----|---------------------|------|-------------|--------------|

SOLUTION; INHALATION

KITABIS PAK

| | | | | |
|-------------|------------------------|------------------|--------------------|--------------|
| <u>AN</u> | PULMOFLOW INC | <u>300MG/5ML</u> | <u>N205433 001</u> | Dec 02, 2014 |
| <u>TOBI</u> | | | | |
| <u>AN</u> | +! MYLAN SPECIALITY LP | <u>300MG/5ML</u> | <u>N050753 001</u> | Dec 22, 1997 |

TOBRAMYCIN

| | | | | |
|-----------|-------------------|------------------|--------------------|--------------|
| <u>AN</u> | AKORN INC | <u>300MG/5ML</u> | <u>A201422 001</u> | May 28, 2014 |
| <u>AN</u> | AMNEAL PHARMS | <u>300MG/5ML</u> | <u>A205501 001</u> | Jul 13, 2015 |
| <u>AN</u> | DR REDDYS LABS SA | <u>300MG/5ML</u> | <u>A207080 001</u> | Jul 09, 2018 |
| <u>AN</u> | LUPIN | <u>300MG/5ML</u> | <u>A208964 001</u> | Mar 22, 2017 |
| <u>AN</u> | MYLAN PHARMS INC | <u>300MG/5ML</u> | <u>A209554 001</u> | Oct 13, 2017 |
| <u>AN</u> | TEVA PHARMS USA | <u>300MG/5ML</u> | <u>A091589 001</u> | Oct 10, 2013 |

BETHKIS

| | | | | |
|----|----------------|-----------|-------------|--------------|
| +! | CHIESI USA INC | 300MG/4ML | N201820 001 | Oct 12, 2012 |
|----|----------------|-----------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

TOBRAMYCIN

SOLUTION/DROPS;OPHTHALMIC

AKTOR

| | | | | |
|-----------|-------|-------------|--------------------|--------------|
| AT | AKORN | 0.3% | A064096 001 | Jan 31, 1996 |
|-----------|-------|-------------|--------------------|--------------|

TOBRAMYCIN

| | | | | |
|-----------|-----------------|-------------|--------------------|--------------|
| AT | BAUSCH AND LOMB | 0.3% | A064052 001 | Nov 29, 1993 |
|-----------|-----------------|-------------|--------------------|--------------|

| | | | | |
|-----------|-------------|-------------|--------------------|--------------|
| AT | FERA PHARMS | 0.3% | A065026 001 | Sep 11, 2001 |
|-----------|-------------|-------------|--------------------|--------------|

| | | | | |
|-----------|-------------------------|-------------|--------------------|--------------|
| AT | SOMERSET THERAPS LLC | 0.3% | A207444 001 | Jun 28, 2017 |
|-----------|-------------------------|-------------|--------------------|--------------|

TOBREX

| | | | | |
|-----------|------------------------------------|-------------|--------------------|--|
| AT | + ! NOVARTIS PHARMS CORP | 0.3% | N050541 001 | |
|-----------|------------------------------------|-------------|--------------------|--|

| | | | | |
|-----------|------------|-------------|--------------------|--------------|
| AT | SANDOZ INC | 0.3% | A062535 001 | Dec 13, 1984 |
|-----------|------------|-------------|--------------------|--------------|

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

TOBRAMYCIN SULFATE

| | | | | |
|-----------|-------|------------------------|--------------------|--------------|
| AP | AKORN | EQ 40MG BASE/ML | A205179 001 | Sep 16, 2014 |
|-----------|-------|------------------------|--------------------|--------------|

| | | | | |
|-----------|-------------------------|------------------------|--------------------|--------------|
| AP | BAXTER HLTHCARE CORP | EQ 40MG BASE/ML | A206965 001 | Jul 01, 2016 |
|-----------|-------------------------|------------------------|--------------------|--------------|

| | | | | |
|-----------|--------------------|------------------------|--------------------|--------------|
| AP | FRESENIUS KABI USA | EQ 10MG BASE/ML | A065122 001 | Nov 29, 2002 |
|-----------|--------------------|------------------------|--------------------|--------------|

| | | | | |
|-----------|----------|------------------------|--------------------|--------------|
| AP | ! | EQ 40MG BASE/ML | A065122 002 | Nov 29, 2002 |
|-----------|----------|------------------------|--------------------|--------------|

| | | | | |
|-----------|----------|---------------------------|--------------------|--------------|
| AP | ! | EQ 1.2GM BASE/VIAL | N050789 001 | Jul 13, 2004 |
|-----------|----------|---------------------------|--------------------|--------------|

| | | | | |
|-----------|------------------|------------------------|--------------------|--------------|
| AP | ! HOSPIRA | EQ 10MG BASE/ML | A063112 001 | Apr 30, 1991 |
|-----------|------------------|------------------------|--------------------|--------------|

| | | | | |
|-----------|----------|------------------------|--------------------|--------------|
| AP | ! | EQ 40MG BASE/ML | A063111 001 | Apr 30, 1991 |
|-----------|----------|------------------------|--------------------|--------------|

| | | | | |
|-----------|----------------|------------------------|--------------------|--------------|
| AP | MYLAN LABS LTD | EQ 40MG BASE/ML | A065407 001 | Mar 11, 2008 |
|-----------|----------------|------------------------|--------------------|--------------|

| | | | | |
|-----------|-----------------|------------------------|--------------------|--------------|
| AP | TEVA PHARMS USA | EQ 40MG BASE/ML | A063100 001 | Jan 30, 1992 |
|-----------|-----------------|------------------------|--------------------|--------------|

| | | | | |
|-----------|-------------------------|------------------------|--------------------|--------------|
| AP | WEST-WARD PHARMS INT | EQ 40MG BASE/ML | A063117 001 | Apr 26, 1991 |
|-----------|-------------------------|------------------------|--------------------|--------------|

| | | | | |
|-----------|-----------------------|---------------------------|--------------------|--------------|
| AP | ! X GEN PHARMS | EQ 1.2GM BASE/VIAL | A065013 001 | Aug 17, 2001 |
|-----------|-----------------------|---------------------------|--------------------|--------------|

| | | | | |
|-----------|-------------------|---------------------------|--------------------|--------------|
| AP | XELLIA PHARMS APS | EQ 1.2GM BASE/VIAL | A205685 001 | Sep 16, 2014 |
|-----------|-------------------|---------------------------|--------------------|--------------|

TOBRAMYCIN SULFATE (PHARMACY BULK)

| | | | | |
|----------|--------------------|-----------------|-------------|--------------|
| ! | FRESENIUS KABI USA | EQ 40MG BASE/ML | A065120 001 | Nov 29, 2002 |
|----------|--------------------|-----------------|-------------|--------------|

TOBRAMYCIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | |
|----------|---------|------------------|-------------|--------------|
| ! | HOSPIRA | EQ 1.2MG BASE/ML | A063081 003 | Jul 31, 1990 |
|----------|---------|------------------|-------------|--------------|

| | | | | |
|----------|--|------------------|-------------|--------------|
| ! | | EQ 1.6MG BASE/ML | A063081 006 | Jun 02, 1993 |
|----------|--|------------------|-------------|--------------|

| | | | | |
|----------|--|--------------------|-------------|--------------|
| ! | | EQ 80MG BASE/100ML | A063081 001 | Jul 31, 1990 |
|----------|--|--------------------|-------------|--------------|

TOFACITINIB CITRATE

TABLET; ORAL

XELJANZ

| | | | | |
|------------|-------------|-------------|-------------|--------------|
| + ! | PF PRISM CV | EQ 5MG BASE | N203214 001 | Nov 06, 2012 |
|------------|-------------|-------------|-------------|--------------|

| | | | | |
|------------|--|--------------|-------------|--------------|
| + ! | | EQ 10MG BASE | N203214 002 | May 30, 2018 |
|------------|--|--------------|-------------|--------------|

TABLET, EXTENDED RELEASE; ORAL

XELJANZ XR

| | | | | |
|------------|------------|--------------|-------------|--------------|
| + ! | PFIZER INC | EQ 11MG BASE | N208246 001 | Feb 23, 2016 |
|------------|------------|--------------|-------------|--------------|

TOLAZAMIDE

TABLET; ORAL

TOLAZAMIDE

| | | | | |
|--|------------------|-------|-------------|--------------|
| | MYLAN PHARMS INC | 250MG | A070259 001 | Jan 02, 1986 |
|--|------------------|-------|-------------|--------------|

| | | | | |
|----------|--|-------|-------------|--------------|
| ! | | 500MG | A070259 003 | Mar 17, 1986 |
|----------|--|-------|-------------|--------------|

TOLBUTAMIDE

TABLET; ORAL

TOLBUTAMIDE

| | | | | |
|----------|------------------|-------|-------------|--|
| ! | MYLAN PHARMS INC | 500MG | A086445 001 | |
|----------|------------------|-------|-------------|--|

TOLCAPONE

TABLET; ORAL

TASMAR

| | | | | |
|-----------|-------------------------------|--------------|--------------------|--------------|
| AB | + ! VALEANT PHARMS LLC | 100MG | N020697 001 | Jan 29, 1998 |
|-----------|-------------------------------|--------------|--------------------|--------------|

TOLCAPONE

| | | | | |
|-----------|--------------------|--------------|--------------------|--------------|
| AB | INGENUS PHARMS LLC | 100MG | A208937 001 | Aug 07, 2018 |
|-----------|--------------------|--------------|--------------------|--------------|

| | | | | |
|-----------|---------------|--------------|--------------------|--------------|
| AB | PAR PHARM INC | 100MG | A204584 001 | Mar 26, 2015 |
|-----------|---------------|--------------|--------------------|--------------|

TOLMETIN SODIUM

CAPSULE; ORAL

TOLMETIN SODIUM

| | | | | |
|-----------|-------|----------------------|--------------------|--------------|
| AB | MYLAN | EQ 400MG BASE | A073393 001 | May 27, 1993 |
|-----------|-------|----------------------|--------------------|--------------|

| | | | | |
|-----------|---------------|----------------------|--------------------|--------------|
| AB | ! TEVA | EQ 400MG BASE | A073290 001 | Nov 27, 1991 |
|-----------|---------------|----------------------|--------------------|--------------|

TABLET; ORAL

TOLMETIN SODIUM

| | | | | |
|----------|-------|---------------|-------------|--------------|
| ! | MYLAN | EQ 600MG BASE | A074473 001 | Aug 30, 1994 |
|----------|-------|---------------|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

TOLTERODINE TARTRATE

CAPSULE, EXTENDED RELEASE;ORAL

DETROL LA

| | | | | | | |
|-----------|---|-------------------------|------------|----------------|------------|--------------|
| <u>AB</u> | + | PHARMACIA AND UPJOHN | <u>2MG</u> | <u>N021228</u> | <u>001</u> | Dec 22, 2000 |
|-----------|---|-------------------------|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|---|------------|----------------|------------|--------------|
| <u>AB</u> | + | ! | <u>4MG</u> | <u>N021228</u> | <u>002</u> | Dec 22, 2000 |
|-----------|---|---|------------|----------------|------------|--------------|

TOLTERODINE TARTRATE

| | | | | | | |
|-----------|--|---------------------|------------|----------------|------------|--------------|
| <u>AB</u> | | HETERO LABS LTD III | <u>2MG</u> | <u>A206419</u> | <u>001</u> | Dec 12, 2017 |
|-----------|--|---------------------|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>4MG</u> | <u>A206419</u> | <u>002</u> | Dec 12, 2017 |
|-----------|--|--|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|------------------|------------|----------------|------------|--------------|
| <u>AB</u> | | MYLAN PHARMS INC | <u>2MG</u> | <u>A201486</u> | <u>001</u> | Oct 31, 2013 |
|-----------|--|------------------|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>4MG</u> | <u>A201486</u> | <u>002</u> | Oct 31, 2013 |
|-----------|--|--|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-----------------|------------|----------------|------------|--------------|
| <u>AB</u> | | TEVA PHARMS USA | <u>2MG</u> | <u>A079141</u> | <u>001</u> | Nov 22, 2016 |
|-----------|--|-----------------|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>4MG</u> | <u>A079141</u> | <u>002</u> | Nov 22, 2016 |
|-----------|--|--|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--------------------|------------|----------------|------------|--------------|
| <u>AB</u> | | TORRENT PHARMS LTD | <u>2MG</u> | <u>A203016</u> | <u>001</u> | Aug 11, 2015 |
|-----------|--|--------------------|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>4MG</u> | <u>A203016</u> | <u>002</u> | Aug 11, 2015 |
|-----------|--|--|------------|----------------|------------|--------------|

TABLET;ORAL

DETROL

| | | | | | | |
|-----------|---|-------------------------|------------|----------------|------------|--------------|
| <u>AB</u> | + | PHARMACIA AND UPJOHN | <u>1MG</u> | <u>N020771</u> | <u>001</u> | Mar 25, 1998 |
|-----------|---|-------------------------|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|---|------------|----------------|------------|--------------|
| <u>AB</u> | + | ! | <u>2MG</u> | <u>N020771</u> | <u>002</u> | Mar 25, 1998 |
|-----------|---|---|------------|----------------|------------|--------------|

TOLTERODINE TARTRATE

| | | | | | | |
|-----------|--|-------------------------|------------|----------------|------------|--------------|
| <u>AB</u> | | IVAX SUB TEVA PHARMS | <u>1MG</u> | <u>A077006</u> | <u>001</u> | Feb 23, 2015 |
|-----------|--|-------------------------|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>2MG</u> | <u>A077006</u> | <u>002</u> | Feb 23, 2015 |
|-----------|--|--|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|---------------------|------------|----------------|------------|--------------|
| <u>AB</u> | | MACLEODS PHARMS LTD | <u>1MG</u> | <u>A203409</u> | <u>001</u> | Aug 31, 2015 |
|-----------|--|---------------------|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>2MG</u> | <u>A203409</u> | <u>002</u> | Aug 31, 2015 |
|-----------|--|--|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|------------------|------------|----------------|------------|--------------|
| <u>AB</u> | | MYLAN PHARMS INC | <u>1MG</u> | <u>A202641</u> | <u>001</u> | Nov 27, 2012 |
|-----------|--|------------------|------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>2MG</u> | <u>A202641</u> | <u>002</u> | Nov 27, 2012 |
|-----------|--|--|------------|----------------|------------|--------------|

TOLVAPTAN

TABLET;ORAL

JYNARQUE

| | | | | | |
|---|---------------------|------|---------|-----|--------------|
| + | OTSUKA PHARM CO LTD | 15MG | N204441 | 001 | Apr 23, 2018 |
|---|---------------------|------|---------|-----|--------------|

| | | | | | |
|---|--|------|---------|-----|--------------|
| + | | 30MG | N204441 | 002 | Apr 23, 2018 |
|---|--|------|---------|-----|--------------|

| | | | | | |
|---|--|------|---------|-----|--------------|
| + | | 45MG | N204441 | 003 | Apr 23, 2018 |
|---|--|------|---------|-----|--------------|

| | | | | | |
|---|--|------|---------|-----|--------------|
| + | | 60MG | N204441 | 004 | Apr 23, 2018 |
|---|--|------|---------|-----|--------------|

| | | | | | |
|---|---|------|---------|-----|--------------|
| + | ! | 90MG | N204441 | 005 | Apr 23, 2018 |
|---|---|------|---------|-----|--------------|

SAMSCA

| | | | | | |
|---|-------------------------|------|---------|-----|--------------|
| + | OTSUKA AMERICA PHARM | 15MG | N022275 | 001 | May 19, 2009 |
|---|-------------------------|------|---------|-----|--------------|

| | | | | | |
|---|---|------|---------|-----|--------------|
| + | ! | 30MG | N022275 | 002 | May 19, 2009 |
|---|---|------|---------|-----|--------------|

TOPIRAMATE

CAPSULE;ORAL

TOPAMAX

| | | | | | | |
|-----------|---|----------------|-------------|----------------|------------|--------------|
| <u>AB</u> | + | JANSSEN PHARMS | <u>15MG</u> | <u>N020844</u> | <u>001</u> | Oct 26, 1998 |
|-----------|---|----------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|---|-------------|----------------|------------|--------------|
| <u>AB</u> | + | ! | <u>25MG</u> | <u>N020844</u> | <u>002</u> | Oct 26, 1998 |
|-----------|---|---|-------------|----------------|------------|--------------|

TOPIRAMATE

| | | | | | | |
|-----------|--|------|-------------|----------------|------------|--------------|
| <u>AB</u> | | TEVA | <u>15MG</u> | <u>A076575</u> | <u>001</u> | Apr 17, 2009 |
|-----------|--|------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>25MG</u> | <u>A076575</u> | <u>002</u> | Apr 17, 2009 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | WATSON LABS | <u>15MG</u> | <u>A077868</u> | <u>001</u> | Apr 15, 2009 |
|-----------|--|-------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>25MG</u> | <u>A077868</u> | <u>002</u> | Apr 15, 2009 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>15MG</u> | <u>A078877</u> | <u>001</u> | Oct 14, 2009 |
|-----------|--|-------------------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>25MG</u> | <u>A078877</u> | <u>002</u> | Oct 14, 2009 |
|-----------|--|--|-------------|----------------|------------|--------------|

CAPSULE, EXTENDED RELEASE;ORAL

TOPIRAMATE

| | | | | | | |
|-----------|--|-------------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>25MG</u> | <u>A207382</u> | <u>001</u> | Nov 24, 2017 |
|-----------|--|-------------------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>50MG</u> | <u>A207382</u> | <u>002</u> | Nov 24, 2017 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|--------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>100MG</u> | <u>A207382</u> | <u>003</u> | Nov 24, 2017 |
|-----------|--|--|--------------|----------------|------------|--------------|

TROKENDI XR

| | | | | | | |
|-----------|---|----------------|-------------|----------------|------------|--------------|
| <u>AB</u> | + | SUPERNU PHARMS | <u>25MG</u> | <u>N201635</u> | <u>001</u> | Aug 16, 2013 |
|-----------|---|----------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|--|-------------|----------------|------------|--------------|
| <u>AB</u> | + | | <u>50MG</u> | <u>N201635</u> | <u>002</u> | Aug 16, 2013 |
|-----------|---|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|--|--------------|----------------|------------|--------------|
| <u>AB</u> | + | | <u>100MG</u> | <u>N201635</u> | <u>003</u> | Aug 16, 2013 |
|-----------|---|--|--------------|----------------|------------|--------------|

QUDEXY XR

| | | | | | |
|---|-------------------|------|---------|-----|--------------|
| + | UPSHER SMITH LABS | 25MG | N205122 | 001 | Mar 11, 2014 |
|---|-------------------|------|---------|-----|--------------|

| | | | | | |
|---|--|------|---------|-----|--------------|
| + | | 50MG | N205122 | 002 | Mar 11, 2014 |
|---|--|------|---------|-----|--------------|

| | | | | | |
|---|--|-------|---------|-----|--------------|
| + | | 100MG | N205122 | 003 | Mar 11, 2014 |
|---|--|-------|---------|-----|--------------|

| | | | | | |
|---|--|-------|---------|-----|--------------|
| + | | 150MG | N205122 | 004 | Mar 11, 2014 |
|---|--|-------|---------|-----|--------------|

| | | | | | |
|---|---|-------|---------|-----|--------------|
| + | ! | 200MG | N205122 | 005 | Mar 11, 2014 |
|---|---|-------|---------|-----|--------------|

TROKENDI XR

| | | | | | |
|---|----------------|-------|---------|-----|--------------|
| + | SUPERNU PHARMS | 200MG | N201635 | 004 | Aug 16, 2013 |
|---|----------------|-------|---------|-----|--------------|

PRESCRIPTION DRUG PRODUCT LIST

TOPIRAMATE

TABLET; ORAL

TOPAMAX

| | | | | | | |
|-----------|---|----------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | JANSSEN PHARMS | <u>25MG</u> | <u>N020505</u> | <u>004</u> | Dec 24, 1996 |
| <u>AB</u> | + | | <u>50MG</u> | <u>N020505</u> | <u>005</u> | Dec 24, 1996 |
| <u>AB</u> | + | ! | <u>100MG</u> | <u>N020505</u> | <u>001</u> | Dec 24, 1996 |
| <u>AB</u> | + | | <u>200MG</u> | <u>N020505</u> | <u>002</u> | Dec 24, 1996 |

TOPIRAMATE

| | | | | | | |
|-----------|--|-------------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | ACCORD HLTHCARE | <u>25MG</u> | <u>A076311</u> | <u>001</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>50MG</u> | <u>A076311</u> | <u>002</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>100MG</u> | <u>A076311</u> | <u>003</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>200MG</u> | <u>A076311</u> | <u>004</u> | Mar 27, 2009 |
| <u>AB</u> | | APOTEX INC | <u>25MG</u> | <u>A077733</u> | <u>001</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>50MG</u> | <u>A077733</u> | <u>002</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>100MG</u> | <u>A077733</u> | <u>003</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>200MG</u> | <u>A077733</u> | <u>004</u> | Mar 27, 2009 |
| <u>AB</u> | | AUROBINDO PHARMA | <u>25MG</u> | <u>A078462</u> | <u>001</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>50MG</u> | <u>A078462</u> | <u>002</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>100MG</u> | <u>A078462</u> | <u>003</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>200MG</u> | <u>A078462</u> | <u>004</u> | Mar 27, 2009 |
| <u>AB</u> | | CIPLA LTD | <u>25MG</u> | <u>A076343</u> | <u>001</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>50MG</u> | <u>A076343</u> | <u>002</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>100MG</u> | <u>A076343</u> | <u>003</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>200MG</u> | <u>A076343</u> | <u>004</u> | Mar 27, 2009 |
| <u>AB</u> | | GLENMARK GENERICS | <u>25MG</u> | <u>A077627</u> | <u>001</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>50MG</u> | <u>A077627</u> | <u>002</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>100MG</u> | <u>A077627</u> | <u>003</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>200MG</u> | <u>A077627</u> | <u>004</u> | Mar 27, 2009 |
| <u>AB</u> | | INVAGEN PHARMS | <u>25MG</u> | <u>A079162</u> | <u>001</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>50MG</u> | <u>A079162</u> | <u>002</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>100MG</u> | <u>A079162</u> | <u>003</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>200MG</u> | <u>A079162</u> | <u>004</u> | Mar 27, 2009 |
| <u>AB</u> | | LUPIN | <u>25MG</u> | <u>A078410</u> | <u>001</u> | Sep 11, 2013 |
| <u>AB</u> | | | <u>50MG</u> | <u>A078410</u> | <u>002</u> | Sep 11, 2013 |
| <u>AB</u> | | | <u>100MG</u> | <u>A078410</u> | <u>003</u> | Sep 11, 2013 |
| <u>AB</u> | | | <u>200MG</u> | <u>A078410</u> | <u>004</u> | Sep 11, 2013 |
| <u>AB</u> | | SUN PHARM INDS LTD | <u>25MG</u> | <u>A076327</u> | <u>001</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>100MG</u> | <u>A076327</u> | <u>002</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>200MG</u> | <u>A076327</u> | <u>003</u> | Mar 27, 2009 |
| <u>AB</u> | | SUN PHARMA GLOBAL | <u>25MG</u> | <u>A090278</u> | <u>001</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>50MG</u> | <u>A090278</u> | <u>002</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>100MG</u> | <u>A090278</u> | <u>003</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>200MG</u> | <u>A090278</u> | <u>004</u> | Mar 27, 2009 |
| <u>AB</u> | | TEVA | <u>25MG</u> | <u>A076317</u> | <u>001</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>50MG</u> | <u>A076317</u> | <u>002</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>100MG</u> | <u>A076317</u> | <u>003</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>200MG</u> | <u>A076317</u> | <u>004</u> | Mar 27, 2009 |
| <u>AB</u> | | TORRENT PHARMS | <u>25MG</u> | <u>A079153</u> | <u>001</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>50MG</u> | <u>A079153</u> | <u>002</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>100MG</u> | <u>A079153</u> | <u>003</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>200MG</u> | <u>A079153</u> | <u>004</u> | Mar 27, 2009 |
| <u>AB</u> | | UNICHEM LABS LTD | <u>25MG</u> | <u>A090162</u> | <u>001</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>50MG</u> | <u>A090162</u> | <u>002</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>100MG</u> | <u>A090162</u> | <u>003</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>200MG</u> | <u>A090162</u> | <u>004</u> | Feb 19, 2013 |
| <u>AB</u> | | UPSHER SMITH LABS | <u>25MG</u> | <u>A078499</u> | <u>001</u> | Jan 07, 2010 |
| <u>AB</u> | | | <u>50MG</u> | <u>A078499</u> | <u>002</u> | Jan 07, 2010 |
| <u>AB</u> | | | <u>100MG</u> | <u>A078499</u> | <u>003</u> | Jan 07, 2010 |
| <u>AB</u> | | | <u>200MG</u> | <u>A078499</u> | <u>004</u> | Jan 07, 2010 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>25MG</u> | <u>A078235</u> | <u>001</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>50MG</u> | <u>A078235</u> | <u>002</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>100MG</u> | <u>A078235</u> | <u>003</u> | Mar 27, 2009 |
| <u>AB</u> | | | <u>200MG</u> | <u>A078235</u> | <u>004</u> | Mar 27, 2009 |

TOPOTECAN HYDROCHLORIDE

CAPSULE; ORAL

HYCANTIN

| | | | | | | |
|--|---|-------------------------|----------------|---------|-----|--------------|
| | + | NOVARTIS PHARMS CORP | EQ 0.25MG BASE | N020981 | 001 | Oct 11, 2007 |
| | + | ! | EQ 1MG BASE | N020981 | 002 | Oct 11, 2007 |

PRESCRIPTION DRUG PRODUCT LIST

TOPOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

HYCAMTIN

| | | | | | |
|-----------|------------|-------------------------|-------------------------|--------------------|--------------|
| <u>AP</u> | <u>+</u> ! | NOVARTIS PHARMS CORP | <u>EQ 4MG BASE/VIAL</u> | <u>N020671 001</u> | May 28, 1996 |
|-----------|------------|-------------------------|-------------------------|--------------------|--------------|

TOPOTECAN HYDROCHLORIDE

| | | | | | |
|-----------|--|--------------------|-------------------------|--------------------|--------------|
| <u>AP</u> | | ACCORD HLTHCARE | <u>EQ 4MG BASE/VIAL</u> | <u>A202351 001</u> | Jun 26, 2013 |
| <u>AP</u> | | ACTAVIS TOTOWA | <u>EQ 4MG BASE/VIAL</u> | <u>A090620 001</u> | Dec 02, 2010 |
| <u>AP</u> | | CIPLA | <u>EQ 4MG BASE/VIAL</u> | <u>A091199 001</u> | Dec 01, 2010 |
| <u>AP</u> | | DR REDDYS LABS LTD | <u>EQ 4MG BASE/VIAL</u> | <u>A201191 001</u> | Mar 09, 2011 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>EQ 4MG BASE/VIAL</u> | <u>A091089 001</u> | Nov 29, 2010 |
| <u>AP</u> | | HONG KONG | <u>EQ 4MG BASE/VIAL</u> | <u>A201166 001</u> | Aug 08, 2012 |
| <u>AP</u> | | MYLAN LABS LTD | <u>EQ 4MG BASE/VIAL</u> | <u>A091542 001</u> | Aug 28, 2012 |
| <u>AP</u> | | NOVAST LABS | <u>EQ 4MG BASE/VIAL</u> | <u>A206962 001</u> | Nov 30, 2016 |
| <u>AP</u> | | SAGENT PHARMS | <u>EQ 4MG BASE/VIAL</u> | <u>A091284 001</u> | Jan 26, 2011 |

SOLUTION; INTRAVENOUS

TOPOTECAN HYDROCHLORIDE

| | | | | | |
|-----------|------------|-----------------|-----------------------------------------|--------------------|--------------|
| <u>AP</u> | | ACCORD HLTHCARE | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A204406 002</u> | Jul 06, 2017 |
| <u>AP</u> | <u>+</u> ! | HOSPIRA INC | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>N200582 001</u> | Feb 02, 2011 |
| <u>AP</u> | | MYLAN LABS LTD | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>A206074 001</u> | Nov 24, 2017 |
| <u>AP</u> | | TEVA PHARMS USA | <u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u> | <u>N022453 001</u> | Dec 20, 2012 |
| | | ACCORD HLTHCARE | EQ 1MG BASE/ML (EQ 1MG BASE/ML) | A204406 001 | Jul 06, 2017 |

TOREMIFENE CITRATE

TABLET; ORAL

FARESTON

| | | | | | |
|-----------|------------|-------------|---------------------|--------------------|--------------|
| <u>AB</u> | <u>+</u> ! | KYOWA KIRIN | <u>EQ 60MG BASE</u> | <u>N020497 001</u> | May 29, 1997 |
|-----------|------------|-------------|---------------------|--------------------|--------------|

TOREMIFENE CITRATE

| | | | | | |
|-----------|--|---------------|---------------------|--------------------|--------------|
| <u>AB</u> | | RISING PHARMS | <u>EQ 60MG BASE</u> | <u>A208813 001</u> | Dec 04, 2018 |
|-----------|--|---------------|---------------------|--------------------|--------------|

TORSEMIDE

TABLET; ORAL

DEMADEX

| | | | | | |
|-----------|------------|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | <u>+</u> | MYLAN SPECIALITY LP | <u>5MG</u> | <u>N020136 001</u> | Aug 23, 1993 |
| <u>AB</u> | <u>+</u> | | <u>10MG</u> | <u>N020136 002</u> | Aug 23, 1993 |
| <u>AB</u> | <u>+</u> ! | | <u>20MG</u> | <u>N020136 003</u> | Aug 23, 1993 |
| <u>AB</u> | <u>+</u> | | <u>100MG</u> | <u>N020136 004</u> | Aug 23, 1993 |

TORSEMIDE

| | | | | | |
|-----------|--|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | | APOTEX INC | <u>5MG</u> | <u>A076894 001</u> | May 31, 2005 |
| <u>AB</u> | | | <u>10MG</u> | <u>A076894 002</u> | May 31, 2005 |
| <u>AB</u> | | | <u>20MG</u> | <u>A076894 003</u> | May 31, 2005 |
| <u>AB</u> | | | <u>100MG</u> | <u>A076894 004</u> | May 31, 2005 |
| <u>AB</u> | | AUROBINDO PHARMA | <u>5MG</u> | <u>A078249 001</u> | Oct 17, 2007 |
| <u>AB</u> | | | <u>10MG</u> | <u>A078249 002</u> | Oct 17, 2007 |
| <u>AB</u> | | | <u>20MG</u> | <u>A078249 003</u> | Oct 17, 2007 |
| <u>AB</u> | | | <u>100MG</u> | <u>A078249 004</u> | Oct 17, 2007 |
| <u>AB</u> | | HETERO LABS LTD III | <u>5MG</u> | <u>A079234 001</u> | Jan 27, 2009 |
| <u>AB</u> | | | <u>10MG</u> | <u>A079234 002</u> | Jan 27, 2009 |
| <u>AB</u> | | | <u>20MG</u> | <u>A079234 003</u> | Jan 27, 2009 |
| <u>AB</u> | | | <u>100MG</u> | <u>A079234 004</u> | Jan 27, 2009 |
| <u>AB</u> | | PAR PHARM | <u>5MG</u> | <u>A076226 001</u> | May 27, 2003 |
| <u>AB</u> | | | <u>10MG</u> | <u>A076226 002</u> | May 27, 2003 |
| <u>AB</u> | | | <u>20MG</u> | <u>A076226 003</u> | May 27, 2003 |
| <u>AB</u> | | | <u>100MG</u> | <u>A076226 004</u> | May 27, 2003 |
| <u>AB</u> | | PLIVA PHARM IND | <u>5MG</u> | <u>A076346 001</u> | May 30, 2003 |
| <u>AB</u> | | | <u>10MG</u> | <u>A076346 002</u> | May 30, 2003 |
| <u>AB</u> | | | <u>20MG</u> | <u>A076346 003</u> | May 30, 2003 |
| <u>AB</u> | | | <u>100MG</u> | <u>A076346 004</u> | Oct 19, 2004 |
| <u>AB</u> | | TEVA | <u>5MG</u> | <u>A076110 001</u> | May 14, 2002 |
| <u>AB</u> | | | <u>10MG</u> | <u>A076110 002</u> | May 14, 2002 |
| <u>AB</u> | | | <u>20MG</u> | <u>A076110 003</u> | May 14, 2002 |
| <u>AB</u> | | | <u>100MG</u> | <u>A076110 004</u> | May 14, 2002 |
| <u>AB</u> | | VINTAGE PHARMS | <u>5MG</u> | <u>A090613 001</u> | Mar 22, 2011 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090613 002</u> | Mar 22, 2011 |
| <u>AB</u> | | | <u>20MG</u> | <u>A090613 003</u> | Mar 22, 2011 |
| <u>AB</u> | | | <u>100MG</u> | <u>A090613 004</u> | Mar 22, 2011 |
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>5MG</u> | <u>A076943 001</u> | Mar 01, 2005 |
| <u>AB</u> | | | <u>10MG</u> | <u>A076943 002</u> | Mar 01, 2005 |
| <u>AB</u> | | | <u>20MG</u> | <u>A076943 003</u> | Mar 01, 2005 |

PRESCRIPTION DRUG PRODUCT LIST

TRABECTEDIN

POWDER; INTRAVENOUS

YONDELIS

+! JANSSEN PRODS 1MG/VIAL N207953 001 Oct 23, 2015

TRAMADOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CONZIP

+! CIPHER PHARMS INC 100MG N022370 001 May 07, 2010

+ 150MG N022370 004 Aug 01, 2011

+ 200MG N022370 002 May 07, 2010

+ 300MG N022370 003 May 07, 2010

TABLET; ORAL

TRAMADOL HYDROCHLORIDE**AB** ACI HEALTHCARE LTD **50MG** **A202075 001** Nov 28, 2011**AB** AMNEAL PHARMS **50MG** **A076003 001** Jun 20, 2002**AB** APOTEX **50MG** **A075981 001** Jul 10, 2002**AB** AUROBINDO PHARMA LTD **50MG** **A203494 001** Mar 31, 2014**AB** CSPEC OUYI PHARM CO **50MG** **A091498 001** Mar 29, 2013**AB** IPCA LABS LTD **50MG** **A201973 001** Nov 16, 2012**AB** MACLEODS PHARMS LTD **50MG** **A205702 001** Sep 25, 2015**AB** MYLAN **50MG** **A075986 001** Jun 21, 2002**AB** PLIVA **50MG** **A075982 001** Jul 01, 2002**AB** SPECGX LLC **50MG** **A075983 001** Jun 25, 2002**AB** SUN PHARM INDS INC **50MG** **A075964 001** Jun 19, 2002**AB** SUN PHARM INDUSTRIES **50MG** **A076100 001** Jun 20, 2002**AB** TEVA **50MG** **A075977 001** Jun 19, 2002**AB** ZYDUS PHARMS USA INC **50MG** **A090404 001** Jan 31, 2011ULTRAM**AB** +! JANSSEN PHARMS **50MG** **N020281 002** Mar 03, 1995

TABLET, EXTENDED RELEASE; ORAL

TRAMADOL HYDROCHLORIDE**AB1** AUROBINDO PHARMA LTD **100MG** **A204421 001** Oct 20, 2015**AB1** **200MG** **A204421 002** Oct 20, 2015**AB1** **300MG** **A204421 003** Oct 20, 2015**AB1** ! LUPIN LTD **100MG** **A200503 001** Aug 29, 2011**AB1** **200MG** **A200503 002** Aug 29, 2011**AB1** **300MG** **A200503 003** Aug 29, 2011**AB1** MYLAN PHARMS INC **100MG** **A205257 001** Dec 22, 2015**AB1** **200MG** **A205257 002** Dec 22, 2015**AB1** **300MG** **A205257 003** Dec 22, 2015**AB1** PAR PHARM INC **100MG** **A078783 001** Nov 13, 2009**AB1** **200MG** **A078783 002** Nov 13, 2009**AB1** **300MG** **A078783 003** Sep 20, 2011**AB1** SUN PHARMA GLOBAL **100MG** **A201384 001** Dec 07, 2011**AB1** **200MG** **A201384 002** Dec 07, 2011**AB1** **300MG** **A201384 003** Dec 07, 2011**AB2** ACTAVIS ELIZABETH **100MG** **A091609 001** Jun 27, 2012**AB2** **200MG** **A091609 002** Jun 27, 2012**AB2** **300MG** **A091609 003** Jun 27, 2012**AB2** ANCHEN PHARMS **100MG** **A200491 001** Jun 27, 2012**AB2** **200MG** **A200491 002** Jun 27, 2012**AB2** **300MG** **A200491 003** Jun 27, 2012**AB2** ! SUN PHARMA GLOBAL **100MG** **A091607 001** Dec 30, 2011**AB2** **200MG** **A091607 002** Dec 30, 2011**AB2** **300MG** **A091607 003** Dec 30, 2011TRAMETINIB DIMETHYL SULFOXIDE

TABLET; ORAL

MEKINIST

+ NOVARTIS PHARMS CORP EQ 0.5MG N204114 001 May 29, 2013

+! EQ 2MG N204114 003 May 29, 2013

TRANDOLAPRIL

TABLET; ORAL

TRANDOLAPRIL**AB** AUROBINDO PHARMA **1MG** **A078438 001** Jun 12, 2007**AB** **2MG** **A078438 002** Jun 12, 2007**AB** **4MG** **A078438 003** Jun 12, 2007**AB** EPIC PHARMA **1MG** **A078508 003** Jun 18, 2008

PRESCRIPTION DRUG PRODUCT LIST

TRANDOLAPRIL

TABLET; ORAL

TRANDOLAPRIL

| | | | | |
|-----------|-------------|------------|--------------------|--------------|
| <u>AB</u> | | <u>2MG</u> | <u>A078508 001</u> | Jun 18, 2008 |
| <u>AB</u> | | <u>4MG</u> | <u>A078508 002</u> | Jun 18, 2008 |
| <u>AB</u> | LUPIN | <u>1MG</u> | <u>A077522 001</u> | Jun 12, 2007 |
| <u>AB</u> | | <u>2MG</u> | <u>A077522 002</u> | Jun 12, 2007 |
| <u>AB</u> | ! | <u>4MG</u> | <u>A077522 003</u> | Jun 12, 2007 |
| <u>AB</u> | TEVA PHARMS | <u>1MG</u> | <u>A077489 001</u> | Dec 12, 2006 |
| <u>AB</u> | | <u>2MG</u> | <u>A077489 002</u> | Dec 12, 2006 |
| <u>AB</u> | | <u>4MG</u> | <u>A077489 003</u> | Dec 12, 2006 |

TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TARKA

| | | | | | |
|-----------|---|--------|------------------|--------------------|--------------|
| <u>AB</u> | + | ABBVIE | <u>1MG;240MG</u> | <u>N020591 003</u> | Oct 22, 1996 |
| <u>AB</u> | + | | <u>2MG;180MG</u> | <u>N020591 001</u> | Oct 22, 1996 |
| <u>AB</u> | + | | <u>2MG;240MG</u> | <u>N020591 004</u> | Oct 22, 1996 |
| <u>AB</u> | + | ! | <u>4MG;240MG</u> | <u>N020591 002</u> | Oct 22, 1996 |

TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE

| | | | | | |
|-----------|--|-------------------|------------------|--------------------|--------------|
| <u>AB</u> | | GLENMARK GENERICS | <u>1MG;240MG</u> | <u>A079135 004</u> | Aug 30, 2010 |
| <u>AB</u> | | | <u>2MG;180MG</u> | <u>A079135 001</u> | May 26, 2010 |
| <u>AB</u> | | | <u>2MG;240MG</u> | <u>A079135 002</u> | May 26, 2010 |
| <u>AB</u> | | | <u>4MG;240MG</u> | <u>A079135 003</u> | May 05, 2010 |

TRANEXAMIC ACID

INJECTABLE; INJECTION

CYKLOKAPRON

| | | | | | | |
|-----------|---|---|----------------------|-----------------|--------------------|--------------|
| <u>AP</u> | + | ! | PHARMACIA AND UPJOHN | <u>100MG/ML</u> | <u>N019281 001</u> | Dec 30, 1986 |
|-----------|---|---|----------------------|-----------------|--------------------|--------------|

TRANEXAMIC ACID

| | | | | | |
|-----------|--|----------------------|-----------------|--------------------|--------------|
| <u>AP</u> | | ACIC PHARMS | <u>100MG/ML</u> | <u>A202436 001</u> | Feb 11, 2014 |
| <u>AP</u> | | AKORN | <u>100MG/ML</u> | <u>A202373 001</u> | Nov 17, 2011 |
| <u>AP</u> | | | <u>100MG/ML</u> | <u>A206594 001</u> | Sep 28, 2017 |
| <u>AP</u> | | | <u>100MG/ML</u> | <u>A206634 001</u> | Jun 09, 2016 |
| <u>AP</u> | | AMNEAL PHARMS CO | <u>100MG/ML</u> | <u>A208840 001</u> | Feb 28, 2017 |
| <u>AP</u> | | AUROBINDO PHARMA LTD | <u>100MG/ML</u> | <u>A205035 001</u> | Jan 14, 2016 |
| <u>AP</u> | | EMCURE PHARMS LTD | <u>100MG/ML</u> | <u>A203521 001</u> | Aug 12, 2014 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>100MG/ML</u> | <u>A091596 001</u> | Mar 02, 2012 |
| <u>AP</u> | | GLAND PHARMA LTD | <u>100MG/ML</u> | <u>A207239 001</u> | Feb 13, 2017 |
| <u>AP</u> | | LUITPOLD | <u>100MG/ML</u> | <u>A201885 001</u> | Aug 10, 2011 |
| <u>AP</u> | | MYLAN INSTITUTIONAL | <u>100MG/ML</u> | <u>A091657 001</u> | Nov 03, 2011 |
| <u>AP</u> | | VIRTUS PHARMS | <u>100MG/ML</u> | <u>A202755 001</u> | Feb 25, 2016 |
| <u>AP</u> | | VIVA HLTHCARE | <u>100MG/ML</u> | <u>A206713 001</u> | Jun 27, 2017 |
| <u>AP</u> | | X-GEN PHARMS INC | <u>100MG/ML</u> | <u>A201580 001</u> | Jun 14, 2013 |
| <u>AP</u> | | ZYDUS PHARMS USA INC | <u>100MG/ML</u> | <u>A205228 001</u> | Jul 17, 2017 |

TABLET; ORAL

LYSTEDA

| | | | | | | |
|-----------|---|---|--------------------|--------------|--------------------|--------------|
| <u>AB</u> | + | ! | FERRING PHARMS INC | <u>650MG</u> | <u>N022430 001</u> | Nov 13, 2009 |
|-----------|---|---|--------------------|--------------|--------------------|--------------|

TRANEXAMIC ACID

| | | | | | |
|-----------|--|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | | ACTAVIS LABS FL INC | <u>650MG</u> | <u>A202093 001</u> | Dec 27, 2012 |
| <u>AB</u> | | APOTEX INC | <u>650MG</u> | <u>A202286 001</u> | Jan 27, 2014 |
| <u>AB</u> | | MYLAN | <u>650MG</u> | <u>A205133 001</u> | Sep 21, 2015 |

TRANLYCYPROMINE SULFATE

TABLET; ORAL

PARNATE

| | | | | | | |
|-----------|---|---|----------------------|---------------------|--------------------|--------------|
| <u>AB</u> | + | ! | CONCORDIA PHARMS INC | <u>EQ 10MG BASE</u> | <u>N012342 003</u> | Aug 16, 1985 |
|-----------|---|---|----------------------|---------------------|--------------------|--------------|

TRANLYCYPROMINE SULFATE

| | | | | | |
|-----------|--|------------------|---------------------|--------------------|--------------|
| <u>AB</u> | | CNTY LINE PHARMS | <u>EQ 10MG BASE</u> | <u>A206856 001</u> | Apr 17, 2018 |
| <u>AB</u> | | PAR PHARM | <u>EQ 10MG BASE</u> | <u>A040640 001</u> | Jun 29, 2006 |

TRAVOPROST

SOLUTION/DROPS; OPHTHALMIC

TRAVATAN Z

| | | | | | | |
|-----------|---|---|----------------------|---------------|--------------------|--------------|
| <u>AT</u> | + | ! | NOVARTIS PHARMS CORP | <u>0.004%</u> | <u>N021994 001</u> | Sep 21, 2006 |
|-----------|---|---|----------------------|---------------|--------------------|--------------|

TRAVOPROST

| | | | | | |
|-----------|---|------------------|---------------|--------------------|--------------|
| <u>AT</u> | | APOTEX INC | <u>0.004%</u> | <u>A203431 001</u> | Jul 10, 2015 |
| <u>AT</u> | | MYLAN PHARMS INC | <u>0.004%</u> | <u>A205050 001</u> | Jul 07, 2017 |
| | ! | PAR PHARM | <u>0.004%</u> | <u>A091340 001</u> | Mar 01, 2013 |

PRESCRIPTION DRUG PRODUCT LIST

TRAZODONE HYDROCHLORIDE

TABLET; ORAL

TRAZODONE HYDROCHLORIDE

| | | | | | |
|-----------|----------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ACCORD HLTHCARE | <u>50MG</u> | <u>A206923</u> | <u>001</u> | Sep 08, 2017 |
| <u>AB</u> | | <u>100MG</u> | <u>A206923</u> | <u>002</u> | Sep 08, 2017 |
| <u>AB</u> | | <u>150MG</u> | <u>A206923</u> | <u>003</u> | Sep 08, 2017 |
| <u>AB</u> | | <u>300MG</u> | <u>A206923</u> | <u>004</u> | Sep 08, 2017 |
| <u>AB</u> | ALVOGEN | <u>50MG</u> | <u>A071636</u> | <u>001</u> | Apr 18, 1988 |
| <u>AB</u> | | <u>100MG</u> | <u>A071514</u> | <u>001</u> | Apr 18, 1988 |
| <u>AB</u> | APOTEX | <u>50MG</u> | <u>A071258</u> | <u>001</u> | Mar 25, 1987 |
| <u>AB</u> | ! APOTEX INC | <u>100MG</u> | <u>A071196</u> | <u>001</u> | Mar 25, 1987 |
| <u>AB</u> | | <u>150MG</u> | <u>A071196</u> | <u>002</u> | Apr 26, 1999 |
| <u>AB</u> | | <u>300MG</u> | <u>A071196</u> | <u>003</u> | Apr 26, 1999 |
| <u>AB</u> | OXFORD PHARMS | <u>50MG</u> | <u>A072192</u> | <u>001</u> | Feb 02, 1989 |
| <u>AB</u> | | <u>100MG</u> | <u>A072193</u> | <u>001</u> | Feb 02, 1989 |
| <u>AB</u> | PLIVA | <u>150MG</u> | <u>A071525</u> | <u>001</u> | Mar 09, 1988 |
| <u>AB</u> | SUN PHARM INDUSTRIES | <u>50MG</u> | <u>A073137</u> | <u>002</u> | Mar 24, 1993 |
| <u>AB</u> | | <u>100MG</u> | <u>A073137</u> | <u>001</u> | Mar 24, 1993 |
| <u>AB</u> | | <u>150MG</u> | <u>A073137</u> | <u>003</u> | Dec 22, 1995 |
| <u>AB</u> | TEVA PHARMS USA | <u>50MG</u> | <u>A071523</u> | <u>001</u> | Dec 11, 1987 |
| <u>AB</u> | | <u>100MG</u> | <u>A071524</u> | <u>001</u> | Dec 11, 1987 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>50MG</u> | <u>A202180</u> | <u>001</u> | Nov 27, 2013 |
| <u>AB</u> | | <u>100MG</u> | <u>A202180</u> | <u>002</u> | Nov 27, 2013 |
| <u>AB</u> | | <u>150MG</u> | <u>A202180</u> | <u>003</u> | Nov 27, 2013 |
| <u>AB</u> | | <u>300MG</u> | <u>A202180</u> | <u>004</u> | Nov 27, 2013 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>50MG</u> | <u>A205253</u> | <u>001</u> | Oct 10, 2017 |
| <u>AB</u> | | <u>100MG</u> | <u>A205253</u> | <u>002</u> | Oct 10, 2017 |
| <u>AB</u> | | <u>150MG</u> | <u>A205253</u> | <u>003</u> | Oct 10, 2017 |
| <u>AB</u> | | <u>300MG</u> | <u>A205253</u> | <u>004</u> | Oct 10, 2017 |

TREPROSTINIL

INJECTABLE; IV (INFUSION), SUBCUTANEOUS

REMODYLIN

| | | | | | |
|-----------|-----------------|-----------------|----------------|------------|--------------|
| <u>AP</u> | ! UNITED THERAP | <u>1MG/ML</u> | <u>N021272</u> | <u>001</u> | May 21, 2002 |
| <u>AP</u> | ! | <u>2.5MG/ML</u> | <u>N021272</u> | <u>002</u> | May 21, 2002 |
| <u>AP</u> | ! | <u>5MG/ML</u> | <u>N021272</u> | <u>003</u> | May 21, 2002 |
| <u>AP</u> | ! | <u>10MG/ML</u> | <u>N021272</u> | <u>004</u> | May 21, 2002 |

TREPROSTINIL

| | | | | | |
|-----------|------------|-----------------|----------------|------------|--------------|
| <u>AP</u> | SANDOZ INC | <u>1MG/ML</u> | <u>A203649</u> | <u>001</u> | Nov 30, 2017 |
| <u>AP</u> | | <u>2.5MG/ML</u> | <u>A203649</u> | <u>002</u> | Nov 30, 2017 |
| <u>AP</u> | | <u>5MG/ML</u> | <u>A203649</u> | <u>003</u> | Nov 30, 2017 |
| <u>AP</u> | | <u>10MG/ML</u> | <u>A203649</u> | <u>004</u> | Nov 30, 2017 |

SOLUTION; INHALATION

TYVASO

| | | | | | |
|---|---------------|----------|---------|-----|--------------|
| ! | UNITED THERAP | 0.6MG/ML | N022387 | 001 | Jul 30, 2009 |
|---|---------------|----------|---------|-----|--------------|

SOLUTION; INTRAVENOUS, SUBCUTANEOUS

REMODYLIN

| | | | | | |
|--|---------------|----------------------|---------|-----|--------------|
| | UNITED THERAP | 20MG/20ML (1MG/ML) | N208276 | 001 | Jul 30, 2018 |
| | | 50MG/20ML (2.5MG/ML) | N208276 | 002 | Jul 30, 2018 |
| | | 100MG/20ML (5MG/ML) | N208276 | 003 | Jul 30, 2018 |
| | | 200MG/20ML (10MG/ML) | N208276 | 004 | Jul 30, 2018 |

TREPROSTINIL DIOLAMINE

TABLET, EXTENDED RELEASE; ORAL

ORENITRAM

| | | | | | |
|---|---------------|-----------------|---------|-----|--------------|
| + | UNITED THERAP | EQ 0.125MG BASE | N203496 | 001 | Dec 20, 2013 |
| + | | EQ 0.25MG BASE | N203496 | 002 | Dec 20, 2013 |
| + | | EQ 1MG BASE | N203496 | 003 | Dec 20, 2013 |
| + | ! | EQ 2.5MG BASE | N203496 | 004 | Dec 20, 2013 |
| + | | EQ 5MG BASE | N203496 | 005 | Oct 07, 2016 |

TRETINOIN

CAPSULE; ORAL

TRETINOIN

| | | | | | |
|-----------|---------------------|-------------|----------------|------------|--------------|
| <u>AB</u> | ANCHEN PHARMS | <u>10MG</u> | <u>A201687</u> | <u>001</u> | Oct 24, 2012 |
| <u>AB</u> | ! BARR LABS INC | <u>10MG</u> | <u>A077684</u> | <u>001</u> | Jun 22, 2007 |
| <u>AB</u> | GLENMARK PHARMS LTD | <u>10MG</u> | <u>A208279</u> | <u>001</u> | Dec 23, 2016 |

CREAM; TOPICAL

AVITA

| | | | | | |
|-----------|------------------|---------------|----------------|------------|--------------|
| <u>AB</u> | MYLAN PHARMS INC | <u>0.025%</u> | <u>N020404</u> | <u>003</u> | Jan 14, 1997 |
|-----------|------------------|---------------|----------------|------------|--------------|

RETIN-A

| | | | | | |
|-----------|-------------------|---------------|----------------|------------|--------------|
| <u>AB</u> | ! VALEANT BERMUDA | <u>0.025%</u> | <u>N019049</u> | <u>001</u> | Sep 16, 1988 |
|-----------|-------------------|---------------|----------------|------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

TRETINOIN

CREAM; TOPICAL

RETIN-A

| | | | | | | |
|-----------|---|-------------------------|-------------|----------------|------------|--|
| <u>AB</u> | + | VALEANT PHARMS NORTH | <u>0.1%</u> | <u>N017340</u> | <u>001</u> | |
|-----------|---|-------------------------|-------------|----------------|------------|--|

TRETINOIN

| | | | | | | |
|-----------|--|---------------------|---------------|----------------|------------|--------------|
| <u>AB</u> | | PERRIGO PHARMA INTL | <u>0.025%</u> | <u>A075264</u> | <u>001</u> | Dec 24, 1998 |
| <u>AB</u> | | | <u>0.1%</u> | <u>A075213</u> | <u>001</u> | Dec 24, 1998 |

RETIN-A

| | | | | | | |
|------------|---|-----------------|--------------|----------------|------------|--|
| <u>AB1</u> | + | VALEANT BERMUDA | <u>0.05%</u> | <u>N017522</u> | <u>001</u> | |
|------------|---|-----------------|--------------|----------------|------------|--|

TRETINOIN

| | | | | | | |
|------------|--|---------------------|--------------|----------------|------------|--------------|
| <u>AB1</u> | | PERRIGO PHARMA INTL | <u>0.05%</u> | <u>A075265</u> | <u>001</u> | Dec 24, 1998 |
|------------|--|---------------------|--------------|----------------|------------|--------------|

RENOVA

| | | | | | | |
|------------|---|-------------------------|--------------|----------------|------------|--------------|
| <u>AB2</u> | + | VALEANT PHARMS NORTH | <u>0.05%</u> | <u>N019963</u> | <u>001</u> | Dec 29, 1995 |
|------------|---|-------------------------|--------------|----------------|------------|--------------|

TRETINOIN

| | | | | | | |
|------------|--|----------------|--------------|----------------|------------|--------------|
| <u>AB2</u> | | ZO SKIN HEALTH | <u>0.05%</u> | <u>A076498</u> | <u>001</u> | Sep 15, 2005 |
|------------|--|----------------|--------------|----------------|------------|--------------|

RENOVA

| | | | | | | |
|--|---|-------------------------|-------|---------|-----|--------------|
| | + | VALEANT PHARMS NORTH | 0.02% | N021108 | 001 | Aug 31, 2000 |
|--|---|-------------------------|-------|---------|-----|--------------|

GEL; TOPICAL

ATRALIN

| | | | | | | |
|-----------|---|-----------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | DOW PHARM | <u>0.05%</u> | <u>N022070</u> | <u>001</u> | Jul 26, 2007 |
|-----------|---|-----------|--------------|----------------|------------|--------------|

RETIN-A

| | | | | | | |
|-----------|---|--------------|--------------|----------------|------------|--|
| <u>AB</u> | + | VALEANT INTL | <u>0.01%</u> | <u>N017955</u> | <u>001</u> | |
|-----------|---|--------------|--------------|----------------|------------|--|

| | | | | | | |
|-----------|---|--|---------------|----------------|------------|--|
| <u>AB</u> | + | | <u>0.025%</u> | <u>N017579</u> | <u>002</u> | |
|-----------|---|--|---------------|----------------|------------|--|

RETIN-A MICRO

| | | | | | | |
|-----------|---|--------------|--------------|----------------|------------|--------------|
| <u>AB</u> | + | VALEANT INTL | <u>0.04%</u> | <u>N020475</u> | <u>002</u> | May 10, 2002 |
|-----------|---|--------------|--------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|--|-------------|----------------|------------|--------------|
| <u>AB</u> | + | | <u>0.1%</u> | <u>N020475</u> | <u>001</u> | Feb 07, 1997 |
|-----------|---|--|-------------|----------------|------------|--------------|

TRETINOIN

| | | | | | | |
|-----------|--|------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | MYLAN PHARMS INC | <u>0.04%</u> | <u>A202567</u> | <u>001</u> | Jul 17, 2013 |
|-----------|--|------------------|--------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|--------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>0.05%</u> | <u>A207955</u> | <u>001</u> | Aug 13, 2015 |
|-----------|--|--|--------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>0.1%</u> | <u>A202026</u> | <u>001</u> | Jul 17, 2013 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|---------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | | PERRIGO PHARMA INTL | <u>0.01%</u> | <u>A075589</u> | <u>001</u> | Jun 11, 2002 |
|-----------|--|---------------------|--------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|---------------|----------------|------------|--------------|
| <u>AB</u> | | | <u>0.025%</u> | <u>A075529</u> | <u>001</u> | Feb 22, 2000 |
|-----------|--|--|---------------|----------------|------------|--------------|

AVITA

| | | | | | | |
|----|--|-------|--------|---------|-----|--------------|
| BT | | MYLAN | 0.025% | N020400 | 001 | Jan 29, 1998 |
|----|--|-------|--------|---------|-----|--------------|

RETIN-A-MICRO

| | | | | | | |
|--|---|--------------|-------|---------|-----|--------------|
| | + | VALEANT INTL | 0.06% | N020475 | 004 | Oct 23, 2017 |
|--|---|--------------|-------|---------|-----|--------------|

| | | | | | | |
|--|---|--|-------|---------|-----|--------------|
| | + | | 0.08% | N020475 | 003 | Jan 28, 2014 |
|--|---|--|-------|---------|-----|--------------|

LOTION; TOPICAL

ALTRENO

| | | | | | | |
|--|---|-----------|-------|---------|-----|--------------|
| | + | DOW PHARM | 0.05% | N209353 | 001 | Aug 23, 2018 |
|--|---|-----------|-------|---------|-----|--------------|

TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

TRIAMCINOLONE ACETONIDE

| | | | | | | |
|-----------|--|----------------|---------------|----------------|------------|--------------|
| <u>AT</u> | | ALKEM LABS LTD | <u>0.025%</u> | <u>A207651</u> | <u>001</u> | Dec 26, 2017 |
|-----------|--|----------------|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AT</u> | | | <u>0.1%</u> | <u>A207651</u> | <u>002</u> | Dec 26, 2017 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AT</u> | | | <u>0.5%</u> | <u>A207651</u> | <u>003</u> | Dec 26, 2017 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|----------------|---------------|----------------|------------|--|
| <u>AT</u> | ! | FOUGERA PHARMS | <u>0.025%</u> | <u>A085692</u> | <u>001</u> | |
|-----------|---|----------------|---------------|----------------|------------|--|

| | | | | | | |
|-----------|---|--|-------------|----------------|------------|--|
| <u>AT</u> | ! | | <u>0.1%</u> | <u>A085692</u> | <u>003</u> | |
|-----------|---|--|-------------|----------------|------------|--|

| | | | | | | |
|-----------|---|--|-------------|----------------|------------|--|
| <u>AT</u> | ! | | <u>0.5%</u> | <u>A085692</u> | <u>002</u> | |
|-----------|---|--|-------------|----------------|------------|--|

| | | | | | | |
|-----------|--|--------------|---------------|----------------|------------|--------------|
| <u>AT</u> | | G AND W LABS | <u>0.025%</u> | <u>A089797</u> | <u>001</u> | May 31, 1991 |
|-----------|--|--------------|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AT</u> | | | <u>0.1%</u> | <u>A089798</u> | <u>001</u> | May 31, 1991 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|---------------------|-------------|----------------|------------|--------------|
| <u>AT</u> | | GLENMARK PHARMS LTD | <u>0.1%</u> | <u>A207117</u> | <u>001</u> | Aug 05, 2016 |
|-----------|--|---------------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|----------------|---------------|----------------|------------|--------------|
| <u>AT</u> | | LANNETT CO INC | <u>0.025%</u> | <u>A040671</u> | <u>001</u> | Jun 09, 2006 |
|-----------|--|----------------|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AT</u> | | | <u>0.1%</u> | <u>A040671</u> | <u>002</u> | Jun 09, 2006 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|----------------|---------------|----------------|------------|--------------|
| <u>AT</u> | | LUPIN ATLANTIS | <u>0.025%</u> | <u>A208763</u> | <u>001</u> | Feb 01, 2017 |
|-----------|--|----------------|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AT</u> | | | <u>0.1%</u> | <u>A208763</u> | <u>002</u> | Feb 01, 2017 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AT</u> | | | <u>0.5%</u> | <u>A208763</u> | <u>003</u> | Feb 01, 2017 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|---------------------|---------------|----------------|------------|--------------|
| <u>AT</u> | | MACLEODS PHARMS LTD | <u>0.025%</u> | <u>A209535</u> | <u>001</u> | May 18, 2018 |
|-----------|--|---------------------|---------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AT</u> | | | <u>0.1%</u> | <u>A209535</u> | <u>002</u> | May 18, 2018 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--------------|
| <u>AT</u> | | | <u>0.5%</u> | <u>A209535</u> | <u>003</u> | May 18, 2018 |
|-----------|--|--|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|---|------------------|---------------|----------------|------------|--|
| <u>AT</u> | + | MYLAN PHARMS INC | <u>0.025%</u> | <u>N011601</u> | <u>003</u> | |
|-----------|---|------------------|---------------|----------------|------------|--|

| | | | | | | |
|-----------|---|--|-------------|----------------|------------|--|
| <u>AT</u> | + | | <u>0.1%</u> | <u>N011601</u> | <u>006</u> | |
|-----------|---|--|-------------|----------------|------------|--|

| | | | | | | |
|-----------|--|------------------|---------------|----------------|------------|--|
| <u>AT</u> | | PERRIGO NEW YORK | <u>0.025%</u> | <u>A086413</u> | <u>002</u> | |
|-----------|--|------------------|---------------|----------------|------------|--|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--|
| <u>AT</u> | | | <u>0.1%</u> | <u>A086413</u> | <u>003</u> | |
|-----------|--|--|-------------|----------------|------------|--|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--|
| <u>AT</u> | | | <u>0.1%</u> | <u>A086414</u> | <u>001</u> | |
|-----------|--|--|-------------|----------------|------------|--|

| | | | | | | |
|-----------|--|--|-------------|----------------|------------|--|
| <u>AT</u> | | | <u>0.5%</u> | <u>A086413</u> | <u>001</u> | |
|-----------|--|--|-------------|----------------|------------|--|

| | | | | | | |
|-----------|--|---------------------|-------------|----------------|------------|--------------|
| <u>AT</u> | | TARO PHARM INDS LTD | <u>0.1%</u> | <u>A040039</u> | <u>001</u> | Nov 26, 1997 |
|-----------|--|---------------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|---------------------|-------------|----------------|------------|--------------|
| <u>AT</u> | | TELIGENT PHARMA INC | <u>0.1%</u> | <u>A208848</u> | <u>001</u> | Sep 18, 2017 |
|-----------|--|---------------------|-------------|----------------|------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

TRIDERM

| | | | | | |
|-----------|------------|---------------|----------------|------------|--------------|
| AT | CROWN LABS | <u>0.025%</u> | <u>A088042</u> | <u>002</u> | Mar 25, 2015 |
| AT | | <u>0.1%</u> | <u>A088042</u> | <u>001</u> | Mar 19, 1984 |
| AT | | <u>0.5%</u> | <u>A088042</u> | <u>003</u> | Mar 25, 2015 |

FOR SUSPENSION, EXTENDED RELEASE; INTRA-ARTICULAR

ZILRETTA

+! FLEXION THERAPS INC 32MG/VIAL N208845 001 Oct 06, 2017

INJECTABLE; INJECTION

KENALOG-40

| | | | | | |
|-----------|--------------|----------------|----------------|------------|--|
| AB | +! APOTHECON | <u>40MG/ML</u> | <u>N014901</u> | <u>001</u> | |
|-----------|--------------|----------------|----------------|------------|--|

TRIAMCINOLONE ACETONIDE

| | | | | | |
|-----------|------------------|----------------|----------------|------------|--------------|
| AB | AMNEAL PHARMS CO | <u>40MG/ML</u> | <u>A207550</u> | <u>001</u> | Dec 11, 2017 |
|-----------|------------------|----------------|----------------|------------|--------------|

| | | | | | |
|-----------|-----------------|----------------|----------------|------------|--------------|
| AB | TEVA PHARMS USA | <u>40MG/ML</u> | <u>A209852</u> | <u>001</u> | Oct 05, 2018 |
|-----------|-----------------|----------------|----------------|------------|--------------|

KENALOG-10

+ APOTHECON 10MG/ML N012041 001

INJECTABLE; INTRAVITREAL

TRIESENCE

+! NOVARTIS PHARMS 40MG/ML (40MG/ML) N022048 001 Nov 29, 2007

CORP

LOTION; TOPICAL

TRIAMCINOLONE ACETONIDE

| | | | | | |
|-----------|-------|---------------|----------------|------------|--------------|
| AT | AKORN | <u>0.025%</u> | <u>A202374</u> | <u>001</u> | May 08, 2013 |
|-----------|-------|---------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-------------|----------------|------------|--------------|
| AT | | <u>0.1%</u> | <u>A202374</u> | <u>002</u> | May 08, 2013 |
|-----------|--|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|----------------|---------------|----------------|------------|--------------|
| AT | FOUGERA PHARMS | <u>0.025%</u> | <u>A040467</u> | <u>001</u> | Apr 21, 2003 |
|-----------|----------------|---------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-------------|----------------|------------|--------------|
| AT | | <u>0.1%</u> | <u>A040467</u> | <u>002</u> | Apr 21, 2003 |
|-----------|--|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|------------------|-------------|----------------|------------|--------------|
| AT | G AND W LABS INC | <u>0.1%</u> | <u>A089129</u> | <u>001</u> | Aug 14, 1986 |
|-----------|------------------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|----------------|-------------|----------------|------------|--------------|
| AT | LANNETT CO INC | <u>0.1%</u> | <u>A040672</u> | <u>002</u> | Dec 13, 2006 |
|-----------|----------------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|---------------------|---------------|----------------|------------|--------------|
| AT | TELIGENT PHARMA INC | <u>0.025%</u> | <u>A204608</u> | <u>001</u> | Jul 07, 2016 |
|-----------|---------------------|---------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-------------|----------------|------------|--------------|
| AT | | <u>0.1%</u> | <u>A204606</u> | <u>001</u> | Jul 07, 2016 |
|-----------|--|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|--------------------|---------------|----------------|------------|--------------|
| AT | ! WOCKHARDT BIO AG | <u>0.025%</u> | <u>A088450</u> | <u>001</u> | Apr 01, 1985 |
|-----------|--------------------|---------------|----------------|------------|--------------|

| | | | | | |
|-----------|---|-------------|----------------|------------|--------------|
| AT | ! | <u>0.1%</u> | <u>A088451</u> | <u>001</u> | Apr 03, 1985 |
|-----------|---|-------------|----------------|------------|--------------|

OINTMENT; TOPICAL

TRIAMCINOLONE ACETATE

| | | | | | |
|-----------|---------------------|---------------|----------------|------------|--------------|
| AT | MACLEODS PHARMS LTD | <u>0.025%</u> | <u>A209828</u> | <u>001</u> | Nov 23, 2018 |
|-----------|---------------------|---------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-------------|----------------|------------|--------------|
| AT | | <u>0.1%</u> | <u>A209828</u> | <u>002</u> | Nov 23, 2018 |
|-----------|--|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-------------|----------------|------------|--------------|
| AT | | <u>0.5%</u> | <u>A209828</u> | <u>003</u> | Nov 23, 2018 |
|-----------|--|-------------|----------------|------------|--------------|

TRIAMCINOLONE ACETONIDE

| | | | | | |
|-----------|----------------|---------------|----------------|------------|--|
| AT | FOUGERA PHARMS | <u>0.025%</u> | <u>A085691</u> | <u>001</u> | |
|-----------|----------------|---------------|----------------|------------|--|

| | | | | | |
|-----------|--|-------------|----------------|------------|--|
| AT | | <u>0.1%</u> | <u>A085691</u> | <u>003</u> | |
|-----------|--|-------------|----------------|------------|--|

| | | | | | |
|-----------|--|-------------|----------------|------------|--|
| AT | | <u>0.5%</u> | <u>A085691</u> | <u>002</u> | |
|-----------|--|-------------|----------------|------------|--|

| | | | | | |
|-----------|--------------|---------------|----------------|------------|--------------|
| AT | G AND W LABS | <u>0.025%</u> | <u>A089795</u> | <u>001</u> | Dec 23, 1988 |
|-----------|--------------|---------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-------------|----------------|------------|--------------|
| AT | | <u>0.1%</u> | <u>A089796</u> | <u>001</u> | Dec 23, 1988 |
|-----------|--|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|------------------|-------------|----------------|------------|--------------|
| AT | G AND W LABS INC | <u>0.5%</u> | <u>A208925</u> | <u>001</u> | Oct 06, 2017 |
|-----------|------------------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|-----------------|-------------|----------------|------------|--------------|
| AT | GLENMARK PHARMS | <u>0.1%</u> | <u>A208320</u> | <u>001</u> | Aug 22, 2017 |
|-----------|-----------------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|---------------------|-------------|----------------|------------|--------------|
| AT | GLENMARK PHARMS LTD | <u>0.5%</u> | <u>A206379</u> | <u>001</u> | Jul 22, 2016 |
|-----------|---------------------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|----------------|-------------|----------------|------------|--------------|
| AT | NOVEL LABS INC | <u>0.1%</u> | <u>A207365</u> | <u>001</u> | Oct 12, 2018 |
|-----------|----------------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|--------------------|---------------|----------------|------------|--|
| AT | ! PERRIGO NEW YORK | <u>0.025%</u> | <u>A087385</u> | <u>002</u> | |
|-----------|--------------------|---------------|----------------|------------|--|

| | | | | | |
|-----------|---|-------------|----------------|------------|--|
| AT | ! | <u>0.1%</u> | <u>A087385</u> | <u>003</u> | |
|-----------|---|-------------|----------------|------------|--|

| | | | | | |
|-----------|---|-------------|----------------|------------|--|
| AT | ! | <u>0.5%</u> | <u>A087385</u> | <u>001</u> | |
|-----------|---|-------------|----------------|------------|--|

| | | | | | |
|-----------|---------------------|-------------|----------------|------------|--------------|
| AT | TARO PHARM INDS LTD | <u>0.1%</u> | <u>A040037</u> | <u>001</u> | Sep 30, 1994 |
|-----------|---------------------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|---------------------|-------------|----------------|------------|--------------|
| AT | TELIGENT PHARMA INC | <u>0.1%</u> | <u>A205373</u> | <u>001</u> | May 13, 2016 |
|-----------|---------------------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|--|-------------|----------------|------------|--------------|
| AT | | <u>0.5%</u> | <u>A208590</u> | <u>001</u> | Mar 03, 2017 |
|-----------|--|-------------|----------------|------------|--------------|

TRIAMCINOLONE ACETONIDE IN ABSORBASE

! CMP PHARMA INC 0.05% A089595 001 Mar 23, 1995

PASTE; DENTAL

TRIAMCINOLONE ACETONIDE

| | | | | | |
|-----------|-------|-------------|----------------|------------|--------------|
| AT | AKORN | <u>0.1%</u> | <u>A206312</u> | <u>001</u> | Aug 11, 2016 |
|-----------|-------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|------------------|-------------|----------------|------------|--------------|
| AT | G AND W LABS INC | <u>0.1%</u> | <u>A205592</u> | <u>001</u> | Jan 12, 2017 |
|-----------|------------------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|------|-------------|----------------|------------|--------------|
| AT | LYNE | <u>0.1%</u> | <u>A040771</u> | <u>001</u> | Jul 01, 2010 |
|-----------|------|-------------|----------------|------------|--------------|

| | | | | | |
|-----------|--------|-------------|----------------|------------|--------------|
| AT | ! TARO | <u>0.1%</u> | <u>A070730</u> | <u>001</u> | Oct 01, 1986 |
|-----------|--------|-------------|----------------|------------|--------------|

SPRAY; TOPICAL

KENALOG

| | | | | | |
|-----------|-----------------------|-------------------|----------------|------------|--|
| AT | +! SUN PHARM INDS INC | <u>0.147MG/GM</u> | <u>N012104</u> | <u>001</u> | |
|-----------|-----------------------|-------------------|----------------|------------|--|

TRIAMCINOLONE ACETONIDE

| | | | | | |
|-----------|-------|-------------------|----------------|------------|--------------|
| AT | AKORN | <u>0.147MG/GM</u> | <u>A207094</u> | <u>001</u> | Dec 07, 2016 |
|-----------|-------|-------------------|----------------|------------|--------------|

| | | | | | |
|-----------|------------------|-------------------|----------------|------------|--------------|
| AT | PERRIGO UK FINCO | <u>0.147MG/GM</u> | <u>A205782</u> | <u>001</u> | Apr 13, 2015 |
|-----------|------------------|-------------------|----------------|------------|--------------|

| | | | | | |
|-----------|---------------|-------------------|----------------|------------|--------------|
| AT | RISING PHARMS | <u>0.147MG/GM</u> | <u>A206786</u> | <u>001</u> | Sep 08, 2017 |
|-----------|---------------|-------------------|----------------|------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

TRIAMCINOLONE HEXACETONIDE

INJECTABLE; INJECTION

ARISTOSPAN

+! SANDOZ INC

5MG/ML

N016466 001

+!

20MG/ML

N016466 002

TRIAMTERENE

CAPSULE; ORAL

DYRENIUM

+ CONCORDIA PHARMS
INC

50MG

N013174 001

+!

100MG

N013174 002

TRIAZOLAM

TABLET; ORAL

HALCIONAB + PHARMACIA AND
UPJOHN0.125MGN017892 003 Apr 26, 1985AB +!0.25MGN017892 001 Nov 15, 1982TRIAZOLAMAB MYLAN PHARMS INC0.125MGA074031 001 Mar 25, 1994AB0.25MGA074031 002 Mar 25, 1994AB WEST-WARD PHARMS
INT0.125MGA074224 001 Jun 01, 1994AB0.25MGA074224 002 Jun 01, 1994TRIENTINE HYDROCHLORIDE

CAPSULE; ORAL

SYPRINEAB +! ATON250MGN019194 001 Nov 08, 1985TRIENTINE HYDROCHLORIDEAB NAVINTA LLC250MGA211251 001 Jan 16, 2019AB

WATSON LABS TEVA

250MGA207567 001 Feb 07, 2018TRIFLUOPERAZINE HYDROCHLORIDE

TABLET; ORAL

TRIFLUOPERAZINE HYDROCHLORIDEAB MYLANEQ 1MG BASEA040209 001 Jul 07, 1997ABEQ 2MG BASEA040209 002 Jul 07, 1997ABEQ 5MG BASEA040209 003 Jul 07, 1997AB !EQ 10MG BASEA040209 004 Jul 07, 1997AB

SANDOZ

EQ 1MG BASEA085785 001ABEQ 2MG BASEA085786 001ABEQ 5MG BASEA085789 001ABEQ 10MG BASEA085788 001TRIFLURIDINE

SOLUTION/DROPS; OPHTHALMIC

TRIFLURIDINEAT HI-TECH PHARMACAL1%A205438 001 Jul 28, 2017AT

SANDOZ INC

1%A074311 001 Oct 06, 1995VIROPTICAT +! MONARCH PHARMS1%N018299 001TRIHEXYPHENIDYL HYDROCHLORIDE

ELIXIR; ORAL

TRIHEXYPHENIDYL HYDROCHLORIDEAA MIKART2MG/5MLA040251 001 Sep 27, 1999AA !

PHARM ASSOC

2MG/5MLA040177 001 Apr 17, 1997

TABLET; ORAL

TRIHEXYPHENIDYL HYDROCHLORIDEAA NATCO PHARMA LTD2MGA091630 001 Nov 17, 2010AA5MGA091630 002 Nov 17, 2010AA

NOVITIUM PHARMA

2MGA040254 001 Dec 24, 1998AA5MGA040254 002 Dec 24, 1998AA !

WATSON LABS

2MGA084363 001AA !5MGA084364 001TRIMETHADIONE

TABLET; ORAL

TRIDIONE

+! ABBVIE

150MG

N005856 009

PRESCRIPTION DRUG PRODUCT LIST

TRIMETHOBENZAMIDE HYDROCHLORIDE

CAPSULE; ORAL

TIGAN

| | | | | | | |
|-----------|------------|-----------------|--------------|----------------|------------|--------------|
| AB | + ! | KING PHARMS LLC | 300MG | N017531 | 006 | Dec 13, 2001 |
|-----------|------------|-----------------|--------------|----------------|------------|--------------|

TRIMETHOBENZAMIDE HYDROCHLORIDE

| | | | | | | |
|-----------|--|--------------|--------------|----------------|------------|--------------|
| AB | | GAVIS PHARMS | 300MG | A076546 | 001 | Aug 20, 2003 |
|-----------|--|--------------|--------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|----------------------|--------------|----------------|------------|--------------|
| AB | | SUN PHARM INDUSTRIES | 300MG | A076570 | 001 | Aug 28, 2003 |
|-----------|--|----------------------|--------------|----------------|------------|--------------|

INJECTABLE; INJECTION

TIGAN

| | | | | | | |
|-----------|------------|----------------------|-----------------|----------------|------------|--|
| AP | + ! | PAR STERILE PRODUCTS | 100MG/ML | N017530 | 001 | |
|-----------|------------|----------------------|-----------------|----------------|------------|--|

TRIMETHOBENZAMIDE HYDROCHLORIDE

| | | | | | | |
|-----------|--|----------|-----------------|----------------|------------|--------------|
| AP | | LUITPOLD | 100MG/ML | A091330 | 001 | Mar 08, 2011 |
|-----------|--|----------|-----------------|----------------|------------|--------------|

TRIMETHOBENZAMIDE HYDROCHLORIDE PRESERVATIVE FREE

| | | | | | | |
|-----------|--|----------|-----------------|----------------|------------|--------------|
| AP | | LUITPOLD | 100MG/ML | A091329 | 001 | Mar 08, 2011 |
|-----------|--|----------|-----------------|----------------|------------|--------------|

TRIMETHOPRIM

TABLET; ORAL

TRIMETHOPRIM

| | | | | | | |
|-----------|------------|--------------|--------------|----------------|------------|--------------|
| AB | + ! | MAYNE PHARMA | 100MG | N018679 | 001 | Jul 30, 1982 |
|-----------|------------|--------------|--------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|----------------|--------------|----------------|------------|--------------|
| AB | | NOVEL LABS INC | 100MG | A091437 | 001 | Jun 15, 2011 |
|-----------|--|----------------|--------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-------------|--------------|----------------|------------|--------------|
| AB | | WATSON LABS | 100MG | A070049 | 001 | Jun 06, 1985 |
|-----------|--|-------------|--------------|----------------|------------|--------------|

TRIMETHOPRIM HYDROCHLORIDE

SOLUTION; ORAL

PRIMSOL

| | | | | | |
|------------|---------|------------------|----------------|------------|--------------|
| + ! | ALLEGIS | EQ 50MG BASE/5ML | N074973 | 001 | Jan 24, 2000 |
|------------|---------|------------------|----------------|------------|--------------|

TRIMIPRAMINE MALEATE

CAPSULE; ORAL

SURMONTIL

| | | | | | | |
|-----------|----------|----------------|---------------------|----------------|------------|--|
| AB | + | ODYSSEY PHARMS | EQ 25MG BASE | N016792 | 001 | |
|-----------|----------|----------------|---------------------|----------------|------------|--|

| | | | | | | |
|-----------|------------|--|---------------------|----------------|------------|--|
| AB | + ! | | EQ 50MG BASE | N016792 | 002 | |
|-----------|------------|--|---------------------|----------------|------------|--|

| | | | | | | |
|-----------|----------|--|----------------------|----------------|------------|--------------|
| AB | + | | EQ 100MG BASE | N016792 | 003 | Sep 15, 1982 |
|-----------|----------|--|----------------------|----------------|------------|--------------|

TRIMIPRAMINE MALEATE

| | | | | | | |
|-----------|--|----------------|---------------------|----------------|------------|--------------|
| AB | | CROSSMEDIKA SA | EQ 25MG BASE | A208127 | 001 | Apr 15, 2016 |
|-----------|--|----------------|---------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|---------------------|----------------|------------|--------------|
| AB | | | EQ 50MG BASE | A208127 | 002 | Apr 15, 2016 |
|-----------|--|--|---------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|----------------------|----------------|------------|--------------|
| AB | | | EQ 100MG BASE | A208127 | 003 | Apr 15, 2016 |
|-----------|--|--|----------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|----------------|---------------------|----------------|------------|--------------|
| AB | | ELITE LABS INC | EQ 25MG BASE | A077361 | 001 | Aug 02, 2006 |
|-----------|--|----------------|---------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|---------------------|----------------|------------|--------------|
| AB | | | EQ 50MG BASE | A077361 | 002 | Aug 02, 2006 |
|-----------|--|--|---------------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|----------------------|----------------|------------|--------------|
| AB | | | EQ 100MG BASE | A077361 | 003 | Aug 02, 2006 |
|-----------|--|--|----------------------|----------------|------------|--------------|

TRIPTORELIN PAMOATE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

TRIPTODUR KIT

| | | | | | |
|------------|------------------|---------------------|----------------|------------|--------------|
| + ! | ARBOR PHARMS LLC | EQ 22.5MG BASE/VIAL | N208956 | 001 | Jun 29, 2017 |
|------------|------------------|---------------------|----------------|------------|--------------|

INJECTABLE; INTRAMUSCULAR

TRELSTAR

| | | | | | |
|------------|--------------------|---------------------|----------------|------------|--------------|
| + ! | ALLERGAN SALES LLC | EQ 3.75MG BASE/VIAL | N020715 | 001 | Jun 15, 2000 |
|------------|--------------------|---------------------|----------------|------------|--------------|

| | | | | | |
|------------|--|----------------------|----------------|------------|--------------|
| + ! | | EQ 11.25MG BASE/VIAL | N021288 | 001 | Jun 29, 2001 |
|------------|--|----------------------|----------------|------------|--------------|

| | | | | | |
|------------|--|---------------------|----------------|------------|--------------|
| + ! | | EQ 22.5MG BASE/VIAL | N022437 | 001 | Mar 10, 2010 |
|------------|--|---------------------|----------------|------------|--------------|

TROPICAMIDE

SOLUTION/DROPS; OPHTHALMIC

MYDRIACYL

| | | | | | | |
|-----------|----------|----------------------|-----------|----------------|------------|--|
| AT | ! | NOVARTIS PHARMS CORP | 1% | A084306 | 001 | |
|-----------|----------|----------------------|-----------|----------------|------------|--|

| | | | | | | |
|-----------|----------|------------|-------------|----------------|------------|--|
| AT | ! | SANDOZ INC | 0.5% | A084305 | 001 | |
|-----------|----------|------------|-------------|----------------|------------|--|

TROPICACYL

| | | | | | | |
|-----------|--|-------|-------------|----------------|------------|--------------|
| AT | | AKORN | 0.5% | A040314 | 001 | Sep 29, 2000 |
|-----------|--|-------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-----------|----------------|------------|--------------|
| AT | | | 1% | A040315 | 001 | Sep 29, 2000 |
|-----------|--|--|-----------|----------------|------------|--------------|

TROPICAMIDE

| | | | | | | |
|-----------|--|-----------------|-------------|----------------|------------|--------------|
| AT | | BAUSCH AND LOMB | 0.5% | A040067 | 001 | Jul 27, 1994 |
|-----------|--|-----------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|--|-----------|----------------|------------|--------------|
| AT | | | 1% | A040064 | 001 | Jul 27, 1994 |
|-----------|--|--|-----------|----------------|------------|--------------|

TROSPIMUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

TROSPIMUM CHLORIDE

| | | | | | | |
|-----------|----------|---------------------|-------------|----------------|------------|--------------|
| AB | ! | ACTAVIS LABS FL INC | 60MG | A091289 | 001 | Oct 12, 2012 |
|-----------|----------|---------------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|-------------|-------------|----------------|------------|--------------|
| AB | | PADDOCK LLC | 60MG | A201291 | 001 | May 24, 2013 |
|-----------|--|-------------|-------------|----------------|------------|--------------|

TABLET; ORAL

TROSPIMUM CHLORIDE

| | | | | | | |
|-----------|--|--------|-------------|----------------|------------|--------------|
| AB | | APOTEX | 20MG | A091513 | 001 | Dec 06, 2011 |
|-----------|--|--------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|----------|-------------------|-------------|----------------|------------|--------------|
| AB | ! | GLENMARK GENERICS | 20MG | A091575 | 001 | Aug 13, 2010 |
|-----------|----------|-------------------|-------------|----------------|------------|--------------|

| | | | | | | |
|-----------|--|---------------------|-------------|----------------|------------|--------------|
| AB | | HERITAGE PHARMS INC | 20MG | A204945 | 001 | Aug 30, 2016 |
|-----------|--|---------------------|-------------|----------------|------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

TROSPIUM CHLORIDE

TABLET; ORAL

TROSPIUM CHLORIDE

| | | | | |
|-----------|----------------|-------------|--------------------|--------------|
| AB | INVAGEN PHARMS | 20MG | A091688 001 | Aug 23, 2016 |
| AB | PADDOCK LLC | 20MG | A091573 001 | Nov 17, 2010 |

TRYPAN BLUESOLUTION; OPHTHALMIC
MEMBRANEBLUE

| | | | | | |
|---|---|------|-------|-------------|--------------|
| + | ! | DORC | 0.15% | N022278 001 | Feb 20, 2009 |
| + | ! | DORC | 0.06% | N021670 001 | Dec 16, 2004 |

ULIPRISTAL ACETATE

TABLET; ORAL

ELLA

| | | | | | | |
|-----------|---|---|----------------|-------------|--------------------|--------------|
| AB | + | ! | LAB HRA PHARMA | 30MG | N022474 001 | Aug 13, 2010 |
|-----------|---|---|----------------|-------------|--------------------|--------------|

LOGILIA

| | | | | | | |
|-----------|--|--|-----------------|-------------|--------------------|--------------|
| AB | | | TEVA PHARMS USA | 30MG | A207952 001 | Feb 13, 2017 |
|-----------|--|--|-----------------|-------------|--------------------|--------------|

UMECLIDINIUM BROMIDE

POWDER; INHALATION

INCRUSE ELLIPTA

| | | | | | |
|---|---|-------------------|---------------------|-------------|--------------|
| + | ! | GLAXO GRP ENGLAND | EQ 62.5MCG BASE/INH | N205382 001 | Apr 30, 2014 |
|---|---|-------------------|---------------------|-------------|--------------|

UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE

POWDER; INHALATION

ANORO ELLIPTA

| | | | | | |
|---|---|-----------------|-------------------------------------------|-------------|--------------|
| + | ! | GLAXOSMITHKLINE | EQ 0.0625MG BASE/INH; EQ 0.025MG BASE/INH | N203975 001 | Dec 18, 2013 |
|---|---|-----------------|-------------------------------------------|-------------|--------------|

UREA, C-14

CAPSULE; ORAL

PYTEST

| | | | | | |
|---|---|-------|------|-------------|--------------|
| + | ! | AVENT | 1uCi | N020617 001 | May 09, 1997 |
|---|---|-------|------|-------------|--------------|

PYTEST KIT

| | | | | | |
|---|---|-------|------|-------------|--------------|
| + | ! | AVENT | 1uCi | N020617 002 | May 09, 1997 |
|---|---|-------|------|-------------|--------------|

URIDINE TRIACETATE

GRANULE; ORAL

VISTOGARD

| | | | | | |
|---|---|-----------------|-------------|-------------|--------------|
| + | ! | WELLSTAT THERAP | 10GM/PACKET | N208159 001 | Dec 11, 2015 |
|---|---|-----------------|-------------|-------------|--------------|

XURIDEN

| | | | | | |
|---|---|-----------------|------------|-------------|--------------|
| + | ! | WELLSTAT THERAP | 2GM/PACKET | N208169 001 | Sep 04, 2015 |
|---|---|-----------------|------------|-------------|--------------|

URSODIOL

CAPSULE; ORAL

ACTIGALL

| | | | | | | |
|-----------|---|---|--------------------|--------------|--------------------|--------------|
| AB | + | ! | ALLERGAN SALES LLC | 300MG | N019594 002 | Dec 31, 1987 |
|-----------|---|---|--------------------|--------------|--------------------|--------------|

URSODIOL

| | | | | | | |
|-----------|--|--|------------------|--------------|--------------------|--------------|
| AB | | | ABHAI LLC | 300MG | A210707 001 | May 17, 2018 |
| AB | | | AMNEAL PHARMS CO | 300MG | A211301 001 | Oct 16, 2018 |
| AB | | | EPIC PHARMA | 300MG | A075517 001 | Mar 14, 2000 |
| AB | | | LANNETT CO INC | 300MG | A079082 001 | Dec 15, 2008 |
| AB | | | MYLAN | 300MG | A090530 001 | Feb 17, 2010 |
| AB | | | TEVA PHARMS | 300MG | A075592 001 | May 25, 2000 |

TABLET; ORAL

URSO 250

| | | | | | | |
|-----------|---|--|--------------------|--------------|--------------------|--------------|
| AB | + | | ALLERGAN SALES LLC | 250MG | N020675 001 | Dec 10, 1997 |
|-----------|---|--|--------------------|--------------|--------------------|--------------|

URSO FORTE

| | | | | | | |
|-----------|---|---|--------------------|--------------|--------------------|--------------|
| AB | + | ! | ALLERGAN SALES LLC | 500MG | N020675 002 | Jul 21, 2004 |
|-----------|---|---|--------------------|--------------|--------------------|--------------|

URSODIOL

| | | | | | | |
|-----------|--|--|-------------------|--------------|--------------------|--------------|
| AB | | | GLENMARK GENERICS | 250MG | A090801 001 | Jul 12, 2011 |
| AB | | | | 500MG | A090801 002 | Jul 12, 2011 |
| AB | | | IMPAX LABS INC | 250MG | A200826 001 | Dec 23, 2011 |
| AB | | | | 500MG | A200826 002 | Dec 23, 2011 |
| AB | | | PAR PHARM | 250MG | A202540 001 | Feb 14, 2013 |
| AB | | | | 500MG | A202540 002 | Feb 14, 2013 |
| AB | | | ZYDUS WORLDWIDE | 250MG | A211145 001 | Oct 30, 2018 |
| AB | | | | 500MG | A211145 002 | Oct 30, 2018 |

PRESCRIPTION DRUG PRODUCT LIST

VALACYCLOVIR HYDROCHLORIDE

TABLET; ORAL

VALACYCLOVIR HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | APOTEX INC | <u>EQ 500MG BASE</u> | <u>A090500 001</u> | Apr 04, 2014 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A090500 002</u> | Apr 04, 2014 |
| <u>AB</u> | AUROBINDO PHARMA | <u>EQ 500MG BASE</u> | <u>A090682 001</u> | May 24, 2010 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A090682 002</u> | May 24, 2010 |
| <u>AB</u> | CIPLA | <u>EQ 500MG BASE</u> | <u>A077135 001</u> | May 24, 2010 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A077135 002</u> | May 24, 2010 |
| <u>AB</u> | HETERO LABS LTD V | <u>EQ 500MG BASE</u> | <u>A203047 001</u> | Apr 08, 2015 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A203047 002</u> | Apr 08, 2015 |
| <u>AB</u> | JUBILANT GENERICS | <u>EQ 500MG BASE</u> | <u>A201506 001</u> | Apr 03, 2012 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A201506 002</u> | Apr 03, 2012 |
| <u>AB</u> | MYLAN PHARMS INC | <u>EQ 500MG BASE</u> | <u>A078518 001</u> | May 24, 2010 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A078518 002</u> | May 24, 2010 |
| <u>AB</u> | SANDOZ | <u>EQ 500MG BASE</u> | <u>A077478 001</u> | May 24, 2010 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A077478 002</u> | May 24, 2010 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>EQ 500MG BASE</u> | <u>A076588 001</u> | Jan 31, 2007 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A076588 002</u> | Jan 31, 2007 |
| <u>AB</u> | TEVA PHARMS | <u>EQ 500MG BASE</u> | <u>A077655 001</u> | May 24, 2010 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A077655 002</u> | May 24, 2010 |
| <u>AB</u> | TIME-CAP LABS INC | <u>EQ 500MG BASE</u> | <u>A079012 001</u> | May 24, 2010 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A079012 002</u> | May 24, 2010 |
| <u>AB</u> | WATSON LABS INC | <u>EQ 500MG BASE</u> | <u>A090370 001</u> | Mar 16, 2011 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A090370 002</u> | Mar 16, 2011 |
| <u>AB</u> | WEST-WARD PHARMS INT | <u>EQ 500MG BASE</u> | <u>A078656 001</u> | May 24, 2010 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A078656 002</u> | May 24, 2010 |
| <u>AB</u> | WOCKHARDT | <u>EQ 500MG BASE</u> | <u>A090216 001</u> | May 24, 2010 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A090216 002</u> | May 24, 2010 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 500MG BASE</u> | <u>A079137 001</u> | Dec 29, 2017 |
| <u>AB</u> | | <u>EQ 1GM BASE</u> | <u>A079137 002</u> | Dec 29, 2017 |

VALTREX

| | | | | |
|-----------|-------------------|----------------------|--------------------|--------------|
| <u>AB</u> | + GLAXOSMITHKLINE | <u>EQ 500MG BASE</u> | <u>N020487 001</u> | Jun 23, 1995 |
| <u>AB</u> | +! | <u>EQ 1GM BASE</u> | <u>N020487 002</u> | Jun 23, 1995 |

VALBENZAZINE TOSYLATE

CAPSULE; ORAL

INGREZZA

+ NEUROCRINE

+!

EQ 40MG BASE

EQ 80MG BASE

N209241 001 Apr 11, 2017

N209241 002 Oct 04, 2017

VALGANCICLOVIR HYDROCHLORIDE

FOR SOLUTION; ORAL

VALCYTE

| | | | | |
|-----------|----------------------|----------------|--------------------|--------------|
| <u>AB</u> | +! HOFFMANN LA ROCHE | <u>50MG/ML</u> | <u>N022257 001</u> | Aug 28, 2009 |
|-----------|----------------------|----------------|--------------------|--------------|

VALGANCICLOVIR HYDROCHLORIDE

| | | | | |
|-----------|---------------------|----------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS LABS FL INC | <u>50MG/ML</u> | <u>A205220 001</u> | Jul 18, 2016 |
|-----------|---------------------|----------------|--------------------|--------------|

TABLET; ORAL

VALCYTE

| | | | | |
|-----------|----------------------|----------------------|--------------------|--------------|
| <u>AB</u> | +! HOFFMANN LA ROCHE | <u>EQ 450MG BASE</u> | <u>N021304 001</u> | Mar 29, 2001 |
|-----------|----------------------|----------------------|--------------------|--------------|

VALGANCICLOVIR HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|----------------------|--------------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 450MG BASE</u> | <u>A204750 001</u> | Mar 31, 2016 |
| <u>AB</u> | CIPLA | <u>EQ 450MG BASE</u> | <u>A209672 001</u> | Nov 09, 2018 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 450MG BASE</u> | <u>A203511 001</u> | Nov 04, 2014 |
| <u>AB</u> | | <u>EQ 450MG BASE</u> | <u>A206876 001</u> | Dec 12, 2017 |
| <u>AB</u> | ENDO PHARMS INC | <u>EQ 450MG BASE</u> | <u>A200790 001</u> | Nov 04, 2014 |
| <u>AB</u> | HETERO LABS LTD V | <u>EQ 450MG BASE</u> | <u>A205166 001</u> | Mar 18, 2016 |

VALPROATE SODIUM

INJECTABLE; INJECTION

DEPACON

| | | | | |
|-----------|-----------|-------------------------|--------------------|--------------|
| <u>AP</u> | +! ABBVIE | <u>EQ 100MG BASE/ML</u> | <u>N020593 001</u> | Dec 30, 1996 |
|-----------|-----------|-------------------------|--------------------|--------------|

VALPROATE SODIUM

| | | | | |
|-----------|--------------------|-------------------------|--------------------|--------------|
| <u>AP</u> | ATHENEX INC | <u>EQ 100MG BASE/ML</u> | <u>A076295 001</u> | Nov 14, 2002 |
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 100MG BASE/ML</u> | <u>A076539 001</u> | Jun 26, 2003 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>EQ 100MG BASE/ML</u> | <u>A078523 001</u> | Feb 17, 2010 |

PRESCRIPTION DRUG PRODUCT LIST

VALPROIC ACID

CAPSULE; ORAL

DEPAKENE

| | | | | | |
|-----------|------------------|--------------|----------------|------------|--|
| <u>AB</u> | <u>+!</u> ABBVIE | <u>250MG</u> | <u>N018081</u> | <u>001</u> | |
|-----------|------------------|--------------|----------------|------------|--|

VALPROIC ACID

| | | | | | |
|-----------|--------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | BIONPHARMA INC | <u>250MG</u> | <u>A073484</u> | <u>001</u> | Jun 29, 1993 |
| <u>AB</u> | CATALENT | <u>250MG</u> | <u>A073229</u> | <u>001</u> | Oct 29, 1991 |
| <u>AB</u> | SUN PHARM INDS LTD | <u>250MG</u> | <u>A091037</u> | <u>001</u> | Feb 22, 2013 |

SYRUP; ORAL

DEPAKENE

| | | | | | |
|-----------|------------------|------------------|----------------|------------|--|
| <u>AA</u> | <u>+!</u> ABBVIE | <u>250MG/5ML</u> | <u>N018082</u> | <u>001</u> | |
|-----------|------------------|------------------|----------------|------------|--|

VALPROIC ACID

| | | | | | |
|-----------|------------------|------------------|----------------|------------|--------------|
| <u>AA</u> | ANI PHARMS INC | <u>250MG/5ML</u> | <u>A073178</u> | <u>001</u> | Aug 25, 1992 |
| <u>AA</u> | ECI PHARMS LLC | <u>250MG/5ML</u> | <u>A090517</u> | <u>001</u> | May 28, 2010 |
| <u>AA</u> | HIGH TECH PHARMA | <u>250MG/5ML</u> | <u>A074060</u> | <u>001</u> | Jan 13, 1995 |
| <u>AA</u> | LANNETT CO INC | <u>250MG/5ML</u> | <u>A077960</u> | <u>001</u> | Oct 13, 2006 |
| <u>AA</u> | PHARM ASSOC | <u>250MG/5ML</u> | <u>A075379</u> | <u>001</u> | Dec 15, 2000 |
| <u>AA</u> | VISTAPHARM | <u>250MG/5ML</u> | <u>A075782</u> | <u>001</u> | Dec 22, 2000 |
| <u>AA</u> | WOCKHARDT BIO AG | <u>250MG/5ML</u> | <u>A070868</u> | <u>001</u> | Jul 01, 1986 |

VALRUBICIN

SOLUTION; INTRAVESICAL

VALSTAR PRESERVATIVE FREE

| | | | | | |
|-----------|------------|----------------|----------------|------------|--------------|
| <u>+!</u> | ENDO PHARM | <u>40MG/ML</u> | <u>N020892</u> | <u>001</u> | Sep 25, 1998 |
|-----------|------------|----------------|----------------|------------|--------------|

VALSARTAN

TABLET; ORAL

DIOVAN

| | | | | | |
|-----------|-------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | <u>+</u> NOVARTIS | <u>40MG</u> | <u>N021283</u> | <u>004</u> | Aug 14, 2002 |
| <u>AB</u> | <u>+</u> | <u>80MG</u> | <u>N021283</u> | <u>001</u> | Jul 18, 2001 |
| <u>AB</u> | <u>+</u> | <u>160MG</u> | <u>N021283</u> | <u>002</u> | Jul 18, 2001 |
| <u>AB</u> | <u>+!</u> | <u>320MG</u> | <u>N021283</u> | <u>003</u> | Jul 18, 2001 |

VALSARTAN

| | | | | | |
|-----------|----------------------|--------------|----------------|------------|--------------|
| <u>AB</u> | ALEMBIC PHARMS LTD | <u>40MG</u> | <u>A091367</u> | <u>001</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>80MG</u> | <u>A091367</u> | <u>002</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>160MG</u> | <u>A091367</u> | <u>003</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>320MG</u> | <u>A091367</u> | <u>004</u> | Jan 05, 2015 |
| <u>AB</u> | AMNEAL PHARMS | <u>40MG</u> | <u>A204011</u> | <u>001</u> | Jan 11, 2016 |
| <u>AB</u> | | <u>80MG</u> | <u>A204011</u> | <u>002</u> | Jan 11, 2016 |
| <u>AB</u> | | <u>160MG</u> | <u>A204011</u> | <u>003</u> | Jan 11, 2016 |
| <u>AB</u> | | <u>320MG</u> | <u>A204011</u> | <u>004</u> | Jan 11, 2016 |
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>40MG</u> | <u>A202223</u> | <u>001</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>80MG</u> | <u>A202223</u> | <u>002</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>160MG</u> | <u>A202223</u> | <u>003</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>320MG</u> | <u>A202223</u> | <u>004</u> | Jan 05, 2015 |
| <u>AB</u> | HETERO LABS LTD V | <u>40MG</u> | <u>A203311</u> | <u>001</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>80MG</u> | <u>A203311</u> | <u>002</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>160MG</u> | <u>A203311</u> | <u>003</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>320MG</u> | <u>A203311</u> | <u>004</u> | Jan 05, 2015 |
| <u>AB</u> | IVAX PHARMS | <u>40MG</u> | <u>A077530</u> | <u>001</u> | Jan 04, 2016 |
| <u>AB</u> | | <u>80MG</u> | <u>A077530</u> | <u>002</u> | Jan 04, 2016 |
| <u>AB</u> | | <u>160MG</u> | <u>A077530</u> | <u>003</u> | Jan 04, 2016 |
| <u>AB</u> | | <u>320MG</u> | <u>A077530</u> | <u>004</u> | Jan 04, 2016 |
| <u>AB</u> | JUBILANT GENERICS | <u>40MG</u> | <u>A203536</u> | <u>001</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>80MG</u> | <u>A203536</u> | <u>002</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>160MG</u> | <u>A203536</u> | <u>003</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>320MG</u> | <u>A203536</u> | <u>004</u> | Jan 05, 2015 |
| <u>AB</u> | LUPIN LTD | <u>40MG</u> | <u>A201677</u> | <u>001</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>80MG</u> | <u>A201677</u> | <u>002</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>160MG</u> | <u>A201677</u> | <u>003</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>320MG</u> | <u>A201677</u> | <u>004</u> | Jan 05, 2015 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>40MG</u> | <u>A202696</u> | <u>001</u> | Sep 16, 2016 |
| <u>AB</u> | | <u>80MG</u> | <u>A202696</u> | <u>002</u> | Sep 16, 2016 |
| <u>AB</u> | | <u>160MG</u> | <u>A202696</u> | <u>003</u> | Sep 16, 2016 |
| <u>AB</u> | | <u>320MG</u> | <u>A202696</u> | <u>004</u> | Sep 16, 2016 |
| <u>AB</u> | MYLAN PHARMS INC | <u>40MG</u> | <u>A090866</u> | <u>001</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>80MG</u> | <u>A090866</u> | <u>002</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>160MG</u> | <u>A090866</u> | <u>003</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>320MG</u> | <u>A090866</u> | <u>004</u> | Jan 05, 2015 |
| <u>AB</u> | OHM LABS INC | <u>40MG</u> | <u>A077492</u> | <u>001</u> | Jun 26, 2014 |
| <u>AB</u> | | <u>80MG</u> | <u>A077492</u> | <u>002</u> | Jun 26, 2014 |
| <u>AB</u> | | <u>160MG</u> | <u>A077492</u> | <u>003</u> | Jun 26, 2014 |
| <u>AB</u> | | <u>320MG</u> | <u>A077492</u> | <u>004</u> | Jun 26, 2014 |

PRESCRIPTION DRUG PRODUCT LIST

VALSARTAN

TABLET; ORAL

VALSARTAN

| | | | | |
|-----------|--------------------|--------------|--------------------|--------------|
| <u>AB</u> | PRINSTON INC | <u>40MG</u> | <u>A204821 001</u> | Jun 09, 2015 |
| <u>AB</u> | | <u>80MG</u> | <u>A204821 002</u> | Jun 09, 2015 |
| <u>AB</u> | | <u>160MG</u> | <u>A204821 003</u> | Jun 09, 2015 |
| <u>AB</u> | | <u>320MG</u> | <u>A204821 004</u> | Jun 09, 2015 |
| <u>AB</u> | SQUARE PHARMS LTD | <u>40MG</u> | <u>A205347 001</u> | Apr 09, 2018 |
| <u>AB</u> | | <u>80MG</u> | <u>A205347 002</u> | Apr 09, 2018 |
| <u>AB</u> | | <u>160MG</u> | <u>A205347 003</u> | Apr 09, 2018 |
| <u>AB</u> | | <u>320MG</u> | <u>A205347 004</u> | Apr 09, 2018 |
| <u>AB</u> | TORRENT PHARMS LTD | <u>40MG</u> | <u>A202728 001</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>80MG</u> | <u>A202728 002</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>160MG</u> | <u>A202728 003</u> | Jan 05, 2015 |
| <u>AB</u> | | <u>320MG</u> | <u>A202728 004</u> | Jan 05, 2015 |
| <u>AB</u> | UNICHEM LABS LTD | <u>40MG</u> | <u>A209261 001</u> | May 04, 2018 |
| <u>AB</u> | | <u>80MG</u> | <u>A209261 002</u> | May 04, 2018 |
| <u>AB</u> | | <u>160MG</u> | <u>A209261 003</u> | May 04, 2018 |
| <u>AB</u> | | <u>320MG</u> | <u>A209261 004</u> | May 04, 2018 |

VANCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

VANCOCIN HYDROCHLORIDE

| | | | | |
|---------------------------------|------------------|----------------------|--------------------|--------------|
| <u>AB</u> | + ANI PHARMS INC | <u>EQ 125MG BASE</u> | <u>N050606 001</u> | Apr 15, 1986 |
| <u>AB</u> | +! | <u>EQ 250MG BASE</u> | <u>N050606 002</u> | Apr 15, 1986 |
| <u>VANCOMYCIN HYDROCHLORIDE</u> | | | | |
| <u>AB</u> | AKORN | <u>EQ 125MG BASE</u> | <u>A065478 001</u> | Apr 09, 2012 |
| <u>AB</u> | | <u>EQ 250MG BASE</u> | <u>A065478 002</u> | Apr 09, 2012 |
| <u>AB</u> | LUPIN LTD | <u>EQ 125MG BASE</u> | <u>A090439 001</u> | Jan 28, 2015 |
| <u>AB</u> | | <u>EQ 250MG BASE</u> | <u>A090439 002</u> | Jan 28, 2015 |
| <u>AB</u> | STRIDES PHARMA | <u>EQ 125MG BASE</u> | <u>A065490 001</u> | Apr 09, 2012 |
| <u>AB</u> | | <u>EQ 250MG BASE</u> | <u>A065490 002</u> | Apr 09, 2012 |
| <u>AB</u> | WATSON LABS | <u>EQ 125MG BASE</u> | <u>A065510 001</u> | Apr 09, 2012 |
| <u>AB</u> | | <u>EQ 250MG BASE</u> | <u>A065510 002</u> | Apr 09, 2012 |

FOR SOLUTION; ORAL

FIRVANQ KIT

| | | | | |
|----|-------------------|-----------------|-------------|--------------|
| +! | RXMTM THERAPS LLC | EQ 25MG BASE/ML | N208910 001 | Jan 26, 2018 |
| +! | | EQ 50MG BASE/ML | N208910 002 | Jan 26, 2018 |

INJECTABLE; INJECTION

VANCOMYCIN HYDROCHLORIDE

| | | | | |
|-----------|----------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>EQ 500MG BASE/VIAL</u> | <u>A205780 001</u> | Mar 31, 2016 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A205780 002</u> | Mar 31, 2016 |
| <u>AP</u> | | <u>EQ 5GM BASE/VIAL</u> | <u>A205779 001</u> | Mar 29, 2016 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A205779 002</u> | Mar 29, 2016 |
| <u>AP</u> | EMCURE PHARMS LTD | <u>EQ 500MG BASE/VIAL</u> | <u>A202275 001</u> | Oct 31, 2013 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A202275 002</u> | Oct 31, 2013 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A202464 001</u> | Oct 09, 2013 |
| <u>AP</u> | | <u>EQ 5GM BASE/VIAL</u> | <u>A202274 001</u> | Oct 31, 2013 |
| <u>AP</u> | ! FRESENIUS KABI USA | <u>EQ 500MG BASE/VIAL</u> | <u>A062663 001</u> | Mar 17, 1987 |
| <u>AP</u> | | <u>EQ 750MG BASE/VIAL</u> | <u>A062663 005</u> | Aug 17, 2016 |
| <u>AP</u> | ! | <u>EQ 1GM BASE/VIAL</u> | <u>A062663 002</u> | Jul 31, 1987 |
| <u>AP</u> | ! | <u>EQ 5GM BASE/VIAL</u> | <u>A062663 003</u> | Jun 03, 1988 |
| <u>AP</u> | ! | <u>EQ 10GM BASE/VIAL</u> | <u>A062663 004</u> | Nov 28, 1997 |
| <u>AP</u> | GLAND PHARMA LTD | <u>EQ 500MG BASE/VIAL</u> | <u>A205694 001</u> | Jan 21, 2016 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A205694 002</u> | Jan 21, 2016 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>EQ 5GM BASE/VIAL</u> | <u>A204360 001</u> | Oct 15, 2018 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A204360 002</u> | Oct 15, 2018 |
| <u>AP</u> | HIKMA PHARMS | <u>EQ 750MG BASE/VIAL</u> | <u>A206616 001</u> | Oct 03, 2018 |
| <u>AP</u> | ! HOSPIRA | <u>EQ 500MG BASE/VIAL</u> | <u>A062911 001</u> | Aug 04, 1988 |
| <u>AP</u> | ! | <u>EQ 500MG BASE/VIAL</u> | <u>A062931 001</u> | Oct 29, 1992 |
| <u>AP</u> | ! | <u>EQ 750MG BASE/VIAL</u> | <u>A062912 002</u> | Jan 07, 2009 |
| <u>AP</u> | ! | <u>EQ 750MG BASE/VIAL</u> | <u>A062933 002</u> | May 27, 2009 |
| <u>AP</u> | ! | <u>EQ 1GM BASE/VIAL</u> | <u>A062912 001</u> | Aug 04, 1988 |
| <u>AP</u> | ! | <u>EQ 1GM BASE/VIAL</u> | <u>A062933 001</u> | Oct 29, 1992 |
| <u>AP</u> | ! | <u>EQ 5GM BASE/VIAL</u> | <u>A063076 001</u> | Dec 21, 1990 |
| <u>AP</u> | HOSPIRA INC | <u>EQ 10GM BASE/VIAL</u> | <u>A065455 001</u> | Apr 29, 2009 |
| <u>AP</u> | MUSTAFA NEVZAT ILAC | <u>EQ 500MG BASE/VIAL</u> | <u>A065401 001</u> | Jun 30, 2008 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A065401 002</u> | Jun 30, 2008 |
| <u>AP</u> | MYLAN LABS LTD | <u>EQ 500MG BASE/VIAL</u> | <u>A065397 001</u> | Dec 30, 2008 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A065397 002</u> | Dec 30, 2008 |
| <u>AP</u> | | <u>EQ 5GM BASE/VIAL</u> | <u>A065432 001</u> | Dec 30, 2008 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A091554 001</u> | Sep 19, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

VANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

VANCOMYCIN HYDROCHLORIDE

| | | | | |
|-----------|-------------------|---------------------------|--------------------|--------------|
| <u>AP</u> | SAGENT PHARMS | <u>EQ 5GM BASE/VIAL</u> | <u>A200837 001</u> | Aug 10, 2012 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A200837 002</u> | Sep 02, 2014 |
| <u>AP</u> | SANDOZ | <u>EQ 500MG BASE/VIAL</u> | <u>A090250 001</u> | Apr 27, 2010 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A090250 002</u> | Apr 27, 2010 |
| <u>AP</u> | SANDOZ INC | <u>EQ 5GM BASE/VIAL</u> | <u>A201048 001</u> | Aug 10, 2012 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A201048 002</u> | Aug 10, 2012 |
| <u>AP</u> | XELLIA PHARMS APS | <u>EQ 5GM BASE/VIAL</u> | <u>A204125 001</u> | Dec 28, 2015 |
| <u>AP</u> | | <u>EQ 10GM BASE/VIAL</u> | <u>A204125 002</u> | Dec 28, 2015 |
| <u>AP</u> | | <u>EQ 500MG BASE/VIAL</u> | <u>A204107 001</u> | Dec 28, 2015 |
| <u>AP</u> | | <u>EQ 1GM BASE/VIAL</u> | <u>A204107 002</u> | Dec 28, 2015 |

VANOCIN HYDROCHLORIDE IN PLASTIC CONTAINER

| | | | | | |
|---|---|-----------------|---------------------|-------------|--------------|
| + | + | BAXTER HLTHCARE | EQ 500MG BASE/100ML | N050671 001 | Apr 29, 1993 |
| + | + | | EQ 750MG BASE/150ML | N050671 002 | Dec 20, 2010 |
| + | + | | EQ 1GM BASE/200ML | N050671 003 | Mar 01, 1999 |

POWDER; INTRAVENOUS

VANCOMYCIN HYDROCHLORIDE

| | | | | | |
|---|---|----------------|---------------------|-------------|--------------|
| + | + | MYLAN LABS LTD | EQ 250MG BASE/VIAL | N209481 001 | Jul 10, 2018 |
| + | + | | EQ 750MG BASE/VIAL | N209481 002 | Jul 10, 2018 |
| + | + | | EQ 1.25GM BASE/VIAL | N209481 003 | Jul 10, 2018 |
| + | + | | EQ 1.5GM BASE/VIAL | N209481 004 | Jul 10, 2018 |

VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER

| | | | | | |
|--|--|--------------|---------------|-------------|--------------|
| | | SAMSON MEDCL | EQ 100GM BASE | A091532 001 | Jan 06, 2016 |
|--|--|--------------|---------------|-------------|--------------|

VANDETANIB

TABLET; ORAL

CAPRELSA

| | | | | | |
|---|---|--------------|-------|-------------|--------------|
| + | + | GENZYME CORP | 100MG | N022405 001 | Apr 06, 2011 |
| + | + | | 300MG | N022405 002 | Apr 06, 2011 |

VARDENAFIL HYDROCHLORIDE

TABLET; ORAL

LEVITRA

| | | | | | |
|-----------|---|----------------|-------------|--------------------|--------------|
| <u>AB</u> | + | BAYER HLTHCARE | <u>5MG</u> | <u>N021400 001</u> | Aug 19, 2003 |
| <u>AB</u> | + | | <u>10MG</u> | <u>N021400 002</u> | Aug 19, 2003 |
| <u>AB</u> | + | | <u>20MG</u> | <u>N021400 004</u> | Aug 19, 2003 |

VARDENAFIL HYDROCHLORIDE

| | | | | | |
|-----------|--|----------------------|--------------|--------------------|--------------|
| <u>AB</u> | | AMNEAL PHARMS CO | <u>5MG</u> | <u>A210738 001</u> | Oct 31, 2018 |
| <u>AB</u> | | | <u>10MG</u> | <u>A210738 002</u> | Oct 31, 2018 |
| <u>AB</u> | | | <u>20MG</u> | <u>A210738 003</u> | Oct 31, 2018 |
| <u>AB</u> | | CROSSMEDIKA SA | <u>2.5MG</u> | <u>A209057 001</u> | Oct 31, 2018 |
| <u>AB</u> | | | <u>5MG</u> | <u>A209057 002</u> | Oct 31, 2018 |
| <u>AB</u> | | | <u>10MG</u> | <u>A209057 003</u> | Oct 31, 2018 |
| <u>AB</u> | | | <u>20MG</u> | <u>A209057 004</u> | Oct 31, 2018 |
| <u>AB</u> | | TEVA PHARMS | <u>2.5MG</u> | <u>A091347 001</u> | May 03, 2012 |
| <u>AB</u> | | | <u>5MG</u> | <u>A091347 002</u> | May 03, 2012 |
| <u>AB</u> | | | <u>10MG</u> | <u>A091347 003</u> | May 03, 2012 |
| <u>AB</u> | | | <u>20MG</u> | <u>A091347 004</u> | May 03, 2012 |
| <u>AB</u> | | ZYDUS PHARMS USA INC | <u>2.5MG</u> | <u>A208960 001</u> | Oct 31, 2018 |
| <u>AB</u> | | | <u>5MG</u> | <u>A208960 002</u> | Oct 31, 2018 |
| <u>AB</u> | | | <u>10MG</u> | <u>A208960 003</u> | Oct 31, 2018 |
| <u>AB</u> | | | <u>20MG</u> | <u>A208960 004</u> | Oct 31, 2018 |

TABLET, ORALLY DISINTEGRATING; ORAL

STAXYN

| | | | | | |
|-----------|---|----------------|-------------|--------------------|--------------|
| <u>AB</u> | + | BAYER HLTHCARE | <u>10MG</u> | <u>N200179 001</u> | Jun 17, 2010 |
|-----------|---|----------------|-------------|--------------------|--------------|

VARDENAFIL HYDROCHLORIDE

| | | | | | |
|-----------|--|--------------------|-------------|--------------------|--------------|
| <u>AB</u> | | ALEMBIC PHARMS LTD | <u>10MG</u> | <u>A208324 001</u> | Nov 16, 2018 |
|-----------|--|--------------------|-------------|--------------------|--------------|

VARENICLINE TARTRATE

TABLET; ORAL

CHANTIX

| | | | | | |
|---|---|-------------|---------------|-------------|--------------|
| + | + | PF PRISM CV | EQ 0.5MG BASE | N021928 001 | May 10, 2006 |
| + | + | | EQ 1MG BASE | N021928 002 | May 10, 2006 |

VASOPRESSIN

SOLUTION; IV (INFUSION)

VASOSTRICT

| | | | | | |
|---|---|----------------------|----------------------------|-------------|--------------|
| + | + | PAR STERILE PRODUCTS | 20UNITS/ML (20UNITS/ML) | N204485 001 | Apr 17, 2014 |
| + | + | | 200UNITS/10ML (20UNITS/ML) | N204485 002 | Dec 17, 2016 |

PRESCRIPTION DRUG PRODUCT LIST

VECURONIUM BROMIDE

INJECTABLE; INJECTION

VECURONIUM BROMIDE

| | | | | |
|-----------|----------------------|------------------|--------------------|--------------|
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>10MG/VIAL</u> | <u>A206670 001</u> | Dec 20, 2018 |
| <u>AP</u> | | <u>20MG/VIAL</u> | <u>A206670 002</u> | Dec 20, 2018 |
| <u>AP</u> | GLAND PHARMA LTD | <u>10MG/VIAL</u> | <u>A205390 001</u> | May 26, 2016 |
| <u>AP</u> | | <u>20MG/VIAL</u> | <u>A205390 002</u> | May 26, 2016 |
| <u>AP</u> | HOSPIRA | <u>10MG/VIAL</u> | <u>A075164 001</u> | Oct 21, 1999 |
| <u>AP</u> | | <u>20MG/VIAL</u> | <u>A075164 002</u> | Oct 21, 1999 |
| <u>AP</u> | MYLAN LABS LTD | <u>10MG/VIAL</u> | <u>A090243 001</u> | May 11, 2010 |
| <u>AP</u> | | <u>20MG/VIAL</u> | <u>A090243 002</u> | May 11, 2010 |
| <u>AP</u> | SAGENT PHARMS | <u>10MG/VIAL</u> | <u>A078274 001</u> | Dec 29, 2008 |
| <u>AP</u> | | <u>20MG/VIAL</u> | <u>A078274 002</u> | Dec 29, 2008 |
| <u>AP</u> | ! SUN PHARMA GLOBAL | <u>10MG/VIAL</u> | <u>A079001 001</u> | Jun 17, 2009 |
| <u>AP</u> | ! | <u>20MG/VIAL</u> | <u>A079001 002</u> | Jun 17, 2009 |
| <u>AP</u> | TEVA PHARMS USA | <u>10MG/VIAL</u> | <u>A074688 001</u> | Aug 25, 1999 |
| <u>AP</u> | | <u>20MG/VIAL</u> | <u>A074688 002</u> | Aug 25, 1999 |
| <u>AP</u> | WEST-WARD PHARMS INT | <u>10MG/VIAL</u> | <u>A075549 001</u> | Jun 13, 2000 |
| <u>AP</u> | | <u>20MG/VIAL</u> | <u>A075549 002</u> | Jun 13, 2000 |

VELAGLUCERASE ALFA

INJECTABLE; INTRAVENOUS

VPRIV

SHIRE HUMAN GENETIC 400 UNITS/VIAL N022575 001 Feb 26, 2010

VEMURAFENIB

TABLET; ORAL

ZELBORAF

+! HOFFMANN LA ROCHE 240MG N202429 001 Aug 17, 2011

VENETOCLAX

TABLET; ORAL

VENCLEXTA

+ ABBVIE INC 10MG N208573 001 Apr 11, 2016

+ 50MG N208573 002 Apr 11, 2016

+! 100MG N208573 003 Apr 11, 2016

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

EFFEXOR XR

| | | | | |
|-----------|----------------|-----------------------|--------------------|--------------|
| <u>AB</u> | + WYETH PHARMS | <u>EQ 37.5MG BASE</u> | <u>N020699 001</u> | Oct 20, 1997 |
| <u>AB</u> | + | <u>EQ 75MG BASE</u> | <u>N020699 002</u> | Oct 20, 1997 |
| <u>AB</u> | +! | <u>EQ 150MG BASE</u> | <u>N020699 004</u> | Oct 20, 1997 |

VENLAFAXINE HYDROCHLORIDE

| | | | | |
|-----------|-----------------------|-----------------------|--------------------|--------------|
| <u>AB</u> | AUROBINDO PHARMA LTD | <u>EQ 37.5MG BASE</u> | <u>A200834 001</u> | Apr 14, 2011 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A200834 002</u> | Apr 14, 2011 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A200834 003</u> | Apr 14, 2011 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 37.5MG BASE</u> | <u>A078421 001</u> | May 06, 2011 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A078421 002</u> | May 06, 2011 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A078421 003</u> | May 06, 2011 |
| <u>AB</u> | INTELLIPHARMACEUTIC S | <u>EQ 37.5MG BASE</u> | <u>A201272 001</u> | Nov 23, 2018 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A201272 002</u> | Nov 23, 2018 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A201272 003</u> | Nov 23, 2018 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>EQ 37.5MG BASE</u> | <u>A204889 001</u> | Oct 05, 2017 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A204889 002</u> | Oct 05, 2017 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A204889 003</u> | Oct 05, 2017 |
| <u>AB</u> | ORCHID HLTHCARE | <u>EQ 37.5MG BASE</u> | <u>A091123 001</u> | Jul 11, 2011 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A091123 002</u> | Jul 11, 2011 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A091123 003</u> | Jul 11, 2011 |
| <u>AB</u> | TEVA | <u>EQ 37.5MG BASE</u> | <u>A076565 001</u> | Jun 28, 2010 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A076565 002</u> | Jun 28, 2010 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A076565 003</u> | Jun 28, 2010 |
| <u>AB</u> | TORRENT PHARMS LLC | <u>EQ 37.5MG BASE</u> | <u>A090899 001</u> | Jun 01, 2011 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A090899 002</u> | Jun 01, 2011 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A090899 003</u> | Jun 01, 2011 |
| <u>AB</u> | VALEANT PHARMS NORTH | <u>EQ 37.5MG BASE</u> | <u>A090071 001</u> | Apr 15, 2011 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A090071 002</u> | Apr 15, 2011 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A090071 003</u> | Apr 15, 2011 |
| <u>AB</u> | WOCKHARDT | <u>EQ 37.5MG BASE</u> | <u>A078865 001</u> | Apr 14, 2011 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A078865 002</u> | Apr 14, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

VENLAFAXINE HYDROCHLORIDE

| | | | | | |
|-----------|-------------------------|-----------------------|----------------|------------|--------------|
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A078865</u> | <u>003</u> | Apr 14, 2011 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 37.5MG BASE</u> | <u>A090174</u> | <u>001</u> | Apr 14, 2011 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A090174</u> | <u>002</u> | Apr 14, 2011 |
| <u>AB</u> | | <u>EQ 150MG BASE</u> | <u>A090174</u> | <u>003</u> | Apr 14, 2011 |

TABLET;ORAL

VENLAFAXINE HYDROCHLORIDE

| | | | | | |
|-----------|---------------------|-----------------------|----------------|------------|--------------|
| <u>AB</u> | ALEMBIC PHARMS LTD | <u>EQ 25MG BASE</u> | <u>A078932</u> | <u>001</u> | Dec 14, 2010 |
| <u>AB</u> | | <u>EQ 37.5MG BASE</u> | <u>A078932</u> | <u>002</u> | Dec 14, 2010 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A078932</u> | <u>003</u> | Dec 14, 2010 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A078932</u> | <u>004</u> | Dec 14, 2010 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A078932</u> | <u>005</u> | Dec 14, 2010 |
| <u>AB</u> | AMNEAL PHARMS | <u>EQ 25MG BASE</u> | <u>A079098</u> | <u>001</u> | May 11, 2010 |
| <u>AB</u> | | <u>EQ 37.5MG BASE</u> | <u>A079098</u> | <u>002</u> | May 11, 2010 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A079098</u> | <u>003</u> | May 11, 2010 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A079098</u> | <u>004</u> | May 11, 2010 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A079098</u> | <u>005</u> | May 11, 2010 |
| <u>AB</u> | AUROBINDO PHARMA | <u>EQ 25MG BASE</u> | <u>A090555</u> | <u>001</u> | Apr 07, 2010 |
| <u>AB</u> | | <u>EQ 37.5MG BASE</u> | <u>A090555</u> | <u>002</u> | Apr 07, 2010 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A090555</u> | <u>003</u> | Apr 07, 2010 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A090555</u> | <u>004</u> | Apr 07, 2010 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A090555</u> | <u>005</u> | Apr 07, 2010 |
| <u>AB</u> | CADILA PHARMS LTD | <u>EQ 25MG BASE</u> | <u>A206250</u> | <u>001</u> | Nov 21, 2018 |
| <u>AB</u> | | <u>EQ 37.5MG BASE</u> | <u>A206250</u> | <u>002</u> | Nov 21, 2018 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A206250</u> | <u>003</u> | Nov 21, 2018 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A206250</u> | <u>004</u> | Nov 21, 2018 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A206250</u> | <u>005</u> | Nov 21, 2018 |
| <u>AB</u> | DR REDDYS LABS LTD | <u>EQ 25MG BASE</u> | <u>A078301</u> | <u>001</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 37.5MG BASE</u> | <u>A078301</u> | <u>002</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A078301</u> | <u>003</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A078301</u> | <u>004</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A078301</u> | <u>005</u> | Jun 13, 2008 |
| <u>AB</u> | HERITAGE PHARMS INC | <u>EQ 25MG BASE</u> | <u>A078554</u> | <u>001</u> | Jan 09, 2009 |
| <u>AB</u> | | <u>EQ 37.5MG BASE</u> | <u>A078554</u> | <u>002</u> | Jan 09, 2009 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A078554</u> | <u>003</u> | Jan 09, 2009 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A078554</u> | <u>004</u> | Jan 09, 2009 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A078554</u> | <u>005</u> | Jan 09, 2009 |
| <u>AB</u> | MYLAN | <u>EQ 25MG BASE</u> | <u>A077166</u> | <u>001</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 37.5MG BASE</u> | <u>A077166</u> | <u>002</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A077166</u> | <u>003</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A077166</u> | <u>004</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A077166</u> | <u>005</u> | Jun 13, 2008 |
| <u>AB</u> | PRINSTON INC | <u>EQ 25MG BASE</u> | <u>A090027</u> | <u>001</u> | Aug 04, 2010 |
| <u>AB</u> | | <u>EQ 37.5MG BASE</u> | <u>A090027</u> | <u>002</u> | Aug 04, 2010 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A090027</u> | <u>003</u> | Aug 04, 2010 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A090027</u> | <u>004</u> | Aug 04, 2010 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A090027</u> | <u>005</u> | Aug 04, 2010 |
| <u>AB</u> | SUN PHARM INDS INC | <u>EQ 25MG BASE</u> | <u>A078627</u> | <u>001</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 37.5MG BASE</u> | <u>A078627</u> | <u>002</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A078627</u> | <u>003</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A078627</u> | <u>004</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A078627</u> | <u>005</u> | Jun 13, 2008 |
| <u>AB</u> | TEVA | <u>EQ 25MG BASE</u> | <u>A076690</u> | <u>001</u> | Aug 03, 2006 |
| <u>AB</u> | | <u>EQ 37.5MG BASE</u> | <u>A076690</u> | <u>002</u> | Aug 03, 2006 |
| <u>AB</u> | ! | <u>EQ 50MG BASE</u> | <u>A076690</u> | <u>003</u> | Aug 03, 2006 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A076690</u> | <u>004</u> | Aug 03, 2006 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A076690</u> | <u>005</u> | Aug 03, 2006 |
| <u>AB</u> | YAOPHARMA CO LTD | <u>EQ 25MG BASE</u> | <u>A202036</u> | <u>001</u> | May 28, 2015 |
| <u>AB</u> | | <u>EQ 37.5MG BASE</u> | <u>A202036</u> | <u>002</u> | May 28, 2015 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A202036</u> | <u>003</u> | May 28, 2015 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A202036</u> | <u>004</u> | May 28, 2015 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A202036</u> | <u>005</u> | May 28, 2015 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>EQ 25MG BASE</u> | <u>A077653</u> | <u>001</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 37.5MG BASE</u> | <u>A077653</u> | <u>002</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 50MG BASE</u> | <u>A077653</u> | <u>003</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 75MG BASE</u> | <u>A077653</u> | <u>004</u> | Jun 13, 2008 |
| <u>AB</u> | | <u>EQ 100MG BASE</u> | <u>A077653</u> | <u>005</u> | Jun 13, 2008 |

PRESCRIPTION DRUG PRODUCT LIST

VENLAFAXINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

VENLAFAXINE HYDROCHLORIDE

| | | | | | |
|-----------|---|-------------------|-----------------------|--------------------|--------------|
| <u>AB</u> | | NOSTRUM LABS INC | <u>EQ 150MG BASE</u> | <u>A205468 002</u> | Mar 24, 2017 |
| <u>AB</u> | | | <u>EQ 225MG BASE</u> | <u>A205468 003</u> | Mar 24, 2017 |
| <u>AB</u> | + | OSMOTICA PHARM | <u>EQ 37.5MG BASE</u> | <u>N022104 001</u> | May 20, 2008 |
| <u>AB</u> | + | | <u>EQ 75MG BASE</u> | <u>N022104 002</u> | May 20, 2008 |
| <u>AB</u> | + | | <u>EQ 150MG BASE</u> | <u>N022104 003</u> | May 20, 2008 |
| <u>AB</u> | + | | <u>EQ 225MG BASE</u> | <u>N022104 004</u> | May 20, 2008 |
| <u>AB</u> | | SUN PHARMA GLOBAL | <u>EQ 37.5MG BASE</u> | <u>A091272 001</u> | Aug 18, 2010 |
| <u>AB</u> | | | <u>EQ 75MG BASE</u> | <u>A091272 002</u> | Aug 18, 2010 |
| <u>AB</u> | | | <u>EQ 150MG BASE</u> | <u>A091272 003</u> | Aug 18, 2010 |
| <u>AB</u> | | | <u>EQ 225MG BASE</u> | <u>A091272 004</u> | Jan 08, 2019 |

VERAPAMIL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

VERAPAMIL HYDROCHLORIDE

| | | | | | |
|-----------|--|-------|--------------|--------------------|--------------|
| <u>AB</u> | | MYLAN | <u>100MG</u> | <u>A078306 001</u> | Aug 09, 2007 |
| <u>AB</u> | | | <u>120MG</u> | <u>A075138 001</u> | Apr 20, 1999 |
| <u>AB</u> | | | <u>180MG</u> | <u>A075138 002</u> | Apr 20, 1999 |
| <u>AB</u> | | | <u>200MG</u> | <u>A078306 002</u> | Aug 09, 2007 |
| <u>AB</u> | | | <u>240MG</u> | <u>A075138 003</u> | Apr 20, 1999 |
| <u>AB</u> | | | <u>300MG</u> | <u>A078306 003</u> | Aug 09, 2007 |

VERELAN

| | | | | | |
|-----------|---|-------------------|--------------|--------------------|--------------|
| <u>AB</u> | + | RECRO GAINESVILLE | <u>120MG</u> | <u>N019614 001</u> | May 29, 1990 |
| <u>AB</u> | + | | <u>180MG</u> | <u>N019614 003</u> | Jan 09, 1992 |
| <u>AB</u> | + | | <u>240MG</u> | <u>N019614 002</u> | May 29, 1990 |

VERELAN PM

| | | | | | |
|-----------|---|-------------------|--------------|--------------------|--------------|
| <u>AB</u> | + | RECRO GAINESVILLE | <u>100MG</u> | <u>N020943 001</u> | Nov 25, 1998 |
| <u>AB</u> | + | | <u>200MG</u> | <u>N020943 002</u> | Nov 25, 1998 |
| <u>AB</u> | + | | <u>300MG</u> | <u>N020943 003</u> | Nov 25, 1998 |

VERELAN

+! RECRO GAINESVILLE

360MG

N019614 004 May 10, 1996

SOLUTION;INTRAVENOUS

VERAPAMIL HYDROCHLORIDE

| | | | | | |
|-----------|---|----------------------|----------------------------|--------------------|--------------|
| <u>AP</u> | | AMNEAL PHARMS CO | <u>5MG/2ML (2.5MG/ML)</u> | <u>A210994 001</u> | Jul 13, 2018 |
| <u>AP</u> | | | <u>10MG/4ML (2.5MG/ML)</u> | <u>A210994 002</u> | Jul 13, 2018 |
| <u>AP</u> | | EXELA PHARMA SCS LLC | <u>5MG/2ML (2.5MG/ML)</u> | <u>N018925 001</u> | Mar 30, 1984 |
| <u>AP</u> | | | <u>10MG/4ML (2.5MG/ML)</u> | <u>N018925 002</u> | Apr 05, 2018 |
| <u>AP</u> | ! | HOSPIRA | <u>5MG/2ML (2.5MG/ML)</u> | <u>A070738 001</u> | May 06, 1987 |
| <u>AP</u> | ! | | <u>5MG/2ML (2.5MG/ML)</u> | <u>A075136 001</u> | Oct 20, 1998 |
| <u>AP</u> | ! | | <u>5MG/2ML (2.5MG/ML)</u> | <u>A070737 001</u> | May 06, 1987 |
| <u>AP</u> | ! | | <u>10MG/4ML (2.5MG/ML)</u> | <u>A070737 002</u> | May 06, 1987 |
| <u>AP</u> | | MICRO LABS | <u>5MG/2ML (2.5MG/ML)</u> | <u>A211370 001</u> | Dec 28, 2018 |
| <u>AP</u> | | | <u>10MG/4ML (2.5MG/ML)</u> | <u>A211370 002</u> | Dec 28, 2018 |
| <u>AP</u> | | SOMERSET THERAPS LLC | <u>5MG/2ML (2.5MG/ML)</u> | <u>A211015 001</u> | Jun 18, 2018 |
| <u>AP</u> | | | <u>5MG/2ML (2.5MG/ML)</u> | <u>A211035 001</u> | Jun 18, 2018 |
| <u>AP</u> | | | <u>10MG/4ML (2.5MG/ML)</u> | <u>A211015 002</u> | Jun 18, 2018 |
| <u>AP</u> | | | <u>10MG/4ML (2.5MG/ML)</u> | <u>A211035 002</u> | Jun 18, 2018 |

TABLET;ORAL

CALAN

| | | | | | |
|-----------|---|---------------|--------------|--------------------|--------------|
| <u>AB</u> | + | GD SEARLE LLC | <u>80MG</u> | <u>N018817 001</u> | Sep 10, 1984 |
| <u>AB</u> | + | | <u>120MG</u> | <u>N018817 002</u> | Sep 10, 1984 |

VERAPAMIL HYDROCHLORIDE

| | | | | | |
|-----------|--|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | | HERITAGE PHARMS INC | <u>40MG</u> | <u>A071881 002</u> | Oct 14, 2015 |
| <u>AB</u> | | | <u>80MG</u> | <u>A071881 003</u> | Apr 05, 1988 |
| <u>AB</u> | | | <u>120MG</u> | <u>A071881 001</u> | Apr 05, 1988 |
| <u>AB</u> | | MYLAN | <u>80MG</u> | <u>A071483 002</u> | Feb 15, 1989 |
| <u>AB</u> | | | <u>120MG</u> | <u>A071483 001</u> | Feb 15, 1989 |
| <u>AB</u> | | WATSON LABS | <u>40MG</u> | <u>A072924 001</u> | Jun 29, 1993 |
| <u>AB</u> | | | <u>80MG</u> | <u>A070995 001</u> | Oct 01, 1986 |
| <u>AB</u> | | | <u>120MG</u> | <u>A070994 001</u> | Oct 01, 1986 |

TABLET, EXTENDED RELEASE;ORAL

CALAN SR

| | | | | | |
|-----------|---|--------|--------------|--------------------|--------------|
| <u>AB</u> | + | PFIZER | <u>120MG</u> | <u>N019152 003</u> | Mar 06, 1991 |
| <u>AB</u> | + | | <u>240MG</u> | <u>N019152 001</u> | Dec 16, 1986 |

VERAPAMIL HYDROCHLORIDE

| | | | | | |
|-----------|---|-------------------|--------------|--------------------|--------------|
| <u>AB</u> | | CADILA PHARMS LTD | <u>180MG</u> | <u>A206173 001</u> | May 05, 2017 |
| <u>AB</u> | | | <u>240MG</u> | <u>A206173 002</u> | May 05, 2017 |
| <u>AB</u> | | GLENMARK GENERICS | <u>120MG</u> | <u>A090700 001</u> | Aug 03, 2011 |
| <u>AB</u> | ! | | <u>180MG</u> | <u>A090700 002</u> | Aug 03, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

VERAPAMIL HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|--------------|--------------------|--------------|
| <u>AB</u> | | <u>240MG</u> | <u>A078906 001</u> | Sep 17, 2009 |
| <u>AB</u> | IVAX SUB TEVA PHARMS | <u>120MG</u> | <u>A073568 002</u> | Oct 10, 1997 |
| <u>AB</u> | | <u>180MG</u> | <u>A074330 001</u> | Jan 31, 1994 |
| <u>AB</u> | | <u>240MG</u> | <u>A073568 001</u> | Jul 31, 1992 |
| <u>AB</u> | MYLAN | <u>120MG</u> | <u>A074587 002</u> | Feb 21, 1997 |
| <u>AB</u> | | <u>180MG</u> | <u>A074587 003</u> | Sep 09, 1997 |
| <u>AB</u> | | <u>240MG</u> | <u>A074587 001</u> | Mar 23, 1996 |
| <u>AB</u> | PAR PHARM | <u>120MG</u> | <u>A075072 001</u> | May 25, 1999 |
| <u>AB</u> | | <u>240MG</u> | <u>A075072 003</u> | May 25, 1999 |
| <u>AB</u> | SUN PHARM INDS INC | <u>120MG</u> | <u>A090529 001</u> | Dec 30, 2011 |
| <u>AB</u> | | <u>180MG</u> | <u>A090529 002</u> | Dec 30, 2011 |
| <u>AB</u> | | <u>240MG</u> | <u>A090529 003</u> | Dec 30, 2011 |

VERTEPORFIN

INJECTABLE;INJECTION

VISUDYNE

+! VALEANT LUXEMBOURG 15MG/VIAL N021119 001 Apr 12, 2000

VIGABATRIN

FOR SOLUTION;ORAL

SABRII

AA +! LUNDBECK PHARMS LLC 500MG/PACKET N022006 001 Aug 21, 2009

VIGABATRIN

| | | | | |
|-----------|--------------------|---------------------|--------------------|--------------|
| <u>AA</u> | AMNEAL PHARMS | <u>500MG/PACKET</u> | <u>A210155 001</u> | Mar 13, 2018 |
| <u>AA</u> | DR REDDYS LABS LTD | <u>500MG/PACKET</u> | <u>A211481 001</u> | Nov 20, 2018 |
| <u>AA</u> | PAR PHARM INC | <u>500MG/PACKET</u> | <u>A208218 001</u> | Apr 27, 2017 |
| <u>AA</u> | TEVA PHARMS USA | <u>500MG/PACKET</u> | <u>A209824 001</u> | Apr 23, 2018 |

VIGADRONE

AA AUCTA PHARMS 500MG/PACKET A210196 001 Jun 21, 2018

TABLET;ORAL

SABRII

AB +! LUNDBECK PHARMS LLC 500MG N020427 001 Aug 21, 2009

VIGABATRIN

AB TEVA PHARMS USA 500MG A209822 001 Jan 14, 2019

VILAZODONE HYDROCHLORIDE

TABLET;ORAL

VIIBRYD

| | | | | |
|----|--------------------|------|-------------|--------------|
| +! | ALLERGAN SALES LLC | 10MG | N022567 001 | Jan 21, 2011 |
| + | | 20MG | N022567 002 | Jan 21, 2011 |
| + | | 40MG | N022567 003 | Jan 21, 2011 |

VINBLASTINE SULFATE

INJECTABLE;INJECTION

VINBLASTINE SULFATE

| | | | | |
|---|-------------------------|-----------|-------------|--------------|
| ! | FRESENIUS KABI USA | 1MG/ML | A089515 001 | Apr 29, 1987 |
| ! | WEST-WARD PHARMS INT | 10MG/VIAL | A089395 001 | Apr 09, 1987 |

VINCRISTINE SULFATE

INJECTABLE;INJECTION

VINCRISTINE SULFATE PFS

| | | | | | |
|-----------|---|-----------------|---------------|--------------------|--------------|
| <u>AP</u> | ! | HOSPIRA | <u>1MG/ML</u> | <u>A071484 001</u> | Apr 19, 1988 |
| <u>AP</u> | | TEVA PHARMS USA | <u>1MG/ML</u> | <u>A075493 001</u> | Sep 01, 1999 |

INJECTABLE, LIPOSOMAL;INTRAVENOUS

MARQIBO KIT

+! TALON THERAP 5MG/5ML (1MG/ML) N202497 001 Aug 09, 2012

VINORELBINE TARTRATE

INJECTABLE;INJECTION

NAVELBINE

AP +! PIERRE FABRE EQ 10MG BASE/ML N020388 001 Dec 23, 1994

VINORELBINE TARTRATE

| | | | | | |
|-----------|--|-------------------------|------------------------|--------------------|--------------|
| <u>AP</u> | | ACTAVIS TOTOWA | <u>EQ 10MG BASE/ML</u> | <u>A078011 001</u> | Jul 22, 2009 |
| <u>AP</u> | | DR REDDYS LABS LTD | <u>EQ 10MG BASE/ML</u> | <u>A202017 001</u> | Sep 12, 2013 |
| <u>AP</u> | | FRESENIUS KABI USA | <u>EQ 10MG BASE/ML</u> | <u>A076849 001</u> | Apr 18, 2005 |
| <u>AP</u> | | HOSPIRA | <u>EQ 10MG BASE/ML</u> | <u>A076827 001</u> | Jun 02, 2005 |
| <u>AP</u> | | JIANGSU HANSON PHARM | <u>EQ 10MG BASE/ML</u> | <u>A091106 001</u> | Sep 26, 2012 |
| <u>AP</u> | | TEVA PHARMS USA | <u>EQ 10MG BASE/ML</u> | <u>A076028 001</u> | Feb 03, 2003 |
| <u>AP</u> | | WEST-WARD PHARMS INT | <u>EQ 10MG BASE/ML</u> | <u>A075992 001</u> | Jun 10, 2003 |

PRESCRIPTION DRUG PRODUCT LIST

VINORELBINE TARTRATE

INJECTABLE; INJECTION

VINORELBINE TARTRATE**AP** **EQ 10MG BASE/ML** **A076461 001** Dec 11, 2003VISMODEGIB

CAPSULE; ORAL

ERIVEDGE

+! GENENTECH

150MG

N203388 001 Jan 30, 2012

VITAMIN A PALMITATE

INJECTABLE; INJECTION

AQUASOL A

+! CASPER PHARMA LLC

EQ 50,000 UNITS BASE/ML

N006823 001

VORAPAXAR SULFATE

TABLET; ORAL

ZONTIVITY

+! ARALEZ PHARMS

EQ 2.08MG BASE

N204886 001 May 08, 2014

VORICONAZOLE

FOR SUSPENSION; ORAL

VFEND**AB** **+!** PF PRISM CV **200MG/5ML** **N021630 001** Dec 19, 2003VORICONAZOLE**AB** AMNEAL PHARMS **200MG/5ML** **A205034 001** Apr 13, 2016**AB** NOVEL LABS INC **200MG/5ML** **A206799 001** May 31, 2016

INJECTABLE; IV (INFUSION)

VFEND**AP** **+!** PF PRISM CV **200MG/VIAL** **N021267 001** May 24, 2002VORICONAZOLE**AP** ALVOGEN INC **200MG/VIAL** **A206398 001** Mar 23, 2016**AP** HAINAN POLY PHARM **200MG/VIAL** **A211661 001** Nov 30, 2018**AP** SANDOZ INC **200MG/VIAL** **A090862 001** May 30, 2012**AP** ZYDUS PHARMS USA **200MG/VIAL** **A208983 001** Jul 16, 2018

INC

POWDER; IV (INFUSION)

VORICONAZOLE

XELLIA PHARMS APS

200MG/VIAL

N208562 001 Mar 09, 2017

TABLET; ORAL

VFEND**AB** **+** PF PRISM CV **50MG** **N021266 001** May 24, 2002**AB** **+!** **200MG** **N021266 002** May 24, 2002VORICONAZOLE**AB** AJANTA PHARMA LTD **50MG** **A206181 001** May 24, 2016**AB** **200MG** **A206181 002** May 24, 2016**AB** AKORN **50MG** **A207049 001** Sep 07, 2016**AB** **200MG** **A207049 002** Sep 07, 2016**AB** AUROBINDO PHARMA **50MG** **A206837 001** Jan 22, 2016

LTD

AB **200MG** **A206837 002** Jan 22, 2016**AB** GLENMARK PHARMS LTD **50MG** **A203503 001** Sep 02, 2015**AB** **200MG** **A203503 002** Sep 02, 2015**AB** MYLAN PHARMS INC **50MG** **A090547 001** Apr 22, 2010**AB** **200MG** **A090547 002** Apr 22, 2010**AB** NOVEL LABS INC **50MG** **A207371 001** May 24, 2016**AB** **200MG** **A207371 002** May 24, 2016**AB** PRINSTON INC **50MG** **A206654 001** Aug 08, 2016**AB** **200MG** **A206654 002** Aug 08, 2016**AB** RISING PHARMS **50MG** **A206762 001** May 24, 2016**AB** **200MG** **A206762 002** May 24, 2016**AB** SANDOZ INC **50MG** **A200265 001** Dec 12, 2011**AB** **200MG** **A200265 002** Dec 12, 2011**AB** TEVA PHARMS **50MG** **A091658 001** Apr 06, 2012**AB** **200MG** **A091658 002** Apr 06, 2012**AB** ZYDUS PHARMS USA **50MG** **A206747 001** May 24, 2016

INC

AB **200MG** **A206747 002** May 24, 2016

PRESCRIPTION DRUG PRODUCT LIST

VORINOSTAT

CAPSULE; ORAL

ZOLINZA

+! MERCK

100MG

N021991 001 Oct 06, 2006

VORTIOXETINE HYDROBROMIDE

TABLET; ORAL

TRINTELLIX

+ TAKEDA PHARMS USA

EQ 5MG BASE

N204447 001 Sep 30, 2013

+

EQ 10MG BASE

N204447 002 Sep 30, 2013

+!

EQ 20MG BASE

N204447 004 Sep 30, 2013

WARFARIN SODIUM

TABLET; ORAL

COUMADINAB + BRISTOL MYERS
SQUIBB1MGN009218 022 Mar 01, 1990AB +2MGN009218 013AB +2.5MGN009218 018AB +3MGN009218 025 Nov 18, 1996AB +4MGN009218 023 Aug 24, 1993AB +5MGN009218 007AB +6MGN009218 026 Nov 18, 1996AB +7.5MGN009218 016AB +!10MGN009218 005JANTOVENAB USL PHARMA1MGA040416 001 Oct 02, 2003AB2MGA040416 002 Oct 02, 2003AB2.5MGA040416 003 Oct 02, 2003AB3MGA040416 004 Oct 02, 2003AB4MGA040416 005 Oct 02, 2003AB5MGA040416 006 Oct 02, 2003AB6MGA040416 007 Oct 02, 2003AB7.5MGA040416 008 Oct 02, 2003AB10MGA040416 009 Oct 02, 2003WARFARIN SODIUMAB AMNEAL PHARMS1MGA202202 001 Mar 04, 2013AB2MGA202202 002 Mar 04, 2013AB2.5MGA202202 003 Mar 04, 2013AB3MGA202202 004 Mar 04, 2013AB4MGA202202 005 Mar 04, 2013AB5MGA202202 006 Mar 04, 2013AB6MGA202202 007 Mar 04, 2013AB7.5MGA202202 008 Mar 04, 2013AB10MGA202202 009 Mar 04, 2013AB

BARR

1MGA040145 001 Mar 26, 1997AB2MGA040145 002 Mar 26, 1997AB2.5MGA040145 003 Mar 26, 1997AB3MGA040145 008 Nov 05, 1998AB4MGA040145 004 Mar 26, 1997AB5MGA040145 005 Mar 26, 1997AB6MGA040145 009 Nov 05, 1998AB7.5MGA040145 006 Mar 26, 1997AB10MGA040145 007 Mar 26, 1997AB

INVAGEN PHARMS

1MGA090935 001 May 25, 2011AB2MGA090935 002 May 25, 2011AB2.5MGA090935 003 May 25, 2011AB3MGA090935 004 May 25, 2011AB4MGA090935 005 May 25, 2011AB5MGA090935 006 May 25, 2011AB6MGA090935 007 May 25, 2011AB7.5MGA090935 008 May 25, 2011AB10MGA090935 009 May 25, 2011AB

IPCA LABS LTD

1MGA200104 001 Jun 27, 2013AB2MGA200104 002 Jun 27, 2013AB2.5MGA200104 003 Jun 27, 2013AB3MGA200104 004 Jun 27, 2013AB4MGA200104 005 Jun 27, 2013AB5MGA200104 006 Jun 27, 2013AB6MGA200104 007 Jun 27, 2013AB7.5MGA200104 008 Jun 27, 2013AB10MGA200104 009 Jun 27, 2013AB

PLIVA

1MGA040616 009 Jul 05, 2006AB2MGA040616 001 Jul 05, 2006

PRESCRIPTION DRUG PRODUCT LIST

WARFARIN SODIUM

TABLET; ORAL

WARFARIN SODIUM

| | | | | |
|-----------|------------------|--------------|--------------------|--------------|
| <u>AB</u> | | <u>2.5MG</u> | <u>A040616 002</u> | Jul 05, 2006 |
| <u>AB</u> | | <u>3MG</u> | <u>A040616 003</u> | Jul 05, 2006 |
| <u>AB</u> | | <u>4MG</u> | <u>A040616 004</u> | Jul 05, 2006 |
| <u>AB</u> | | <u>5MG</u> | <u>A040616 005</u> | Jul 05, 2006 |
| <u>AB</u> | | <u>6MG</u> | <u>A040616 006</u> | Jul 05, 2006 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A040616 007</u> | Jul 05, 2006 |
| <u>AB</u> | | <u>10MG</u> | <u>A040616 008</u> | Jul 05, 2006 |
| <u>AB</u> | TARO PHARM | <u>1MG</u> | <u>A040301 002</u> | Jul 15, 1999 |
| <u>AB</u> | | <u>2MG</u> | <u>A040301 003</u> | Jul 15, 1999 |
| <u>AB</u> | | <u>2.5MG</u> | <u>A040301 004</u> | Jul 15, 1999 |
| <u>AB</u> | | <u>3MG</u> | <u>A040301 005</u> | Jul 15, 1999 |
| <u>AB</u> | | <u>4MG</u> | <u>A040301 006</u> | Jul 15, 1999 |
| <u>AB</u> | | <u>5MG</u> | <u>A040301 007</u> | Jul 15, 1999 |
| <u>AB</u> | | <u>6MG</u> | <u>A040301 008</u> | Jul 15, 1999 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A040301 009</u> | Jul 15, 1999 |
| <u>AB</u> | | <u>10MG</u> | <u>A040301 001</u> | Jul 15, 1999 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>1MG</u> | <u>A040663 001</u> | May 30, 2006 |
| <u>AB</u> | | <u>2MG</u> | <u>A040663 002</u> | May 30, 2006 |
| <u>AB</u> | | <u>2.5MG</u> | <u>A040663 003</u> | May 30, 2006 |
| <u>AB</u> | | <u>3MG</u> | <u>A040663 004</u> | May 30, 2006 |
| <u>AB</u> | | <u>4MG</u> | <u>A040663 005</u> | May 30, 2006 |
| <u>AB</u> | | <u>5MG</u> | <u>A040663 006</u> | May 30, 2006 |
| <u>AB</u> | | <u>6MG</u> | <u>A040663 007</u> | May 30, 2006 |
| <u>AB</u> | | <u>7.5MG</u> | <u>A040663 008</u> | May 30, 2006 |
| <u>AB</u> | | <u>10MG</u> | <u>A040663 009</u> | May 30, 2006 |

XENON XE-133

GAS; INHALATION

XENON XE 133

| | | | | |
|--|---------------------|------------|-------------|--------------|
| | LANTHEUS MEDCL | 10mCi/VIAL | N017284 001 | |
| | | 20mCi/VIAL | N017284 002 | |
| | MALLINKRODT NUCLEAR | 10mCi/VIAL | N018327 001 | Mar 09, 1982 |
| | | 20mCi/VIAL | N018327 002 | Mar 09, 1982 |

ZAFIRLUKAST

TABLET; ORAL

ACCOLATE

| | | | | | |
|-----------|---|---------------|-------------|--------------------|--------------|
| <u>AB</u> | + | PAR PHARM INC | <u>10MG</u> | <u>N020547 003</u> | Sep 17, 1999 |
| <u>AB</u> | + | ! | <u>20MG</u> | <u>N020547 001</u> | Sep 26, 1996 |

ZAFIRLUKAST

| | | | | | |
|-----------|--|--------------------|-------------|--------------------|--------------|
| <u>AB</u> | | DR REDDYS LABS LTD | <u>10MG</u> | <u>A090372 001</u> | Nov 18, 2010 |
| <u>AB</u> | | | <u>20MG</u> | <u>A090372 002</u> | Nov 18, 2010 |

ZALEPLON

CAPSULE; ORAL

SONATA

| | | | | | |
|-----------|---|--------|-------------|--------------------|--------------|
| <u>AB</u> | + | PFIZER | <u>5MG</u> | <u>N020859 001</u> | Aug 13, 1999 |
| <u>AB</u> | + | ! | <u>10MG</u> | <u>N020859 002</u> | Aug 13, 1999 |

ZALEPLON

| | | | | | |
|-----------|--|-------------------------|-------------|--------------------|--------------|
| <u>AB</u> | | AUROBINDO PHARMA | <u>5MG</u> | <u>A078829 001</u> | Jun 06, 2008 |
| <u>AB</u> | | | <u>10MG</u> | <u>A078829 002</u> | Jun 06, 2008 |
| <u>AB</u> | | CIPLA LTD | <u>5MG</u> | <u>A077505 001</u> | Jun 20, 2008 |
| <u>AB</u> | | | <u>10MG</u> | <u>A077505 002</u> | Jun 20, 2008 |
| <u>AB</u> | | HIKMA PHARMS | <u>5MG</u> | <u>A078147 001</u> | Nov 25, 2008 |
| <u>AB</u> | | | <u>10MG</u> | <u>A078147 002</u> | Nov 25, 2008 |
| <u>AB</u> | | ORCHID HLTHCARE | <u>5MG</u> | <u>A090374 001</u> | Sep 17, 2009 |
| <u>AB</u> | | | <u>10MG</u> | <u>A090374 002</u> | Sep 17, 2009 |
| <u>AB</u> | | TEVA PHARMS | <u>5MG</u> | <u>A077239 001</u> | Jun 06, 2008 |
| <u>AB</u> | | | <u>10MG</u> | <u>A077239 002</u> | Jun 06, 2008 |
| <u>AB</u> | | UNICHEM | <u>5MG</u> | <u>A078989 001</u> | Jun 06, 2008 |
| <u>AB</u> | | | <u>10MG</u> | <u>A078989 002</u> | Jun 06, 2008 |
| <u>AB</u> | | WEST-WARD PHARMS INT | <u>5MG</u> | <u>A077237 001</u> | Jun 06, 2008 |
| <u>AB</u> | | | <u>10MG</u> | <u>A077237 002</u> | Jun 06, 2008 |

ZANAMIVIR

POWDER; INHALATION

RELENZA

| | | | | | |
|--|---|-----------------|-----|-------------|--------------|
| | + | GLAXOSMITHKLINE | 5MG | N021036 001 | Jul 26, 1999 |
|--|---|-----------------|-----|-------------|--------------|

PRESCRIPTION DRUG PRODUCT LIST

ZICONOTIDE ACETATE

INJECTABLE; INTRATHECAL

PRIALT

| | | | | | |
|---|---|---------------------|------------------------|-------------|--------------|
| + | ! | TERSERA THERAPS LLC | 100MCG/1ML (100MCG/ML) | N021060 002 | Dec 28, 2004 |
| + | ! | | 500MCG/20ML (25MCG/ML) | N021060 001 | Dec 28, 2004 |
| + | ! | | 500MCG/5ML (100MCG/ML) | N021060 004 | Dec 28, 2004 |

ZIDOVUDINE

CAPSULE; ORAL

RETROVIR

| | | | | | | |
|-----------|---|---|---------------|--------------|--------------------|--------------|
| AB | + | ! | VIIV HLTHCARE | 100MG | N019655 001 | Mar 19, 1987 |
|-----------|---|---|---------------|--------------|--------------------|--------------|

ZIDOVUDINE

| | | | | | | |
|-----------|--|--|----------------------|--------------|--------------------|--------------|
| AB | | | AUROBINDO PHARMA LTD | 100MG | A078128 001 | Mar 27, 2006 |
| AB | | | CIPLA LTD | 100MG | A078349 001 | May 23, 2007 |

INJECTABLE; INJECTION

RETROVIR

| | | | | | | |
|-----------|---|---|---------------|----------------|--------------------|--------------|
| AP | + | ! | VIIV HLTHCARE | 10MG/ML | N019951 001 | Feb 02, 1990 |
|-----------|---|---|---------------|----------------|--------------------|--------------|

ZIDOVUDINE

| | | | | | | |
|-----------|--|--|----------|----------------|--------------------|--------------|
| AP | | | LUITPOLD | 10MG/ML | A091457 001 | May 06, 2010 |
|-----------|--|--|----------|----------------|--------------------|--------------|

SYRUP; ORAL

RETROVIR

| | | | | | | |
|-----------|---|---|---------------|-----------------|--------------------|--------------|
| AA | + | ! | VIIV HLTHCARE | 50MG/5ML | N019910 001 | Sep 28, 1989 |
|-----------|---|---|---------------|-----------------|--------------------|--------------|

ZIDOVUDINE

| | | | | | | |
|-----------|--|--|-----------|-----------------|--------------------|--------------|
| AA | | | AUROBINDO | 50MG/5ML | A077268 001 | Sep 19, 2005 |
| AA | | | CIPLA LTD | 50MG/5ML | A077981 001 | Jun 26, 2008 |

TABLET; ORAL

ZIDOVUDINE

| | | | | | | |
|-----------|---|--|----------------------|--------------|--------------------|--------------|
| AB | | | AUROBINDO | 300MG | A077267 001 | Sep 19, 2005 |
| AB | | | CIPLA | 300MG | A090561 001 | Oct 27, 2010 |
| AB | ! | | HETERO LABS LTD III | 300MG | A090092 001 | Apr 25, 2008 |
| AB | | | MYLAN PHARMS INC | 300MG | A078922 001 | Feb 14, 2008 |
| AB | | | WEST-WARD PHARMS INT | 300MG | A076844 001 | Sep 19, 2005 |

ZILEUTON

TABLET; ORAL

ZYFLO

| | | | | | |
|---|---|----------------|-------|-------------|--------------|
| + | ! | CHIESI USA INC | 600MG | N020471 003 | Dec 09, 1996 |
|---|---|----------------|-------|-------------|--------------|

TABLET, EXTENDED RELEASE; ORAL

ZILEUTON

| | | | | | | |
|-----------|--|--|---------------|--------------|--------------------|--------------|
| AB | | | RISING PHARMS | 600MG | A204929 001 | Mar 17, 2017 |
|-----------|--|--|---------------|--------------|--------------------|--------------|

ZYFLO CR

| | | | | | | |
|-----------|---|---|----------------|--------------|--------------------|--------------|
| AB | + | ! | CHIESI USA INC | 600MG | N022052 001 | May 30, 2007 |
|-----------|---|---|----------------|--------------|--------------------|--------------|

ZINC ACETATE

CAPSULE; ORAL

GALZIN

| | | | | | |
|---|---|------|--------------|-------------|--------------|
| + | | TEVA | EQ 25MG ZINC | N020458 001 | Jan 28, 1997 |
| + | ! | | EQ 50MG ZINC | N020458 002 | Jan 28, 1997 |

ZINC CHLORIDE

INJECTABLE; INJECTION

ZINC CHLORIDE IN PLASTIC CONTAINER

| | | | | | |
|---|---|---------|----------------|-------------|--------------|
| + | ! | HOSPIRA | EQ 1MG ZINC/ML | N018959 001 | Jun 26, 1986 |
|---|---|---------|----------------|-------------|--------------|

ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL

GEODON

| | | | | | | |
|-----------|---|---|--------|---------------------|--------------------|--------------|
| AB | + | ! | PFIZER | EQ 20MG BASE | N020825 001 | Feb 05, 2001 |
| AB | + | | | EQ 40MG BASE | N020825 002 | Feb 05, 2001 |
| AB | + | | | EQ 60MG BASE | N020825 003 | Feb 05, 2001 |
| AB | + | | | EQ 80MG BASE | N020825 004 | Feb 05, 2001 |

ZIPRASIDONE HYDROCHLORIDE

| | | | | | | |
|-----------|--|--|----------------------|---------------------|--------------------|--------------|
| AB | | | APOTEX INC | EQ 20MG BASE | A077561 001 | Mar 02, 2012 |
| AB | | | | EQ 40MG BASE | A077561 002 | Mar 02, 2012 |
| AB | | | | EQ 60MG BASE | A077561 003 | Mar 02, 2012 |
| AB | | | | EQ 80MG BASE | A077561 004 | Mar 02, 2012 |
| AB | | | AUROBINDO PHARMA LTD | EQ 20MG BASE | A204117 001 | Dec 27, 2016 |
| AB | | | | EQ 40MG BASE | A204117 002 | Dec 27, 2016 |
| AB | | | | EQ 60MG BASE | A204117 003 | Dec 27, 2016 |
| AB | | | | EQ 80MG BASE | A204117 004 | Dec 27, 2016 |
| AB | | | DR REDDYS LABS INC | EQ 20MG BASE | A077565 001 | Mar 02, 2012 |
| AB | | | | EQ 40MG BASE | A077565 002 | Mar 02, 2012 |

PRESCRIPTION DRUG PRODUCT LIST

ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL

ZIPRASIDONE HYDROCHLORIDE

| | | | | |
|-----------|-------------------------|---------------------|--------------------|--------------|
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A077565 003</u> | Mar 02, 2012 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A077565 004</u> | Mar 02, 2012 |
| <u>AB</u> | LUPIN PHARMS | <u>EQ 20MG BASE</u> | <u>A077560 001</u> | Mar 02, 2012 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077560 002</u> | Mar 02, 2012 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A077560 003</u> | Mar 02, 2012 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A077560 004</u> | Mar 02, 2012 |
| <u>AB</u> | MACLEODS PHARMS LTD | <u>EQ 20MG BASE</u> | <u>A204375 001</u> | Feb 17, 2017 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A204375 002</u> | Feb 17, 2017 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A204375 003</u> | Feb 17, 2017 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A204375 004</u> | Feb 17, 2017 |
| <u>AB</u> | SANDOZ INC | <u>EQ 20MG BASE</u> | <u>A077562 001</u> | Jun 01, 2012 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A077562 002</u> | Jun 01, 2012 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A077562 003</u> | Jun 01, 2012 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A077562 004</u> | Jun 01, 2012 |
| <u>AB</u> | WOCKHARDT LTD | <u>EQ 20MG BASE</u> | <u>A090348 001</u> | Sep 05, 2012 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A090348 002</u> | Sep 05, 2012 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A090348 003</u> | Sep 05, 2012 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A090348 004</u> | Sep 05, 2012 |
| <u>AB</u> | ZYDUS PHARMS USA INC | <u>EQ 20MG BASE</u> | <u>A208988 001</u> | Aug 22, 2017 |
| <u>AB</u> | | <u>EQ 40MG BASE</u> | <u>A208988 002</u> | Aug 22, 2017 |
| <u>AB</u> | | <u>EQ 60MG BASE</u> | <u>A208988 003</u> | Aug 22, 2017 |
| <u>AB</u> | | <u>EQ 80MG BASE</u> | <u>A208988 004</u> | Aug 22, 2017 |

ZIPRASIDONE MESYLATE

INJECTABLE; INTRAMUSCULAR

GEODON

+! PFIZER

EQ 20MG BASE/ML

N020919 001 Jun 21, 2002

ZOLEDRONIC ACID

INJECTABLE; IV (INFUSION)

RECLAST

| | | | | |
|-----------|-------------------------|--------------------------|--------------------|--------------|
| <u>AP</u> | +! NOVARTIS | <u>EQ 5MG BASE/100ML</u> | <u>N021817 001</u> | Apr 16, 2007 |
| | <u>ZOLEDRONIC</u> | | | |
| <u>AP</u> | GLAND PHARMA LTD | <u>EQ 4MG BASE/100ML</u> | <u>A205749 001</u> | Jun 29, 2018 |
| | <u>ZOLEDRONIC ACID</u> | | | |
| <u>AP</u> | ACCORD HLTHCARE | <u>EQ 4MG BASE/5ML</u> | <u>A205279 001</u> | Nov 28, 2016 |
| <u>AP</u> | ACTAVIS INC | <u>EQ 4MG BASE/5ML</u> | <u>A202472 001</u> | Mar 04, 2013 |
| <u>AP</u> | AKORN | <u>EQ 5MG BASE/100ML</u> | <u>A200918 001</u> | Aug 21, 2014 |
| <u>AP</u> | AKORN INC | <u>EQ 4MG BASE/5ML</u> | <u>A202548 001</u> | May 22, 2014 |
| <u>AP</u> | APOTEX INC | <u>EQ 5MG BASE/100ML</u> | <u>A204367 001</u> | Dec 24, 2015 |
| <u>AP</u> | AUROBINDO PHARMA LTD | <u>EQ 4MG BASE/5ML</u> | <u>A207751 001</u> | Sep 26, 2016 |
| <u>AP</u> | | <u>EQ 5MG BASE/100ML</u> | <u>A209125 001</u> | Dec 08, 2017 |
| <u>AP</u> | BPI LABS LLC | <u>EQ 4MG BASE/5ML</u> | <u>A207341 001</u> | Dec 29, 2017 |
| <u>AP</u> | BRECKENRIDGE PHARM | <u>EQ 4MG BASE/5ML</u> | <u>A091170 001</u> | Mar 04, 2013 |
| <u>AP</u> | | <u>EQ 4MG BASE/5ML</u> | <u>A202571 001</u> | May 07, 2013 |
| <u>AP</u> | | <u>EQ 5MG BASE/100ML</u> | <u>A202163 001</u> | Aug 05, 2013 |
| <u>AP</u> | CIPLA | <u>EQ 4MG BASE/100ML</u> | <u>A210174 001</u> | Oct 27, 2017 |
| <u>AP</u> | DR REDDYS LABS LTD | <u>EQ 4MG BASE/5ML</u> | <u>A091186 001</u> | Mar 04, 2013 |
| <u>AP</u> | | <u>EQ 4MG BASE/100ML</u> | <u>A204344 001</u> | Nov 19, 2018 |
| <u>AP</u> | | <u>EQ 5MG BASE/100ML</u> | <u>A091363 001</u> | Mar 29, 2013 |
| <u>AP</u> | EMCURE PHARMS LTD | <u>EQ 4MG BASE/5ML</u> | <u>A201783 001</u> | Mar 12, 2013 |
| <u>AP</u> | | <u>EQ 5MG BASE/100ML</u> | <u>A201801 001</u> | Mar 29, 2013 |
| <u>AP</u> | FRESENIUS KABI USA | <u>EQ 4MG BASE/5ML</u> | <u>A091516 001</u> | Apr 23, 2015 |
| <u>AP</u> | GLAND PHARMA LTD | <u>EQ 4MG BASE/5ML</u> | <u>A202930 001</u> | Aug 05, 2013 |
| <u>AP</u> | | <u>EQ 5MG BASE/100ML</u> | <u>A204217 001</u> | Aug 18, 2016 |
| <u>AP</u> | HIKMA FARMACEUTICA | <u>EQ 4MG BASE/5ML</u> | <u>A202182 001</u> | Jun 03, 2013 |
| <u>AP</u> | HOSPIRA INC | <u>EQ 4MG BASE/5ML</u> | <u>A090621 001</u> | Mar 19, 2015 |
| <u>AP</u> | | <u>EQ 5MG BASE/100ML</u> | <u>A202837 001</u> | Apr 05, 2013 |
| <u>AP</u> | INFORLIFE | <u>EQ 4MG BASE/100ML</u> | <u>N203231 001</u> | Aug 02, 2013 |
| <u>AP</u> | | <u>EQ 5MG BASE/100ML</u> | <u>A202828 001</u> | Sep 23, 2013 |
| <u>AP</u> | MYLAN LABS LTD | <u>EQ 4MG BASE/5ML</u> | <u>A202650 001</u> | Mar 04, 2013 |
| <u>AP</u> | | <u>EQ 5MG BASE/100ML</u> | <u>A203841 001</u> | Feb 14, 2017 |
| <u>AP</u> | | <u>EQ 5MG BASE/100ML</u> | <u>A205254 001</u> | Oct 27, 2017 |
| <u>AP</u> | SAGENT PHARMS | <u>EQ 4MG BASE/5ML</u> | <u>A091493 001</u> | Nov 24, 2014 |
| <u>AP</u> | USV NORTH AMERICA | <u>EQ 4MG BASE/5ML</u> | <u>A202923 001</u> | Sep 04, 2014 |
| | <u>ZOMETA</u> | | | |
| <u>AP</u> | +! NOVARTIS | <u>EQ 4MG BASE/5ML</u> | <u>N021223 002</u> | Mar 07, 2003 |
| <u>AP</u> | +! | <u>EQ 4MG BASE/100ML</u> | <u>N021223 003</u> | Jun 17, 2011 |

PRESCRIPTION DRUG PRODUCT LIST

ZOLEDRONIC ACID

SOLUTION;IV (INFUSION)

ZOLEDRONIC ACID

HOSPIRA INC

EQ 4MG BASE/100ML (EQ 0.04MG BASE/ML)

N204016 001 Dec 28, 2015

ZOLMITRIPTAN

SPRAY;NASAL

ZOMIG

+ ASTRAZENECA

2.5MG/SPRAY

N021450 003 Sep 16, 2013

+!

5MG/SPRAY

N021450 004 Sep 30, 2003

TABLET;ORAL

ZOLMITRIPTAN

| | | | | |
|-----------|------------------------------------|--------------|--------------------|--------------|
| AB | AJANTA PHARMA LTD | 2.5MG | A204041 001 | May 20, 2016 |
| AB | | 5MG | A204041 002 | May 20, 2016 |
| AB | ALEMBIC PHARMS LTD | 2.5MG | A204232 001 | Sep 30, 2015 |
| AB | | 5MG | A204232 002 | Sep 30, 2015 |
| AB | AUROBINDO PHARMA LTD | 2.5MG | A207021 001 | May 11, 2016 |
| AB | | 5MG | A207021 002 | May 11, 2016 |
| AB | GLENMARK GENERICS | 2.5MG | A201779 001 | May 14, 2013 |
| AB | | 5MG | A201779 002 | May 14, 2013 |
| AB | INVAGEN PHARMS | 2.5MG | A204284 001 | Apr 09, 2014 |
| AB | | 5MG | A204284 002 | Apr 09, 2014 |
| AB | JUBILANT GENERICS | 2.5MG | A202279 001 | Nov 20, 2014 |
| AB | | 5MG | A202279 002 | Nov 20, 2014 |
| AB | MACLEODS PHARMS LTD | 2.5MG | A203772 001 | Sep 30, 2015 |
| AB | | 5MG | A203772 002 | Sep 30, 2015 |
| AB | PLD ACQUISITIONS LLC | 2.5MG | A207867 001 | Feb 27, 2017 |
| AB | | 5MG | A207867 002 | Feb 27, 2017 |
| AB | TEVA PHARMS USA | 2.5MG | A090861 001 | Mar 04, 2014 |
| AB | | 5MG | A090861 002 | Mar 04, 2014 |
| AB | TWI PHARMS | 2.5MG | A206973 001 | Jun 30, 2017 |
| AB | | 5MG | A206973 002 | Jun 30, 2017 |
| AB | ZYDUS PHARMS USA INC | 2.5MG | A203019 001 | Jul 11, 2018 |
| AB | | 5MG | A203019 002 | Jul 11, 2018 |
| | ZOMIG | | | |
| AB | + IPR | 2.5MG | N020768 001 | Nov 25, 1997 |
| AB | +! | 5MG | N020768 002 | Nov 25, 1997 |
| | TABLET, ORALLY DISINTEGRATING;ORAL | | | |
| | ZOLMITRIPTAN | | | |
| AB | ALEMBIC PHARMS LTD | 2.5MG | A205074 001 | Dec 01, 2016 |
| AB | | 5MG | A205074 002 | Dec 01, 2016 |
| AB | GLENMARK GENERICS | 2.5MG | A202560 001 | May 14, 2013 |
| AB | | 5MG | A202560 002 | May 14, 2013 |
| AB | JUBILANT GENERICS | 2.5MG | A202956 001 | Sep 17, 2015 |
| AB | | 5MG | A202956 002 | Sep 17, 2015 |
| AB | MACLEODS PHARMS LTD | 2.5MG | A204336 001 | Oct 22, 2015 |
| AB | | 5MG | A204336 002 | Oct 22, 2015 |
| AB | ZYDUS PHARMS USA INC | 2.5MG | A202890 001 | May 15, 2013 |
| AB | | 5MG | A202890 002 | May 15, 2013 |
| | ZOMIG-ZMT | | | |
| AB | + ASTRAZENECA | 2.5MG | N021231 001 | Feb 13, 2001 |
| AB | +! | 5MG | N021231 002 | Sep 17, 2001 |
| | <u>ZOLPIDEM TARTRATE</u> | | | |
| | SPRAY, METERED;ORAL | | | |
| | ZOLPIMIST | | | |
| | +! MAGNA PHARMS | 5MG/SPRAY | | |
| | TABLET;ORAL | | | |
| | AMBIEN | | | |
| AB | + SANOFI AVENTIS US | 5MG | N019908 001 | Dec 16, 1992 |
| AB | +! | 10MG | N019908 002 | Dec 16, 1992 |
| | ZOLPIDEM TARTRATE | | | |
| AB | ACME LABS | 5MG | A077214 001 | Apr 23, 2007 |
| AB | | 10MG | A077214 002 | Apr 23, 2007 |
| AB | APOTEX INC | 5MG | A077884 001 | Apr 23, 2007 |
| AB | | 10MG | A077884 002 | Apr 23, 2007 |
| AB | AUROBINDO PHARMA | 5MG | A078413 001 | May 04, 2007 |
| AB | | 10MG | A078413 002 | May 04, 2007 |
| AB | CIPLA LTD | 5MG | A077388 001 | Jul 30, 2012 |
| AB | | 10MG | A077388 002 | Jul 30, 2012 |

PRESCRIPTION DRUG PRODUCT LIST

ZOLPIDEM TARTRATE

TABLET;ORAL

ZOLPIDEM TARTRATE

| | | | | |
|-----------|--------------------|-------------|--------------------|--------------|
| <u>AB</u> | INVAGEN PHARMS | <u>5MG</u> | <u>A078184 001</u> | Sep 07, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A078184 002</u> | Sep 07, 2007 |
| <u>AB</u> | MYLAN | <u>5MG</u> | <u>A076578 001</u> | Apr 23, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A076578 002</u> | Apr 23, 2007 |
| <u>AB</u> | SANDOZ INC | <u>5MG</u> | <u>A077322 001</u> | Apr 23, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A077322 002</u> | Apr 23, 2007 |
| <u>AB</u> | SUN PHARM INDS INC | <u>5MG</u> | <u>A077359 001</u> | Apr 23, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A077359 002</u> | Apr 23, 2007 |
| <u>AB</u> | TEVA | <u>5MG</u> | <u>A076410 001</u> | Apr 23, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A076410 002</u> | Apr 23, 2007 |
| <u>AB</u> | TORRENT PHARMS | <u>5MG</u> | <u>A077903 001</u> | Aug 17, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A077903 002</u> | Aug 17, 2007 |
| <u>AB</u> | VINTAGE | <u>5MG</u> | <u>A078616 001</u> | Nov 21, 2008 |
| <u>AB</u> | | <u>10MG</u> | <u>A078616 002</u> | Nov 21, 2008 |
| <u>AB</u> | WOCKHARDT | <u>5MG</u> | <u>A078426 001</u> | May 15, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A078426 002</u> | May 15, 2007 |
| <u>AB</u> | YUNG SHIN PHARM | <u>5MG</u> | <u>A077990 001</u> | Apr 23, 2007 |
| <u>AB</u> | | <u>10MG</u> | <u>A077990 002</u> | Apr 23, 2007 |

TABLET;SUBLINGUAL

EDLUAR

| | | | | |
|-----------|-----------------------|-------------|--------------------|--------------|
| <u>AB</u> | + MYLAN SPECIALITY LP | <u>5MG</u> | <u>N021997 001</u> | Mar 13, 2009 |
| <u>AB</u> | +! | <u>10MG</u> | <u>N021997 002</u> | Mar 13, 2009 |

INTERMEZZO

| | | | | |
|-----------|-----------------|---------------|--------------------|--------------|
| <u>AB</u> | + PURDUE PHARMA | <u>1.75MG</u> | <u>N022328 001</u> | Nov 23, 2011 |
| <u>AB</u> | +! | <u>3.5MG</u> | <u>N022328 002</u> | Nov 23, 2011 |

ZOLPIDEM TARTRATE

| | | | | |
|-----------|--------------------|---------------|--------------------|--------------|
| <u>AB</u> | DR REDDYS LABS INC | <u>1.75MG</u> | <u>A204503 001</u> | Nov 18, 2016 |
| <u>AB</u> | | <u>3.5MG</u> | <u>A204503 002</u> | Nov 18, 2016 |
| <u>AB</u> | MYLAN PHARMS INC | <u>5MG</u> | <u>A202657 001</u> | Aug 08, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A202657 002</u> | Aug 08, 2016 |
| <u>AB</u> | NOVEL LABS INC | <u>1.75MG</u> | <u>A204299 001</u> | Jun 03, 2015 |
| <u>AB</u> | | <u>3.5MG</u> | <u>A204299 002</u> | Jun 03, 2015 |
| <u>AB</u> | PAR FORM | <u>5MG</u> | <u>A201509 001</u> | Aug 01, 2016 |
| <u>AB</u> | | <u>10MG</u> | <u>A201509 002</u> | Aug 01, 2016 |
| <u>AB</u> | PAR PHARM INC | <u>1.75MG</u> | <u>A204229 001</u> | Sep 11, 2017 |
| <u>AB</u> | | <u>3.5MG</u> | <u>A204229 002</u> | Sep 11, 2017 |

TABLET, EXTENDED RELEASE;ORAL

AMBIEN CR

| | | | | |
|-----------|---------------------|---------------|--------------------|--------------|
| <u>AB</u> | + SANOFI AVENTIS US | <u>6.25MG</u> | <u>N021774 002</u> | Sep 02, 2005 |
| <u>AB</u> | +! | <u>12.5MG</u> | <u>N021774 001</u> | Sep 02, 2005 |

ZOLPIDEM TARTRATE

| | | | | |
|-----------|---------------------|---------------|--------------------|--------------|
| <u>AB</u> | ACTAVIS ELIZABETH | <u>6.25MG</u> | <u>A078179 002</u> | Oct 13, 2010 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A078179 001</u> | Jun 06, 2011 |
| <u>AB</u> | ACTAVIS LABS FL INC | <u>6.25MG</u> | <u>A090153 001</u> | Mar 25, 2013 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A090153 002</u> | Mar 25, 2013 |
| <u>AB</u> | ANCHEN PHARMS | <u>6.25MG</u> | <u>A078148 002</u> | Apr 14, 2011 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A078148 001</u> | Dec 03, 2010 |
| <u>AB</u> | APOTEX INC | <u>6.25MG</u> | <u>A200266 001</u> | Sep 10, 2013 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A200266 002</u> | Sep 10, 2013 |
| <u>AB</u> | LUPIN LTD | <u>6.25MG</u> | <u>A078970 001</u> | Sep 11, 2013 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A078970 002</u> | Sep 11, 2013 |
| <u>AB</u> | SANDOZ | <u>6.25MG</u> | <u>A090107 001</u> | Jul 01, 2011 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A090107 002</u> | Jul 01, 2011 |
| <u>AB</u> | SUN PHARMA GLOBAL | <u>6.25MG</u> | <u>A204170 001</u> | Jan 24, 2017 |
| <u>AB</u> | | <u>12.5MG</u> | <u>A204170 002</u> | Jan 24, 2017 |

ZONISAMIDE

CAPSULE;ORAL

ZONEGRAN

| | | | | |
|-----------|-----------------------|--------------|--------------------|--------------|
| <u>AB</u> | + SUNOVION PHARMS INC | <u>25MG</u> | <u>N020789 003</u> | Aug 22, 2003 |
| <u>AB</u> | +! | <u>100MG</u> | <u>N020789 001</u> | Mar 27, 2000 |

ZONISAMIDE

| | | | | |
|-----------|-------------------|--------------|--------------------|--------------|
| <u>AB</u> | APOTEX INC | <u>25MG</u> | <u>A077642 001</u> | Dec 22, 2005 |
| <u>AB</u> | | <u>50MG</u> | <u>A077642 002</u> | Dec 22, 2005 |
| <u>AB</u> | | <u>100MG</u> | <u>A077642 003</u> | Dec 22, 2005 |
| <u>AB</u> | BLUEPHARMA | <u>25MG</u> | <u>A077813 001</u> | Aug 16, 2006 |
| <u>AB</u> | | <u>50MG</u> | <u>A077813 002</u> | Aug 16, 2006 |
| <u>AB</u> | | <u>100MG</u> | <u>A077813 003</u> | Aug 16, 2006 |
| <u>AB</u> | GLENMARK GENERICS | <u>25MG</u> | <u>A077651 001</u> | Jan 30, 2006 |
| <u>AB</u> | | <u>50MG</u> | <u>A077651 002</u> | Jan 30, 2006 |

PRESCRIPTION DRUG PRODUCT LIST

ZONISAMIDE

CAPSULE; ORAL

ZONISAMIDE

| | | | | |
|-----------|---------------------|--------------|--------------------|--------------|
| <u>AB</u> | | <u>100MG</u> | <u>A077651 003</u> | Jan 30, 2006 |
| <u>AB</u> | INVAGEN PHARMS | <u>25MG</u> | <u>A077869 001</u> | May 31, 2006 |
| <u>AB</u> | | <u>50MG</u> | <u>A077869 002</u> | May 31, 2006 |
| <u>AB</u> | | <u>100MG</u> | <u>A077869 003</u> | May 31, 2006 |
| <u>AB</u> | MYLAN | <u>25MG</u> | <u>A077637 001</u> | Dec 22, 2005 |
| <u>AB</u> | | <u>50MG</u> | <u>A077637 002</u> | Dec 22, 2005 |
| <u>AB</u> | | <u>100MG</u> | <u>A077637 003</u> | Dec 22, 2005 |
| <u>AB</u> | SUN PHARM INDS (IN) | <u>25MG</u> | <u>A077634 001</u> | Mar 17, 2006 |
| <u>AB</u> | | <u>50MG</u> | <u>A077634 002</u> | Mar 17, 2006 |
| <u>AB</u> | | <u>100MG</u> | <u>A077634 003</u> | Mar 17, 2006 |
| <u>AB</u> | WOCKHARDT | <u>25MG</u> | <u>A077636 003</u> | Jul 27, 2006 |
| <u>AB</u> | | <u>50MG</u> | <u>A077636 002</u> | Jul 27, 2006 |
| <u>AB</u> | | <u>100MG</u> | <u>A077636 001</u> | Dec 22, 2005 |
| <u>AB</u> | ZYDUS PHARMS USA | <u>25MG</u> | <u>A077625 001</u> | Oct 16, 2006 |
| <u>AB</u> | | <u>50MG</u> | <u>A077625 002</u> | Oct 16, 2006 |
| <u>AB</u> | | <u>100MG</u> | <u>A077625 003</u> | Oct 16, 2006 |

OTC DRUG PRODUCT LIST

ACETAMINOPHEN

SUPPOSITORY; RECTAL

ACEPHEN

| | | |
|--------------|-------|--------------------------|
| G AND W LABS | 120MG | N018060 001 |
| | 325MG | A072344 001 Mar 27, 1992 |
| | 650MG | A072237 001 Mar 27, 1992 |

ACETAMINOPHEN

| | | |
|-----------------------|-------|--------------------------|
| PERRIGO NEW YORK | 120MG | A070607 001 Apr 06, 1987 |
| | 650MG | A070608 001 Dec 01, 1986 |
| + TARO PHARM INDS LTD | 120MG | N018337 003 Sep 12, 1983 |
| + | 325MG | N018337 002 |
| +! | 650MG | N018337 001 |

INFANTS' FEVERALL

| | | |
|-----------------------|------|--------------------------|
| + TARO PHARM INDS LTD | 80MG | N018337 004 Aug 26, 1992 |
|-----------------------|------|--------------------------|

NEOPAP

| | | |
|------------|-------|-------------|
| POLYMEDICA | 120MG | N016401 001 |
|------------|-------|-------------|

TABLET, EXTENDED RELEASE; ORAL

ACETAMINOPHEN

| | | |
|----------------------|-------|--------------------------|
| AUROBINDO PHARMA LTD | 650MG | A207229 001 Nov 09, 2016 |
| HERITAGE PHARMA | 650MG | A207035 001 May 31, 2018 |
| OHM LABS | 650MG | A076200 001 Mar 19, 2002 |
| PERRIGO | 650MG | A075077 001 Feb 25, 2000 |
| SUN PHARM INDS LTD | 650MG | A078569 001 Dec 14, 2011 |

TYLENOL

| | | |
|-------------------------|-------|--------------------------|
| +! J AND J CONSUMER INC | 650MG | N019872 001 Jun 08, 1994 |
| +! | 650MG | N019872 002 Jan 11, 2001 |

ACETAMINOPHEN; ASPIRIN; CAFFEINE

TABLET; ORAL

ACETAMINOPHEN, ASPIRIN AND CAFFEINE

| | | |
|---------|--------------------|--------------------------|
| PERRIGO | 250MG; 250MG; 65MG | A075794 001 Nov 26, 2001 |
|---------|--------------------|--------------------------|

EXCEDRIN (MIGRAINE)

| | | |
|-------------------------|--------------------|--------------------------|
| +! GLAXOSMITHKLINE CONS | 250MG; 250MG; 65MG | N020802 001 Jan 14, 1998 |
|-------------------------|--------------------|--------------------------|

ADAPALENE

GEL; TOPICAL

DIFFERIN

| | | |
|---------------------|------|--------------------------|
| +! GALDERMA LABS LP | 0.1% | N020380 002 Jul 08, 2016 |
|---------------------|------|--------------------------|

ALCOHOL; CHLORHEXIDINE GLUCONATE

SOLUTION; TOPICAL

AVAGARD

| | | |
|-------|---------|--------------------------|
| +! 3M | 61%; 1% | N021074 001 Jun 07, 2001 |
|-------|---------|--------------------------|

ALUMINUM HYDROXIDE; MAGNESIUM TRISILICATE

TABLET, CHEWABLE; ORAL

GAVISCON

| | | |
|---------------------|------------|--------------------------|
| + SANOFI AVENTIS US | 80MG; 20MG | N018685 001 Dec 09, 1983 |
|---------------------|------------|--------------------------|

ASPIRIN

CAPSULE; ORAL

VAZALORE

| | | |
|---------------|-------|--------------------------|
| +! PLX PHARMA | 325MG | N203697 001 Jan 14, 2013 |
|---------------|-------|--------------------------|

AVOBENZONE; ECAMSULE; OCTOCRYLENE

CREAM; TOPICAL

ANTHELIOS SX

| | | |
|---------------|-------------|--------------------------|
| +! LOREAL USA | 2%; 2%; 10% | N021502 001 Jul 21, 2006 |
|---------------|-------------|--------------------------|

CAPITAL SOLEIL 15

| | | |
|---------------|-------------|--------------------------|
| +! LOREAL USA | 2%; 3%; 10% | N021501 001 Oct 02, 2006 |
|---------------|-------------|--------------------------|

AVOBENZONE; ECAMSULE; OCTOCRYLENE; TITANIUM DIOXIDE

CREAM; TOPICAL

ANTHELIOS 20

| | | |
|---------------|-----------------|--------------------------|
| +! LOREAL USA | 2%; 2%; 10%; 2% | N021471 001 Oct 05, 2006 |
|---------------|-----------------|--------------------------|

ANTHELIOS 40

| | | |
|---------------|-----------------|--------------------------|
| +! LOREAL USA | 2%; 3%; 10%; 5% | N022009 001 Mar 31, 2008 |
|---------------|-----------------|--------------------------|

| | | |
|----|-----------------|--------------------------|
| +! | 2%; 3%; 10%; 5% | N022009 002 Oct 29, 2009 |
|----|-----------------|--------------------------|

OTC DRUG PRODUCT LIST

BENTOQUATAM

LOTION; TOPICAL

IVY BLOCK

+! STAND HOMEOPATH 5% N020532 001 Aug 26, 1996

BRIMONIDINE TARTRATE

SOLUTION/DROPS; OPHTHALMIC

LUMIFY

+! BAUSCH AND LOMB INC 0.025% N208144 001 Dec 22, 2017

BUDESONIDE

SPRAY, METERED; NASAL

BUDESONIDE

APOTEX INC 0.032MG/SPRAY A078949 002 Nov 20, 2015

RHINOCORT ALLERGY

+! ASTRAZENECA PHARMS 0.032MG/SPRAY N020746 003 Mar 23, 2015

BUTENAFINE HYDROCHLORIDE

CREAM; TOPICAL

BUTENAFINE HYDROCHLORIDE

TARO PHARMS 1% A205181 001 Nov 16, 2017

LOTRIMIN ULTRA

+! BAYER HEALTHCARE 1% N021307 001 Dec 07, 2001

LLC

CALCIUM CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE

TABLET, CHEWABLE; ORAL

FAMOTIDINE, CALCIUM CARBONATE, AND MAGNESIUM HYDROXIDE

PERRIGO R AND D 800MG;10MG;165MG A077355 001 Feb 06, 2008

800MG;10MG;165MG A204782 001 Aug 29, 2016

PEPCID COMPLETE

+! J AND J CONSUMER 800MG;10MG;165MG N020958 001 Oct 16, 2000

INC

CETIRIZINE HYDROCHLORIDE

CAPSULE; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

APOTEX INC 10MG A207235 001 Aug 12, 2016

AUROBINDO PHARMA 10MG A209107 001 Jul 20, 2018

LTD

+ BIONPHARMA INC 5MG N022429 001 Jul 23, 2009

+! 10MG N022429 004 Jul 23, 2009

STRIDES PHARMA 10MG A205291 001 Jul 21, 2017

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

AUROBINDO PHARMA 10MG A209107 002 Jul 20, 2018

LTD

+ BIONPHARMA INC 5MG N022429 003 Jul 23, 2009

+! 10MG N022429 002 Jul 23, 2009

SYRUP; ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

AMNEAL PHARMS 5MG/5ML A090765 002 Oct 07, 2009

APNAR PHARMA LP 5MG/5ML A091327 001 Oct 17, 2011

AUROBINDO PHARMA 5MG/5ML A090750 002 Feb 02, 2010

BIO PHARM INC 5MG/5ML A090474 002 Mar 30, 2009

LANNETT CO INC 5MG/5ML A091130 001 Apr 22, 2011

PERRIGO R AND D 5MG/5ML A204226 001 Sep 09, 2013

5MG/5ML A090254 002 Apr 09, 2008

TARO 5MG/5ML A090182 002 Apr 22, 2008

5MG/5ML A201546 001 May 20, 2011

TRIS PHARMA INC 5MG/5ML A090572 001 Nov 16, 2012

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

AMNEAL PHARMS 5MG/5ML A090765 001 Oct 07, 2009

APNAR PHARMA LP 5MG/5ML A091327 002 Oct 17, 2011

AUROBINDO PHARMA 5MG/5ML A090750 001 Feb 02, 2010

BIO PHARM INC 5MG/5ML A090474 001 Mar 30, 2009

LANNETT CO INC 5MG/5ML A091130 002 Apr 22, 2011

PERRIGO R AND D 5MG/5ML A090254 001 Apr 09, 2008

TARO 5MG/5ML A090182 001 Apr 22, 2008

5MG/5ML A201546 002 May 20, 2011

TRIS PHARMA INC 5MG/5ML A090572 002 Nov 16, 2012

CHILDREN'S ZYRTEC ALLERGY

+! J AND J CONSUMER 5MG/5ML N022155 002 Nov 16, 2007

INC

CHILDREN'S ZYRTEC HIVES RELIEF

+! J AND J CONSUMER 5MG/5ML N022155 001 Nov 16, 2007

INC

OTC DRUG PRODUCT LIST

CETIRIZINE HYDROCHLORIDE

TABLET; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

| | | | | |
|----------------------|------|---------|-----|--------------|
| AMNEAL PHARMS NY | 5MG | A078780 | 001 | Jan 21, 2010 |
| | 10MG | A078780 | 004 | Jan 21, 2010 |
| APOTEX INC | 5MG | A078317 | 001 | Dec 27, 2007 |
| | 10MG | A078317 | 002 | Dec 27, 2007 |
| AUROBINDO PHARMA LTD | 5MG | A090760 | 001 | Aug 05, 2015 |
| | 10MG | A090760 | 003 | Aug 05, 2015 |
| CIPLA LTD | 5MG | A077318 | 001 | Jul 25, 2013 |
| | 10MG | A077318 | 002 | Jul 25, 2013 |
| CONTRACT PHARMACAL | 5MG | A076047 | 001 | Dec 27, 2007 |
| | 10MG | A076047 | 002 | Dec 27, 2007 |
| DR REDDYS LABS LTD | 5MG | A078343 | 004 | Jan 15, 2008 |
| | 10MG | A078343 | 003 | Jan 15, 2008 |
| GRANULES INDIA LTD | 10MG | A209274 | 001 | Dec 22, 2017 |
| IPCA LABS LTD | 5MG | A202277 | 002 | Mar 11, 2014 |
| | 10MG | A202277 | 004 | Mar 11, 2014 |
| MARKSANS PHARMA | 5MG | A078933 | 001 | Jun 15, 2010 |
| | 10MG | A078933 | 002 | Jun 15, 2010 |
| MYLAN | 5MG | A076677 | 001 | Dec 27, 2007 |
| | 10MG | A076677 | 002 | Dec 27, 2007 |
| ORCHID HLTHCARE | 5MG | A078862 | 001 | Feb 19, 2009 |
| | 10MG | A078862 | 002 | Feb 19, 2009 |
| PERRIGO R AND D | 5MG | A078336 | 001 | Dec 27, 2007 |
| | 10MG | A078336 | 002 | Dec 27, 2007 |
| PLD ACQUISITIONS | 5MG | A077946 | 001 | Dec 27, 2007 |
| | 10MG | A077946 | 002 | Dec 27, 2007 |
| SUN PHARM INDS INC | 5MG | A077499 | 001 | Dec 27, 2007 |
| | 10MG | A077499 | 002 | Dec 27, 2007 |
| SUN PHARM INDS LTD | 5MG | A077498 | 001 | Dec 27, 2007 |
| | 10MG | A077498 | 002 | Dec 27, 2007 |
| TARO | 5MG | A078072 | 001 | Jul 22, 2009 |
| | 5MG | A078072 | 003 | Jul 22, 2009 |
| TORRENT PHARMS LLC | 5MG | A079191 | 001 | Apr 15, 2010 |
| | 10MG | A079191 | 004 | Apr 15, 2010 |
| UNICHEM | 5MG | A078680 | 003 | Jun 26, 2009 |
| | 10MG | A078680 | 004 | Jun 26, 2009 |
| UNIQUE PHARM LABS | 5MG | A077829 | 001 | Aug 26, 2009 |
| | 10MG | A077829 | 004 | Aug 26, 2009 |
| WOCKHARDT | 5MG | A078427 | 003 | Dec 28, 2007 |
| | 10MG | A078427 | 004 | Dec 28, 2007 |

CETIRIZINE HYDROCHLORIDE HIVES

| | | | | |
|--------------------|------|---------|-----|--------------|
| DR REDDYS LABS LTD | 5MG | A078343 | 001 | Jan 15, 2008 |
| | 10MG | A078343 | 002 | Jan 15, 2008 |
| IPCA LABS LTD | 5MG | A202277 | 001 | Mar 11, 2014 |
| | 10MG | A202277 | 003 | Mar 11, 2014 |
| MARKSANS PHARMA | 5MG | A078933 | 003 | Jun 15, 2010 |
| | 10MG | A078933 | 004 | Jun 15, 2010 |
| MYLAN | 5MG | A076677 | 004 | Dec 27, 2007 |
| | 10MG | A076677 | 003 | Dec 27, 2007 |
| ORCHID HLTHCARE | 5MG | A078862 | 003 | Feb 19, 2009 |
| | 10MG | A078862 | 004 | Feb 19, 2009 |
| PERRIGO R AND D | 5MG | A078336 | 003 | Dec 27, 2007 |
| | 10MG | A078336 | 004 | Dec 27, 2007 |
| SUN PHARM INDS INC | 5MG | A077499 | 003 | Dec 27, 2007 |
| | 10MG | A077499 | 004 | Dec 27, 2007 |
| SUN PHARM INDS LTD | 5MG | A077498 | 003 | Dec 27, 2007 |
| | 10MG | A077498 | 004 | Dec 27, 2007 |
| UNICHEM | 5MG | A078680 | 001 | Jun 26, 2009 |
| | 10MG | A078680 | 002 | Jun 26, 2009 |
| UNIQUE PHARM LABS | 5MG | A077829 | 003 | Aug 26, 2009 |
| ! | 10MG | A077829 | 002 | Aug 26, 2009 |

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

| | | | | |
|----------------------|------|---------|-----|--------------|
| AMNEAL PHARMS NY | 5MG | A078780 | 003 | Jan 21, 2010 |
| | 10MG | A078780 | 002 | Jan 21, 2010 |
| AUROBINDO PHARMA LTD | 5MG | A090760 | 002 | Aug 05, 2015 |
| | 10MG | A090760 | 004 | Aug 05, 2015 |
| TARO | 10MG | A078072 | 002 | Jul 22, 2009 |
| | 10MG | A078072 | 004 | Jul 22, 2009 |
| TORRENT PHARMS LLC | 5MG | A079191 | 003 | Apr 15, 2010 |

OTC DRUG PRODUCT LIST

CETIRIZINE HYDROCHLORIDE

TABLET;ORAL

CETIRIZINE HYDROCHLORIDE HIVES RELIEF
10MG

A079191 002 Apr 15, 2010

ZYRTEC ALLERGY

+! J AND J CONSUMER
INC

N019835 004 Nov 16, 2007

TABLET, CHEWABLE;ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

JUBILANT GENERICS 5MG

A091116 001 Feb 19, 2015

10MG

A091116 002 Feb 19, 2015

NOVEL LABS INC 5MG

A206793 001 Mar 08, 2016

10MG

A206793 002 Mar 08, 2016

SANDOZ 5MG

A078692 001 Feb 14, 2008

! 10MG

A078692 002 Feb 14, 2008

SUN PHARMA GLOBAL 5MG

A090142 001 Aug 30, 2011

10MG

A090142 002 Aug 30, 2011

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

JUBILANT GENERICS 5MG

A091116 003 Feb 19, 2015

10MG

A091116 004 Feb 19, 2015

SUN PHARMA GLOBAL 5MG

A090142 003 Aug 30, 2011

10MG

A090142 004 Aug 30, 2011

TABLET, ORALLY DISINTEGRATING;ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

PERRIGO R AND D 10MG

A205490 001 Sep 02, 2015

ZYRTEC ALLERGY

+! J AND J CONSUMER
INC

N022578 001 Sep 03, 2010

CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

IVAX SUB TEVA 5MG;120MG

A077170 001 Feb 25, 2008

PHARMS

PERRIGO R AND D 5MG;120MG

A210719 001 Nov 16, 2018

PLD ACQUISITIONS 5MG;120MG

A077991 001 Mar 05, 2008

SUN PHARM INDS LTD 5MG;120MG

A090922 001 Sep 28, 2012

ZYRTEC-D 12 HOUR

+! J AND J CONSUMER
INC

N021150 002 Nov 09, 2007

CHLORHEXIDINE GLUCONATE

AEROSOL, METERED;TOPICAL

EXIDINE

+! XTTRIUM 4%

N019127 001 Dec 24, 1984

CLOTH;TOPICAL

CHLORHEXIDINE GLUCONATE

+! SAGE PRODS 2%

N021669 001 Apr 25, 2005

READYPREP CHG

MEDLINE INDUSTRIES 2%

N207964 001 Nov 20, 2018

SOLUTION;TOPICAL

BRIAN CARE

SOAPCO 4%

A071419 001 Dec 17, 1987

CHG SCRUB

ECOLAB 4%

N019258 002 Jul 22, 1986

CIDA-STAT

ECOLAB 2%

N019258 001 Jul 22, 1986

EXIDINE

+! XTTRIUM 2%

N019422 001 Dec 17, 1985

4%

N019125 001 Dec 24, 1984

HIBICLENS

+! MOLNLYCKE HLTH 4%

N017768 001

HIBISTAT

+! MOLNLYCKE HLTH 0.5%

N018300 001

SPONGE;TOPICAL

BIOSCRUB

GRIFFEN 4%

N019822 001 Mar 31, 1989

CHLORHEXIDINE GLUCONATE

! BECTON DICKINSON 4%

A072525 001 Oct 24, 1989

OTC DRUG PRODUCT LIST

CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL

SOLUTION; TOPICAL

SOLUPREP

+! 3M HEALTH CARE 2%;70% N208288 001 Aug 08, 2018

SPONGE; TOPICAL

CHLORAPREP ONE-STEP

+! BECTON DICKINSON CO 2%;70% (3ML) N020832 001 Jul 14, 2000

+! 2%;70% (10.5ML) N020832 004 Aug 20, 2003

+! 2%;70% (26ML) N020832 006 Nov 21, 2006

+! 2%;70% (1ML) N020832 008 Oct 23, 2008

CHLORAPREP ONE-STEP FREPP

+! BECTON DICKINSON CO 2%;70% (1.5ML) N020832 003 Apr 26, 2002

CHLORAPREP WITH TINT

+! BECTON DICKINSON CO 2%;70% (26ML) N020832 002 May 03, 2005

+! 2%;70% (10.5ML) N020832 005 Apr 03, 2006

+! 2%;70% (3ML) N020832 007 Oct 10, 2006

SWAB; TOPICAL

CHLORAPREP ONE-STEP SEPP

+! BECTON DICKINSON CO 2%;70% (0.67ML) N021555 001 Oct 07, 2002

CHLORAPREP SINGLE SWABSTICK

+! BECTON DICKINSON CO 2%;70% (1.75ML) N021555 002 May 10, 2005

CHLORAPREP TRIPLE SWABSTICK

+! BECTON DICKINSON CO 2%;70% (5.25ML) N021555 003 Jun 10, 2009

PREVANTICS MAXI SWABSTICK

+! PROF DSPLS 3.15%;70% (5.1ML) N021524 003 Jun 03, 2005

PREVANTICS SWAB

+! PROF DSPLS 3.15%;70% (1ML) N021524 001 Jun 03, 2005

PREVANTICS SWABSTICK

+! PROF DSPLS 3.15%;70% (1.6ML) N021524 002 Jun 03, 2005

CHLORPHENIRAMINE MALEATE

TABLET, EXTENDED RELEASE; ORAL

CHLOR-TRIMETON

+ BAYER HEALTHCARE 12MG N007638 002

LLC

CHLORPHENIRAMINE MALEATE

! AVANTHI INC 12MG A040829 001 May 13, 2009

CHLORPHENIRAMINE MALEATE; IBUPROFEN; PHENYLEPHRINE HYDROCHLORIDE

TABLET; ORAL

ADVIL ALLERGY AND CONGESTION RELIEF

+! PFIZER 4MG;200MG;10MG N022113 001 Dec 21, 2011

ADVIL MULTI-SYMPTOM COLD & FLU

+! PFIZER 4MG;200MG;10MG N022113 002 Apr 28, 2017

CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

SUSPENSION; ORAL

CHILDREN'S ADVIL ALLERGY SINUS

+! PFIZER 1MG/5ML;100MG/5ML;15MG/5ML N021587 001 Feb 24, 2004

TABLET; ORAL

ADVIL ALLERGY SINUS

+! PFIZER 2MG;200MG;30MG N021441 001 Dec 19, 2002

CIMETIDINE

TABLET; ORAL

CIMETIDINE

APOTEX

100MG

A074948 001 Jun 19, 1998

200MG

A074948 002 Jul 26, 2002

IVAX SUB TEVA

200MG

A075345 001 Jun 16, 1999

PHARMS

L PERRIGO CO

200MG

A075285 001 Oct 29, 1998

TAGAMET HB

+! MEDTECH PRODUCTS 200MG N020238 002 Aug 21, 1996

CLEMASTINE FUMARATE

TABLET; ORAL

CLEMASTINE FUMARATE

! L PERRIGO CO 1.34MG A074512 001 Nov 22, 1995

CLOTRIMAZOLE

CREAM; VAGINAL

CLOTRIMAZOLE

! ACTAVIS MID 1% A074165 001 Jul 16, 1993

ATLANTIC

TARO

1%

A072641 001 Dec 04, 1995

OTC DRUG PRODUCT LIST

CLOTRIMAZOLE

CREAM;VAGINAL

TRIVAGIZOLE 3

TARO

2%

N021143 001 Apr 12, 2000

CROMOLYN SODIUM

SPRAY, METERED;NASAL

CROMOLYN SODIUM

!

BAUSCH AND LOMB

5.2MG/SPRAY

A075702 001 Jul 03, 2001

PERRIGO

5.2MG/SPRAY

A075427 001 Dec 12, 2001

DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN

TABLET, EXTENDED RELEASE;ORAL

GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE

ACTAVIS LABS FL

30MG;600MG

A091070 001 Aug 31, 2015

60MG;1.2GM

A091070 002 Aug 31, 2015

AMNEAL PHARMS

30MG;600MG

A209692 001 Nov 01, 2018

60MG;1.2GM

A209692 002 Nov 01, 2018

AUROBINDO PHARMA

30MG;600MG

A206941 001 Mar 17, 2017

LTD

60MG;1.2GM

A206941 002 Mar 17, 2017

PERRIGO R AND D

30MG;600MG

A207602 002 Mar 05, 2018

60MG;1.2GM

A207602 001 Mar 05, 2018

MUCINEX DM

+

RB HLTH

30MG;600MG

N021620 002 Apr 29, 2004

+!

60MG;1.2GM

N021620 001 Apr 29, 2004

DEXTROMETHORPHAN POLISTIREX

SUSPENSION, EXTENDED RELEASE;ORAL

DELSYM

+!

RB HLTH

EQ 30MG HYDROBROMIDE/5ML

N018658 001 Oct 08, 1982

DEXTROMETHORPHAN POLISTIREX

AMNEAL PHARMS LLC

EQ 30MG HYDROBROMIDE/5ML

A203133 001 Jul 28, 2017

TRIS PHARMA INC

EQ 30MG HYDROBROMIDE/5ML

A091135 001 May 25, 2012

DIPHENHYDRAMINE CITRATE; IBUPROFEN

TABLET;ORAL

ADVIL PM

+!

PFIZER

38MG;200MG

N021394 001 Dec 21, 2005

IBUPROFEN AND DIPHENHYDRAMINE CITRATE

DR REDDYS LABS LTD

38MG;200MG

A090619 001 Jul 08, 2009

PERRIGO R AND D

38MG;200MG

A079113 001 Dec 22, 2008

DIPHENHYDRAMINE HYDROCHLORIDE; IBUPROFEN

CAPSULE;ORAL

ADVIL PM

+!

PFIZER

25MG;EQ 200MG FREE ACID AND POTASSIUM
SALT

N021393 001 Dec 21, 2005

IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE

BIONPHARMA INC

25MG;EQ 200MG FREE ACID AND POTASSIUM
SALT

A090397 001 Nov 22, 2010

STRIDES PHARMA

25MG;EQ 200MG FREE ACID AND POTASSIUM
SALT

A200888 001 Mar 05, 2012

DIPHENHYDRAMINE HYDROCHLORIDE; NAPROXEN SODIUM

TABLET;ORAL

ALEVE PM

+!

BAYER HLTHCARE

25MG;220MG

N205352 001 Jan 17, 2014

NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE

AMNEAL PHARMS CO

25MG;220MG

A209726 001 Oct 23, 2018

DOCOSANOL

CREAM;TOPICAL

ABREVA

+!

GLAXOSMITHKLINE

10%

N020941 001 Jul 25, 2000

DOCOSANOL

ACTAVIS LABS UT INC

10%

A208754 001 Nov 19, 2018

DOXYLAMINE SUCCINATE

TABLET;ORAL

DOXYLAMINE SUCCINATE

LNK

25MG

A040564 001 Aug 27, 2004

PERRIGO

25MG

A040167 001 Sep 18, 1996

UNISOM

+!

CHATTEM

25MG

N018066 001

OTC DRUG PRODUCT LIST

EPINEPHRINE

AEROSOL, METERED; INHALATION

PRIMATENE MIST

+! ARMSTRONG PHARMS 0.125MG/INH N205920 001 Nov 07, 2018

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE; ORAL

ESOMEPRAZOLE MAGNESIUM

AUROBINDO PHARMA EQ 20MG BASE A209339 001 Oct 16, 2017

LTD

DR REDDYS LABS LTD EQ 20MG BASE A207673 001 May 15, 2018

PERRIGO R AND D EQ 20MG BASE A207193 001 Aug 18, 2017

NEXIUM 24HR

+! ASTRAZENECA LP EQ 20MG BASE N204655 001 Mar 28, 2014

TABLET, DELAYED RELEASE; ORAL

NEXIUM 24HR

+! ASTRAZENECA LP EQ 20MG BASE N207920 001 Nov 23, 2015

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

AUROBINDO PHARMA 10MG A206531 001 Apr 26, 2016

LTD

20MG A206531 002 Apr 26, 2016

DR REDDYS LABS LTD 10MG A075758 001 Aug 17, 2001

20MG A077367 001 Sep 25, 2006

IVAX SUB TEVA 10MG A075512 001 Jul 26, 2001

PHARMS

MYLAN 10MG A075674 001 Dec 21, 2001

PERRIGO 10MG A075400 001 Mar 18, 2005

PERRIGO R AND D 20MG A077351 001 Sep 25, 2006

SUN PHARM INDS LTD 10MG A090283 001 Nov 17, 2009

20MG A090283 002 Nov 17, 2009

TEVA 10MG A075312 001 May 31, 2001

WOCKHARDT 10MG A077146 001 Mar 07, 2005

20MG A090837 001 Aug 04, 2010

PEPCID AC

+ J AND J CONSUMER 10MG N020325 001 Apr 28, 1995

INC

10MG N020902 001 Aug 05, 1999

+! 20MG N020325 002 Sep 23, 2003

TABLET, CHEWABLE; ORAL

FAMOTIDINE

PERRIGO 10MG A075715 001 Aug 22, 2003

PEPCID AC

+! J AND J CONSUMER 20MG N020801 002 Dec 17, 2007

INC

FEXOFENADINE HYDROCHLORIDE

SUSPENSION; ORAL

CHILDREN'S ALLEGRA ALLERGY

+! SANOFI AVENTIS US 30MG/5ML N201373 001 Jan 24, 2011

CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

ACTAVIS MID 30MG/5ML A203330 001 Nov 18, 2014

ATLANTIC

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

! ACTAVIS MID 30MG/5ML A203330 002 Nov 18, 2014

ATLANTIC

TABLET; ORAL

ALLEGRA ALLERGY

+ SANOFI AVENTIS US 60MG N020872 007 Jan 24, 2011

+! 180MG N020872 010 Jan 24, 2011

CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

AUROLIFE PHARMA LLC 30MG A202039 001 Nov 19, 2014

DR REDDYS LABS LTD 30MG A076502 004 Apr 12, 2011

HETERO LABS LTD V 30MG A204097 001 Aug 19, 2016

MYLAN 30MG A077081 004 Jul 21, 2011

SUN PHARM INDS 30MG A091567 002 Feb 06, 2012

TEVA 30MG A076447 004 Apr 13, 2011

WOCKHARDT LTD 30MG A079112 002 Feb 08, 2012

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

DR REDDYS LABS LTD 30MG A076502 005 Apr 12, 2011

MYLAN 30MG A077081 005 Jul 21, 2011

SUN PHARM INDS 30MG A091567 001 Feb 06, 2012

TEVA 30MG A076447 005 Apr 13, 2011

OTC DRUG PRODUCT LIST

FEXOFENADINE HYDROCHLORIDE

TABLET; ORAL

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

WOCKHARDT LTD 30MG A079112 001 Feb 08, 2012

FEXOFENADINE HYDROCHLORIDE ALLERGY

AUROLIFE PHARMA LLC 60MG A202039 002 Nov 19, 2014

180MG A202039 003 Nov 19, 2014

DR REDDYS LABS LTD 60MG A076502 006 Apr 12, 2011

180MG A076502 008 Apr 12, 2011

HETERO LABS LTD V 60MG A204097 002 Aug 19, 2016

180MG A204097 003 Aug 19, 2016

MYLAN 60MG A077081 006 Jul 21, 2011

180MG A077081 008 Jul 21, 2011

SCIEGEN PHARMS INC 60MG A204507 002 Sep 16, 2015

180MG A204507 003 Sep 16, 2015

SUN PHARM INDS 60MG A091567 004 Feb 06, 2012

180MG A091567 006 Feb 06, 2012

TEVA 60MG A076447 006 Apr 13, 2011

180MG A076447 008 Apr 13, 2011

UNIQUE PHARM LABS 180MG A210137 001 Aug 13, 2018

WOCKHARDT LTD 60MG A079112 004 Feb 08, 2012

180MG A079112 006 Feb 08, 2012

FEXOFENADINE HYDROCHLORIDE HIVES

DR REDDYS LABS LTD 60MG A076502 007 Apr 12, 2011

180MG A076502 009 Apr 12, 2011

MYLAN 60MG A077081 007 Jul 21, 2011

180MG A077081 009 Jul 21, 2011

SCIEGEN PHARMS INC 60MG A204507 004 Sep 16, 2015

180MG A204507 005 Sep 16, 2015

SUN PHARM INDS 60MG A091567 003 Feb 06, 2012

180MG A091567 005 Feb 06, 2012

TEVA 60MG A076447 007 Apr 13, 2011

WOCKHARDT LTD 60MG A079112 003 Feb 08, 2012

180MG A079112 005 Feb 08, 2012

FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION

+! SANOFI AVENTIS US 60MG;120MG N020786 002 Jan 24, 2011

ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION

+! SANOFI AVENTIS US 180MG;240MG N021704 002 Jan 24, 2011

FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

AUROBINDO PHARMA 60MG;120MG A209116 001 Oct 30, 2017

LTD

DR REDDYS LABS LTD 60MG;120MG A076667 001 Nov 18, 2014

180MG;240MG A079043 002 Jun 22, 2011

SUN PHARMA GLOBAL 60MG;120MG A090818 001 Jan 29, 2015

FLUTICASONE FUROATE

SPRAY, METERED; NASAL

FLONASE SENSIMIST ALLERGY RELIEF

+! GLAXOSMITHKLINE 0.0275MG/SPRAY N022051 002 Aug 02, 2016

CONS

FLUTICASONE PROPIONATE

SPRAY, METERED; NASAL

FLONASE ALLERGY RELIEF

+! GLAXOSMITHKLINE 0.05MG/SPRAY N205434 001 Jul 23, 2014

CONS

FLUTICASONE PROPIONATE

APOTEX INC 0.05MG/SPRAY A208150 001 Feb 29, 2016

WEST-WARD PHARMS 0.05MG/SPRAY A207957 001 May 26, 2016

INT

GUAIFENESIN

TABLET, EXTENDED RELEASE; ORAL

GUAIFENESIN

ACTAVIS LABS FL 1.2GM A091009 002 Sep 03, 2015

AMNEAL PHARMS 600MG A207342 001 Jul 11, 2018

1.2GM A207342 002 Jul 11, 2018

GUARDIAN DRUG 600MG A209215 001 Sep 06, 2017

1.2GM A209215 002 Sep 06, 2017

OHM LABS INC 600MG A209254 001 Jul 16, 2018

1.2GM A209254 002 Jul 16, 2018

PERRIGO R AND D 600MG A078912 001 Nov 23, 2011

OTC DRUG PRODUCT LIST

GUAIFENESIN

TABLET, EXTENDED RELEASE;ORAL

MUCINEX

| | | | | | |
|---|---------|-------|---------|-----|--------------|
| + | RB HLTH | 600MG | N021282 | 001 | Jul 12, 2002 |
| + | ! | 1.2GM | N021282 | 002 | Dec 18, 2002 |

GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE

| | | | | |
|--------------------|-------------|---------|-----|--------------|
| ACTAVIS LABS FL | 600MG;60MG | A091071 | 001 | May 27, 2015 |
| | 1.2GM;120MG | A091071 | 002 | May 27, 2015 |
| DR REDDYS LABS LTD | 600MG;60MG | A208369 | 001 | Dec 29, 2017 |
| | 1.2GM;120MG | A208369 | 002 | Dec 29, 2017 |

MUCINEX D

| | | | | | |
|---|---------|-------------|---------|-----|--------------|
| + | RB HLTH | 600MG;60MG | N021585 | 001 | Jun 22, 2004 |
| + | ! | 1.2GM;120MG | N021585 | 002 | Jun 22, 2004 |

IBUPROFEN

CAPSULE;ORAL

ADVIL LIQUI-GELS

| | | | | | | |
|---|---|--------|---------------------------------------|---------|-----|--------------|
| + | ! | PFIZER | EQ 200MG FREE ACID AND POTASSIUM SALT | N020402 | 001 | Apr 20, 1995 |
|---|---|--------|---------------------------------------|---------|-----|--------------|

ADVIL MIGRAINE LIQUI-GELS

| | | | | | | |
|---|---|--------|---------------------------------------|---------|-----|--------------|
| + | ! | PFIZER | EQ 200MG FREE ACID AND POTASSIUM SALT | N020402 | 002 | Mar 16, 2000 |
|---|---|--------|---------------------------------------|---------|-----|--------------|

IBUPROFEN

| | | | | |
|----------------------|---------------------------------------|---------|-----|--------------|
| AMNEAL PHARMS | EQ 200MG FREE ACID AND POTASSIUM SALT | A202300 | 001 | Dec 23, 2011 |
| ASCENT PHARMS INC | EQ 200MG FREE ACID AND POTASSIUM SALT | A206999 | 001 | Dec 21, 2017 |
| AUROBINDO PHARMA LTD | EQ 200MG FREE ACID AND POTASSIUM SALT | A207753 | 001 | Jun 29, 2018 |
| BIONPHARMA INC | EQ 200MG FREE ACID AND POTASSIUM SALT | A078682 | 001 | Mar 24, 2009 |
| HUMANWELL PURACAP | EQ 200MG FREE ACID AND POTASSIUM SALT | A206568 | 001 | Jun 21, 2016 |
| MARKSANS PHARMA | EQ 200MG FREE ACID AND POTASSIUM SALT | A079205 | 001 | Jun 26, 2009 |
| P AND L DEV LLC | EQ 200MG FREE ACID AND POTASSIUM SALT | A077338 | 001 | Jul 10, 2009 |
| SOFGEN PHARMS | EQ 200MG FREE ACID AND POTASSIUM SALT | A203599 | 001 | Sep 07, 2016 |
| STRIDES PHARMA | EQ 200MG FREE ACID AND POTASSIUM SALT | A204469 | 001 | Mar 28, 2018 |

MIDOL LIQUID GELS

| | | | | | | |
|---|---|----------------|-------|---------|-----|--------------|
| + | ! | BIONPHARMA INC | 200MG | N021472 | 001 | Oct 18, 2002 |
|---|---|----------------|-------|---------|-----|--------------|

SUSPENSION;ORAL

CHILDREN'S ADVIL

| | | | | |
|--------|-----------|---------|-----|--------------|
| PFIZER | 100MG/5ML | N020589 | 001 | Jun 27, 1996 |
|--------|-----------|---------|-----|--------------|

CHILDREN'S ADVIL-FLAVORED

| | | | | |
|--------|-----------|---------|-----|--------------|
| PFIZER | 100MG/5ML | N020589 | 002 | Nov 07, 1997 |
|--------|-----------|---------|-----|--------------|

CHILDREN'S ELIXSURE

| | | | | |
|---------------------|-----------|---------|-----|--------------|
| MOBERG PHARMA NORTH | 100MG/5ML | N021604 | 001 | Jan 07, 2004 |
|---------------------|-----------|---------|-----|--------------|

CHILDREN'S IBUPROFEN

| | | | | |
|---------|-----------|---------|-----|--------------|
| PERRIGO | 100MG/5ML | A074937 | 001 | Dec 22, 1998 |
|---------|-----------|---------|-----|--------------|

CHILDREN'S MOTRIN

| | | | | | | |
|---|---|----------------------|-----------|---------|-----|--------------|
| + | ! | J AND J CONSUMER INC | 100MG/5ML | N020516 | 001 | Jun 16, 1995 |
|---|---|----------------------|-----------|---------|-----|--------------|

IBUPROFEN

| | | | | |
|----------------------|-----------|---------|-----|--------------|
| ACTAVIS MID ATLANTIC | 100MG/5ML | A074916 | 001 | Apr 30, 1999 |
| APTAPHARMA INC | 100MG/5ML | A210602 | 001 | Nov 23, 2018 |
| ARISE PHARMS | 100MG/5ML | A200457 | 001 | Aug 18, 2011 |
| AUROBINDO PHARMA LTD | 100MG/5ML | A209179 | 001 | Apr 17, 2018 |
| GUARDIAN DRUG | 100MG/5ML | A210149 | 001 | Aug 17, 2018 |
| TARO | 100MG/5ML | A209207 | 001 | Jun 27, 2017 |

SUSPENSION/DROPS;ORAL

CHILDREN'S MOTRIN

| | | | | | | |
|---|---|----------------------|---------|---------|-----|--------------|
| + | ! | J AND J CONSUMER INC | 40MG/ML | N020603 | 001 | Jun 10, 1996 |
|---|---|----------------------|---------|---------|-----|--------------|

IBUPROFEN

| | | | | |
|-----------------|---------|---------|-----|--------------|
| GUARDIAN DRUG | 40MG/ML | A210755 | 001 | Sep 26, 2018 |
| L PERRIGO CO | 40MG/ML | A075217 | 001 | Dec 16, 1998 |
| TRIS PHARMA INC | 40MG/ML | A079058 | 001 | Aug 31, 2009 |

PEDIATRIC ADVIL

| | | | | | | |
|---|---|--------|-------------|---------|-----|--------------|
| + | ! | PFIZER | 100MG/2.5ML | N020812 | 001 | Jan 30, 1998 |
|---|---|--------|-------------|---------|-----|--------------|

TABLET;ORAL

ADVIL

| | | | | |
|--------|-------|---------|-----|--------------|
| PFIZER | 200MG | N018989 | 001 | May 18, 1984 |
|--------|-------|---------|-----|--------------|

IBU-TAB 200

| | | | | |
|------|-------|---------|-----|--------------|
| ALRA | 200MG | A071057 | 001 | Aug 11, 1988 |
|------|-------|---------|-----|--------------|

IBUPROFEN

| | | | | |
|---------------|-------|---------|-----|--------------|
| AMNEAL PHARMS | 200MG | A079233 | 001 | Mar 18, 2014 |
|---------------|-------|---------|-----|--------------|

OTC DRUG PRODUCT LIST

IBUPROFEN

TABLET;ORAL

IBUPROFEN

| | | | | |
|----------------------|-------|---------|-----|--------------|
| AMNEAL PHARMS NY | 200MG | A071333 | 001 | Feb 17, 1987 |
| | 200MG | A072199 | 001 | May 23, 1988 |
| AUROBINDO PHARMA LTD | 200MG | A208865 | 001 | Nov 08, 2017 |
| AVEMA PHARMA | 200MG | A076460 | 001 | Nov 26, 2003 |
| CONTRACT PHARMACAL | 200MG | A072299 | 001 | Jul 01, 1988 |
| DR REDDYS LA | 200MG | A075661 | 001 | Dec 12, 2001 |
| DR REDDYS LABS INC | 200MG | A076117 | 001 | Nov 20, 2001 |
| GRANULES INDIA | 200MG | A079174 | 001 | Dec 10, 2010 |
| GRANULES INDIA LTD | 200MG | A202312 | 001 | Oct 07, 2016 |
| LNK | 200MG | A075010 | 001 | Mar 01, 1999 |
| | 200MG | A075139 | 001 | Mar 01, 1999 |
| MARKSANS PHARMA | 200MG | A091237 | 001 | Feb 08, 2011 |
| | 200MG | A091239 | 001 | Feb 01, 2011 |
| MCNEIL | 200MG | A073019 | 001 | Mar 30, 1994 |
| MERRO PHARM | 200MG | A070985 | 001 | Oct 02, 1987 |
| OHM | 200MG | A071163 | 001 | Jul 15, 1986 |
| PAR PHARM | 200MG | A070481 | 001 | Sep 24, 1986 |
| PERRIGO | 200MG | A072096 | 001 | Dec 08, 1987 |
| | 200MG | A075995 | 001 | Mar 14, 2002 |
| PERRIGO R AND D | 200MG | A077349 | 001 | Jun 21, 2005 |
| SHANDONG XINHUA | 200MG | A206990 | 001 | Aug 21, 2018 |
| | 200MG | A207095 | 001 | May 05, 2017 |
| STRIDES PHARMA | 200MG | A079129 | 001 | Mar 28, 2011 |
| | 200MG | A091355 | 001 | Apr 04, 2011 |
| | 200MG | A206989 | 001 | Jun 29, 2018 |
| | 200MG | A207052 | 001 | May 30, 2017 |
| VINTAGE PHARMS | 200MG | A071229 | 001 | Apr 01, 1987 |
| | 200MG | A071639 | 001 | Feb 02, 1988 |

IBUPROHM

OHM LABS 200MG A071214 001 Dec 01, 1986

JUNIOR STRENGTH ADVIL

PFIZER 100MG N020267 002 Dec 13, 1996

JUNIOR STRENGTH IBUPROFEN

L PERRIGO CO 100MG A075367 001 Apr 22, 1999

JUNIOR STRENGTH MOTRIN

J AND J CONSUMER INC 100MG N020602 001 Jun 10, 1996

MOTRIN IB

+! J AND J CONSUMER INC 200MG N019012 003 Dec 17, 1990

TAB-PROFEN

PERRIGO 200MG A072095 001 Dec 08, 1987

TABLET, CHEWABLE;ORAL

CHILDREN'S ADVIL

PFIZER 50MG N020944 001 Dec 18, 1998

IBUPROFEN

! PERRIGO 100MG A076359 002 Jan 16, 2004

JUNIOR STRENGTH ADVIL

PFIZER 100MG N020944 002 Dec 18, 1998

IBUPROFEN SODIUM

TABLET;ORAL

ADVIL

+! PFIZER CONS EQ 200MG BASE N201803 001 Jun 12, 2012
HLTHCARE

IBUPROFEN SODIUM

PERRIGO R AND D EQ 200MG BASE A206581 001 Aug 03, 2015

IBUPROFEN; PHENYLEPHRINE HYDROCHLORIDE

TABLET;ORAL

ADVIL CONGESTION RELIEF

+! PFIZER 200MG;10MG N022565 001 May 27, 2010

IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE

PERRIGO R AND D 200MG;10MG A203200 001 Jul 03, 2014

OTC DRUG PRODUCT LIST

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE; ORAL

ADVIL COLD AND SINUS

| | | | | | | |
|---|---|--------|------------------------------------------------|---------|-----|--------------|
| + | ! | PFIZER | EQ 200MG FREE ACID AND POTASSIUM SALT; 30MG | N021374 | 001 | May 30, 2002 |
|---|---|--------|------------------------------------------------|---------|-----|--------------|

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

| | | | | | | |
|--|--|-------------------------|------------------------------------------------|---------|-----|--------------|
| | | AUROBINDO PHARMA LTD | EQ 200MG FREE ACID AND POTASSIUM SALT; 30MG | A209235 | 001 | Dec 01, 2017 |
|--|--|-------------------------|------------------------------------------------|---------|-----|--------------|

SUSPENSION; ORAL

CHILDREN'S ADVIL COLD

| | | | | | | |
|--|--|--------|---------------------|---------|-----|--------------|
| | | PFIZER | 100MG/5ML; 15MG/5ML | N021373 | 001 | Apr 18, 2002 |
|--|--|--------|---------------------|---------|-----|--------------|

CHILDREN'S MOTRIN COLD

| | | | | | | |
|---|---|-------------------------|---------------------|---------|-----|--------------|
| + | ! | J AND J CONSUMER INC | 100MG/5ML; 15MG/5ML | N021128 | 001 | Aug 01, 2000 |
|---|---|-------------------------|---------------------|---------|-----|--------------|

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

| | | | | | | |
|--|--|---------|---------------------|---------|-----|--------------|
| | | PERRIGO | 100MG/5ML; 15MG/5ML | A076478 | 001 | Nov 05, 2003 |
|--|--|---------|---------------------|---------|-----|--------------|

TABLET; ORAL

ADVIL COLD AND SINUS

| | | | | | | |
|---|---|--------|-------------|---------|-----|--------------|
| + | ! | PFIZER | 200MG; 30MG | N019771 | 001 | Sep 19, 1989 |
|---|---|--------|-------------|---------|-----|--------------|

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

| | | | | | | |
|--|--|--------------------|-------------|---------|-----|--------------|
| | | DR REDDYS LABS LTD | 200MG; 30MG | A077628 | 001 | Aug 14, 2006 |
|--|--|--------------------|-------------|---------|-----|--------------|

IBUPROHM COLD AND SINUS

| | | | | | | |
|--|--|----------|-------------|---------|-----|--------------|
| | | OHM LABS | 200MG; 30MG | A074567 | 001 | Apr 17, 2001 |
|--|--|----------|-------------|---------|-----|--------------|

SINE-AID IB

| | | | | | | |
|--|--|-------------------------|-------------|---------|-----|--------------|
| | | J AND J CONSUMER INC | 200MG; 30MG | N019899 | 001 | Dec 31, 1992 |
|--|--|-------------------------|-------------|---------|-----|--------------|

INSULIN RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN R PEN

| | | | | | | |
|---|---|-------|--------------|---------|-----|--------------|
| + | ! | LILLY | 100 UNITS/ML | N018780 | 005 | Aug 06, 1998 |
|---|---|-------|--------------|---------|-----|--------------|

NOVOLIN R

| | | | | | | |
|---|---|------------------|--------------|---------|-----|--------------|
| + | ! | NOVO NORDISK INC | 100 UNITS/ML | N019938 | 001 | Jun 25, 1991 |
|---|---|------------------|--------------|---------|-----|--------------|

INSULIN RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN R

| | | | | | | |
|---|---|-------|--------------|---------|-----|--------------|
| + | ! | LILLY | 100 UNITS/ML | N018780 | 001 | Oct 28, 1982 |
|---|---|-------|--------------|---------|-----|--------------|

INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN 70/30

| | | | | | | |
|---|---|-------|--------------------------|---------|-----|--------------|
| + | ! | LILLY | 30 UNITS/ML; 70 UNITS/ML | N019717 | 001 | Apr 25, 1989 |
|---|---|-------|--------------------------|---------|-----|--------------|

HUMULIN 70/30 PEN

| | | | | | | |
|---|---|-------|--------------------------|---------|-----|--------------|
| + | ! | LILLY | 30 UNITS/ML; 70 UNITS/ML | N019717 | 002 | Aug 06, 1998 |
|---|---|-------|--------------------------|---------|-----|--------------|

NOVOLIN 70/30

| | | | | | | |
|---|---|------------------|--------------------------|---------|-----|--------------|
| + | ! | NOVO NORDISK INC | 30 UNITS/ML; 70 UNITS/ML | N019991 | 001 | Jun 25, 1991 |
|---|---|------------------|--------------------------|---------|-----|--------------|

INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN N

| | | | | | | |
|---|---|-------|--------------|---------|-----|--------------|
| + | ! | LILLY | 100 UNITS/ML | N018781 | 001 | Oct 28, 1982 |
|---|---|-------|--------------|---------|-----|--------------|

NOVOLIN N

| | | | | | | |
|---|---|------------------|--------------|---------|-----|--------------|
| + | ! | NOVO NORDISK INC | 100 UNITS/ML | N019959 | 001 | Jul 01, 1991 |
|---|---|------------------|--------------|---------|-----|--------------|

IODINE POVACRYLEX; ISOPROPYL ALCOHOL

SPONGE; TOPICAL

DURAPREP

| | | | | | | |
|---|---|----|---------------------------|---------|-----|--------------|
| + | ! | 3M | EQ 0.7% IODINE; 74% (6ML) | N021586 | 001 | Sep 29, 2006 |
|---|---|----|---------------------------|---------|-----|--------------|

| | | | | | | |
|---|---|--|----------------------------|---------|-----|--------------|
| + | ! | | EQ 0.7% IODINE; 74% (26ML) | N021586 | 002 | Sep 29, 2006 |
|---|---|--|----------------------------|---------|-----|--------------|

KETOCONAZOLE

SHAMPOO; TOPICAL

NIZORAL A-D

| | | | | | | |
|---|---|---------------------|----|---------|-----|--------------|
| + | ! | JOHNSON AND JOHNSON | 1% | N020310 | 001 | Oct 10, 1997 |
|---|---|---------------------|----|---------|-----|--------------|

KETOTIFEN FUMARATE

SOLUTION/DROPS; OPHTHALMIC

ALAWAY

| | | | | | | |
|---|---|-----------------|----------------|---------|-----|--------------|
| + | ! | BAUSCH AND LOMB | EQ 0.025% BASE | N021996 | 001 | Dec 01, 2006 |
|---|---|-----------------|----------------|---------|-----|--------------|

| | | | | | | |
|---|--|--|----------------|---------|-----|--------------|
| + | | | EQ 0.035% BASE | N021996 | 002 | Feb 11, 2015 |
|---|--|--|----------------|---------|-----|--------------|

KETOTIFEN FUMARATE

| | | | | | | |
|--|--|-------|----------------|---------|-----|--------------|
| | | AKORN | EQ 0.025% BASE | A077958 | 001 | Jul 26, 2007 |
|--|--|-------|----------------|---------|-----|--------------|

| | | | | | | |
|---|--|------------------|----------------|---------|-----|--------------|
| ! | | ALCON PHARMS LTD | EQ 0.025% BASE | A077200 | 001 | Sep 02, 2008 |
|---|--|------------------|----------------|---------|-----|--------------|

OTC DRUG PRODUCT LIST

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

| | | | | |
|--------------------|------|---------|-----|--------------|
| DR REDDYS LABS LTD | 15MG | A202194 | 001 | May 18, 2012 |
| LANNETT CO INC | 15MG | A207157 | 001 | Sep 29, 2017 |
| MYLAN PHARMS INC | 15MG | A203187 | 001 | Jun 01, 2016 |
| NATCO PHARMA LTD | 15MG | A203306 | 001 | Jan 13, 2016 |
| PERRIGO R AND D | 15MG | A202319 | 001 | May 18, 2012 |
| WOCKHARDT LTD | 15MG | A202727 | 001 | May 18, 2012 |

PREVACID 24 HR

| | | | | |
|--------------------|------|---------|-----|--------------|
| +! GLAXOSMITHKLINE | 15MG | N022327 | 001 | May 18, 2009 |
|--------------------|------|---------|-----|--------------|

CONS

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

LANSOPRAZOLE

| | | | | |
|---------------|------|---------|-----|--------------|
| DEXCEL PHARMA | 15MG | N208025 | 001 | Jun 07, 2016 |
|---------------|------|---------|-----|--------------|

LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION;ORAL

XYZAL ALLERGY 24HR

| | | | | |
|----------------------|-----------|---------|-----|--------------|
| +! SANOFI AVENTIS US | 2.5MG/5ML | N209090 | 001 | Jan 31, 2017 |
|----------------------|-----------|---------|-----|--------------|

TABLET;ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

| | | | | |
|--------------------|-----|---------|-----|--------------|
| DR REDDYS LABS LTD | 5MG | A210375 | 001 | Jan 19, 2018 |
| MICRO LABS | 5MG | A211551 | 001 | Nov 20, 2018 |

XYZAL ALLERGY 24HR

| | | | | |
|----------------------|-----|---------|-----|--------------|
| +! SANOFI AVENTIS US | 5MG | N209089 | 001 | Jan 31, 2017 |
|----------------------|-----|---------|-----|--------------|

LEVONORGESTREL

TABLET;ORAL

ATHENTIA NEXT

| | | | | |
|----------------------|-------|---------|-----|--------------|
| AUROBINDO PHARMA LTD | 1.5MG | A206867 | 001 | Dec 08, 2015 |
|----------------------|-------|---------|-----|--------------|

FALLBACK SOLO

| | | | | |
|-----------|-------|---------|-----|--------------|
| LUPIN LTD | 1.5MG | A201446 | 001 | Jun 19, 2014 |
|-----------|-------|---------|-----|--------------|

HER STYLE

| | | | | |
|-------------|-------|---------|-----|--------------|
| NOVAST LABS | 1.5MG | A207976 | 001 | Mar 11, 2016 |
|-------------|-------|---------|-----|--------------|

LEVONORGESTREL

| | | | | |
|---------------------|--------|---------|-----|--------------|
| AMNEAL PHARMS | 1.5MG | A204044 | 001 | Jul 03, 2018 |
| APOTEX INC | 1.5MG | A205329 | 001 | Sep 18, 2018 |
| FDN CONSUMER | 1.5MG | A200670 | 001 | Jul 12, 2012 |
| GLENMARK PHARMS LTD | 1.5MG | A207044 | 001 | Mar 25, 2016 |
| LOTUS PHARM CO LTD | 0.75MG | A202684 | 001 | Sep 02, 2016 |
| MYLAN LABS LTD | 0.75MG | A202740 | 001 | Sep 02, 2016 |
| | 1.5MG | A202739 | 001 | Oct 31, 2014 |
| NOVEL LABS INC | 1.5MG | A202508 | 001 | Feb 22, 2013 |
| OC PHARMA | 1.5MG | A202380 | 001 | May 29, 2015 |
| ! PERRIGO R AND D | 0.75MG | A090740 | 001 | Dec 30, 2010 |
| | 1.5MG | A202334 | 001 | Aug 20, 2014 |
| RECKITT BENCKISER | 1.5MG | A202246 | 001 | Jun 05, 2015 |

OPCICON ONE-STEP

| | | | | |
|--------------------|-------|---------|-----|--------------|
| SUN PHARM INDS LTD | 1.5MG | A202635 | 001 | Sep 11, 2014 |
|--------------------|-------|---------|-----|--------------|

PLAN B ONE-STEP

| | | | | |
|-----------------|-------|---------|-----|--------------|
| +! FDN CONSUMER | 1.5MG | N021998 | 001 | Jul 10, 2009 |
|-----------------|-------|---------|-----|--------------|

LOPERAMIDE HYDROCHLORIDE

CAPSULE;ORAL

LOPERAMIDE HYDROCHLORIDE

| | | | | |
|------------------|-----|---------|-----|--------------|
| + BIONPHARMA INC | 1MG | N021855 | 001 | Aug 04, 2005 |
| +! | 2MG | N021855 | 002 | Aug 04, 2005 |

SOLUTION;ORAL

IMODIUM A-D

| | | | | |
|-------------------------|---------|---------|-----|--------------|
| +! J AND J CONSUMER INC | 1MG/5ML | N019487 | 001 | Mar 01, 1988 |
|-------------------------|---------|---------|-----|--------------|

LOPERAMIDE HYDROCHLORIDE

| | | | | |
|------------------|---------|---------|-----|--------------|
| HI TECH PHARMA | 1MG/5ML | A074352 | 001 | Nov 17, 1995 |
| PERRIGO | 1MG/5ML | A073243 | 001 | Jan 21, 1992 |
| WOCKHARDT BIO AG | 1MG/5ML | A074730 | 001 | Aug 28, 1997 |

SUSPENSION;ORAL

IMODIUM A-D

| | | | | |
|-------------------------|-----------|---------|-----|--------------|
| +! J AND J CONSUMER INC | 1MG/7.5ML | N019487 | 002 | Jul 08, 2004 |
|-------------------------|-----------|---------|-----|--------------|

LOPERAMIDE HYDROCHLORIDE

| | | | | |
|-----------------|-----------|---------|-----|--------------|
| PERRIGO R AND D | 1MG/7.5ML | A091292 | 001 | May 20, 2011 |
|-----------------|-----------|---------|-----|--------------|

OTC DRUG PRODUCT LIST

LOPERAMIDE HYDROCHLORIDE

TABLET;ORAL

IMODIUM A-D

| | | | | | |
|----|-------------------------|-----|---------|-----|--------------|
| +! | J AND J CONSUMER INC | 2MG | N019860 | 001 | Nov 22, 1989 |
|----|-------------------------|-----|---------|-----|--------------|

LOPERAMIDE HYDROCHLORIDE

| | | | | | |
|--|-------------------------|-----|---------|-----|--------------|
| | AUROBINDO PHARMA LTD | 2MG | A206548 | 001 | Dec 15, 2015 |
|--|-------------------------|-----|---------|-----|--------------|

| | | | | | |
|--|--------------|-----|---------|-----|--------------|
| | L PERRIGO CO | 2MG | A075232 | 001 | Jan 06, 2000 |
|--|--------------|-----|---------|-----|--------------|

| | | | | | |
|--|-----|-----|---------|-----|--------------|
| | LNK | 2MG | A076497 | 001 | Jun 10, 2003 |
|--|-----|-----|---------|-----|--------------|

| | | | | | |
|--|----------|-----|---------|-----|--------------|
| | OHM LABS | 2MG | A074091 | 001 | Dec 10, 1992 |
|--|----------|-----|---------|-----|--------------|

LOPERAMIDE HYDROCHLORIDE; SIMETHICONE

TABLET;ORAL

IMODIUM MULTI-SYMPOM RELIEF

| | | | | | |
|----|-------------------------|-----------|---------|-----|--------------|
| +! | J AND J CONSUMER INC | 2MG;125MG | N021140 | 001 | Nov 30, 2000 |
|----|-------------------------|-----------|---------|-----|--------------|

LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE

| | | | | | |
|--|-----------------|-----------|---------|-----|--------------|
| | PERRIGO R AND D | 2MG;125MG | A209837 | 001 | Sep 05, 2018 |
|--|-----------------|-----------|---------|-----|--------------|

| | | | | | |
|--|--------------------|-----------|---------|-----|--------------|
| | SUN PHARM INDS LTD | 2MG;125MG | A077500 | 001 | Sep 06, 2006 |
|--|--------------------|-----------|---------|-----|--------------|

TABLET, CHEWABLE;ORAL

LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE

| | | | | | |
|---|---------|-----------|---------|-----|--------------|
| ! | PERRIGO | 2MG;125MG | A076029 | 001 | Aug 30, 2002 |
|---|---------|-----------|---------|-----|--------------|

LORATADINE

CAPSULE;ORAL

CLARITIN

| | | | | | |
|----|-------------------------|------|---------|-----|--------------|
| +! | BAYER HEALTHCARE LLC | 10MG | N021952 | 001 | Jun 16, 2008 |
|----|-------------------------|------|---------|-----|--------------|

LORATADINE

| | | | | | |
|--|----------------|------|---------|-----|--------------|
| | BIONPHARMA INC | 10MG | A202538 | 001 | Dec 21, 2018 |
|--|----------------|------|---------|-----|--------------|

| | | | | | |
|--|-----------------|------|---------|-----|--------------|
| | MARKSANS PHARMA | 10MG | A206214 | 001 | Sep 23, 2016 |
|--|-----------------|------|---------|-----|--------------|

SUSPENSION;ORAL

LORATADINE

| | | | | | |
|----|------|--------|---------|-----|--------------|
| +! | TARO | 1MG/ML | N021734 | 001 | Oct 04, 2005 |
|----|------|--------|---------|-----|--------------|

SYRUP;ORAL

CLARITIN

| | | | | | |
|----|-------------------------|--------|---------|-----|--------------|
| +! | BAYER HEALTHCARE LLC | 1MG/ML | N020641 | 002 | Nov 27, 2002 |
|----|-------------------------|--------|---------|-----|--------------|

LORATADINE

| | | | | | |
|--|-------------------------|--------|---------|-----|--------------|
| | AUROBINDO PHARMA LTD | 1MG/ML | A208931 | 001 | Jun 29, 2018 |
|--|-------------------------|--------|---------|-----|--------------|

| | | | | | |
|--|----------------|--------|---------|-----|--------------|
| | LANNETT CO INC | 1MG/ML | A077421 | 001 | Jun 29, 2006 |
|--|----------------|--------|---------|-----|--------------|

| | | | | | |
|--|---------|--------|---------|-----|--------------|
| | PERRIGO | 1MG/ML | A075728 | 001 | Aug 20, 2004 |
|--|---------|--------|---------|-----|--------------|

| | | | | | |
|--|------|--------|---------|-----|--------------|
| | TARO | 1MG/ML | A076805 | 001 | Aug 20, 2004 |
|--|------|--------|---------|-----|--------------|

| | | | | | |
|--|------------|--------|---------|-----|--------------|
| | TARO PHARM | 1MG/ML | A201865 | 001 | Jul 31, 2015 |
|--|------------|--------|---------|-----|--------------|

| | | | | | |
|--|------|--------|---------|-----|--------------|
| | TEVA | 1MG/ML | A075505 | 001 | Nov 07, 2003 |
|--|------|--------|---------|-----|--------------|

| | | | | | |
|--|------------------|--------|---------|-----|--------------|
| | WOCKHARDT BIO AG | 1MG/ML | A075815 | 001 | Aug 20, 2004 |
|--|------------------|--------|---------|-----|--------------|

TABLET;ORAL

CLARITIN

| | | | | | |
|----|-------------------------|------|---------|-----|--------------|
| +! | BAYER HEALTHCARE LLC | 10MG | N019658 | 002 | Nov 27, 2002 |
|----|-------------------------|------|---------|-----|--------------|

CLARITIN HIVES RELIEF

| | | | | | |
|----|-------------------------|------|---------|-----|--------------|
| +! | BAYER HEALTHCARE LLC | 10MG | N019658 | 003 | Nov 19, 2003 |
|----|-------------------------|------|---------|-----|--------------|

LORATADINE

| | | | | | |
|--|------------|------|---------|-----|--------------|
| | APOTEX INC | 10MG | A076471 | 001 | Feb 14, 2006 |
|--|------------|------|---------|-----|--------------|

| | | | | | |
|--|-------------------------|------|---------|-----|--------------|
| | AUROBINDO PHARMA LTD | 10MG | A208314 | 001 | Apr 16, 2018 |
|--|-------------------------|------|---------|-----|--------------|

| | | | | | |
|--|-------|------|---------|-----|--------------|
| | MYLAN | 10MG | A075790 | 001 | Nov 07, 2008 |
|--|-------|------|---------|-----|--------------|

| | | | | | |
|--|--|------|---------|-----|--------------|
| | | 10MG | A076154 | 001 | Aug 20, 2003 |
|--|--|------|---------|-----|--------------|

| | | | | | |
|--|--|------|---------|-----|--------------|
| | | 10MG | A078447 | 001 | Aug 12, 2011 |
|--|--|------|---------|-----|--------------|

| | | | | | |
|--|---------|------|---------|-----|--------------|
| | PERRIGO | 10MG | A076301 | 001 | Jun 25, 2004 |
|--|---------|------|---------|-----|--------------|

| | | | | | |
|--|-------------------------|------|---------|-----|--------------|
| | PLD ACQUISITIONS LLC | 10MG | A075209 | 001 | Jan 21, 2003 |
|--|-------------------------|------|---------|-----|--------------|

| | | | | | |
|--|--------------------|------|---------|-----|--------------|
| | SUN PHARM INDS LTD | 10MG | A076134 | 001 | Aug 18, 2003 |
|--|--------------------|------|---------|-----|--------------|

TABLET, CHEWABLE;ORAL

CHILDREN'S CLARITIN

| | | | | | |
|----|-------------------------|-----|---------|-----|--------------|
| +! | BAYER HEALTHCARE LLC | 5MG | N021891 | 001 | Aug 23, 2006 |
|----|-------------------------|-----|---------|-----|--------------|

CLARITIN

| | | | | | |
|---|-------------------------|------|---------|-----|--------------|
| + | BAYER HEALTHCARE LLC | 10MG | N021891 | 002 | Nov 21, 2018 |
|---|-------------------------|------|---------|-----|--------------|

LORATADINE

| | | | | | |
|--|-------------------|-----|---------|-----|--------------|
| | SUN PHARMA GLOBAL | 5MG | A210088 | 001 | Apr 16, 2018 |
|--|-------------------|-----|---------|-----|--------------|

OTC DRUG PRODUCT LIST

LORATADINE

TABLET, ORALLY DISINTEGRATING;ORAL

ALAVERT

PFIZER 10MG N021375 001 Dec 19, 2002

CLARITIN HIVES RELIEF REDITAB

+! BAYER HEALTHCARE 10MG N020704 003 Nov 19, 2003
LLC

CLARITIN REDITABS

+! BAYER HEALTHCARE 5MG N021993 001 Dec 12, 2006
LLC

+! 10MG N020704 002 Nov 27, 2002

LORATADINE

ACTAVIS LABS FL INC 10MG A075990 001 Nov 03, 2003

AUROBINDO PHARMA 10MG A208477 001 Apr 11, 2018

LTD

PERRIGO PHARMA INTL 10MG A076011 001 Sep 29, 2003

PFIZER 10MG A075822 001 Feb 10, 2003

LORATADINE REDIDOSE

SUN PHARM INDS LTD 10MG A077153 001 Apr 11, 2007

LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

CLARITIN-D

+! BAYER HEALTHCARE 5MG;120MG N019670 002 Nov 27, 2002
LLC

CLARITIN-D 24 HOUR

+! BAYER HEALTHCARE 10MG;240MG N020470 002 Nov 27, 2002
LLC

LORATADINE AND PSEUDOEPHEDRINE SULFATE

ACTAVIS LABS FL INC 10MG;240MG A075706 001 Feb 21, 2003

PERRIGO PHARMA INTL 5MG;120MG A076050 001 Jan 30, 2003

10MG;240MG A075989 001 Mar 04, 2004

SUN PHARM INDS LTD 10MG;240MG A076557 001 Sep 22, 2004

MENTHOL; METHYL SALICYLATE

PATCH;TOPICAL

SALONPAS

+! HISAMITSU PHARM CO 3%;10% N022029 001 Feb 20, 2008

+ 3%;10% N022029 002 Nov 05, 2012

MICONAZOLE NITRATE

CREAM;TOPICAL, VAGINAL

MICONAZOLE 3 COMBINATION PACK

PERRIGO 2%,4% A076357 001 Mar 30, 2004

MONISTAT 3 COMBINATION PACK

+ MEDTECH PRODUCTS 2%,4% N021261 003 Jun 17, 2003

MONISTAT 3 COMBINATION PACK (PREFILLED)

+! MEDTECH PRODUCTS 2%,4% N021261 001 Feb 02, 2001

CREAM;VAGINAL

MICONAZOLE 3

TARO 4% A076773 001 Mar 02, 2005

MICONAZOLE 7

ACTAVIS MID 2% A074164 001 Mar 29, 1996

ATLANTIC

MICONAZOLE NITRATE

G AND W LABS INC 2% A074366 001 Feb 22, 1996

PERRIGO 2% A074760 001 May 15, 1997

PERRIGO R AND D 4% A091366 001 Jan 15, 2010

TARO PHARMS 2% A074444 001 Jan 13, 1997

MONISTAT 3

+! MEDTECH PRODUCTS 4% N020827 001 Mar 30, 1998

MONISTAT 7

+! MEDTECH PRODUCTS 2% N017450 002 Feb 15, 1991

CREAM, SUPPOSITORY;TOPICAL, VAGINAL

M-ZOLE 3 COMBINATION PACK

ACTAVIS MID 2%,200MG A074926 001 Apr 16, 1999

ATLANTIC

MICONAZOLE NITRATE

PERRIGO R AND D 2%,1.2GM A079114 001 Jun 02, 2010

MICONAZOLE NITRATE COMBINATION PACK

PERRIGO 2%,200MG A075329 001 Apr 20, 1999

MONISTAT 1 COMBINATION PACK

+! MEDTECH PRODUCTS 2%,1.2GM N021308 001 Jun 29, 2001

MONISTAT 3 COMBINATION PACK

+! MEDTECH PRODUCTS 2%,200MG N020670 002 Apr 16, 1996

OTC DRUG PRODUCT LIST

MICONAZOLE NITRATE

CREAM, SUPPOSITORY;TOPICAL, VAGINAL

MONISTAT 7 COMBINATION PACK

+! MEDTECH PRODUCTS 2%,100MG N020288 002 Apr 26, 1993

SUPPOSITORY;VAGINAL

MICONAZOLE NITRATE

ACTAVIS PHARMA 100MG A073507 001 Nov 19, 1993

G AND W LABS 100MG A074414 001 Apr 30, 1997

! PERRIGO 100MG A074395 001 Mar 20, 1997

MONISTAT 7

+! MEDTECH PRODUCTS 100MG N018520 002 Feb 15, 1991

MINOXIDIL

AEROSOL, FOAM;TOPICAL

MEN'S ROGAINE

+! JOHNSON AND JOHNSON 5% N021812 001 Jan 20, 2006

MINOXIDIL

PERRIGO ISRAEL 5% A091344 001 Apr 28, 2011

MINOXIDIL (FOR MEN)

TARO PHARM 5% A209074 001 Dec 31, 2018

WATSON LABS INC 5% A208092 001 Feb 17, 2017

MINOXIDIL (FOR WOMEN)

WATSON LABS INC 5% A208092 002 Jul 27, 2017

WOMEN'S ROGAINE

+! JOHNSON AND JOHNSON 5% N021812 002 Feb 28, 2014

SOLUTION;TOPICAL

MINOXIDIL (FOR MEN)

ACTAVIS MID 2% A074588 001 Apr 05, 1996

ATLANTIC

HI TECH PHARMA 2% A074731 001 Dec 24, 1996

L PERRIGO CO 2% A075357 001 Jul 30, 1999

WOCKHARDT BIO AG 2% A074767 001 Feb 28, 1997

MINOXIDIL (FOR WOMEN)

HI TECH PHARMA 2% A074731 002 May 11, 2005

L PERRIGO CO 2% A075357 002 Jul 30, 1999

MINOXIDIL EXTRA STRENGTH (FOR MEN)

ACTAVIS MID 5% A075518 001 Nov 17, 2000

ATLANTIC

AVACOR PRODS 5% A075619 001 Nov 17, 2000

PERRIGO 5% A075598 001 Jun 13, 2001

PERRIGO NEW YORK 5% A075737 001 Mar 15, 2002

WOCKHARDT BIO AG 5% A075438 001 Feb 27, 2003

ROGAINE (FOR MEN)

+! JOHNSON AND JOHNSON 2% N019501 002 Feb 09, 1996

ROGAINE (FOR WOMEN)

+! JOHNSON AND JOHNSON 2% N019501 003 Feb 09, 1996

ROGAINE EXTRA STRENGTH (FOR MEN)

+! JOHNSON AND JOHNSON 5% N020834 001 Nov 14, 1997

THEROXIDIL

EI INC 2% A078176 001 Nov 09, 2007

5% A076239 001 Aug 24, 2004

NAPHAZOLINE HYDROCHLORIDE; PHENIRAMINE MALEATE

SOLUTION/DROPS;OPHTHALMIC

NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE

AKORN INC 0.025%;0.3% A202795 001 Jan 24, 2013

ALTAIRE PHARMS INC 0.02675%;0.315% A078208 001 Sep 27, 2010

NAPHCN-A

+! ALCON 0.025%;0.3% N020226 001 Jun 08, 1994

OPCON-A

+! BAUSCH AND LOMB 0.02675%;0.315% N020065 001 Jun 08, 1994

VISINE

+! JOHNSON AND JOHNSON 0.025%;0.3% N020485 001 Jan 31, 1996

NAPROXEN SODIUM

CAPSULE;ORAL

NAPROXEN SODIUM

+! BIONPHARMA INC EQ 200MG BASE N021920 001 Feb 17, 2006

CATALENT EQ 200MG BASE A202807 001 Jan 04, 2019

PURACAP PHARM LLC EQ 200MG BASE A208363 001 Mar 15, 2018

TABLET;ORAL

ALEVE

+! BAYER EQ 200MG BASE N020204 002 Jan 11, 1994

OTC DRUG PRODUCT LIST

NAPROXEN SODIUM

TABLET;ORAL

NAPROXEN SODIUM

| | | | | |
|----------------------|---------------|---------|-----|--------------|
| AMNEAL PHARMS NY | EQ 200MG BASE | A079096 | 001 | Dec 16, 2008 |
| AUROBINDO PHARMA LTD | EQ 200MG BASE | A205497 | 001 | Mar 18, 2016 |
| CONTRACT PHARMACAL | EQ 200MG BASE | A074635 | 001 | Jan 13, 1997 |
| DR REDDYS LABS INC | EQ 200MG BASE | A075168 | 001 | Jul 28, 1998 |
| GRANULES INDIA | EQ 200MG BASE | A091353 | 001 | Sep 20, 2011 |
| LNK INTL INC | EQ 200MG BASE | A204872 | 001 | Jan 23, 2017 |
| MARKSANS PHARMA | EQ 200MG BASE | A090545 | 001 | Mar 16, 2011 |
| NOVELGENIX THERAPS | EQ 200MG BASE | A207612 | 001 | Nov 16, 2018 |
| PERRIGO | EQ 200MG BASE | A074661 | 001 | Jan 13, 1997 |
| SUN PHARM INDS LTD | EQ 200MG BASE | A091183 | 001 | May 20, 2011 |

NAPROXEN SODIUM; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

ALEVE-D SINUS & COLD

| | | | | |
|----------|-------------|---------|-----|--------------|
| +! BAYER | 200MG;120MG | N021076 | 001 | Nov 29, 1999 |
|----------|-------------|---------|-----|--------------|

NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE

| | | | | |
|--------------------|---------------------|---------|-----|--------------|
| DR REDDYS LABS INC | EQ 220MG BASE;120MG | A077381 | 001 | Sep 27, 2006 |
| PERRIGO | EQ 200MG BASE;120MG | A076518 | 001 | Mar 17, 2004 |

NICOTINE

FILM, EXTENDED RELEASE;TRANSDERMAL

HABITROL

| | | | | |
|---------------------|-----------|---------|-----|--------------|
| + DR REDDYS LABS SA | 7MG/24HR | N020076 | 004 | Nov 12, 1999 |
| + | 14MG/24HR | N020076 | 005 | Nov 12, 1999 |
| +! | 21MG/24HR | N020076 | 006 | Nov 12, 1999 |

NICODERM CQ

| | | | | |
|---------------------|-----------|---------|-----|--------------|
| + SANOFI AVENTIS US | 7MG/24HR | N020165 | 006 | Aug 02, 1996 |
| + | 14MG/24HR | N020165 | 005 | Aug 02, 1996 |
| +! | 21MG/24HR | N020165 | 004 | Aug 02, 1996 |

NICOTINE

| | | | | |
|-------|-----------|---------|-----|--------------|
| AVEVA | 7MG/24HR | A074612 | 002 | Jul 28, 2003 |
| | 14MG/24HR | A074612 | 003 | Oct 20, 1997 |
| | 21MG/24HR | A074612 | 001 | Oct 20, 1997 |

NICOTINE POLACRILEX

GUM, CHEWING;BUCCAL

NICORETTE

| | | | | |
|-------------------|-------------|---------|-----|--------------|
| + GLAXOSMITHKLINE | EQ 2MG BASE | N018612 | 002 | Feb 09, 1996 |
| + | EQ 2MG BASE | N018612 | 004 | Sep 25, 2000 |
| +! | EQ 4MG BASE | N020066 | 002 | Feb 09, 1996 |
| + | EQ 4MG BASE | N020066 | 004 | Sep 25, 2000 |

NICORETTE (MINT)

| | | | | |
|-------------------|-------------|---------|-----|--------------|
| + GLAXOSMITHKLINE | EQ 2MG BASE | N018612 | 003 | Dec 23, 1998 |
| + | EQ 4MG BASE | N020066 | 003 | Dec 23, 1998 |

NICOTINE POLACRILEX

| | | | | |
|---------------------|-------------|---------|-----|--------------|
| ACTAVIS LABS NY INC | EQ 2MG BASE | A074507 | 001 | Mar 15, 1999 |
| | EQ 2MG BASE | A076569 | 001 | Jul 29, 2004 |
| | EQ 2MG BASE | A078699 | 001 | Dec 29, 2008 |
| | EQ 2MG BASE | A079216 | 001 | Jul 08, 2009 |
| | EQ 2MG BASE | A204794 | 001 | May 10, 2016 |
| | EQ 4MG BASE | A074707 | 001 | Mar 19, 1999 |
| | EQ 4MG BASE | A076568 | 002 | Jul 29, 2004 |
| | EQ 4MG BASE | A078697 | 001 | Dec 29, 2008 |
| | EQ 4MG BASE | A079038 | 001 | Jul 08, 2009 |
| | EQ 4MG BASE | A079219 | 001 | Jul 08, 2009 |
| | EQ 4MG BASE | A204833 | 001 | Feb 26, 2016 |
| L PERRIGO CO | EQ 2MG BASE | A076775 | 001 | Sep 16, 2004 |
| | EQ 2MG BASE | A076776 | 001 | Sep 16, 2004 |
| | EQ 2MG BASE | A076777 | 001 | Sep 16, 2004 |
| | EQ 4MG BASE | A076778 | 001 | Sep 16, 2004 |
| | EQ 4MG BASE | A076779 | 001 | Sep 16, 2004 |
| | EQ 4MG BASE | A076789 | 001 | Sep 16, 2004 |
| PERRIGO R AND D | EQ 2MG BASE | A078325 | 001 | Oct 30, 2006 |
| | EQ 2MG BASE | A078547 | 001 | May 24, 2007 |
| | EQ 2MG BASE | A078967 | 001 | Apr 23, 2008 |
| | EQ 2MG BASE | A091349 | 001 | Jul 20, 2011 |
| | EQ 2MG BASE | A206394 | 001 | Dec 15, 2016 |
| | EQ 4MG BASE | A078326 | 001 | Oct 30, 2006 |
| | EQ 4MG BASE | A078546 | 001 | May 24, 2007 |
| | EQ 4MG BASE | A078968 | 001 | Apr 23, 2008 |

OTC DRUG PRODUCT LIST

NICOTINE POLACRILEX

GUM, CHEWING;BUCCAL

NICOTINE POLACRILEX

EQ 4MG BASE

A091354 001 Jul 20, 2011

EQ 4MG BASE

A206393 001 Dec 15, 2016

WATSON LABS

EQ 2MG BASE

A079044 001 Jul 08, 2009

TROCHE/LOZENGE;ORAL

NICORETTE

+ GLAXOSMITHKLINE
CONS

EQ 2MG BASE

N021330 001 Oct 31, 2002

+

EQ 2MG BASE

N022360 001 May 18, 2009

+!

EQ 4MG BASE

N021330 002 Oct 31, 2002

+!

EQ 4MG BASE

N022360 002 May 18, 2009

NICOTINE POLACRILEX

PERRIGO R AND D

EQ 2MG BASE

A077007 001 Jan 31, 2006

EQ 2MG BASE

A090711 001 Jul 10, 2009

EQ 2MG BASE

A090821 001 Jul 10, 2009

EQ 2MG BASE

A203690 001 Oct 09, 2012

EQ 4MG BASE

A077007 002 Jan 31, 2006

EQ 4MG BASE

A090711 002 Jul 10, 2009

EQ 4MG BASE

A090821 002 Jul 10, 2009

EQ 4MG BASE

A203690 002 Oct 09, 2012

WATSON LABS INC

EQ 2MG BASE

A209206 001 Jun 26, 2018

EQ 4MG BASE

A209206 002 Jun 26, 2018

WATSON LABS TEVA

EQ 2MG BASE

A209519 001 Jul 02, 2018

EQ 4MG BASE

A209519 002 Jul 02, 2018

NIZATIDINE

TABLET;ORAL

AXID AR

+! PFIZER

75MG

N020555 001 May 09, 1996

NONOXYNOL-9

SPONGE;VAGINAL

TODAY

+! MAYER LABS INC

1GM

N018683 001 Apr 01, 1983

OMEPRAZOLE

TABLET, DELAYED RELEASE;ORAL

OMEPRAZOLE

+! DEXCEL PHARMA 20MG

N022032 001 Dec 04, 2007

DR REDDYS LABS LTD 20MG

A207740 001 Nov 05, 2018

SUN PHARM INDS LTD 20MG

A207891 001 Oct 12, 2018

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

OMEPRAZOLE

+ DEXCEL PHARMA 20MG

N209400 001 Jul 05, 2017

OMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE;ORAL

OMEPRAZOLE MAGNESIUM

! DR REDDYS LABS LTD EQ 20MG BASE

A078878 001 Jun 05, 2009

SPIL EQ 20MG BASE

A210593 001 Jul 20, 2018

TABLET, DELAYED RELEASE;ORAL

OMEPRAZOLE MAGNESIUM

AUROBINDO PHARMA EQ 20MG BASE

A206877 001 Jun 06, 2018

LTD

PERRIGO R AND D EQ 20MG BASE

A204152 001 Jul 30, 2015

PRILOSEC OTC

+! ASTRAZENECA PHARMS EQ 20MG BASE

N021229 001 Jun 20, 2003

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

ACTAVIS ELIZABETH 20MG;1.1GM

A204137 001 Jul 15, 2016

AUROLIFE PHARMA LLC 20MG;1.1GM

A204923 001 Nov 07, 2016

PAR PHARM 20MG;1.1GM

A201946 001 Jul 15, 2016

PERRIGO R AND D 20MG;1.1GM

A201361 001 Jul 15, 2016

ZYDUS PHARMS USA 20MG;1.1GM

A203345 001 Mar 16, 2018

INC

ZEGERID OTC

+! BAYER HEALTHCARE 20MG;1.1GM

N022281 001 Dec 01, 2009

LLC

FOR SUSPENSION;ORAL

ZEGERID OTC

+! BAYER HEALTHCARE 20MG/PACKET;1.68GM/PACKET

N022283 001 Jun 17, 2013

LLC

OTC DRUG PRODUCT LIST

ORLISTAT

CAPSULE;ORAL

ALLI

| | | | | | | |
|---|---|-----------------|------|---------|-----|--------------|
| + | ! | GLAXOSMITHKLINE | 60MG | N021887 | 001 | Feb 07, 2007 |
| | | CONS | | | | |

OXYBUTYNIN

FILM, EXTENDED RELEASE;TRANSDERMAL

OXYTROL FOR WOMEN

| | | | | | | |
|---|---|--------------------|------------|---------|-----|--------------|
| + | ! | ALLERGAN SALES LLC | 3.9MG/24HR | N202211 | 001 | Jan 25, 2013 |
|---|---|--------------------|------------|---------|-----|--------------|

OXYMETAZOLINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

VISINE L.R.

| | | | | | | |
|---|---|---------------------|--------|---------|-----|--------------|
| + | ! | JOHNSON AND JOHNSON | 0.025% | N019407 | 001 | Mar 31, 1989 |
|---|---|---------------------|--------|---------|-----|--------------|

PERMETHRIN

LOTION;TOPICAL

NIX

| | | | | | | |
|---|---|------------------|----|---------|-----|--------------|
| + | ! | MEDTECH PRODUCTS | 1% | N019918 | 001 | May 02, 1990 |
|---|---|------------------|----|---------|-----|--------------|

PERMETHRIN

ACTAVIS MID

1%

A075014 001 Mar 28, 2000

ATLANTIC

PERRIGO NEW YORK

1%

A076090 001 Dec 20, 2001

POLYETHYLENE GLYCOL 3350

FOR SOLUTION;ORAL

GLYCOLAX

LANNETT CO INC

17GM/PACKET

A090600 001 Oct 06, 2009

17GM/SCOOPFUL

A090600 002 Oct 06, 2009

MIRALAX

| | | | | | | |
|---|---|------------------|---------------|---------|-----|--------------|
| + | ! | BAYER HEALTHCARE | 17GM/SCOOPFUL | N022015 | 001 | Oct 06, 2006 |
| | | LLC | | | | |

POLYETHYLENE GLYCOL 3350

ANI PHARMS INC

17GM/SCOOPFUL

A202850 001 Dec 15, 2015

APNAR PHARMA LP

17GM/SCOOPFUL

A202071 001 Dec 28, 2012

AUROBINDO PHARMA

17GM/SCOOPFUL

A209017 001 Apr 09, 2018

LTD

MYLAN

17GM/PACKET

A078915 001 Oct 06, 2009

17GM/SCOOPFUL

A078915 002 Oct 06, 2009

NEXGEN PHARMA

17GM/SCOOPFUL

A090812 001 Oct 07, 2009

NOVEL LABS INC

17GM/SCOOPFUL

A091077 001 Oct 06, 2009

NUVO PHARMS INC

17GM/SCOOPFUL

A206105 001 Oct 28, 2016

PAR PHARM

17GM/SCOOPFUL

A079214 001 Jan 31, 2013

PERRIGO R AND D

17GM/PACKET

A090685 001 Oct 06, 2009

17GM/SCOOPFUL

A090685 002 Oct 06, 2009

STRIDES PHARMA

17GM/SCOOPFUL

A203928 001 Aug 24, 2016

17GM/PACKET

A203928 002 Aug 24, 2016

POTASSIUM IODIDE

SOLUTION;ORAL

POTASSIUM IODIDE

MISSION PHARMACAL

65MG/ML

A206211 001 Mar 24, 2016

CO

THYROSHIELD

| | | | | | | |
|---|---|-----------------|---------|---------|-----|--------------|
| ! | ! | ARCO PHARMS LLC | 65MG/ML | A077218 | 001 | Jan 12, 2005 |
|---|---|-----------------|---------|---------|-----|--------------|

A077218 001 Jan 12, 2005

TABLET;ORAL

IOSAT

| | | | | | | |
|---|--|-------|------|---------|-----|--------------|
| + | | ANBEX | 65MG | N018664 | 002 | May 12, 2011 |
|---|--|-------|------|---------|-----|--------------|

N018664 002 May 12, 2011

| | | | | | | |
|---|---|--|-------|---------|-----|--------------|
| + | ! | | 130MG | N018664 | 001 | Oct 14, 1982 |
|---|---|--|-------|---------|-----|--------------|

N018664 001 Oct 14, 1982

THYROSAFE

| | | | | | | |
|---|--|-------|------|---------|-----|--------------|
| ! | | RECIP | 65MG | A076350 | 001 | Sep 10, 2002 |
|---|--|-------|------|---------|-----|--------------|

A076350 001 Sep 10, 2002

POVIDONE-IODINE

SOLUTION;TOPICAL

POVIDONE IODINE

| | | | | | | |
|---|---|---------------------|----|---------|-----|--------------|
| + | ! | ALLEGIANCE HLTHCARE | 1% | N019522 | 001 | Mar 31, 1989 |
|---|---|---------------------|----|---------|-----|--------------|

N019522 001 Mar 31, 1989

SPONGE;TOPICAL

E-Z SCRUB 201

| | | | | | | |
|---|---|------------------|-----|---------|-----|--------------|
| + | ! | BECTON DICKINSON | 20% | N019240 | 001 | Nov 29, 1985 |
|---|---|------------------|-----|---------|-----|--------------|

N019240 001 Nov 29, 1985

E-Z SCRUB 241

| | | | | | | |
|---|---|------------------|-----|---------|-----|--------------|
| + | ! | BECTON DICKINSON | 10% | N019476 | 001 | Jan 07, 1987 |
|---|---|------------------|-----|---------|-----|--------------|

N019476 001 Jan 07, 1987

OTC DRUG PRODUCT LIST

PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

PSEUDOEPHEDRINE HYDROCHLORIDE

| | | | | |
|-------------------------|-------|---------|-----|--------------|
| AUROBINDO PHARMA LTD | 120MG | A209008 | 001 | Jun 09, 2017 |
| L PERRIGO CO | 120MG | A075153 | 001 | Feb 26, 1999 |
| SUN PHARM INDS LTD | 120MG | A077442 | 001 | Sep 28, 2005 |
| SUDAFED 12 HOUR | | | | |
| ! MCNEIL CONS | 120MG | A073585 | 001 | Oct 31, 1991 |
| SUDAFED 24 HOUR | | | | |
| +! J AND J CONSUMER INC | 240MG | N020021 | 002 | Dec 15, 1992 |

PURIFIED WATER

SOLUTION;OPHTHALMIC

PUR-WASH

| | | | | |
|-------------------|-------|---------|-----|--------------|
| +! NIAGARA PHARMS | 98.3% | N022305 | 001 | Sep 01, 2011 |
|-------------------|-------|---------|-----|--------------|

RANITIDINE HYDROCHLORIDE

TABLET;ORAL

RANITIDINE HYDROCHLORIDE

| | | | | |
|----------------------|---------------|---------|-----|--------------|
| APOTEX INC | EQ 75MG BASE | A075167 | 001 | May 04, 2000 |
| | EQ 150MG BASE | A200172 | 001 | May 31, 2012 |
| AUROBINDO PHARMA LTD | EQ 75MG BASE | A207579 | 001 | Nov 13, 2017 |
| | EQ 150MG BASE | A207578 | 001 | Nov 13, 2017 |
| DR REDDYS LABS LTD | EQ 75MG BASE | A075294 | 001 | Mar 28, 2000 |
| | EQ 150MG BASE | A078192 | 001 | Aug 31, 2007 |
| GRANULES INDIA LTD | EQ 150MG BASE | A210243 | 001 | Aug 20, 2018 |
| | EQ 150MG BASE | A210243 | 002 | Aug 20, 2018 |
| IVAX SUB TEVA PHARMS | EQ 75MG BASE | A075296 | 001 | Jan 14, 2000 |
| MYLAN | EQ 75MG BASE | A075497 | 001 | Jan 14, 2000 |
| PERRIGO | EQ 75MG BASE | A076195 | 001 | Aug 30, 2002 |
| PERRIGO R AND D | EQ 150MG BASE | A091429 | 001 | May 11, 2011 |
| | EQ 150MG BASE | A091429 | 002 | May 11, 2011 |
| STRIDES PHARMA | EQ 75MG BASE | A201745 | 001 | Feb 29, 2012 |
| | EQ 150MG BASE | A200536 | 001 | Jun 28, 2011 |
| STRIDES VIVIMED | EQ 75MG BASE | A209160 | 001 | Mar 05, 2018 |
| | EQ 150MG BASE | A209161 | 001 | Feb 22, 2018 |
| WOCKHARDT | EQ 75MG BASE | A076760 | 001 | Feb 24, 2006 |
| ZANTAC 150 | | | | |
| +! SANOFI US | EQ 150MG BASE | N021698 | 001 | Aug 31, 2004 |
| + | EQ 150MG BASE | N021698 | 002 | Mar 13, 2007 |
| ZANTAC 75 | | | | |
| + SANOFI US | EQ 75MG BASE | N020520 | 001 | Dec 19, 1995 |

SODIUM CHLORIDE

AEROSOL, METERED;INHALATION

BRONCHO SALINE

| | | | | |
|------------|------|---------|-----|--------------|
| +! BLAIREX | 0.9% | N019912 | 001 | Sep 03, 1992 |
|------------|------|---------|-----|--------------|

SODIUM FLUORIDE; TRICLOSAN

PASTE;DENTAL

COLGATE TOTAL

| | | | | |
|----------------------|------------|---------|-----|--------------|
| +! COLGATE PALMOLIVE | 0.24%;0.3% | N020231 | 001 | Jul 11, 1997 |
|----------------------|------------|---------|-----|--------------|

TERBINAFINE

GEL;TOPICAL

LAMISIL AT

| | | | | |
|-------------------------|----|---------|-----|--------------|
| +! GLAXOSMITHKLINE CONS | 1% | N021958 | 001 | Jul 24, 2006 |
|-------------------------|----|---------|-----|--------------|

TERBINAFINE HYDROCHLORIDE

CREAM;TOPICAL

LAMISIL

| | | | | |
|--------------------|----|---------|-----|--------------|
| +! GLAXOSMITHKLINE | 1% | N020980 | 001 | Mar 09, 1999 |
|--------------------|----|---------|-----|--------------|

TERBINAFINE HYDROCHLORIDE

TARO

| | | | | |
|--|----|---------|-----|--------------|
| | 1% | A077511 | 001 | Jul 02, 2007 |
|--|----|---------|-----|--------------|

SOLUTION;TOPICAL

LAMISIL AT

| | | | | |
|-------------------------|----|---------|-----|--------------|
| +! GLAXOSMITHKLINE CONS | 1% | N021124 | 001 | Mar 17, 2000 |
|-------------------------|----|---------|-----|--------------|

SPRAY;TOPICAL

LAMISIL AT

| | | | | |
|-------------------------|----|---------|-----|--------------|
| +! GLAXOSMITHKLINE CONS | 1% | N021124 | 002 | Mar 17, 2000 |
|-------------------------|----|---------|-----|--------------|

OTC DRUG PRODUCT LISTTIOCONAZOLE

OINTMENT; VAGINAL

TIOCONAZOLE

PERRIGO

6.5%

A075915 001 Nov 21, 2001

VAGISTAT-1

+! COMBE

6.5%

N020676 001 Feb 11, 1997

TRIAMCINOLONE ACETONIDE

SPRAY, METERED; NASAL

NASACORT ALLERGY 24 HOUR

+! SANOFI AVENTIS US

0.055MG/SPRAY

N020468 002 Oct 11, 2013

TRIAMCINOLONE ACETONIDE

PERRIGO ISRAEL

0.055MG/SPRAY

A078104 002 Nov 14, 2014

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

ANTICOAGULANT 4% SODIUM CITRATE SOLUTION USP

INJECTABLE; INJECTION

NONE

HAEMONETICS

N760305

Jun 30, 1978

MANUFACTURING INC

ANTICOAGULANT CITRATE DEXTROSE SOLUTION (ACD)

INJECTABLE; INJECTION

CITRA LABS LLC

N020037

Aug 26, 2003

ACD-A SOLUTION

TERUMO BCT INC

A010228

Feb 25, 2002

ADSOL WITH ACD-A

FENWAL INC

N000922

Aug 29, 2002

ANTICOAGULANT CITRATE DEXTROSE SOLUTION FORMULA A

HAEMONETICS CORP

A980728

Feb 06, 2002

AS3 SOLUTION/ACD-A

TERUMO BCT INC

N001214

May 29, 2002

NONE

HAEMONETICS

A710497

Nov 06, 1987

MANUFACTURING INC

ANTICOAGULANT CITRATE DEXTROSE SOLUTION USP

INJECTABLE; INJECTION

NONE

FENWAL INC

N160918

Mar 17, 1978

ANTICOAGULANT CITRATE DEXTROSE SOLUTION USP (ACD-A)

INJECTABLE; INJECTION

NONE

ARTERIOCYTE MEDICAL

N160767

May 11, 2012

SYSTEMS, INC

ANTICOAGULANT CITRATE PHOSPHATE 2X DEXTROSE SOLUTION (CP2D)

INJECTABLE; INJECTION

CITRATE PHOSPHATE DOUBLE DEXTROSE/ADDITIVE SOLUTION 3

HAEMONETICS CORP

N000127

Jan 18, 2002

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION

INJECTABLE; INJECTION

NONE

TERUMO MEDICAL CORP

N820528

Nov 03, 1982

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION (CPDA)

INJECTABLE; INJECTION

CPDA-1 BLOOD-PACK UNIT (PL 146 PLASTIC) 250, 450, 500 ML BLOOD PACK UNITS

FENWAL INC

N770420

May 12, 1978

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION USP

INJECTABLE; INJECTION

BLOOD PACK UNIT CPDA-1 IN PLASTIC CONTAINER

FENWAL INC

N940404

Jul 28, 1994

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE-1 SOLUTION

INJECTABLE; INJECTION

NONE

HAEMONETICS

N800077

Nov 06, 1980

MANUFACTURING INC

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION

INJECTABLE; INJECTION

ADSOL IN PLASTIC CONTAINER
FENWAL INC

N900223 Dec 27, 1991

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION (CPD)

INJECTABLE; INJECTION

CPD ANTICOAGULANT IN PL 2209 PLASTIC CONTAINER
FENWAL INC

N900224 Dec 27, 1991

MACOPRODUCTIONS SAS CPD/AS-1: MACOPHARMA LEUCOFLEX MTL1 LEUKOREDUCTION SYSTEM FOR BLOOD
COMPONENTS KNOWN AS MTL1-WB

MACOPRODUCTIONS SAS

N040083 Nov 21, 2005

NONE

TERUMO BCT INC

A070025 Jan 06, 2008

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP

INJECTABLE; INJECTION

NONE

FENWAL INC

N170401 Dec 06, 1977

N811012 Jun 28, 1983

HAEMONETICS

N800222 Aug 23, 1982

MANUFACTURING INC

TERUMO MEDICAL CORP

N781211 Jun 10, 1981

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP WITH: AS-1:
DEXTROSE USP; SODIUM CHLORIDE USP; MANNITOL USP; ADENINE

INJECTABLE; INJECTION

ADSOL RED BLOOD CELL PRESERVATIVE SOLUTION
FENWAL INC

N811104 May 16, 1983

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP WITH: AS-5:
DEXTROSE USP; SODIUM CHLORIDE USP; MANNITOL USP; ADENINE

INJECTABLE; INJECTION

OPTISOL RED BLOOD CELL PRESERVATIVE SOLUTION
TERUMO MEDICAL CORP

N880217 Oct 07, 1988

ANTICOAGULANT CITRATE PHOSPHATE DOUBLE DEXTROSE SOLUTION WITH:
AS-3: CITRIC ACID USP; MONOBASIC SODIUM PHOSPHATE USP; SODIUM CHLORIDE USP; ADENINE;
DEXTROSE USP; SODIUM CITRATE USP

INJECTABLE; INJECTION

AS-3 NUTRICEL ADDITIVE SYSTEM

HAEMONETICS 0.042GM/100ML;0.276GM/100ML;

N820915 Oct 19, 1984

MANUFACTURING INC 0.410GM/100ML;0.30GM/100ML;

1.10GM/100ML;0.588GM/100ML

ANTICOAGULANT CITRATE PHOSPHATE DOUBLE DEXTROSE SOLUTION WITH:
AS-2: CITRIC ACID USP; DIBASIC SODIUM PHOSPHATE USP; SODIUM CHLORIDE USP; ADENINE;
DEXTROSE USP; SODIUM CITRATE USP

INJECTABLE; INJECTION

AS-2 NUTRICEL ADDITIVE SYSTEM

MEDSEP CORP 0.042GM/100ML;0.285GM/100ML;

N820915 Sep 22, 1983

0.718GM/100ML;0.017GM/100ML;

0.396GM/100ML;0.588GM/100ML

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

ANTICOAGULANT SODIUM CITRATE 4% SOLUTION

INJECTABLE; INJECTION

NONE

HAEMONETICS

N980123

Mar 03, 2000

CORPORATION

TERUMO BCT

N125608

Jun 26, 2018

ANTICOAGULANT SODIUM CITRATE SOLUTION

INJECTABLE; INJECTION

TRICITRASOL

CYTOSOL

N010409

Jul 10, 2003

LABORATORIES INC

ANTICOAGULANT SODIUM CITRATE SOLUTION USP

INJECTABLE; INJECTION

NONE

FENWAL INC

N770923

Jan 20, 1978

TERUMO MEDICAL CORP

N781214

Feb 08, 1980

CORD BLOOD STERILE COLLECTION BAG, ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION (CPD)

STERILE CORD BLOOD COLLECTION UNIT

NONE

MACOPHARMA

N125552

Dec 21, 2016

DEXTRAN 1 IN SODIUM CHLORIDE 0.6%

INJECTABLE; INJECTION

PROMIT

MEDA AB

N830715

Oct 30, 1984

DEXTRAN 40, 10% IN DEXTROSE 5%

INJECTABLE; INJECTION

LMD IN GLASS BOTTLE

HOSPIRA INC

10GM/100ML;5GM/100ML

A720563

Oct 30, 1992

DEXTRAN 40, 10% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION

LMD IN PLASTIC CONTAINER

HOSPIRA INC

10GM/100ML;0.9GM/100ML

A720562

Oct 30, 1992

HETASTARCH 6% IN LACTATED ELECTROLYTE INJECTION

INJECTABLE; INJECTION

HEXTEND

BIOTIME INC

6GM/100ML

N200952

Mar 31, 1999

HETASTARCH 6% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION

6% HETASTARCH IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

HOSPIRA INC

6GM/100ML;0.9GM/100ML

A740193

Jan 30, 1995

HESPAN IN PLASTIC CONTAINER

B BRAUN MEDICAL INC

6GM/100ML;0.9GM/100ML

N890105

Apr 04, 1991

NONE

TEVA PARENTERAL

6GM/100ML;0.9GM/100ML

A740592

Nov 12, 1998

MEDICINES INC

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

HYDROXYETHYL STARCH 130/0.4 IN 6% SODIUM CHLORIDE 0.9%

STORAGE/PROCESSING SOLUTION ONLY; SHOULD NEVER BE
INFUSED DIRECTLY TO THE PATIENT.

NONE

| | | | |
|------------------|-----------------------|---------|--------------|
| B. BRAUN MEDICAL | | A110013 | Jan 09, 2015 |
| VOLUVEN | | | |
| FRESENIUS KABI | 6GM/100ML;0.9GM/100ML | N070012 | Dec 27, 2007 |
| DEUTSCHLAND GMBH | | | |

ISOPLATE SOLUTION IN THE 500 ML EXCEL CONTAINER

STORAGE/PROCESSING SOLUTION ONLY; SHOULD NEVER BE
INFUSED DIRECTLY TO THE PATIENT.

ISOPLATE SOLUTION

| | | | |
|------------------|--|--------|--------------|
| HAEMONETICS CORP | | N90067 | Mar 05, 2013 |
|------------------|--|--------|--------------|

LEUKOCYTE REDUCTION FILTRATION SYSTEM FOR WHOLE BLOOD WITH CPD ANTICOAGULANT AND
SOLX ADDITIVE

INJECTABLE; INJECTION

| | | | |
|---------------------|--|---------|--------------|
| LEUKOSEP HWB-600-XL | | | |
| HAEMONETICS CORP | | N110059 | Apr 25, 2013 |

RED BLOOD CELL PROCESSING SOLUTION

STORAGE/PROCESSING SOLUTION ONLY; SHOULD NEVER BE
INFUSED DIRECTLY TO THE PATIENT.

REJUVESOL

| | | | |
|----------------|--|---------|--------------|
| CITRA LABS LLC | | N950522 | Feb 26, 1997 |
|----------------|--|---------|--------------|

SODIUM CHLORIDE; SODIUM ACETATE; SODIUM CITRATE DIHYDRATE; SODIUM PHOSPHATE,
DIABASIC ANHYDROUS; SODIUM PHOSPHATE MONOBASIC, MONOHYDRATE

STORAGE/PROCESSING SOLUTION ONLY; SHOULD NEVER BE
INFUSED DIRECTLY TO THE PATIENT.

INTERSOL SOLUTION

| | | | |
|-------------|---------------------------------------------------------------------|---------|--------------|
| FENWAL INC. | 2.26G/500ML; 2.21G/500ML; 1.59G/500ML; 1.53G/500ML; 0.465G/500ML | N080041 | Dec 09, 2009 |
|-------------|---------------------------------------------------------------------|---------|--------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ABARELIXINJECTABLE; INTRAMUSCULAR
PLENAXIS

SPECIALITY EUROPEAN 100MG/VIAL N021320 001 Nov 25, 2003

ACAMPROSATE CALCIUMTABLET, DELAYED RELEASE; ORAL
CAMPRAL

+ FOREST LABS 333MG ** N021431 001 Jul 29, 2004

ACEBUTOLOL HYDROCHLORIDE

CAPSULE; ORAL

ACEBUTOLOL HYDROCHLORIDE

WATSON LABS EQ 200MG BASE A074007 001 Oct 18, 1995

EQ 400MG BASE A074007 002 Oct 18, 1995

SECTRAL

+ PROMIUS PHARMA EQ 200MG BASE N018917 001 Dec 28, 1984

+ EQ 400MG BASE N018917 003 Dec 28, 1984

ACETAMINOPHEN

INJECTABLE; INJECTION

INJECTAPAP

ORTHO MCNEIL PHARM 100MG/ML N017785 001 Mar 07, 1986

SUPPOSITORY; RECTAL

ACEPHEN

G AND W LABS 120MG A072218 001 Mar 27, 1992

325MG N018060 003 Dec 18, 1986

650MG N018060 002

ACETAMINOPHEN

ABLE 120MG A073106 001 Feb 27, 1995

325MG A073107 001 Feb 27, 1995

650MG A073108 001 Feb 27, 1995

ACINO PRODS 120MG A071010 001 May 12, 1987

650MG A071011 001 May 12, 1987

TYLENOL

J AND J CONSUMER INC 120MG N017756 002

650MG N017756 001

TABLET, EXTENDED RELEASE; ORAL

ACETAMINOPHEN

SUN PHARM INDS LTD 650MG A090205 001 Nov 18, 2009

ACETAMINOPHEN; ASPIRIN; CODEINE PHOSPHATE

CAPSULE; ORAL

ACETAMINOPHEN, ASPIRIN, AND CODEINE PHOSPHATE

MIKART 150MG; 180MG; 15MG A081095 001 Oct 26, 1990

150MG; 180MG; 30MG A081096 001 Oct 26, 1990

150MG; 180MG; 60MG A081097 001 Oct 26, 1990

CODEINE, ASPIRIN, APAP FORMULA NO. 2

SCHERER LABS 150MG; 180MG; 15MG A085640 001

CODEINE, ASPIRIN, APAP FORMULA NO. 3

SCHERER LABS 150MG; 180MG; 30MG A085639 001

CODEINE, ASPIRIN, APAP FORMULA NO. 4

SCHERER LABS 150MG; 180MG; 60MG A085638 001

ACETAMINOPHEN; BUTALBITAL

CAPSULE; ORAL

BANCAP

FOREST PHARMS 325MG; 50MG A088889 001 Jan 16, 1986

BUCET

MALLINCKRODT 650MG; 50MG A088991 001 Jun 28, 1985

PHRENILIN FORTE

VALEANT 650MG; 50MG A088831 001 Jun 19, 1985

TENCON

MALLINCKRODT 650MG; 50MG A089405 001 May 15, 1990

TRIAPRIN

DUNHALL 325MG; 50MG A089268 001 Jul 02, 1987

TABLET; ORAL

BUTALBITAL AND ACETAMINOPHEN

HALSEY 325MG; 50MG A089568 001 Oct 05, 1988

WATSON LABS 325MG; 50MG A087550 001 Oct 19, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; BUTALBITAL

TABLET; ORAL

BUTAPAP

| | | | | |
|-----------|----------------|---------|-----|--------------|
| MIKART | 650MG; 50MG | A089988 | 001 | Oct 26, 1992 |
| PHRENILIN | | | | |
| VALEANT | 325MG; 50MG ** | A087811 | 001 | Jun 19, 1985 |
| SEDAPAP | | | | |
| MAYRAND | 650MG; 50MG | A088944 | 001 | Oct 17, 1985 |

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

ANOQUAN

| | | | | |
|----------------------------------------|----------------------|---------|-----|--------------|
| SHIRE | 325MG; 50MG; 40MG | A087628 | 001 | Oct 01, 1986 |
| BUTALBITAL, ACETAMINOPHEN AND CAFFEINE | | | | |
| GILBERT LABS | 325MG; 50MG; 40MG ** | A088825 | 001 | Dec 05, 1984 |
| GRAHAM DM | 325MG; 50MG; 40MG | A088743 | 001 | Jul 18, 1985 |
| | 325MG; 50MG; 40MG | A088765 | 001 | Mar 27, 1985 |
| | 325MG; 50MG; 40MG | A089067 | 001 | Apr 19, 1985 |
| HIKMA PHARMS | 500MG; 50MG; 40MG | A040261 | 001 | Oct 28, 1998 |
| MALLINCKRODT | 325MG; 50MG; 40MG | A088758 | 001 | Mar 27, 1985 |

ESGIC-PLUS

| | | | | |
|--------|-------------------|---------|-----|--------------|
| MIKART | 500MG; 50MG; 40MG | A040085 | 001 | Mar 28, 1996 |
|--------|-------------------|---------|-----|--------------|

FEMCET

| | | | | |
|--------------|-------------------|---------|-----|--------------|
| MALLINCKRODT | 325MG; 50MG; 40MG | A089102 | 001 | Jun 19, 1985 |
|--------------|-------------------|---------|-----|--------------|

MEDIGESIC PLUS

| | | | | |
|---------|-------------------|---------|-----|--------------|
| US CHEM | 325MG; 50MG; 40MG | A089115 | 001 | Jan 14, 1986 |
|---------|-------------------|---------|-----|--------------|

TRIAD

| | | | | |
|--------------|-------------------|---------|-----|--------------|
| MALLINCKRODT | 325MG; 50MG; 40MG | A089023 | 001 | Jun 19, 1985 |
|--------------|-------------------|---------|-----|--------------|

TABLET; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

| | | | | |
|----------------------|-------------------|---------|-----|--------------|
| ABLE | 325MG; 50MG; 40MG | A040390 | 001 | Jul 23, 2001 |
| | 500MG; 50MG; 40MG | A040394 | 001 | Jul 23, 2001 |
| GILBERT LABS | 325MG; 50MG; 40MG | A087629 | 001 | Nov 13, 1984 |
| HIKMA PHARMS | 500MG; 50MG; 40MG | A040336 | 001 | Aug 18, 1999 |
| MIKART | 750MG; 50MG; 40MG | A040496 | 001 | Dec 23, 2003 |
| MIRROR PHARMS LLC | 500MG; 50MG; 40MG | A040883 | 001 | Dec 23, 2008 |
| NOVAST LABS | 325MG; 50MG; 40MG | A040864 | 001 | Dec 01, 2008 |
| SUN PHARM INDUSTRIES | 325MG; 50MG; 40MG | A040601 | 001 | Jul 29, 2005 |
| VINTAGE PHARMS | 500MG; 50MG; 40MG | A040513 | 001 | Aug 25, 2003 |
| WATSON LABS | 325MG; 50MG; 40MG | A089536 | 001 | Feb 16, 1988 |
| | 500MG; 50MG; 40MG | A040267 | 001 | Jul 30, 1998 |

ESGIC

| | | | | |
|---------------|-------------------|---------|-----|--------------|
| FOREST PHARMS | 325MG; 50MG; 40MG | A089660 | 001 | Dec 23, 1988 |
|---------------|-------------------|---------|-----|--------------|

ESGIC-PLUS

| | | | | |
|--------|-------------------|---------|-----|--------------|
| MIKART | 500MG; 50MG; 40MG | A089451 | 001 | May 23, 1988 |
|--------|-------------------|---------|-----|--------------|

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE

| | | | | |
|-------------------------------------|-------------------------|---------|-----|--------------|
| ABLE | 325MG; 50MG; 40MG; 30MG | A076528 | 001 | Aug 21, 2003 |
| HIKMA INTL PHARMS | 325MG; 50MG; 40MG; 30MG | A075618 | 001 | Mar 23, 2001 |
| PHRENILIN WITH CAFFEINE AND CODEINE | | | | |
| VALEANT | 325MG; 50MG; 40MG; 30MG | A074911 | 001 | Aug 22, 2001 |

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE

| | | | | |
|-------------------|---------------------|---------|-----|--------------|
| MIKART | 356.4MG; 30MG; 16MG | A040109 | 001 | Aug 26, 1997 |
| WRASER PHARMS LLC | 356.4MG; 30MG; 16MG | A040688 | 001 | Apr 03, 2007 |
| DHC PLUS | | | | |
| PHARM RES ASSOC | 356.4MG; 30MG; 16MG | A088584 | 001 | Mar 04, 1986 |
| SYNALGOS-DC-A | | | | |
| LEITNER PHARMS | 356.4MG; 30MG; 16MG | A089166 | 001 | May 14, 1986 |

TABLET; ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE

| | | | | |
|----------------------|---------------------|---------|-----|--------------|
| BOCA PHARMA LLC | 712.8MG; 60MG; 32MG | A040701 | 001 | Apr 03, 2007 |
| MIKART | 712.8MG; 60MG; 32MG | A040316 | 001 | Apr 28, 1999 |
| WEST-WARD PHARM CORP | 712.8MG; 60MG; 32MG | A040637 | 001 | Sep 22, 2006 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; CLEMASTINE FUMARATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET; ORAL

TAVIST ALLERGY/SINUS/HEADACHE

NOVARTIS 500MG;EQ 0.25MG BASE;30MG N021082 001 Mar 01, 2001

ACETAMINOPHEN; CODEINE PHOSPHATE

CAPSULE; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

TEVA 300MG;15MG A088537 001 Jun 04, 1984

300MG;30MG A088324 001 Dec 29, 1983

300MG;60MG A088599 001 Jun 01, 1984

PHENAPHEN W/ CODEINE NO. 2

ROBINS AH 325MG;15MG A084444 001

PHENAPHEN W/ CODEINE NO. 3

ROBINS AH 325MG;30MG A084445 001

PHENAPHEN W/ CODEINE NO. 4

ROBINS AH 325MG;60MG A084446 001

PROVAL #3

SOLVAY 325MG;30MG A085685 001

TYLENOL W/ CODEINE NO. 3

ORTHO MCNEIL PHARM 300MG;30MG A087422 001

TYLENOL W/ CODEINE NO. 4

ORTHO MCNEIL PHARM 300MG;60MG A087421 001

SOLUTION; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

ACI HEALTHCARE LTD 120MG/5ML;12MG/5ML A086366 001

ACTAVIS MID ATLANTIC 120MG/5ML;12MG/5ML A085861 001

DAVA PHARMS INC 120MG/5ML;12MG/5ML A040098 001 Sep 20, 1996

TYLENOL W/ CODEINE

ORTHO MCNEIL PHARM 120MG/5ML;12MG/5ML A085057 001

SUSPENSION; ORAL

CAPITAL AND CODEINE

ACTAVIS MID ATLANTIC 120MG/5ML;12MG/5ML A085883 001

VALEANT PHARMS LLC 120MG/5ML;12MG/5ML A086024 001

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

ABLE 300MG;30MG A040452 001 Aug 01, 2002

300MG;60MG A040459 001 Aug 01, 2002

AM THERAP 300MG;15MG A089478 001 Mar 03, 1987

300MG;15MG A089481 001 Mar 03, 1987

300MG;30MG A089479 001 Mar 03, 1987

300MG;30MG A089482 001 Mar 03, 1987

300MG;60MG A089480 001 Mar 03, 1987

300MG;60MG A089483 001 Mar 03, 1987

DURAMED PHARMS BARR 300MG;15MG A040223 001 Nov 18, 1997

300MG;15MG A088353 001 Feb 06, 1984

300MG;30MG A040223 002 Nov 18, 1997

300MG;30MG A088354 001 Feb 06, 1984

300MG;60MG A040223 003 Nov 18, 1997

300MG;60MG A088355 001 Feb 06, 1984

EVERYLIFE 325MG;30MG A085217 001

FOSUN PHARMA 300MG;30MG A081250 001 Jul 16, 1992

300MG;60MG A081249 001 Jul 16, 1992

HALSEY 300MG;15MG A083871 001

300MG;30MG A083872 001

300MG;60MG A086549 001

KV PHARM 300MG;30MG A085288 001

300MG;60MG A085365 001

325MG;15MG A085364 001

325MG;45MG ** A085363 001

LEDERLE 300MG;30MG A087141 001

MIKART 300MG;30MG A089238 001 Feb 25, 1986

300MG;60MG A089244 001 Feb 25, 1986

650MG;30MG A089231 001 Mar 03, 1986

650MG;60MG A089363 001 Sep 09, 1991

MUTUAL PHARM 300MG;15MG A085795 001

300MG;30MG A085794 001

300MG;60MG A087653 001 Apr 13, 1982

PURACAP PHARM 300MG;30MG A087762 001 Dec 10, 1982

PUREPAC PHARM 300MG;30MG A086681 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

| | | | |
|----------------------------------|----------------|-------------|--------------|
| | 300MG;30MG | A089080 001 | Jul 17, 1986 |
| | 300MG;60MG | A086683 001 | |
| RHODES PHARMS | 300MG;15MG | A089673 002 | Feb 10, 1988 |
| | 300MG;30MG | A089673 003 | Feb 10, 1988 |
| | 300MG;60MG | A089673 001 | Feb 10, 1988 |
| ROXANE | 300MG;15MG | A084659 001 | |
| | 300MG;30MG | A084656 001 | |
| | 300MG;60MG | A084667 001 | |
| | 500MG;15MG | A089511 001 | Apr 25, 1989 |
| | 500MG;30MG | A089512 001 | Apr 25, 1989 |
| | 500MG;60MG | A089513 001 | Apr 25, 1989 |
| SANDOZ | 300MG;15MG | A087433 001 | |
| | 300MG;30MG | A085291 002 | |
| | 300MG;30MG | A085917 001 | |
| | 300MG;60MG | A085964 001 | |
| | 300MG;60MG | A087423 001 | |
| SUPERPHARM | 300MG;15MG | A089183 001 | Oct 18, 1985 |
| | 300MG;30MG | A089184 001 | Oct 18, 1985 |
| | 300MG;30MG | A089253 001 | May 19, 1986 |
| | 300MG;60MG | A089185 001 | Oct 18, 1985 |
| | 300MG;60MG | A089254 001 | May 19, 1986 |
| USL PHARMA | 300MG;30MG | A087919 001 | Jun 22, 1982 |
| | 300MG;60MG | A087920 001 | Jun 22, 1982 |
| VALEANT PHARM INTL | 300MG;30MG | A085896 001 | |
| VITARINE | 300MG;30MG | A085676 001 | |
| WARNER CHILCOTT | 300MG;15MG | A085992 001 | |
| | 300MG;30MG | A085218 002 | |
| | 300MG;60MG | A087306 001 | |
| WATSON LABS | 300MG;15MG | A087277 001 | May 26, 1982 |
| | 300MG;15MG | A089997 001 | Dec 28, 1994 |
| | 300MG;30MG | A087276 001 | May 26, 1982 |
| | 300MG;30MG | A089998 001 | Dec 28, 1994 |
| | 300MG;60MG | A087275 001 | May 26, 1982 |
| | 300MG;60MG | A089999 001 | Dec 28, 1994 |
| WATSON LABS FLORIDA | 300MG;15MG | A040443 001 | Jan 22, 2003 |
| | 300MG;30MG | A040443 002 | Jan 22, 2003 |
| | 300MG;60MG | A040443 003 | Jan 22, 2003 |
| WHITEWORTH TOWN PLSN | 300MG;30MG | A084360 001 | |
| | 300MG;60MG | A085607 001 | |
| CAPITAL AND CODEINE | | | |
| CARNRICK | 325MG;30MG | A083643 001 | |
| CODRIX | | | |
| WATSON LABS FLORIDA | 500MG;15MG | A040447 001 | Feb 26, 2003 |
| | 500MG;30MG | A040441 001 | Mar 27, 2003 |
| | 500MG;60MG | A040488 001 | Mar 28, 2003 |
| EMPRACET W/ CODEINE PHOSPHATE #3 | | | |
| GLAXOSMITHKLINE | 300MG;30MG | A083951 001 | |
| EMPRACET W/ CODEINE PHOSPHATE #4 | | | |
| GLAXOSMITHKLINE | 300MG;60MG | A083951 002 | |
| PAPA-DEINE #3 | | | |
| VANGARD | 300MG;30MG | A088037 001 | Mar 20, 1984 |
| PAPA-DEINE #4 | | | |
| VANGARD | 300MG;60MG | A088715 001 | Mar 20, 1984 |
| PHENAPHEN-650 W/ CODEINE | | | |
| ROBINS AH | 650MG;30MG | A085856 001 | |
| TYLENOL W/ CODEINE | | | |
| ORTHO MCNEIL PHARM | 325MG;7.5MG ** | A085056 001 | |
| | 325MG;15MG ** | A085056 002 | |
| | 325MG;30MG ** | A085056 003 | |
| | 325MG;60MG ** | A085056 004 | |
| TYLENOL W/ CODEINE NO. 1 | | | |
| JANSSEN PHARMS | 300MG;7.5MG | A085055 001 | |
| TYLENOL W/ CODEINE NO. 2 | | | |
| JANSSEN PHARMS | 300MG;15MG | A085055 002 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

DRIXORAL PLUS

SCHERING PLOUGH 500MG;3MG;60MG N019453 001 May 22, 1987

ACETAMINOPHEN; HYDROCODONE BITARTRATE

CAPSULE;ORAL

ACETAMINOPHEN AND HYDROCODONE BITARTRATE

CENT PHARMS 500MG;5MG A088898 001 Mar 27, 1985

ALLAY

IVAX PHARMS 500MG;5MG A089907 001 Jan 13, 1989

BANCAP HC

FOREST PHARMS 500MG;5MG A087961 001 Mar 17, 1983

CO-GESIC

CENT PHARMS 500MG;5MG A089360 001 Mar 02, 1988

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

MALLINCKRODT 500MG;5MG A088956 001 Jul 19, 1985

500MG;5MG A089006 001 Aug 09, 1985

MIKART 500MG;5MG A081067 001 Nov 30, 1989

500MG;5MG A081068 001 Nov 30, 1989

500MG;5MG A081069 001 Nov 30, 1989

500MG;5MG A081070 001 Nov 30, 1989

500MG;5MG A089008 001 Feb 21, 1986

LORCET-HD

MALLINCKRODT 500MG;5MG A087336 001 Jul 08, 1982

SOLUTION;ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

MALLINCKRODT 500MG/15ML;7.5MG/15ML A040418 001 Jun 27, 2001

MALLINCKRODT INC 500MG/15ML;10MG/15ML A040508 001 Aug 29, 2003

MIKART 500MG/15ML;5MG/15ML A081226 001 Oct 27, 1992

500MG/15ML;5MG/15ML A089557 001 Apr 29, 1992

500MG/15ML;7.5MG/15ML A081051 001 Aug 28, 1992

NESHER PHARMS 500MG/15ML;7.5MG/15ML A040366 001 Jan 23, 2002

PHARM ASSOC 500MG/15ML;7.5MG/15ML A040182 001 Mar 13, 1998

VINTAGE PHARMS 500MG/15ML;7.5MG/15ML A040520 001 Oct 30, 2003

ZYFREL

CYPRESS PHARM INC 325MG/15ML;7.5MG/15ML A090468 001 Apr 14, 2016

TABLET;ORAL

ANEXSIA

MALLINCKRODT 500MG;5MG A089160 001 Apr 23, 1987

750MG;10MG A040468 001 Oct 31, 2002

ANEXSIA 7.5/650

MALLINCKRODT 650MG;7.5MG A089725 001 Sep 30, 1987

CO-GESIC

UCB INC 500MG;5MG A087757 001 May 03, 1982

DURADYNE DHC

FOREST PHARMS 500MG;5MG A087809 001 Mar 17, 1983

HY-PHEN

ASCHER 500MG;5MG A087677 001 May 03, 1982

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

ABLE 325MG;5MG A040478 001 Nov 08, 2002

325MG;7.5MG A040464 001 Oct 23, 2002

325MG;10MG A040464 002 Oct 23, 2002

500MG;5MG A040477 001 Nov 06, 2002

500MG;7.5MG A040490 001 May 21, 2003

500MG;10MG A040473 001 Nov 06, 2002

650MG;7.5MG A040474 001 Jan 02, 2003

650MG;10MG A040476 001 Oct 23, 2002

750MG;7.5MG A040469 001 Oct 25, 2002

AMNEAL PHARMS NY 500MG;5MG A040729 001 Aug 25, 2006

500MG;7.5MG A040748 001 Aug 25, 2006

500MG;10MG A040813 001 Feb 23, 2007

650MG;7.5MG A040754 001 Aug 25, 2006

650MG;10MG A040757 001 Aug 25, 2006

750MG;7.5MG A040769 001 Aug 28, 2006

APIL 500MG;10MG A040148 002 Feb 14, 1997

BARR 500MG;2.5MG A040307 001 Jul 26, 2000

500MG;5MG A040308 001 Jul 26, 2000

500MG;5MG A088577 001 Dec 21, 1984

500MG;7.5MG A040307 002 Jul 26, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

| | | | | |
|--------------------|-------------|---------|-----|--------------|
| | 500MG;10MG | A040309 | 001 | Jul 26, 2000 |
| | 650MG;7.5MG | A040307 | 003 | Jul 26, 2000 |
| | 650MG;10MG | A040307 | 004 | Jul 26, 2000 |
| | 750MG;7.5MG | A040308 | 002 | Jul 26, 2000 |
| CARACO | 500MG;5MG | A090265 | 001 | Dec 23, 2008 |
| | 500MG;7.5MG | A090265 | 002 | Dec 23, 2008 |
| | 500MG;10MG | A090265 | 003 | Dec 23, 2008 |
| | 650MG;7.5MG | A090380 | 001 | Dec 23, 2008 |
| | 650MG;10MG | A090380 | 002 | Dec 23, 2008 |
| | 660MG;10MG | A090380 | 003 | Dec 23, 2008 |
| | 750MG;7.5MG | A090380 | 004 | Dec 23, 2008 |
| HALSEY | 500MG;5MG | A089554 | 001 | Jun 12, 1987 |
| IVAX PHARMS | 500MG;5MG | A089696 | 001 | Apr 21, 1988 |
| MALLINCKRODT | 500MG;5MG | A040084 | 002 | Jun 01, 1995 |
| | 500MG;7.5MG | A040201 | 001 | Feb 27, 1998 |
| | 500MG;10MG | A040201 | 002 | Feb 27, 1998 |
| | 650MG;10MG | A040084 | 004 | Oct 16, 1996 |
| | 660MG;10MG | A040084 | 003 | Jul 29, 1996 |
| | 750MG;7.5MG | A040084 | 001 | Jun 01, 1995 |
| MIKART | 500MG;2.5MG | A089698 | 001 | Aug 25, 1989 |
| | 500MG;5MG | A089271 | 001 | Jul 16, 1986 |
| | 500MG;5MG | A089697 | 001 | Jan 28, 1992 |
| | 500MG;7.5MG | A089699 | 001 | Aug 25, 1989 |
| | 650MG;5MG | A040849 | 001 | Jun 09, 2010 |
| | 650MG;7.5MG | A089689 | 001 | Jun 29, 1988 |
| | 650MG;10MG | A081223 | 001 | May 29, 1992 |
| MUTUAL PHARM | 500MG;5MG | A040236 | 001 | Sep 25, 1997 |
| | 650MG;7.5MG | A040240 | 002 | Nov 26, 1997 |
| | 650MG;10MG | A040240 | 001 | Nov 26, 1997 |
| | 750MG;7.5MG | A040236 | 002 | Sep 25, 1997 |
| RANBAXY | 500MG;5MG | A040825 | 001 | Aug 16, 2007 |
| | 500MG;10MG | A040824 | 001 | Aug 16, 2007 |
| RANBAXY LABS LTD | 750MG;7.5MG | A040822 | 001 | Aug 16, 2007 |
| SANDOZ | 500MG;5MG | A040149 | 001 | Jan 27, 1997 |
| | 750MG;7.5MG | A040149 | 002 | Jan 27, 1997 |
| SUN PHARM INDS LTD | 325MG;10MG | A040826 | 001 | Aug 16, 2007 |
| UCB INC | 500MG;10MG | A040210 | 001 | Aug 13, 1997 |
| | 650MG;7.5MG | A040134 | 001 | Nov 21, 1996 |
| USL PHARMA | 500MG;5MG | A089290 | 001 | May 29, 1987 |
| | 500MG;5MG | A089291 | 001 | May 29, 1987 |
| VINTAGE PHARMS | 500MG;2.5MG | A040144 | 002 | Apr 25, 1997 |
| | 500MG;5MG | A089831 | 001 | Sep 07, 1988 |
| | 500MG;5MG | A089971 | 001 | Dec 02, 1988 |
| | 500MG;7.5MG | A040144 | 001 | Feb 22, 1996 |
| | 500MG;10MG | A040356 | 001 | May 31, 2000 |
| | 650MG;7.5MG | A040155 | 001 | Apr 14, 1997 |
| | 650MG;10MG | A040143 | 001 | Feb 22, 1996 |
| | 660MG;10MG | A040358 | 001 | May 31, 2000 |
| | 750MG;7.5MG | A040157 | 001 | Apr 12, 1996 |
| VINTAGE PHARMS LLC | 500MG;5MG | A040281 | 001 | Sep 30, 1998 |
| | 500MG;7.5MG | A040280 | 001 | Sep 30, 1998 |
| | 650MG;7.5MG | A040280 | 002 | Sep 30, 1998 |
| | 650MG;10MG | A040280 | 003 | Sep 30, 1998 |
| | 750MG;7.5MG | A040281 | 002 | Sep 30, 1998 |
| WATSON LABS | 325MG;7.5MG | A040248 | 001 | Apr 28, 2000 |
| | 325MG;10MG | A040248 | 002 | Apr 28, 2000 |
| | 500MG;2.5MG | A040123 | 003 | Mar 04, 1996 |
| | 500MG;2.5MG | A081079 | 001 | Aug 30, 1991 |
| | 500MG;5MG | A040122 | 001 | Mar 04, 1996 |
| | 500MG;5MG | A089883 | 001 | Dec 01, 1988 |
| | 500MG;7.5MG | A040123 | 004 | Mar 04, 1996 |
| | 500MG;7.5MG | A081080 | 001 | Aug 30, 1991 |
| | 650MG;7.5MG | A040094 | 001 | Sep 29, 1995 |
| | 650MG;7.5MG | A040123 | 001 | Mar 04, 1996 |
| | 650MG;10MG | A040094 | 002 | Sep 29, 1995 |
| | 650MG;10MG | A040123 | 002 | Mar 04, 1996 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

| | | | | |
|---------------------|-------------|---------|-----|--------------|
| | 660MG;10MG | A040094 | 003 | Aug 08, 2000 |
| | 750MG;7.5MG | A040122 | 002 | Mar 04, 1996 |
| | 750MG;7.5MG | A081083 | 001 | Aug 30, 1991 |
| | 750MG;10MG | A040094 | 004 | Mar 22, 1999 |
| WATSON LABS FLORIDA | 500MG;5MG | A040493 | 001 | May 28, 2003 |
| | 660MG;10MG | A040495 | 001 | May 28, 2003 |
| | 750MG;7.5MG | A040494 | 001 | May 28, 2003 |
| LORTAB | | | | |
| UCB INC | 500MG;5MG | A087722 | 001 | Jul 09, 1982 |
| | 500MG;10MG | A040100 | 001 | Jan 26, 1996 |
| NORCET | | | | |
| ABANA | 500MG;5MG | A088871 | 001 | May 15, 1986 |
| TYCOLET | | | | |
| ORTHO MCNEIL PHARM | 500MG;5MG | A089385 | 001 | Aug 27, 1986 |
| VICODIN | | | | |
| ABBOTT | 500MG;5MG | A085667 | 001 | |
| ABBVIE | 500MG;5MG | A088058 | 001 | Jan 07, 1983 |
| VICODIN ES | | | | |
| ABBVIE | 750MG;7.5MG | A089736 | 001 | Dec 09, 1988 |
| VICODIN HP | | | | |
| ABBVIE | 660MG;10MG | A040117 | 001 | Sep 23, 1996 |
| ZYDONE | | | | |
| VINTAGE PHARMS LLC | 400MG;5MG | A040288 | 001 | Nov 27, 1998 |
| | 400MG;7.5MG | A040288 | 002 | Nov 27, 1998 |
| | 400MG;10MG | A040288 | 003 | Nov 27, 1998 |

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

CAPSULE; ORAL

OXYCODONE AND ACETAMINOPHEN

| | | | | |
|---------------------|-----------|---------|-----|--------------|
| ACTAVIS ELIZABETH | 500MG;5MG | A040199 | 001 | Dec 30, 1998 |
| BARR | 500MG;5MG | A040304 | 001 | Oct 02, 2000 |
| DURAMED PHARMS BARR | 500MG;5MG | A040289 | 001 | Mar 16, 1999 |
| HALSEY | 500MG;5MG | A089994 | 001 | May 04, 1989 |
| MALLINCKRODT | 500MG;5MG | A040257 | 001 | Aug 04, 1998 |
| MUTUAL PHARM | 500MG;5MG | A040219 | 001 | Jan 22, 1998 |
| VINTAGE PHARMS | 500MG;5MG | A040106 | 001 | Jul 30, 1996 |
| VINTAGE PHARMS LLC | 500MG;5MG | A040303 | 001 | Dec 30, 1999 |
| WATSON LABS | 500MG;5MG | A040234 | 001 | Oct 30, 1997 |
| ROXILOX | | | | |
| ROXANE | 500MG;5MG | A040061 | 001 | Jul 03, 1995 |
| TYLOX | | | | |
| JANSSEN PHARMS | 500MG;5MG | A088790 | 001 | Dec 12, 1984 |
| TYLOX-325 | | | | |
| ORTHO MCNEIL PHARM | 325MG;5MG | A088246 | 001 | Nov 08, 1984 |

SOLUTION; ORAL

OXYCODONE AND ACETAMINOPHEN

| | | | | |
|-------------------------------------------|-------------------|---------|-----|--------------|
| SPECGX LLC | 325MG/5ML;5MG/5ML | A040680 | 001 | Sep 29, 2006 |
| OXYCODONE HYDROCHLORIDE AND ACETAMINOPHEN | | | | |
| VINTAGE PHARMS | 325MG/5ML;5MG/5ML | A203573 | 001 | Dec 18, 2014 |
| ROXICET | | | | |
| WEST-WARD PHARMS INT | 325MG/5ML;5MG/5ML | A089351 | 001 | Dec 03, 1986 |

TABLET; ORAL

OXYCODONE 2.5/APAP 500

| | | | | |
|-----------------------------|-------------|---------|-----|--------------|
| BRISTOL MYERS SQUIBB | 500MG;2.5MG | A085910 | 001 | |
| OXYCODONE 5/APAP 500 | | | | |
| BRISTOL MYERS SQUIBB | 500MG;5MG | A085911 | 001 | |
| OXYCODONE AND ACETAMINOPHEN | | | | |
| ACTAVIS ELIZABETH | 325MG;5MG | A040203 | 001 | Mar 15, 1999 |
| | 325MG;7.5MG | A040800 | 001 | Apr 03, 2012 |
| | 325MG;10MG | A040800 | 002 | Apr 03, 2012 |
| AMNEAL PHARMS NY | 500MG;7.5MG | A040789 | 001 | Nov 27, 2007 |
| | 650MG;10MG | A040789 | 002 | Nov 27, 2007 |
| BARR | 325MG;5MG | A087406 | 001 | |
| DURAMED PHARMS BARR | 325MG;5MG | A040272 | 001 | Jun 30, 1998 |
| MALLINCKRODT | 500MG;7.5MG | A040550 | 001 | Jun 30, 2004 |
| | 650MG;10MG | A040550 | 002 | Jun 30, 2004 |
| MAYNE PHARMA INC | 500MG;7.5MG | A090177 | 005 | Oct 20, 2008 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE AND ACETAMINOPHEN

| | | | | |
|--------------------------------|-------------|---------|-----|--------------|
| | 650MG;10MG | A090177 | 006 | Oct 20, 2008 |
| MIKART | 400MG;2.5MG | A040679 | 001 | May 16, 2006 |
| | 400MG;5MG | A040687 | 001 | Apr 27, 2006 |
| | 400MG;7.5MG | A040698 | 001 | Apr 27, 2006 |
| | 400MG;10MG | A040692 | 001 | Apr 27, 2006 |
| | 500MG;10MG | A040676 | 001 | Apr 19, 2006 |
| WATSON LABS | 500MG;7.5MG | A040371 | 001 | Dec 29, 2000 |
| | 650MG;10MG | A040371 | 002 | Dec 29, 2000 |
| PERCOCET | | | | |
| VINTAGE PHARMS LLC | 325MG;5MG | A085106 | 002 | |
| | 500MG;7.5MG | A040341 | 001 | Jul 26, 1999 |
| | 650MG;10MG | A040341 | 002 | Jul 26, 1999 |
| ROXICET 5/500 | | | | |
| ROXANE | 500MG;5MG | A089775 | 001 | Jan 12, 1989 |
| TABLET, EXTENDED RELEASE; ORAL | | | | |
| XARTEMIS XR | | | | |
| + MALLINCKRODT INC | 325MG;7.5MG | N204031 | 001 | Mar 11, 2014 |

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

CAPSULE; ORAL

TYLOX

| | | | | |
|--------------------|--------------------|---------|-----|--|
| ORTHO MCNEIL PHARM | 500MG;4.5MG;0.38MG | A085375 | 001 | |
|--------------------|--------------------|---------|-----|--|

ACETAMINOPHEN; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

ACETAMINOPHEN AND PENTAZOCINE HYDROCHLORIDE

| | | | | |
|-------------------|--------------------|---------|-----|--------------|
| GAVIS PHARMS | 650MG;EQ 25MG BASE | A076202 | 001 | Aug 02, 2002 |
| WATSON LABS | 650MG;EQ 25MG BASE | A074699 | 001 | Mar 24, 2000 |
| TALACEN | | | | |
| SANOFI AVENTIS US | 650MG;EQ 25MG BASE | N018458 | 001 | Sep 23, 1982 |

ACETAMINOPHEN; PROPOXYPHENE HYDROCHLORIDE

TABLET; ORAL

DARVOCET

| | | | | |
|----------------------------------------------|--------------|---------|-----|--------------|
| AAIPHARMA LLC | 325MG;32.5MG | N016844 | 001 | |
| DOLENE AP-65 | | | | |
| LEDERLE | 650MG;65MG | A085100 | 001 | |
| PROPOXYPHENE HYDROCHLORIDE AND ACETAMINOPHEN | | | | |
| MYLAN | 325MG;32MG | A083689 | 001 | |
| | 650MG;65MG | A083978 | 001 | |
| SANDOZ | 650MG;65MG | A089959 | 001 | Jul 18, 1989 |
| VINTAGE PHARMS | 650MG;65MG | A040507 | 001 | Jul 30, 2003 |
| WATSON LABS | 650MG;65MG | A040139 | 001 | Dec 16, 1996 |
| WYGESIC | | | | |
| CARACO | 650MG;65MG | A084999 | 001 | |

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET; ORAL

DARVOCET A500

| | | | | |
|------------------------------------------|-------------|---------|-----|--------------|
| XANODYNE PHARM | 500MG;100MG | A076429 | 001 | Sep 10, 2003 |
| DARVOCET-N 100 | | | | |
| XANODYNE PHARM | 650MG;100MG | N017122 | 002 | |
| DARVOCET-N 50 | | | | |
| XANODYNE PHARM | 325MG;50MG | N017122 | 001 | |
| PROPACET 100 | | | | |
| TEVA | 650MG;100MG | A070107 | 001 | Jun 12, 1985 |
| PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN | | | | |
| ABLE | 650MG;100MG | A075838 | 001 | Jul 11, 2001 |
| ACTAVIS ELIZABETH | 650MG;100MG | A070910 | 001 | Jan 02, 1987 |
| CORNERSTONE | 325MG;100MG | A076743 | 001 | May 07, 2004 |
| | 500MG;100MG | A076750 | 001 | Jun 28, 2004 |
| HALSEY | 325MG;50MG | A072105 | 001 | May 13, 1988 |
| | 650MG;100MG | A072106 | 001 | May 13, 1988 |
| IVAX SUB TEVA PHARMS | 650MG;100MG | A070146 | 001 | Aug 02, 1985 |
| MALLINCKRODT | 650MG;100MG | A075738 | 001 | Feb 02, 2001 |
| MIRROR PHARMS | 650MG;100MG | A077821 | 001 | Feb 11, 2008 |
| MUTUAL PHARM | 325MG;50MG | A070115 | 001 | Jun 12, 1985 |
| | 650MG;100MG | A070116 | 001 | Jun 12, 1985 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET; ORAL

PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN

| | | | | |
|---------------------|-------------|---------|-----|--------------|
| | 650MG;100MG | A070615 | 001 | Mar 21, 1986 |
| | 650MG;100MG | A070771 | 001 | Mar 21, 1986 |
| | 650MG;100MG | A070775 | 001 | Mar 21, 1986 |
| MYLAN | 650MG;100MG | A072195 | 001 | Feb 16, 1988 |
| MYLAN PHARMS INC | 650MG;100MG | A070145 | 001 | Jun 12, 1985 |
| SANDOZ | 650MG;100MG | A070443 | 001 | Jan 23, 1986 |
| SUPERPHARM | 650MG;100MG | A071319 | 001 | Jan 06, 1987 |
| TEVA | 650MG;100MG | A070732 | 001 | Jan 03, 1986 |
| | 650MG;100MG | A074119 | 001 | Dec 19, 1994 |
| VINTAGE PHARMS | 325MG;50MG | A074843 | 002 | Feb 15, 2001 |
| | 650MG;100MG | A074843 | 001 | Feb 12, 1997 |
| WATSON LABS | 325MG;50MG | A070398 | 001 | Dec 18, 1986 |
| | 650MG;100MG | A070399 | 001 | Dec 18, 1986 |
| WATSON LABS FLORIDA | 500MG;100MG | A077196 | 001 | Jun 28, 2005 |
| | 650MG;100MG | A076609 | 001 | Nov 16, 2004 |
| WOCKHARDT LTD | 325MG;50MG | A077677 | 001 | Mar 16, 2007 |
| | 650MG;100MG | A077677 | 002 | Mar 16, 2007 |

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

| | | | | |
|--------------------|--------------|---------|-----|--------------|
| CSPC OUYI PHARM CO | 325MG;37.5MG | A076914 | 001 | Jul 26, 2006 |
|--------------------|--------------|---------|-----|--------------|

ACETAZOLAMIDE

TABLET; ORAL

ACETAZOLAMIDE

| | | | | |
|----------------------|----------|---------|-----|--------------|
| ALRA | 250MG | A083320 | 001 | |
| ASCOT | 250MG | A087686 | 001 | Oct 20, 1982 |
| SUN PHARM INDUSTRIES | 250MG | A089753 | 001 | Jun 22, 1988 |
| VANGARD | 250MG | A087654 | 001 | Feb 05, 1982 |
| WATSON LABS | 250MG | A084498 | 002 | |
| | 250MG | A088882 | 001 | Oct 22, 1985 |
| DIAMOX | | | | |
| + TEVA BRANDED PHARM | 125MG ** | N008943 | 001 | |
| + | 250MG ** | N008943 | 002 | |

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION

ACETAZOLAMIDE SODIUM

| | | | | |
|---------------|-----------------------|---------|-----|--------------|
| HOSPIRA | EQ 500MG BASE/VIAL | A040108 | 001 | Oct 30, 1995 |
| DIAMOX | | | | |
| + TEVA WOMENS | EQ 500MG BASE/VIAL ** | N009388 | 001 | |

ACETIC ACID, GLACIAL

SOLUTION/DROPS; OTIC

ACETASOL

| | | | | |
|----------------------|----|---------|-----|--|
| ACTAVIS MID ATLANTIC | 2% | A087146 | 001 | |
| ACETIC ACID | | | | |
| KV PHARM | 2% | A085493 | 001 | |
| ORLEX | | | | |
| WARNER CHILCOTT | 2% | A086845 | 001 | |

ACETIC ACID, GLACIAL; ALUMINUM ACETATE

SOLUTION/DROPS; OTIC

ACETIC ACID 2% IN AQUEOUS ALUMINUM ACETATE

| | | | | |
|-----------------|----------|---------|-----|--------------|
| BAUSCH AND LOMB | 2%;0.79% | A040063 | 001 | Feb 25, 1994 |
| BOROFAIR | | | | |
| PHARMAFAIR | 2%;0.79% | A088606 | 001 | Aug 21, 1985 |
| DOMEBORO | | | | |
| BAYER PHARMS | 2%;0.79% | A084476 | 001 | |

ACETIC ACID, GLACIAL; DESONIDE

SOLUTION/DROPS; OTIC

TRIDESILON

| | | | | |
|--------------|----------|---------|-----|--|
| BAYER PHARMS | 2%;0.05% | N017914 | 001 | |
|--------------|----------|---------|-----|--|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS;OTIC

ACETIC ACID W/ HYDROCORTISONE

KV PHARM 2%;1% A085492 001

HYDROCORTISONE AND ACETIC ACID

BAUSCH AND LOMB 2%;1% A040097 001 Oct 31, 1994

WOCKHARDT 2%;1% A040168 001 Aug 30, 1996

ORLEX HC

WARNER CHILCOTT 2%;1% A086844 001

ACETIC ACID, GLACIAL; HYDROCORTISONE; NEOMYCIN SULFATE

SUSPENSION/DROPS;OTIC

NEO-CORT-DOME

BAYER PHARMS 2%;1%;EQ 0.35% BASE N050238 001

ACETOHEXAMIDE

TABLET;ORAL

ACETOHEXAMIDE

ANI PHARMS INC 250MG A070869 001 Feb 09, 1987

500MG A070870 001 Feb 09, 1987

USL PHARMA 250MG A070753 001 Nov 03, 1986

500MG A070754 001 Nov 03, 1986

WATSON LABS TEVA 250MG A071893 001 Nov 25, 1987

500MG A071894 001 Nov 25, 1987

DYMELOR

LILLY 250MG N013378 002

500MG N013378 001

ACETOPHENAZINE MALEATE

TABLET;ORAL

TINDAL

SCHERING 20MG N012254 002

ACETRIZOATE SODIUM

SOLUTION;INTRAUTERINE

SALPIX

ORTHO MCNEIL PHARM 53% N009008 001

ACETYLCHOLINE CHLORIDE

FOR SOLUTION;OPHTHALMIC

MIOCHOL

NOVARTIS 20MG/VIAL N016211 001

ACETYLCYSTEINE

SOLUTION;INHALATION, ORAL

ACETYLCYSTEINE

HOSPIRA 10% A071364 001 May 01, 1989

20% A071365 001 May 01, 1989

ROXANE 10% A072323 001 Apr 30, 1992

10% A072621 001 Sep 30, 1992

20% A072324 001 Apr 30, 1992

20% A072622 001 Sep 30, 1992

MUCOMYST

+ APOTHECON 10% ** N013601 002

+ 20% ** N013601 001

MUCOSIL-10

DEY 10% A070575 001 Oct 14, 1986

MUCOSIL-20

DEY 20% A070576 001 Oct 14, 1986

ACETYLCYSTEINE; ISOPROTERENOL HYDROCHLORIDE

SOLUTION;INHALATION

MUCOMYST W/ ISOPROTERENOL

MEAD JOHNSON 10%;0.05% N017366 001

ACETYLDIGITOXIN

TABLET;ORAL

ACYLANID

NOVARTIS 0.1MG N009436 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACITRETIN

CAPSULE; ORAL

ACITRETIN

| | | | |
|------------------|--------|-------------|--------------|
| MYLAN PHARMS INC | 17.5MG | A203707 001 | Sep 10, 2015 |
| | 22.5MG | A203707 002 | Sep 10, 2015 |

ACRISORCIN

CREAM; TOPICAL

AKRINOL

| | | | |
|----------|--------|-------------|--|
| SCHERING | 2MG/GM | N012470 001 | |
|----------|--------|-------------|--|

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

| | | | |
|----------------------|-------|-------------|--------------|
| ACTAVIS ELIZABETH | 200MG | A074906 001 | Aug 26, 1997 |
| CHARTWELL MOLECULES | 200MG | A074872 001 | Apr 22, 1997 |
| IVAX SUB TEVA PHARMS | 200MG | A074674 001 | Apr 22, 1997 |
| LEK PHARM | 200MG | A074750 001 | Apr 22, 1997 |
| MYLAN | 200MG | A074727 001 | Apr 22, 1997 |
| | 200MG | A074977 001 | Apr 13, 1998 |
| RANBAXY | 200MG | A074975 001 | Sep 30, 1998 |
| ROXANE | 200MG | A074570 002 | Apr 22, 1997 |
| TEVA | 200MG | A074828 001 | Apr 22, 1997 |
| TEVA PHARMS | 200MG | A074914 001 | Nov 26, 1997 |
| WATSON LABS | 200MG | A075101 001 | Apr 15, 1998 |

TABLET; ORAL

ACYCLOVIR

| | | | |
|----------------------|----------|-------------|--------------|
| ACTAVIS ELIZABETH | 400MG | A074870 001 | Jun 05, 1997 |
| | 800MG | A074870 002 | Jun 05, 1997 |
| CHARTWELL MOLECULES | 400MG | A074834 001 | Apr 24, 1997 |
| | 800MG | A074834 002 | Apr 24, 1997 |
| IVAX SUB TEVA PHARMS | 400MG | A074836 001 | Apr 22, 1997 |
| | 800MG | A074836 002 | Apr 22, 1997 |
| LEK PHARM | 400MG | A074658 001 | Apr 22, 1997 |
| | 800MG | A074658 002 | Apr 22, 1997 |
| MYLAN | 400MG | A074976 001 | Apr 13, 1998 |
| | 800MG | A074976 002 | Apr 13, 1998 |
| MYLAN PHARMS INC | 400MG | A075211 001 | Sep 28, 1998 |
| | 800MG | A075211 002 | Sep 28, 1998 |
| SUN PHARM INDS LTD | 400MG | A074980 001 | Sep 30, 1998 |
| | 800MG | A074980 002 | Sep 30, 1998 |
| TEVA | 200MG ** | A074556 001 | Apr 22, 1997 |
| TEVA PHARMS | 400MG | A075021 001 | Mar 18, 1998 |
| | 800MG | A075021 002 | Mar 18, 1998 |

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR

| | | | |
|--------|-----------------|-------------|--------------|
| ABBVIE | EQ 50MG BASE/ML | A075114 001 | Jul 26, 1999 |
|--------|-----------------|-------------|--------------|

ACYCLOVIR IN SODIUM CHLORIDE 0.9% PRESERVATIVE FREE

| | | | |
|--------------------|--------------------|-------------|--------------|
| EUROHLTH INTL SARL | EQ 500MG BASE/VIAL | A074885 001 | Dec 19, 1997 |
| | EQ 1GM BASE/VIAL | A074885 002 | Dec 19, 1997 |

ACYCLOVIR SODIUM

| | | | |
|--------------------|--------------------|-------------|--------------|
| APOTHECON | EQ 500MG BASE/VIAL | A074897 001 | Feb 27, 1998 |
| | EQ 1GM BASE/VIAL | A074897 002 | Feb 27, 1998 |
| ATHENEX INC | EQ 500MG BASE/VIAL | A074596 002 | Apr 22, 1997 |
| | EQ 1GM BASE/VIAL | A074596 001 | Apr 22, 1997 |
| EUROHLTH INTL SARL | EQ 500MG BASE/VIAL | A074913 001 | Oct 15, 1997 |
| | EQ 1GM BASE/VIAL | A074913 002 | Oct 15, 1997 |
| FRESENIUS KABI USA | EQ 500MG BASE/VIAL | A075015 001 | Apr 30, 1998 |
| HIKMA PHARMS | EQ 500MG BASE/VIAL | A205771 001 | Feb 29, 2016 |
| | EQ 1GM BASE/VIAL | A205771 002 | Feb 29, 2016 |
| HOSPIRA | EQ 25MG BASE/ML | A074720 001 | Apr 22, 1997 |
| | EQ 50MG BASE/ML | A075065 001 | Feb 25, 1999 |
| | EQ 500MG BASE/VIAL | A074663 001 | Apr 22, 1997 |
| | EQ 500MG BASE/VIAL | A074758 001 | Apr 22, 1997 |
| | EQ 1GM BASE/VIAL | A074663 002 | Apr 22, 1997 |
| | EQ 1GM BASE/VIAL | A074758 002 | Apr 22, 1997 |
| MYLAN LABS LTD | EQ 500MG BASE/VIAL | A203927 001 | Mar 29, 2017 |
| | EQ 1GM BASE/VIAL | A203927 002 | Mar 29, 2017 |
| TEVA PARENTERAL | EQ 50MG BASE/ML | A075627 001 | Mar 28, 2001 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR SODIUM

| | | | | |
|---|----------------------|-----------------------|-------------|--------------|
| | | EQ 500MG BASE/VIAL | A074969 001 | Aug 26, 1997 |
| | | EQ 1GM BASE/VIAL | A074969 002 | Aug 26, 1997 |
| | ZYDUS PHARMS USA INC | EQ 500MG BASE/VIAL | A206606 001 | Jun 13, 2017 |
| | | EQ 1GM BASE/VIAL | A206606 002 | Jun 13, 2017 |
| | ZOVIRAX | | | |
| + | GLAXOSMITHKLINE | EQ 250MG BASE/VIAL ** | N018603 003 | Aug 30, 1983 |
| + | | EQ 500MG BASE/VIAL ** | N018603 001 | Oct 22, 1982 |
| + | | EQ 1GM BASE/VIAL ** | N018603 002 | Jun 29, 1989 |

ADAPALENE

SOLUTION; TOPICAL

DIFFERIN

| | | | | |
|---|------------------|---------|-------------|--------------|
| + | GALDERMA LABS LP | 0.1% ** | N020338 001 | May 31, 1996 |
|---|------------------|---------|-------------|--------------|

ADENOSINE

INJECTABLE; INJECTION

ADENOCARD

| | | | | |
|---|----------|--------|-------------|--------------|
| + | ASTELLAS | 3MG/ML | N019937 002 | Oct 30, 1989 |
|---|----------|--------|-------------|--------------|

ADENOSINE

| | | | | |
|--|----------------------|--------|-------------|--------------|
| | TEVA PHARMS USA | 3MG/ML | A076564 001 | Jun 16, 2004 |
| | | 3MG/ML | A078676 001 | Jul 31, 2008 |
| | WEST-WARD PHARMS INT | 3MG/ML | A076501 001 | Jun 16, 2004 |
| | WOCKHARDT | 3MG/ML | A090220 001 | Jul 20, 2009 |

SOLUTION; INTRAVENOUS

ADENOSCAN

| | | | | |
|---|----------|-----------------------|-------------|--------------|
| + | ASTELLAS | 60MG/20ML (3MG/ML) ** | N020059 001 | May 18, 1995 |
| + | | 90MG/30ML (3MG/ML) ** | N020059 002 | May 18, 1995 |

ALATROFLOXACIN MESYLATE

INJECTABLE; INJECTION

TROVAN PRESERVATIVE FREE

| | | | | |
|--|--------|--------------------|-------------|--------------|
| | PFIZER | EQ 200MG BASE/VIAL | N020760 001 | Dec 18, 1997 |
| | | EQ 300MG BASE/VIAL | N020760 002 | Dec 18, 1997 |

ALBENDAZOLE

TABLET, CHEWABLE; ORAL

ALBENZA

| | | | | |
|--|-------------------|-------|-------------|--------------|
| | AMEDRA PHARMS LLC | 200MG | N207844 001 | Jun 11, 2015 |
|--|-------------------|-------|-------------|--------------|

ALBUMIN CHROMATED CR-51 SERUM

INJECTABLE; INJECTION

CHROMALBIN

| | | | | |
|--|---------|-------------|-------------|--|
| | ISO TEX | 100uCi/VIAL | N017835 001 | |
| | | 250uCi/VIAL | N017835 002 | |
| | | 500uCi/VIAL | N017835 003 | |

ALBUMIN IODINATED I-125 SERUM

INJECTABLE; INJECTION

RADIO-IODINATED (I 125) SERUM ALBUMIN (HUMAN)

| | | | | |
|--|--------------|------------|-------------|--|
| | BAYER PHARMS | 2.5uCi/AMP | N017846 001 | |
|--|--------------|------------|-------------|--|

RADIOIODINATED SERUM ALBUMIN (HUMAN) IHSA I 125

| | | | | |
|--|--------------|------------|-------------|--|
| | MALLINCKRODT | 6.67uCi/ML | N017844 003 | |
| | | 10uCi/ML | N017844 001 | |
| | | 100uCi/ML | N017844 002 | |

ALBUMIN IODINATED I-131 SERUM

INJECTABLE; INJECTION

MEGATOPE

| | | | | |
|--|---------|-----------|-------------|--|
| | ISO TEX | 2mCi/VIAL | N017837 003 | |
| | | 5uCi/AMP | N017837 004 | |
| | | 20uCi/AMP | N017837 005 | |

ALBUTEROL

AEROSOL, METERED; INHALATION

ALBUTEROL

| | | | | |
|--|----------------------|------------|-------------|--------------|
| | ARMSTRONG PHARMS | 0.09MG/INH | A072273 001 | Aug 14, 1996 |
| | GENPHARM | 0.09MG/INH | A073045 001 | Aug 19, 1997 |
| | IVAX SUB TEVA PHARMS | 0.09MG/INH | A073272 001 | Dec 28, 1995 |
| | PLIVA | 0.09MG/INH | A074072 001 | Aug 01, 1996 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALBUTEROL

AEROSOL, METERED; INHALATION

PROVENTIL

SCHERING 0.09MG/INH N017559 001

VENTOLIN

GLAXOSMITHKLINE 0.09MG/INH N018473 001

ALBUTEROL SULFATE

CAPSULE; INHALATION

VENTOLIN ROTACAPS

GLAXOSMITHKLINE EQ 0.2MG BASE N019489 001 May 04, 1988

SOLUTION; INHALATION

ALBUTEROL SULFATE

ACTAVIS MID ATLANTIC EQ 0.083% BASE A073533 001 Sep 26, 1995

APOTEX INC EQ 0.021% BASE A078623 001 Apr 05, 2010

EQ 0.042% BASE A078623 002 Apr 05, 2010

EQ 0.083% BASE A075717 001 Feb 02, 2007

EQ 0.5% BASE A076391 001 Apr 01, 2003

BAUSCH AND LOMB EQ 0.083% BASE A075358 001 Mar 29, 2000

COPLEY PHARM EQ 0.083% BASE A073495 001 May 28, 1993

EQ 0.5% BASE A073307 001 Nov 27, 1991

HI TECH PHARMA EQ 0.083% BASE A075063 001 Feb 09, 1999

LANDELA PHARM EQ 0.083% BASE A077569 001 Apr 04, 2006

MYLAN SPECLT EQ 0.083% BASE ** A072652 001 Feb 21, 1992

ROXANE EQ 0.083% BASE A075129 001 Feb 13, 2001

TEVA PHARMS EQ 0.083% BASE A075343 001 Nov 09, 1999

WATSON LABS INC EQ 0.083% BASE A076370 001 Nov 24, 2003

WOCKHARDT EU OPERATN EQ 0.083% BASE A075394 001 Nov 22, 1999

PROVENTIL

+ SCHERING EQ 0.083% BASE ** N019243 002 Jan 14, 1987

+ EQ 0.5% BASE ** N019243 001 Jan 14, 1987

VENTOLIN

+ GLAXOSMITHKLINE EQ 0.083% BASE ** N019773 001 Apr 23, 1992

EQ 0.5% BASE ** N019269 002 Jan 16, 1987

SYRUP; ORAL

ALBUTEROL SULFATE

ACTAVIS MID ATLANTIC EQ 2MG BASE/5ML A075262 001 Mar 30, 1999

MOVA EQ 2MG BASE/5ML A074302 001 Sep 30, 1994

WATSON LABS EQ 2MG BASE/5ML A073165 001 Apr 29, 1993

PROVENTIL

+ SCHERING EQ 2MG BASE/5ML ** N018062 001 Jan 19, 1983

VENTOLIN

GLAXOSMITHKLINE EQ 2MG BASE/5ML ** N019621 001 Jun 10, 1987

TABLET; ORAL

ALBUTEROL SULFATE

AM THERAP EQ 2MG BASE A072449 001 Dec 05, 1989

EQ 4MG BASE A072450 001 Dec 05, 1989

COPLEY PHARM EQ 2MG BASE A072966 001 Nov 22, 1991

EQ 4MG BASE A072967 001 Nov 22, 1991

DAVA PHARMS INC EQ 2MG BASE A072860 002 Dec 20, 1989

EQ 4MG BASE A072860 001 Dec 20, 1989

PLIVA EQ 2MG BASE A072316 001 Dec 05, 1989

EQ 4MG BASE A072317 001 Dec 05, 1989

TEVA EQ 2MG BASE A072619 001 Dec 05, 1989

EQ 2MG BASE A072779 001 Jun 25, 1993

EQ 2MG BASE A072938 001 Mar 30, 1990

EQ 4MG BASE A072620 001 Dec 05, 1989

EQ 4MG BASE A072780 001 Jun 25, 1993

EQ 4MG BASE A072939 001 Mar 30, 1990

UCB INC EQ 2MG BASE A073120 001 Sep 29, 1992

EQ 4MG BASE A073121 001 Sep 29, 1992

WARNER CHILCOTT EQ 2MG BASE A072817 001 Jan 09, 1990

EQ 4MG BASE A072818 001 Jan 09, 1990

WATSON LABS EQ 2MG BASE A072629 001 Jan 31, 1991

EQ 2MG BASE A072764 001 Aug 28, 1991

EQ 4MG BASE A072630 001 Jan 31, 1991

EQ 4MG BASE A072765 001 Aug 28, 1991

YAOPHARMA CO LTD EQ 2MG BASE A072151 001 Dec 05, 1989

EQ 4MG BASE A072152 001 Dec 05, 1989

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALBUTEROL SULFATE

TABLET; ORAL

PROVENTIL

+ SCHERING

EQ 2MG BASE **

N017853 001 May 07, 1982

+

EQ 4MG BASE **

N017853 002 May 07, 1982

VENTOLIN

GLAXOSMITHKLINE

EQ 2MG BASE

N019112 001 Jul 10, 1986

EQ 4MG BASE

N019112 002 Jul 10, 1986

TABLET, EXTENDED RELEASE; ORAL

PROVENTIL

SCHERING

EQ 4MG BASE

N019383 001 Jul 13, 1987

VOLMAX

MURO

EQ 4MG BASE

N019604 002 Dec 23, 1992

EQ 8MG BASE

N019604 001 Dec 23, 1992

VOSPIRE ER

DAVA PHARMS INC

EQ 8MG BASE

A076130 003 Sep 26, 2002

ALBUTEROL SULFATE; IPRATROPIUM BROMIDE

AEROSOL, METERED; INHALATION

COMBIVENT

BOEHRINGER INGELHEIM EQ 0.09MG BASE/INH;0.018MG/INH

N020291 001 Oct 24, 1996

SOLUTION; INHALATION

ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE

APOTEX INC

EQ 0.083% BASE;0.017%

A077117 001 Dec 31, 2007

FOSUN PHARMA

EQ 0.083% BASE;0.017%

A076867 001 Dec 21, 2006

TEVA PHARMS

EQ 0.083% BASE;0.017%

A076724 001 Dec 31, 2007

DUONEB

+ MYLAN SPECIALITY LP

EQ 0.083% BASE;0.017% **

N020950 001 Mar 21, 2001

ALCLOMETASONE DIPROPIONATE

CREAM; TOPICAL

ACLOVATE

+ FOUGERA PHARMS

0.05% **

N018707 001 Dec 14, 1982

OINTMENT; TOPICAL

ACLOVATE

+ FOUGERA PHARMS

0.05% **

N018702 001 Dec 14, 1982

ALCOHOL

INJECTABLE; INJECTION

ALCOHOL 5% IN DEXTROSE 5%

MILES

5ML/100ML

A083483 001

ALCOHOL; DEXTROSE

INJECTABLE; INJECTION

ALCOHOL 10% AND DEXTROSE 5%

B BRAUN

10ML/100ML;5GM/100ML

N004589 006

ALCOHOL 5% AND DEXTROSE 5%

B BRAUN

5ML/100ML;5GM/100ML

N004589 004

ALCOHOL 5% IN D5-W

HOSPIRA

5ML/100ML;5GM/100ML

A083263 001

ALCOHOL 5% IN DEXTROSE 5% IN WATER

BAXTER HLTHCARE

5ML/100ML;5GM/100ML

A083256 001

ALENDRONATE SODIUM

SOLUTION; ORAL

FOSAMAX

+ MERCK

EQ 70MG BASE/75ML **

N021575 001 Sep 17, 2003

TABLET; ORAL

ALENDRONATE SODIUM

MYLAN

EQ 35MG BASE

A076584 003 Aug 04, 2008

EQ 35MG BASE

A078638 001 Aug 04, 2008

EQ 70MG BASE

A076584 004 Aug 04, 2008

EQ 70MG BASE

A078638 002 Aug 04, 2008

TEVA PHARMS

EQ 35MG BASE

A076184 002 Aug 04, 2008

EQ 70MG BASE

A076184 001 Feb 06, 2008

UPSHER SMITH LABS

EQ 5MG BASE

A075871 001 Apr 22, 2009

EQ 10MG BASE

A075871 002 Apr 22, 2009

EQ 35MG BASE

A075871 004 Apr 22, 2009

EQ 40MG BASE

A075871 003 Apr 22, 2009

EQ 70MG BASE

A075871 005 Apr 22, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALENDRONATE SODIUM

TABLET; ORAL

FOSAMAX

| | | | | |
|---|------------------|-----------------|-------------|--------------|
| + | MERCK AND CO INC | EQ 5MG BASE ** | N020560 003 | Apr 25, 1997 |
| + | | EQ 10MG BASE ** | N020560 001 | Sep 29, 1995 |
| + | | EQ 35MG BASE ** | N020560 004 | Oct 20, 2000 |
| + | | EQ 40MG BASE ** | N020560 002 | Sep 29, 1995 |

ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

ALFUZOSIN HYDROCHLORIDE

WOCKHARDT LTD

10MG

A090221 001 Aug 10, 2012

ALGLUCERASE

INJECTABLE; INJECTION

CEREDASE

GENZYME

10 UNITS/ML

N020057 004 May 08, 1992

80 UNITS/ML

N020057 003 Apr 05, 1991

ALISKIREN HEMIFUMARATE

CAPSULE, PELLET; ORAL

TEKTURNA

+ NODEN PHARMA

EQ 37.5MG BASE

N210709 001 Nov 14, 2017

ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE

TABLET; ORAL

TEKAMLO

NOVARTIS

EQ 150MG BASE; EQ 5MG BASE

N022545 001 Aug 26, 2010

EQ 150MG BASE; EQ 10MG BASE

N022545 002 Aug 26, 2010

EQ 300MG BASE; EQ 5MG BASE

N022545 003 Aug 26, 2010

EQ 300MG BASE; EQ 10MG BASE

N022545 004 Aug 26, 2010

ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMTURNIDE

NOVARTIS

EQ 150MG BASE; EQ 5MG BASE; 12.5MG

N200045 001 Dec 21, 2010

EQ 300MG BASE; EQ 5MG BASE; 12.5MG

N200045 002 Dec 21, 2010

EQ 300MG BASE; EQ 5MG BASE; 25MG

N200045 003 Dec 21, 2010

EQ 300MG BASE; EQ 10MG BASE; 12.5MG

N200045 004 Dec 21, 2010

EQ 300MG BASE; EQ 10MG BASE; 25MG

N200045 005 Dec 21, 2010

ALISKIREN HEMIFUMARATE; VALSARTAN

TABLET; ORAL

VALTURNA

NOVARTIS

EQ 150MG BASE; 160MG

N022217 001 Sep 16, 2009

EQ 300MG BASE; 320MG

N022217 002 Sep 16, 2009

ALKAVERVIR

TABLET; ORAL

VERILOID

3M

2MG

N007336 002

3MG

N007336 003

ALLOPURINOL

TABLET; ORAL

ALLOPURINOL

FOSUN PHARMA

100MG

A070268 001 Dec 31, 1985

MUTUAL PHARM

100MG

A070466 001 Dec 24, 1985

300MG

A070467 001 Dec 24, 1985

PURACAP PHARM

100MG

A070150 001 Dec 10, 1985

300MG

A070147 001 Dec 10, 1985

PUREPAC PHARM

100MG

A070579 001 Apr 14, 1986

300MG

A070580 001 Apr 14, 1986

SANDOZ

300MG

A070269 001 Dec 31, 1985

SUPERPHARM

100MG

A070950 001 Nov 30, 1988

300MG

A070951 001 Nov 30, 1988

WATSON LABS

100MG

N018241 001 Nov 16, 1984

100MG

N018785 001 Sep 28, 1984

300MG

N018241 002 Nov 16, 1984

300MG

N018785 002 Sep 28, 1984

LOPURIN

ABBOTT

100MG

N018297 001

300MG

N018297 002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALPRAZOLAM

SOLUTION; ORAL

ALPRAZOLAM

ROXANE

0.5MG/5ML

A074314 001 Oct 31, 1993

TABLET; ORAL

ALPRAZOLAM

ANI PHARMS INC

0.25MG

A074085 001 Feb 16, 1994

0.5MG

A074085 002 Feb 16, 1994

1MG

A074085 003 Feb 16, 1994

2MG

A074085 004 Feb 26, 1996

IVAX SUB TEVA PHARMS

0.25MG

A074294 001 Jul 29, 1994

0.5MG

A074294 002 Jul 29, 1994

1MG

A074294 003 Jul 29, 1994

2MG

A074294 004 Jul 29, 1994

MYLAN PHARMS INC

0.25MG

A074046 001 Oct 19, 1993

0.5MG

A074046 002 Oct 19, 1993

1MG

A074046 003 Oct 19, 1993

2MG

A074046 004 May 07, 1997

ROXANE

0.25MG

A074199 001 Oct 19, 1993

0.5MG

A074199 002 Oct 19, 1993

1MG

A074199 003 Oct 19, 1993

WATSON LABS

0.25MG

A074456 001 Aug 31, 1995

0.25MG

A074479 001 Jan 21, 1997

0.5MG

A074456 002 Aug 31, 1995

0.5MG

A074479 002 Jan 21, 1997

1MG

A074456 003 Aug 31, 1995

1MG

A074479 003 Jan 21, 1997

TABLET, EXTENDED RELEASE; ORAL

ALPRAZOLAM

ACTAVIS LABS FL INC

0.5MG

A077198 001 May 13, 2010

1MG

A077198 002 May 13, 2010

2MG

A077198 003 May 13, 2010

3MG

A077198 004 May 13, 2010

ANI PHARMS INC

0.5MG

A077979 001 Feb 28, 2007

1MG

A077979 002 Feb 28, 2007

2MG

A077979 003 Feb 28, 2007

3MG

A077979 004 Feb 28, 2007

IMPAX LABS

0.5MG

A077968 004 May 24, 2007

1MG

A077968 003 May 24, 2007

2MG

A077968 002 May 24, 2007

3MG

A077968 001 May 24, 2007

IMPAX LABS INC

0.5MG

A077996 001 Jan 31, 2007

1MG

A077996 002 Jan 31, 2007

2MG

A077996 003 Jan 31, 2007

3MG

A077996 004 Jan 31, 2007

MYLAN

0.5MG

A077391 002 Jan 26, 2006

1MG

A077391 003 Jan 26, 2006

2MG

A077391 004 Jan 26, 2006

3MG

A077391 001 Jan 26, 2006

SANDOZ INC

0.5MG

A077777 001 Jun 30, 2006

1MG

A077777 002 Jun 30, 2006

2MG

A077777 003 Jun 30, 2006

3MG

A077777 004 Jun 30, 2006

VINTAGE PHARMS

0.5MG

A078442 001 Oct 15, 2007

1MG

A078442 002 Oct 15, 2007

2MG

A078442 003 Oct 15, 2007

3MG

A078442 004 Oct 15, 2007

TABLET, ORALLY DISINTEGRATING; ORAL

NIRAVAM

+ UCB INC

0.25MG **

N021726 001 Jan 19, 2005

+

0.5MG **

N021726 002 Jan 19, 2005

+

1MG **

N021726 003 Jan 19, 2005

+

2MG **

N021726 004 Jan 19, 2005

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ALPROSTADIL

INJECTABLE; INJECTION

CAVERJECT

| | | | |
|------------------------|--------------|-------------|--------------|
| PFIZER | 0.005MG/ML | N020755 001 | Oct 31, 1997 |
| | 0.01MG/ML | N020755 002 | Oct 01, 1997 |
| | 0.02MG/ML | N020755 003 | Oct 01, 1997 |
| + PHARMACIA AND UPJOHN | 0.005MG/VIAL | N020379 003 | Jun 27, 1996 |
| EDEX | | | |
| AUXILIUM PHARMS LLC | 0.005MG/VIAL | N020649 001 | Jun 12, 1997 |

ALSEROXYLON

TABLET; ORAL

RAUTENSIN

| | | | |
|----------|-----|-------------|--|
| NOVARTIS | 2MG | N009215 001 | |
|----------|-----|-------------|--|

RAUWILOID

| | | | |
|----|-----|-------------|--|
| 3M | 2MG | N008867 001 | |
|----|-----|-------------|--|

ALUMINUM HYDROXIDE; MAGNESIUM TRISILICATE

TABLET, CHEWABLE; ORAL

ALUMINUM HYDROXIDE AND MAGNESIUM TRISILICATE

| | | | |
|---------------------|-------------|-------------|--------------|
| PENNEX | 80MG; 20MG | A089449 001 | Nov 27, 1987 |
| FOAMCOAT | | | |
| GUARDIAN DRUG | 80MG; 20MG | A071793 001 | Sep 04, 1987 |
| FOAMICON | | | |
| NOVARTIS | 80MG; 20MG | A072687 001 | Jun 28, 1989 |
| GAVISCON | | | |
| + SANOFI AVENTIS US | 160MG; 40MG | N018685 002 | Dec 09, 1983 |

AMANTADINE HYDROCHLORIDE

CAPSULE; ORAL

AMANTADINE HYDROCHLORIDE

| | | | |
|-------------------|-------|-------------|--------------|
| ACTAVIS ELIZABETH | 100MG | A077659 001 | Feb 23, 2006 |
| LANNETT CO INC | 100MG | A209221 001 | Jun 15, 2017 |
| WATSON LABS | 100MG | A071382 001 | Jan 21, 1987 |

SYMADINE

| | | | |
|--------|-------|-------------|--------------|
| SOLVAY | 100MG | A071000 001 | Sep 04, 1986 |
|--------|-------|-------------|--------------|

SYMMETREL

| | | | |
|---------------|----------|-------------|--|
| + ENDO PHARMS | 100MG ** | N016020 001 | |
|---------------|----------|-------------|--|

SYRUP; ORAL

AMANTADINE HYDROCHLORIDE

| | | | |
|------------------|----------|-------------|--------------|
| G AND W LABS INC | 50MG/5ML | A072655 001 | Oct 30, 1990 |
| LANNETT CO INC | 50MG/5ML | A076352 001 | Sep 10, 2004 |
| TEVA PHARMS | 50MG/5ML | A073115 001 | Aug 23, 1991 |
| VINTAGE | 50MG/5ML | A077992 001 | Dec 12, 2006 |

SYMMETREL

| | | | |
|---------------|-------------|-------------|--|
| + ENDO PHARMS | 50MG/5ML ** | N016023 002 | |
|---------------|-------------|-------------|--|

TABLET; ORAL

SYMMETREL

| | | | |
|---------------|----------|-------------|--|
| + ENDO PHARMS | 100MG ** | N018101 001 | |
|---------------|----------|-------------|--|

AMBENONIUM CHLORIDE

TABLET; ORAL

MYTELASE

| | | | |
|-------------------|------|-------------|--|
| SANOFI AVENTIS US | 10MG | N010155 002 | |
|-------------------|------|-------------|--|

AMCINONIDE

CREAM; TOPICAL

CYCLOCORT

| | | | |
|------------|--------|-------------|--|
| + ASTELLAS | 0.025% | N018116 001 | |
| + ASTELLAS | 0.1% | N018116 002 | |

LOTION; TOPICAL

CYCLOCORT

| | | | |
|------------|------|-------------|--------------|
| + ASTELLAS | 0.1% | N019729 001 | Jun 13, 1988 |
|------------|------|-------------|--------------|

OINTMENT; TOPICAL

CYCLOCORT

| | | | |
|------------|------|-------------|--|
| + ASTELLAS | 0.1% | N018498 001 | |
|------------|------|-------------|--|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMDINOCILLIN

INJECTABLE; INJECTION

COACTIN

| | | | |
|-------|------------|-------------|--------------|
| ROCHE | 250MG/VIAL | N050565 001 | Dec 21, 1984 |
| | 500MG/VIAL | N050565 002 | Dec 21, 1984 |
| | 1GM/VIAL | N050565 003 | Dec 21, 1984 |

AMIFOSTINE

INJECTABLE; INJECTION

ETHYOL

| | | | |
|-------------------|------------|-------------|--------------|
| CLINIGEN HLTHCARE | 375MG/VIAL | N020221 002 | Sep 10, 1999 |
|-------------------|------------|-------------|--------------|

AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKACIN SULFATE

| | | | |
|----------------------|-------------------|-------------|--------------|
| ABBOTT | EQ 250MG BASE/ML | A063265 001 | Nov 30, 1994 |
| | EQ 250MG BASE/ML | A063266 001 | Oct 31, 1994 |
| HOSPIRA | EQ 50MG BASE/ML | A063263 001 | Nov 30, 1994 |
| | EQ 50MG BASE/ML | A063350 001 | Jul 30, 1993 |
| | EQ 62.5MG BASE/ML | A063283 001 | Oct 31, 1994 |
| | EQ 250MG BASE/ML | A063264 001 | Nov 30, 1994 |
| | EQ 250MG BASE/ML | A063350 002 | Jul 30, 1993 |
| | EQ 250MG BASE/ML | A064098 001 | Jun 26, 1995 |
| IGI LABS INC | EQ 250MG BASE/ML | A064099 001 | Jun 20, 1995 |
| | EQ 50MG BASE/ML | A063167 001 | Dec 14, 1995 |
| | EQ 250MG BASE/ML | A063169 001 | Dec 14, 1995 |
| TEVA PHARMS USA | EQ 50MG BASE/ML | A064045 001 | Sep 28, 1993 |
| WEST-WARD PHARMS INT | EQ 50MG BASE/ML | A063274 001 | May 18, 1992 |
| | EQ 250MG BASE/ML | A063275 001 | May 18, 1992 |

AMIKACIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | |
|---------|---------------------|-------------|--------------|
| HOSPIRA | EQ 500MG BASE/100ML | A064146 001 | Apr 02, 1997 |
|---------|---------------------|-------------|--------------|

AMIKIN

| | | | |
|-----------------------------------------------------|---------------------|-------------|--------------|
| APOTHECON | EQ 50MG BASE/ML | A062311 001 | |
| | EQ 50MG BASE/ML | A062562 001 | Sep 20, 1984 |
| + | EQ 50MG BASE/ML ** | N050495 001 | |
| | EQ 250MG BASE/ML | A062311 002 | |
| + | EQ 250MG BASE/ML | A062562 002 | Sep 20, 1984 |
| | EQ 250MG BASE/ML ** | N050495 002 | |
| AMIKIN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | |
| APOTHECON | EQ 5MG BASE/ML | N050618 002 | Nov 30, 1987 |
| | EQ 10MG BASE/ML | N050618 001 | Nov 30, 1987 |

AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

| | | | |
|------------------|------------------------|---------------------------|--------------|
| TEVA | EQ 5MG ANHYDROUS; 50MG | A070795 001 | Apr 17, 1988 |
| WATSON LABS | EQ 5MG ANHYDROUS; 50MG | A073334 001 | Jul 19, 1991 |
| YAOPHARMA CO LTD | EQ 5MG ANHYDROUS; 50MG | A073357 001 | Nov 27, 1991 |
| HYDRO-RIDE | | | |
| PAR PHARM | EQ 5MG ANHYDROUS; 50MG | A070347 001 | Dec 25, 1990 |
| MODURETIC 5-50 | | | |
| + | MERCK | EQ 5MG ANHYDROUS; 50MG ** | N018201 001 |

AMINO ACIDS

INJECTABLE; INJECTION

AMINESS 5.2% ESSENTIAL AMINO ACIDS W/ HISTADINE

| | | | |
|---------|--------------------|-------------|--------------|
| HOSPIRA | 5.2% (5.2GM/100ML) | N018901 001 | Apr 06, 1984 |
|---------|--------------------|-------------|--------------|

AMINOSYN 10% (PH6)

| | | | |
|-----------------|------------------|-------------|--------------|
| ICU MEDICAL INC | 10% (10GM/100ML) | N017673 008 | Nov 18, 1985 |
|-----------------|------------------|-------------|--------------|

AMINOSYN 3.5%

| | | | |
|-----------------|--------------------|-------------|--|
| ICU MEDICAL INC | 3.5% (3.5GM/100ML) | N017789 004 | |
|-----------------|--------------------|-------------|--|

AMINOSYN 3.5% IN PLASTIC CONTAINER

| | | | |
|--------|--------------------|-------------|--------------|
| ABBOTT | 3.5% (3.5GM/100ML) | N018804 001 | May 15, 1984 |
| | 3.5% (3.5GM/100ML) | N018875 001 | Aug 08, 1984 |

AMINOSYN 5%

| | | | |
|-----------------|----------------|-------------|--|
| ICU MEDICAL INC | 5% (5GM/100ML) | N017673 001 | |
|-----------------|----------------|-------------|--|

AMINOSYN 7%

| | | | |
|-----------------|----------------|-------------|--|
| ICU MEDICAL INC | 7% (7GM/100ML) | N017673 002 | |
|-----------------|----------------|-------------|--|

AMINOSYN 7% (PH6)

| | | | |
|-----------------|----------------|-------------|--------------|
| ICU MEDICAL INC | 7% (7GM/100ML) | N017673 006 | Nov 18, 1985 |
|-----------------|----------------|-------------|--------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS

INJECTABLE; INJECTION

| | | | | |
|------------------------------------------------|----------------------|---------|-----|--------------|
| AMINOSYN 8.5% (PH6) | | | | |
| ICU MEDICAL INC | 8.5% (8.5GM/100ML) | N017673 | 007 | Nov 18, 1985 |
| AMINOSYN II 10% | | | | |
| ICU MEDICAL INC | 10% (10GM/100ML) | N019438 | 005 | Apr 03, 1986 |
| AMINOSYN II 3.5% | | | | |
| ICU MEDICAL INC | 3.5% (3.5GM/100ML) | N019438 | 001 | Apr 03, 1986 |
| AMINOSYN II 3.5% IN PLASTIC CONTAINER | | | | |
| ABBOTT | 3.5% (3.5GM/100ML) | N019491 | 001 | Oct 10, 1986 |
| AMINOSYN II 5% | | | | |
| ICU MEDICAL INC | 5% (5GM/100ML) | N019438 | 002 | Apr 03, 1986 |
| AMINOSYN II 7% | | | | |
| ICU MEDICAL INC | 7% (7GM/100ML) | N019438 | 003 | Apr 03, 1986 |
| AMINOSYN II 8.5% | | | | |
| ICU MEDICAL INC | 8.5% (8.5GM/100ML) | N019438 | 004 | Apr 03, 1986 |
| AMINOSYN-HBC 7% | | | | |
| ICU MEDICAL INC | 7% (7GM/100ML) | N019374 | 001 | Jul 12, 1985 |
| AMINOSYN-HBC 7% IN PLASTIC CONTAINER | | | | |
| ABBOTT | 7% (7GM/100ML) | N019400 | 001 | Jul 23, 1986 |
| AMINOSYN-HF 8% | | | | |
| ICU MEDICAL INC | 8% (8GM/100ML) | A020345 | 001 | Apr 04, 1996 |
| AMINOSYN-RF 5.2% | | | | |
| ICU MEDICAL INC | 5.2% (5.2GM/100ML) | N018429 | 001 | |
| BRANCHAMIN 4% | | | | |
| BAXTER HLTHCARE | 4% (4GM/100ML) | N018678 | 001 | Sep 28, 1984 |
| BRANCHAMIN 4% IN PLASTIC CONTAINER | | | | |
| BAXTER HLTHCARE | 4% (4GM/100ML) | N018684 | 001 | Sep 28, 1984 |
| FREAMINE 8.5% | | | | |
| B BRAUN | 8.5% (8.5GM/100ML) | N016822 | 001 | |
| FREAMINE II 8.5% | | | | |
| B BRAUN | 8.5% (8.5GM/100ML) | N016822 | 002 | |
| HEPATASOL 8% | | | | |
| BAXTER HLTHCARE | 8% (8GM/100ML) | A020360 | 001 | Apr 04, 1996 |
| NEOPHAM 6.4% | | | | |
| HOSPIRA | 6.4% (6.4GM/100ML) | N018792 | 001 | Jan 17, 1984 |
| NOVAMINE 11.4% | | | | |
| HOSPIRA INC | 11.4% (11.4GM/100ML) | N017957 | 003 | Aug 09, 1982 |
| NOVAMINE 15% | | | | |
| HOSPIRA INC | 15% (75GM/500ML) | N017957 | 004 | Nov 28, 1986 |
| NOVAMINE 15% SULFITE FREE IN PLASTIC CONTAINER | | | | |
| BAXTER HLTHCARE | 15% (15GM/100ML) ** | N020107 | 001 | Feb 05, 1993 |
| NOVAMINE 8.5% | | | | |
| HOSPIRA INC | 8.5% (8.5GM/100ML) | N017957 | 002 | Aug 09, 1982 |
| RENAMIN W/O ELECTROLYTES | | | | |
| BAXTER HLTHCARE | 6.5% (6.5GM/100ML) | N017493 | 007 | Oct 15, 1982 |
| TRAVASOL 10% W/O ELECTROLYTES | | | | |
| BAXTER HLTHCARE | 10% (10GM/100ML) | N017493 | 006 | |
| TRAVASOL 5.5% W/O ELECTROLYTES | | | | |
| BAXTER HLTHCARE | 5.5% (5.5GM/100ML) | N017493 | 004 | |
| TRAVASOL 8.5% W/O ELECTROLYTES | | | | |
| BAXTER HLTHCARE | 8.5% (8.5GM/100ML) | N017493 | 005 | |

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | | | |
|-----------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------|---------|-----|--------------|
| AMINOSYN II 3.5% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER | | | | |
| ABBOTT | 3.5%; 36.8MG/100ML; 25GM/100ML; 51MG/100ML ; 22.4MG/100ML; 261MG/100ML; 205MG/100ML | N019714 | 001 | Sep 12, 1988 |
| HOSPIRA INC | 3.5%; 36.8MG/100ML; 25GM/100ML; 51MG/100ML ; 22.4MG/100ML; 261MG/100ML; 205MG/100ML | N019683 | 001 | Nov 07, 1988 |
| AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER | | | | |
| ABBOTT | 4.25%; 36.8MG/100ML; 20GM/100ML; 51MG/100M L; 22.4MG/100ML; 261MG/100ML; 205MG/100ML | N019714 | 002 | Sep 12, 1988 |
| HOSPIRA INC | 4.25%; 36.8MG/100ML; 20GM/100ML; 51MG/100M L; 22.4MG/100ML; 261MG/100ML; 205MG/100ML | N019683 | 002 | Nov 07, 1988 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER

| | | | |
|--------|------------------------------------------|-------------|--------------|
| ABBOTT | 4.25%;36.8MG/100ML;25GM/100ML;51MG/100ML | N019714 004 | Sep 12, 1988 |
| | L;22.4MG/100ML;261MG/100ML;205MG/100ML | | |

| | | | |
|-------------|------------------------------------------|-------------|--------------|
| HOSPIRA INC | 4.25%;36.8MG/100ML;25GM/100ML;51MG/100ML | N019683 003 | Nov 07, 1988 |
| | L;22.4MG/100ML;261MG/100ML;205MG/100ML | | |

AMINOSYN II 5% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER

| | | | |
|--------|-----------------------------------------|-------------|--------------|
| ABBOTT | 5%;36.8MG/100ML;25GM/100ML;51MG/100ML;2 | N019714 003 | Sep 12, 1988 |
| | 2.4MG/100ML;261MG/100ML;205MG/100ML | | |

| | | | |
|-------------|-----------------------------------------|-------------|--------------|
| HOSPIRA INC | 5%;36.8MG/100ML;25GM/100ML;51MG/100ML;2 | N019683 004 | Nov 07, 1988 |
| | 2.4MG/100ML;261MG/100ML;205MG/100ML | | |

AMINO ACIDS; DEXTROSE

INJECTABLE; INJECTION

AMINOSYN 3.5% W/ DEXTROSE 25% IN PLASTIC CONTAINER

| | | | |
|--------|-----------------|-------------|--------------|
| ABBOTT | 3.5%;25GM/100ML | N019118 001 | Oct 11, 1984 |
|--------|-----------------|-------------|--------------|

AMINOSYN 3.5% W/ DEXTROSE 5% IN PLASTIC CONTAINER

| | | | |
|--------|----------------|-------------|--------------|
| ABBOTT | 3.5%;5GM/100ML | N019120 001 | Oct 11, 1984 |
|--------|----------------|-------------|--------------|

AMINOSYN 4.25% W/ DEXTROSE 25% IN PLASTIC CONTAINER

| | | | |
|--------|------------------|-------------|--------------|
| ABBOTT | 4.25%;25GM/100ML | N019119 001 | Oct 11, 1984 |
|--------|------------------|-------------|--------------|

AMINOSYN II 3.5% IN DEXTROSE 25% IN PLASTIC CONTAINER

| | | | |
|--------|-----------------|-------------|--------------|
| ABBOTT | 3.5%;25GM/100ML | N019505 002 | Nov 07, 1986 |
|--------|-----------------|-------------|--------------|

| | | | |
|--|-----------------|-------------|--------------|
| | 3.5%;25GM/100ML | N019713 006 | Sep 09, 1988 |
|--|-----------------|-------------|--------------|

| | | | |
|---------|-----------------|-------------|--------------|
| HOSPIRA | 3.5%;25GM/100ML | N019681 001 | Nov 01, 1988 |
|---------|-----------------|-------------|--------------|

AMINOSYN II 3.5% IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | |
|--------|----------------|-------------|--------------|
| ABBOTT | 3.5%;5GM/100ML | N019506 001 | Nov 07, 1986 |
|--------|----------------|-------------|--------------|

| | | | |
|--|----------------|-------------|--------------|
| | 3.5%;5GM/100ML | N019713 002 | Sep 09, 1988 |
|--|----------------|-------------|--------------|

| | | | |
|---------|----------------|-------------|--------------|
| HOSPIRA | 3.5%;5GM/100ML | N019681 002 | Nov 01, 1988 |
|---------|----------------|-------------|--------------|

AMINOSYN II 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER

| | | | |
|--------|------------------|-------------|--------------|
| ABBOTT | 4.25%;10GM/100ML | N019713 001 | Sep 09, 1988 |
|--------|------------------|-------------|--------------|

| | | | |
|---------|------------------|-------------|--------------|
| HOSPIRA | 4.25%;10GM/100ML | N019681 004 | Nov 01, 1988 |
|---------|------------------|-------------|--------------|

AMINOSYN II 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER

| | | | |
|--------|------------------|-------------|--------------|
| ABBOTT | 4.25%;20GM/100ML | N019713 004 | Sep 09, 1988 |
|--------|------------------|-------------|--------------|

| | | | |
|---------|------------------|-------------|--------------|
| HOSPIRA | 4.25%;20GM/100ML | N019681 005 | Nov 01, 1988 |
|---------|------------------|-------------|--------------|

AMINOSYN II 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER

| | | | |
|--------|------------------|-------------|--------------|
| ABBOTT | 4.25%;25GM/100ML | N019504 002 | Nov 07, 1986 |
|--------|------------------|-------------|--------------|

| | | | |
|--|------------------|-------------|--------------|
| | 4.25%;25GM/100ML | N019713 005 | Sep 09, 1988 |
|--|------------------|-------------|--------------|

| | | | |
|---------|------------------|-------------|--------------|
| HOSPIRA | 4.25%;25GM/100ML | N019681 003 | Nov 01, 1988 |
|---------|------------------|-------------|--------------|

AMINOSYN II 5% IN DEXTROSE 25% IN PLASTIC CONTAINER

| | | | |
|--------|---------------|-------------|--------------|
| ABBOTT | 5%;25GM/100ML | N019565 001 | Dec 17, 1986 |
|--------|---------------|-------------|--------------|

| | | | |
|--|---------------|-------------|--------------|
| | 5%;25GM/100ML | N019713 003 | Sep 09, 1988 |
|--|---------------|-------------|--------------|

| | | | |
|---------|---------------|-------------|--------------|
| HOSPIRA | 5%;25GM/100ML | N019681 006 | Nov 01, 1988 |
|---------|---------------|-------------|--------------|

TRAVASOL 2.75% IN DEXTROSE 10% IN PLASTIC CONTAINER

| | | | |
|-----------------|------------------|-------------|--------------|
| BAXTER HLTHCARE | 2.75%;10GM/100ML | N019520 002 | Sep 23, 1988 |
|-----------------|------------------|-------------|--------------|

TRAVASOL 2.75% IN DEXTROSE 15% IN PLASTIC CONTAINER

| | | | |
|-----------------|------------------|-------------|--------------|
| BAXTER HLTHCARE | 2.75%;15GM/100ML | N019520 003 | Sep 23, 1988 |
|-----------------|------------------|-------------|--------------|

TRAVASOL 2.75% IN DEXTROSE 20% IN PLASTIC CONTAINER

| | | | |
|-----------------|------------------|-------------|--------------|
| BAXTER HLTHCARE | 2.75%;20GM/100ML | N019520 004 | Sep 23, 1988 |
|-----------------|------------------|-------------|--------------|

TRAVASOL 2.75% IN DEXTROSE 25% IN PLASTIC CONTAINER

| | | | |
|-----------------|------------------|-------------|--------------|
| BAXTER HLTHCARE | 2.75%;25GM/100ML | N019520 005 | Sep 23, 1988 |
|-----------------|------------------|-------------|--------------|

TRAVASOL 2.75% IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | |
|-----------------|-----------------|-------------|--------------|
| BAXTER HLTHCARE | 2.75%;5GM/100ML | N019520 001 | Sep 23, 1988 |
|-----------------|-----------------|-------------|--------------|

TRAVASOL 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER

| | | | |
|-----------------|------------------|-------------|--------------|
| BAXTER HLTHCARE | 4.25%;10GM/100ML | N019520 007 | Sep 23, 1988 |
|-----------------|------------------|-------------|--------------|

TRAVASOL 4.25% IN DEXTROSE 15% IN PLASTIC CONTAINER

| | | | |
|-----------------|------------------|-------------|--------------|
| BAXTER HLTHCARE | 4.25%;15GM/100ML | N019520 008 | Sep 23, 1988 |
|-----------------|------------------|-------------|--------------|

TRAVASOL 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER

| | | | |
|-----------------|------------------|-------------|--------------|
| BAXTER HLTHCARE | 4.25%;20GM/100ML | N019520 009 | Sep 23, 1988 |
|-----------------|------------------|-------------|--------------|

TRAVASOL 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER

| | | | |
|-----------------|------------------|-------------|--------------|
| BAXTER HLTHCARE | 4.25%;25GM/100ML | N019520 010 | Sep 23, 1988 |
|-----------------|------------------|-------------|--------------|

TRAVASOL 4.25% IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | |
|-----------------|-----------------|-------------|--------------|
| BAXTER HLTHCARE | 4.25%;5GM/100ML | N019520 006 | Sep 23, 1988 |
|-----------------|-----------------|-------------|--------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM ACETATE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 4.25% W/ ELECT AND ADJUSTED PHOSPHATE IN DEXTROSE 10% IN PLASTIC CONTAINER

| | | | |
|--------|------------------------------------------------------------------------------------------|-------------|--------------|
| ABBOTT | 4.25%;10GM/100ML;51MG/100ML;176.5MG/100 ML;22.4MG/100ML;104.5MG/100ML;205MG/100 ML | N019712 002 | Sep 08, 1988 |
|--------|------------------------------------------------------------------------------------------|-------------|--------------|

| | | | |
|-------------|------------------------------------------------------------------------------------------|-------------|--------------|
| HOSPIRA INC | 4.25%;10GM/100ML;51MG/100ML;176.5MG/100 ML;22.4MG/100ML;104.5MG/100ML;205MG/100 ML | N019682 003 | Nov 01, 1988 |
|-------------|------------------------------------------------------------------------------------------|-------------|--------------|

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 3.5% W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER

| | | | |
|--------|---------------------------------------------------------------------|-------------|--------------|
| ABBOTT | 3.5%;25GM/100ML;51MG/100ML;22.4MG/100ML ;261MG/100ML;205MG/100ML | N019564 002 | Dec 16, 1986 |
|--------|---------------------------------------------------------------------|-------------|--------------|

AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER

| | | | |
|--------|----------------------------------------------------------------------|-------------|--------------|
| ABBOTT | 4.25%;25GM/100ML;51MG/100ML;22.4MG/100M L;261MG/100ML;205MG/100ML | N019564 004 | Dec 16, 1986 |
|--------|----------------------------------------------------------------------|-------------|--------------|

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | |
|--------|-------------------------------------------------------------------|-------------|--------------|
| ABBOTT | 3.5%;5GM/100ML;30MG/100ML;97MG/100ML;12 0MG/100ML;49.3MG/100ML | N019564 001 | Dec 16, 1986 |
|--------|-------------------------------------------------------------------|-------------|--------------|

| | | | |
|--|-------------------------------------------------------------------|-------------|--------------|
| | 3.5%;5GM/100ML;30MG/100ML;97MG/100ML;12 0MG/100ML;49.3MG/100ML | N019712 001 | Sep 08, 1988 |
|--|-------------------------------------------------------------------|-------------|--------------|

| | | | |
|-------------|-------------------------------------------------------------------|-------------|--------------|
| HOSPIRA INC | 3.5%;5GM/100ML;30MG/100ML;97MG/100ML;12 0MG/100ML;49.3MG/100ML | N019682 001 | Nov 01, 1988 |
|-------------|-------------------------------------------------------------------|-------------|--------------|

AMINOSYN II 4.25% M IN DEXTROSE 10% IN PLASTIC CONTAINER

| | | | |
|--------|---------------------------------------------------------------------|-------------|--------------|
| ABBOTT | 4.25%;10GM/100ML;30MG/100ML;97MG/100ML; 120MG/100ML;49.3MG/100ML | N019564 003 | Dec 16, 1986 |
|--------|---------------------------------------------------------------------|-------------|--------------|

| | | | |
|-------------|--------------------------------------------------------------------|-------------|--------------|
| HOSPIRA INC | 4.25%;5GM/100ML;30MG/100ML;97MG/100ML;1 20MG/100ML;49.3MG/100ML | N019682 002 | Nov 01, 1988 |
|-------------|--------------------------------------------------------------------|-------------|--------------|

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 10% IN PLASTIC CONTAINER

| | | | |
|-----------------|---------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE | 2.75%;10GM/100ML;51MG/100ML;261MG/100ML ;216MG/100ML;112MG/100ML | N020147 002 | Oct 23, 1995 |
|-----------------|---------------------------------------------------------------------|-------------|--------------|

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 15% IN PLASTIC CONTAINER

| | | | |
|-----------------|---------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE | 2.75%;15GM/100ML;51MG/100ML;261MG/100ML ;216MG/100ML;112MG/100ML | N020147 003 | Oct 23, 1995 |
|-----------------|---------------------------------------------------------------------|-------------|--------------|

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 20% IN PLASTIC CONTAINER

| | | | |
|-----------------|---------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE | 2.75%;20GM/100ML;51MG/100ML;261MG/100ML ;216MG/100ML;112MG/100ML | N020147 004 | Oct 23, 1995 |
|-----------------|---------------------------------------------------------------------|-------------|--------------|

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER

| | | | |
|-----------------|---------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE | 2.75%;25GM/100ML;51MG/100ML;261MG/100ML ;216MG/100ML;112MG/100ML | N020147 005 | Oct 23, 1995 |
|-----------------|---------------------------------------------------------------------|-------------|--------------|

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | |
|-----------------|--------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE | 2.75%;5GM/100ML;51MG/100ML;261MG/100ML; 216MG/100ML;112MG/100ML | N020147 001 | Oct 23, 1995 |
|-----------------|--------------------------------------------------------------------|-------------|--------------|

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 10% IN PLASTIC CONTAINER

| | | | |
|-----------------|--------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE | 4.25%;10GM/100ML;51MG/100ML;261MG/100ML ;297MG/100ML;77MG/100ML | N020147 007 | Oct 23, 1995 |
|-----------------|--------------------------------------------------------------------|-------------|--------------|

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 15% IN PLASTIC CONTAINER

| | | | |
|-----------------|--------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE | 4.25%;15GM/100ML;51MG/100ML;261MG/100ML ;297MG/100ML;77MG/100ML | N020147 008 | Oct 23, 1995 |
|-----------------|--------------------------------------------------------------------|-------------|--------------|

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 20% IN PLASTIC CONTAINER

| | | | |
|-----------------|--------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE | 4.25%;20GM/100ML;51MG/100ML;261MG/100ML ;297MG/100ML;77MG/100ML | N020147 009 | Oct 23, 1995 |
|-----------------|--------------------------------------------------------------------|-------------|--------------|

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER

| | | | |
|-----------------|--------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE | 4.25%;25GM/100ML;51MG/100ML;261MG/100ML ;297MG/100ML;77MG/100ML | N020147 010 | Oct 23, 1995 |
|-----------------|--------------------------------------------------------------------|-------------|--------------|

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | |
|-----------------|-------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE | 4.25%;5GM/100ML;51MG/100ML;261MG/100ML; 297MG/100ML;77MG/100ML | N020147 006 | Oct 23, 1995 |
|-----------------|-------------------------------------------------------------------|-------------|--------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 3.5% M IN PLASTIC CONTAINER

| | | | |
|--------|-----------------------------------------------------------|-------------|--------------|
| ABBOTT | 3.5%; 21MG/100ML; 40MG/100ML; 128MG/100ML; 234MG/100ML | N018804 002 | May 15, 1984 |
| | 3.5%; 21MG/100ML; 40MG/100ML; 128MG/100ML; 234MG/100ML | N018875 002 | Aug 08, 1984 |

AMINO ACIDS; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 3.5% M

| | | | |
|-----------------|--------------------------------------------|-------------|--|
| ICU MEDICAL INC | 3.5%; 21MG/100ML; 128MG/100ML; 234MG/100ML | N017789 005 | |
|-----------------|--------------------------------------------|-------------|--|

AMINO ACIDS; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M IN PLASTIC CONTAINER

| | | | |
|--------|------------------------------------------------------------|-------------|--------------|
| ABBOTT | 3.5%; 32MG/100ML; 128MG/100ML; 222MG/100ML ; 49MG/100ML | N019493 001 | Oct 16, 1986 |
|--------|------------------------------------------------------------|-------------|--------------|

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM ACETATE; POTASSIUM CHLORIDE; SODIUM ACETATE

INJECTABLE; INJECTION

VEINAMINE 8%

| | | | |
|-------------|----------------------------------------------------------|-------------|--|
| HOSPIRA INC | 8%; 61MG/100ML; 211MG/100ML; 56MG/100ML; 38 8MG/100ML | N017957 001 | |
|-------------|----------------------------------------------------------|-------------|--|

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 10% W/ ELECTROLYTES

| | | | |
|-----------------|-----------------------------------------------------------|-------------|--------------|
| ICU MEDICAL INC | 10%; 102MG/100ML; 45MG/100ML; 522MG/100ML; 410MG/100ML | N019437 004 | Apr 03, 1986 |
|-----------------|-----------------------------------------------------------|-------------|--------------|

AMINOSYN II 7% W/ ELECTROLYTES

| | | | |
|-----------------|-----------------------------------------------------------|-------------|--------------|
| ICU MEDICAL INC | 7%; 102MG/100ML; 45MG/100ML; 522MG/100ML; 4 10MG/100ML | N019437 006 | Apr 03, 1986 |
|-----------------|-----------------------------------------------------------|-------------|--------------|

AMINOSYN II 8.5% W/ ELECTROLYTES

| | | | |
|-----------------|-------------------------------------------------------------|-------------|--------------|
| ICU MEDICAL INC | 8.5%; 102MG/100ML; 45MG/100ML; 522MG/100ML ; 410MG/100ML | N019437 005 | Apr 03, 1986 |
|-----------------|-------------------------------------------------------------|-------------|--------------|

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC

INJECTABLE; INJECTION

AMINOSYN II 8.5% W/ELECTROLYTES

| | | | |
|-----------------|-------------------------------------------------------------|-------------|--------------|
| ICU MEDICAL INC | 8.5%; 102MG/100ML; 492MG/100ML; 60MG/100ML ; 425MG/100ML | N019437 008 | Oct 25, 2002 |
|-----------------|-------------------------------------------------------------|-------------|--------------|

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M

| | | | |
|-----------------|----------------------------------------------------------|-------------|--------------|
| ICU MEDICAL INC | 3.5%; 30MG/100ML; 97MG/100ML; 120MG/100ML; 49MG/100ML | N019437 007 | Apr 03, 1986 |
|-----------------|----------------------------------------------------------|-------------|--------------|

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TRAVASOL 3.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER

| | | | |
|-----------------|------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE | 3.5%; 51MG/100ML; 131MG/100ML; 218MG/100ML ; 35MG/100ML | N020177 001 | Oct 23, 1995 |
|-----------------|------------------------------------------------------------|-------------|--------------|

TRAVASOL 3.5% W/ ELECTROLYTES

| | | | |
|-----------------|------------------------------------------------------------|-------------|--|
| BAXTER HLTHCARE | 3.5%; 51MG/100ML; 131MG/100ML; 218MG/100ML ; 35MG/100ML | N017493 003 | |
|-----------------|------------------------------------------------------------|-------------|--|

TRAVASOL 5.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER

| | | | |
|-----------------|--------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE | 5.5%; 102MG/100ML; 522MG/100ML; 431MG/100M L; 224MG/100ML | N020173 001 | Oct 27, 1995 |
|-----------------|--------------------------------------------------------------|-------------|--------------|

TRAVASOL 5.5% W/ ELECTROLYTES

| | | | |
|-----------------|--------------------------------------------------------------|-------------|--|
| BAXTER HLTHCARE | 5.5%; 102MG/100ML; 522MG/100ML; 431MG/100M L; 224MG/100ML | N017493 001 | |
|-----------------|--------------------------------------------------------------|-------------|--|

TRAVASOL 8.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER

| | | | |
|-----------------|--------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE | 8.5%; 102MG/100ML; 522MG/100ML; 594MG/100M L; 154MG/100ML | N020173 002 | Oct 27, 1995 |
|-----------------|--------------------------------------------------------------|-------------|--------------|

TRAVASOL 8.5% W/ ELECTROLYTES

| | | | |
|-----------------|--------------------------------------------------------------|-------------|--|
| BAXTER HLTHCARE | 8.5%; 102MG/100ML; 522MG/100ML; 594MG/100M L; 154MG/100ML | N017493 002 | |
|-----------------|--------------------------------------------------------------|-------------|--|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 7% W/ ELECTROLYTES

ICU MEDICAL INC 7%;102MG/100ML;522MG/100ML;410MG/100ML N017789 002

AMINOSYN 8.5% W/ ELECTROLYTES

ICU MEDICAL INC 8.5%;102MG/100ML;522MG/100ML;410MG/100ML N017673 005
LAMINOCAPROIC ACID

INJECTABLE; INJECTION

AMICAR

+ CLOVER PHARMS 250MG/ML ** N015229 002

AMINOCAPROIC ACID

ABRAXIS PHARM 250MG/ML A070522 001 Jun 17, 1986

BAXTER HLTHCARE 250MG/ML N018590 001 Oct 29, 1982

HOSPIRA 250MG/ML A070888 001 Jun 16, 1988

SYRUP; ORAL

AMINOCAPROIC ACID

AKORN 1.25GM/5ML A074759 001 Sep 02, 1998

TABLET; ORAL

AMINOCAPROIC

AKORN 500MG A075602 001 May 24, 2001

AMINOGLUTETHIMIDE

TABLET; ORAL

CYTADREN

NOVARTIS 250MG N018202 001

AMINOHIPPURATE SODIUM

INJECTABLE; INJECTION

AMINOHIPPURATE SODIUM

MERCCK 20% N005619 001

AMINOPHYLLINE

ENEMA; RECTAL

SOMOPHYLLIN

FISONS 300MG/5ML N018232 001 Apr 02, 1982

INJECTABLE; INJECTION

AMINOPHYLLIN

GD SEARLE LLC 25MG/ML A087243 001 May 24, 1982

25MG/ML A087621 001 May 24, 1982

AMINOPHYLLINE

ABRAXIS PHARM 25MG/ML A084568 001

25MG/ML A087200 001

25MG/ML A087250 001 Jan 06, 1982

25MG/ML A087886 001 Aug 30, 1983

25MG/ML A088407 001 Jan 25, 1984

25MG/ML A087239 001

ELKINS SINN 25MG/ML A087601 001 Jul 23, 1982

HOSPIRA 25MG/ML A087209 001 Feb 01, 1982

INTL MEDICATION 25MG/ML A087867 001 Nov 10, 1983

25MG/ML A087868 001 Nov 10, 1983

25MG/ML A086606 001

KING PHARMS 25MG/ML A087240 001

LUITPOLD 25MG/ML A087431 001

PHARMA SERVE NY 25MG/ML A087387 001 Jun 03, 1983

25MG/ML A087392 001 Dec 15, 1983

SMITH AND NEPHEW 25MG/ML A088429 001 May 30, 1985

25MG/ML A088749 001 May 30, 1985

TEVA PARENTERAL 25MG/ML A081142 001 Sep 25, 1991

AMINOPHYLLINE IN SODIUM CHLORIDE 0.45%

HOSPIRA 100MG/100ML A088147 002 May 03, 1983

200MG/100ML A088147 003 May 03, 1983

AMINOPHYLLINE IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

HOSPIRA 100MG/100ML N018924 001 Dec 12, 1984

200MG/100ML N018924 002 Dec 12, 1984

400MG/100ML N018924 003 Dec 12, 1984

500MG/100ML N018924 004 Dec 12, 1984

SOLUTION; ORAL

AMINOPHYLLINE

MORTON GROVE 105MG/5ML A088156 001 Dec 05, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINOPHYLLINE

SOLUTION; ORAL

AMINOPHYLLINE

ROXANE 105MG/5ML A088126 001 Aug 19, 1983

AMINOPHYLLINE DYE FREE

ACTAVIS MID ATLANTIC 105MG/5ML A087727 001 Apr 16, 1982

SOMOPHYLLIN

FISONS 105MG/5ML A086466 001

SOMOPHYLLIN-DF

FISONS 105MG/5ML A087045 001

SUPPOSITORY; RECTAL

TRUPHYLLINE

G AND W LABS 250MG A085498 001 Mar 23, 1983

500MG A085498 002 Jan 03, 1983

TABLET; ORAL

AMINOPHYLLIN

GD SEARLE LLC 100MG N002386 002

200MG N002386 003

AMINOPHYLLINE

ANI PHARMS INC 100MG A085261 004

200MG A085261 002

ASCOT 100MG A087522 001 Feb 12, 1982

200MG A087523 001 Feb 12, 1982

BARR 100MG A088297 001 Aug 19, 1983

200MG A088298 001 Aug 19, 1983

DURAMED PHARMS BARR 100MG A088182 001 Mar 31, 1983

200MG A088183 001 Mar 31, 1983

HALSEY 100MG A084674 001

HIKMA INTL PHARMS 100MG A084540 001

200MG A085003 001

IMPAX LABS 100MG A084574 001

200MG A084576 001

KV PHARM 100MG A085284 001

200MG A085289 001

LANNETT 100MG A084588 001

200MG A084588 002

PAL PAK 100MG A084533 001

PANRAY 100MG A084552 001

200MG A084552 002

PUREPAC PHARM 100MG A084699 001

200MG A085333 001

ROXANE 100MG A087500 001 Feb 09, 1982

200MG A087501 001 Feb 09, 1982

VALEANT PHARM INTL 200MG A084563 001

VANGARD 100MG A088314 001 Oct 03, 1983

200MG A088319 001 Oct 03, 1983

VINTAGE PHARMS 100MG A085409 001

200MG A085410 001

WATSON LABS 100MG A085567 001

200MG A085564 001

TABLET, DELAYED RELEASE; ORAL

AMINOPHYLLINE

IMPAX LABS 100MG A084577 001

200MG A084575 001

TABLICAPS 100MG A084632 002

VALE 100MG A084531 001

200MG A084530 001

TABLET, EXTENDED RELEASE; ORAL

PHYLLOCONTIN

PHARM RES ASSOC 225MG A086760 001

AMINOSALICYLATE SODIUM

POWDER; ORAL

P.A.S. SODIUM

CENTURY PHARMS 4GM/PACKET A080947 001

SODIUM AMINOSALICYLATE

HEXCEL 100% A080097 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMINOSALICYLATE SODIUM

TABLET; ORAL

PARASAL SODIUM

| | | | |
|--------|-------|---------|-----|
| PANRAY | 500MG | N006811 | 006 |
| | 1GM | N006811 | 011 |

SODIUM P.A.S.

| | | | |
|---------|-------|---------|-----|
| LANNETT | 500MG | A080138 | 002 |
|---------|-------|---------|-----|

TEEBACIN

| | | | |
|----------------------|-------|---------|-----|
| CONSOLIDATED MIDLAND | 500MG | N007320 | 002 |
|----------------------|-------|---------|-----|

AMINOSALICYLATE SODIUM; AMINOSALICYLIC ACID

TABLET; ORAL

NEOPASALATE

| | | | |
|---------------------|-------------|---------|-----|
| MEDPOINTE PHARM HLC | 846MG;112MG | A080059 | 002 |
|---------------------|-------------|---------|-----|

AMINOSALICYLIC ACID

TABLET; ORAL

PARASAL

| | | | |
|--------|-------|---------|-----|
| PANRAY | 500MG | N006811 | 001 |
| | 1GM | N006811 | 002 |

AMINOSALICYLIC ACID RESIN COMPLEX

POWDER; ORAL

REZIPAS

| | | | |
|----------------------|------------------|---------|-----|
| BRISTOL MYERS SQUIBB | EQ 500MG BASE/GM | N009052 | 001 |
|----------------------|------------------|---------|-----|

AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION

AMIODARONE HYDROCHLORIDE

| | | | | |
|----------------------|---------|---------|-----|--------------|
| AKORN | 50MG/ML | A076232 | 001 | Jul 05, 2006 |
| BEDFORD | 50MG/ML | A076018 | 001 | Oct 15, 2002 |
| BEDFORD LABS | 50MG/ML | A076299 | 001 | Oct 24, 2002 |
| BEN VENUE | 50MG/ML | A076088 | 001 | Oct 15, 2002 |
| HOSPIRA | 50MG/ML | A076108 | 001 | Oct 15, 2002 |
| INTL MEDICATION SYS | 50MG/ML | N021594 | 001 | Feb 04, 2004 |
| PAR STERILE PRODUCTS | 50MG/ML | A076394 | 001 | Apr 25, 2003 |
| TEVA PHARMS USA | 50MG/ML | A076163 | 001 | Sep 05, 2003 |

CORDARONE

| | | | | |
|--------------------|------------|---------|-----|--------------|
| + WYETH PHARMS INC | 50MG/ML ** | N020377 | 001 | Aug 03, 1995 |
|--------------------|------------|---------|-----|--------------|

NEXTERONE

| | | | | |
|-------------------|---------|---------|-----|--------------|
| + BAXTER HLTHCARE | 50MG/ML | N022325 | 001 | Dec 24, 2008 |
|-------------------|---------|---------|-----|--------------|

TABLET; ORAL

AMIODARONE HYDROCHLORIDE

| | | | | |
|-------|-------|---------|-----|--------------|
| MYLAN | 200MG | A075188 | 001 | Feb 24, 1999 |
| TEVA | 200MG | A074895 | 001 | Apr 16, 1999 |

CORDARONE

| | | | | |
|----------------|----------|---------|-----|--------------|
| + WYETH PHARMS | 200MG ** | N018972 | 001 | Dec 24, 1985 |
|----------------|----------|---------|-----|--------------|

AMITRIPTYLINE HYDROCHLORIDE

CONCENTRATE; ORAL

ENDEP

| | | | |
|-------|---------|---------|-----|
| ROCHE | 40MG/ML | A085749 | 001 |
|-------|---------|---------|-----|

INJECTABLE; INJECTION

AMITRIPTYLINE HYDROCHLORIDE

| | | | |
|-------------|---------|---------|-----|
| WATSON LABS | 10MG/ML | A085594 | 001 |
|-------------|---------|---------|-----|

ELAVIL

| | | | |
|-------------|---------|---------|-----|
| ASTRAZENECA | 10MG/ML | N012704 | 001 |
|-------------|---------|---------|-----|

TABLET; ORAL

AMITID

| | | | |
|----------------------|-------|---------|-----|
| BRISTOL MYERS SQUIBB | 10MG | A086454 | 001 |
| | 25MG | A086454 | 002 |
| | 50MG | A086454 | 003 |
| | 75MG | A086454 | 004 |
| | 100MG | A086454 | 005 |

AMITRIL

| | | | |
|-----------------|-------|---------|-----|
| WARNER CHILCOTT | 10MG | A083939 | 001 |
| | 25MG | A083937 | 001 |
| | 50MG | A083938 | 002 |
| | 75MG | A084957 | 001 |
| | 100MG | A085093 | 001 |
| | 150MG | A086295 | 001 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

AMITRIPTYLINE HYDROCHLORIDE

| | | | | |
|----------------|-------|---------|-----|--------------|
| AM THERAP | 25MG | A088672 | 001 | Nov 20, 1984 |
| | 50MG | A088673 | 001 | Nov 20, 1984 |
| | 75MG | A088674 | 001 | Nov 20, 1984 |
| | 100MG | A088675 | 001 | Nov 20, 1984 |
| ANI PHARMS INC | 10MG | A085031 | 002 | |
| | 25MG | A085031 | 001 | |
| | 50MG | A085031 | 003 | |
| | 75MG | A085031 | 004 | |
| COPLEY PHARM | 10MG | A088421 | 001 | Apr 30, 1984 |
| | 25MG | A088422 | 001 | Apr 30, 1984 |
| | 50MG | A088423 | 001 | Apr 30, 1984 |
| | 75MG | A088424 | 001 | Apr 30, 1984 |
| | 100MG | A088425 | 001 | Apr 30, 1984 |
| | 150MG | A088426 | 001 | Apr 30, 1984 |
| HALSEY | 10MG | A085923 | 001 | |
| | 25MG | A085922 | 001 | |
| | 50MG | A085925 | 001 | |
| | 50MG | A087557 | 001 | Mar 05, 1982 |
| | 75MG | A085926 | 001 | May 20, 1983 |
| | 100MG | A085927 | 001 | May 20, 1983 |
| LEDERLE | 10MG | A086744 | 001 | |
| | 10MG | A087366 | 001 | Jan 04, 1982 |
| | 25MG | A086746 | 001 | |
| | 25MG | A087367 | 001 | May 03, 1982 |
| | 50MG | A086743 | 001 | |
| | 50MG | A087181 | 001 | Jan 04, 1982 |
| | 75MG | A086745 | 001 | |
| | 75MG | A087369 | 001 | Jan 04, 1982 |
| | 100MG | A086747 | 001 | |
| | 100MG | A087368 | 001 | May 03, 1982 |
| | 150MG | A087370 | 001 | Jan 04, 1982 |
| MUTUAL PHARM | 10MG | A085744 | 001 | |
| | 25MG | A085627 | 001 | |
| | 50MG | A085745 | 001 | |
| | 75MG | A085743 | 001 | |
| | 100MG | A085742 | 002 | May 11, 1982 |
| | 150MG | A089423 | 001 | Feb 17, 1987 |
| PAR PHARM | 10MG | A088697 | 001 | Sep 25, 1984 |
| | 25MG | A088698 | 001 | Sep 25, 1984 |
| | 50MG | A088699 | 001 | Sep 25, 1984 |
| | 75MG | A088700 | 001 | Sep 25, 1984 |
| | 100MG | A088701 | 001 | Sep 25, 1984 |
| | 150MG | A088702 | 001 | Sep 25, 1984 |
| PLIVA | 10MG | A088883 | 001 | Sep 26, 1984 |
| | 25MG | A088884 | 001 | Sep 26, 1984 |
| | 50MG | A088885 | 001 | Sep 26, 1984 |
| | 75MG | A088886 | 001 | Sep 26, 1984 |
| | 100MG | A088887 | 001 | Sep 26, 1984 |
| | 150MG | A088888 | 001 | Sep 26, 1984 |
| PUREPAC PHARM | 10MG | A088075 | 001 | Sep 16, 1983 |
| | 10MG | A088084 | 001 | Jul 18, 1983 |
| | 25MG | A088076 | 001 | May 20, 1983 |
| | 25MG | A088085 | 001 | Jul 18, 1983 |
| | 50MG | A088077 | 001 | Sep 16, 1983 |
| | 50MG | A088105 | 001 | Jul 18, 1983 |
| | 75MG | A088078 | 001 | Sep 16, 1983 |
| | 75MG | A088106 | 001 | Jul 18, 1983 |
| | 100MG | A088079 | 001 | Sep 16, 1983 |
| | 100MG | A088107 | 001 | Jul 18, 1983 |
| ROXANE | 10MG | A086002 | 001 | |
| | 10MG | A086144 | 001 | |
| | 25MG | A085944 | 001 | |
| | 25MG | A086145 | 001 | |
| | 50MG | A085945 | 001 | |
| | 50MG | A086143 | 001 | |
| | 75MG | A086004 | 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

AMITRIPTYLINE HYDROCHLORIDE

| | | | | |
|--------------------|-------------|---------|-----|--------------|
| | 75MG | A086147 | 001 | |
| | 100MG | A086003 | 001 | |
| | 100MG | A086146 | 001 | |
| | 150MG | A086090 | 001 | |
| | 150MG | A086148 | 001 | |
| SUN PHARM INDS INC | 10MG | A040816 | 002 | Jun 27, 2008 |
| | 25MG | A040816 | 001 | Jun 27, 2008 |
| | 50MG | A040816 | 003 | Jun 27, 2008 |
| | 75MG | A040816 | 004 | Jun 27, 2008 |
| | 100MG | A040816 | 005 | Jun 27, 2008 |
| | 150MG | A040816 | 006 | Jun 27, 2008 |
| SUPERPHARM | 10MG | A088853 | 001 | Nov 13, 1984 |
| | 25MG | A088854 | 001 | Nov 13, 1984 |
| | 50MG | A088855 | 001 | Nov 13, 1984 |
| | 75MG | A088856 | 001 | Nov 13, 1984 |
| | 100MG | A088857 | 001 | Nov 13, 1984 |
| TEVA | 10MG | A086610 | 001 | |
| | 25MG | A086859 | 001 | |
| | 50MG | A086857 | 001 | |
| | 75MG | A086860 | 001 | |
| | 100MG | A085836 | 001 | |
| | 100MG | A086854 | 001 | |
| | 150MG | A086853 | 001 | |
| UCB INC | 10MG | A085864 | 001 | |
| | 25MG | A085935 | 001 | |
| | 50MG | A085936 | 001 | |
| | 75MG | A086337 | 001 | |
| | 100MG | A086336 | 001 | |
| | 150MG | A086335 | 001 | |
| USL PHARMA | 25MG | A087775 | 001 | Feb 10, 1982 |
| VANGARD | 10MG | A087632 | 001 | Feb 01, 1982 |
| | 50MG | A087616 | 001 | Feb 08, 1982 |
| | 75MG | A087617 | 001 | Feb 05, 1982 |
| | 100MG | A087639 | 001 | Feb 08, 1982 |
| WATSON LABS | 10MG | A085816 | 001 | |
| | 10MG | A088620 | 001 | Mar 02, 1984 |
| | 25MG | A085817 | 001 | |
| | 25MG | A088621 | 001 | Mar 02, 1984 |
| | 50MG | A085815 | 001 | |
| | 50MG | A088622 | 001 | Mar 02, 1984 |
| | 75MG | A085819 | 001 | |
| | 75MG | A088633 | 001 | Mar 02, 1984 |
| | 100MG | A085820 | 001 | |
| | 100MG | A088634 | 001 | Mar 02, 1984 |
| | 150MG | A085821 | 001 | |
| | 150MG | A088635 | 001 | Mar 02, 1984 |
| WEST WARD | 10MG | A087647 | 001 | Mar 05, 1982 |
| | 25MG | A087278 | 001 | |
| ELAVIL | | | | |
| + | ASTRAZENECA | 10MG | ** | N012703 001 |
| + | | 25MG | ** | N012703 003 |
| + | | 50MG | ** | N012703 004 |
| + | | 75MG | ** | N012703 005 |
| + | | 100MG | ** | N012703 006 |
| + | | 150MG | ** | N012703 007 |
| ENDEP | | | | |
| ROCHE | 10MG | A083639 | 001 | |
| | 25MG | A083639 | 002 | |
| | 50MG | A083639 | 003 | |
| | 75MG | A083639 | 004 | |
| | 100MG | A083639 | 005 | |
| | 150MG | A085303 | 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE; CHLORDIAZEPOXIDE

TABLET;ORAL

CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE

| | | | | |
|-----------------------|-----------------------|---------|-----|--------------|
| FRONTIDA BIOPHARM | EQ 12.5MG BASE;5MG | A070765 | 001 | Dec 10, 1986 |
| | EQ 25MG BASE;10MG | A070766 | 001 | Dec 10, 1986 |
| PAR PHARM | EQ 12.5MG BASE;5MG | A072277 | 001 | May 09, 1988 |
| | EQ 25MG BASE;10MG | A072278 | 001 | May 09, 1988 |
| USL PHARMA | EQ 12.5MG BASE;5MG | A070477 | 001 | Jan 12, 1988 |
| | EQ 25MG BASE;10MG | A070478 | 001 | Jan 12, 1988 |
| WATSON LABS | EQ 25MG BASE;10MG | A072053 | 001 | Dec 16, 1988 |
| WATSON LABS TEVA | EQ 12.5MG BASE;5MG | A072052 | 001 | Dec 16, 1988 |
| LIMBITROL | | | | |
| + HERITAGE PHARMS INC | EQ 12.5MG BASE;5MG ** | N016949 | 001 | |
| LIMBITROL DS | | | | |
| + HERITAGE PHARMS INC | EQ 25MG BASE;10MG ** | N016949 | 002 | |

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET;ORAL

| | | | | |
|----------------------------------------------|-------------|---------|-----|--------------|
| ETRAFON 2-10 | | | | |
| SCHERING | 10MG;2MG ** | N014713 | 007 | |
| ETRAFON 2-25 | | | | |
| SCHERING | 25MG;2MG ** | N014713 | 004 | |
| ETRAFON-A | | | | |
| SCHERING | 10MG;4MG ** | N014713 | 002 | |
| ETRAFON-FORTE | | | | |
| SCHERING | 25MG;4MG ** | N014713 | 006 | |
| PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE | | | | |
| FOSUN PHARMA | 10MG;2MG | A071062 | 001 | Nov 27, 1987 |
| | 10MG;4MG | A071862 | 001 | Dec 21, 1987 |
| | 25MG;2MG | A071063 | 001 | Nov 27, 1987 |
| | 25MG;4MG | A071064 | 001 | Nov 27, 1987 |
| | 50MG;4MG | A071863 | 001 | Dec 21, 1987 |
| IVAX SUB TEVA PHARMS | 10MG;2MG | A070935 | 001 | Sep 11, 1986 |
| | 10MG;4MG | A070937 | 001 | Sep 11, 1986 |
| | 25MG;2MG | A070936 | 001 | Sep 11, 1986 |
| | 25MG;4MG | A070938 | 001 | Sep 11, 1986 |
| | 50MG;4MG | A070939 | 001 | Sep 12, 1986 |
| PAR PHARM | 10MG;2MG | A070565 | 001 | Sep 11, 1986 |
| | 10MG;4MG | A070620 | 001 | Sep 11, 1986 |
| | 25MG;2MG | A070621 | 001 | Sep 11, 1986 |
| | 25MG;4MG | A070595 | 001 | Sep 11, 1986 |
| | 50MG;4MG | A070574 | 001 | Sep 11, 1986 |
| SUN PHARM INDUSTRIES | 10MG;2MG | A071077 | 001 | Nov 12, 1986 |
| | 10MG;4MG | A071078 | 001 | Nov 12, 1986 |
| | 25MG;2MG | A070297 | 001 | Nov 12, 1986 |
| | 25MG;4MG | A071079 | 001 | Nov 12, 1986 |
| WATSON LABS | 10MG;2MG | A070373 | 001 | Aug 25, 1986 |
| | 10MG;2MG | A072539 | 001 | Feb 15, 1989 |
| | 10MG;2MG | A073007 | 001 | Oct 17, 1991 |
| | 10MG;4MG | A070375 | 001 | Aug 25, 1986 |
| | 10MG;4MG | A072540 | 001 | Feb 15, 1989 |
| | 10MG;4MG | A073009 | 001 | Oct 17, 1991 |
| | 25MG;2MG | A070374 | 001 | Aug 25, 1986 |
| | 25MG;2MG | A072541 | 001 | Feb 15, 1989 |
| | 25MG;2MG | A073008 | 001 | Oct 17, 1991 |
| | 25MG;4MG | A070376 | 001 | Aug 25, 1986 |
| | 25MG;4MG | A072134 | 001 | Feb 15, 1989 |
| | 25MG;4MG | A073010 | 001 | Oct 17, 1991 |
| | 50MG;4MG | A070377 | 001 | Nov 04, 1986 |
| | 50MG;4MG | A071558 | 001 | Mar 02, 1987 |
| | 50MG;4MG | A072135 | 001 | Feb 15, 1989 |
| TRIAVIL 2-10 | | | | |
| NEW RIVER | 10MG;2MG ** | N014715 | 004 | |
| TRIAVIL 2-25 | | | | |
| NEW RIVER | 25MG;2MG ** | N014715 | 002 | |
| TRIAVIL 4-10 | | | | |
| NEW RIVER | 10MG;4MG ** | N014715 | 003 | |
| TRIAVIL 4-25 | | | | |
| NEW RIVER | 25MG;4MG ** | N014715 | 005 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET; ORAL

TRIAVIL 4-50

NEW RIVER

50MG; 4MG **

N014715 006

AMLEXANOX

PASTE; DENTAL

APHTHASOL

ULURU

5%

N020511 001 Dec 17, 1996

PATCH; TOPICAL

AMLEXANOX

ULURU

2MG

N021727 001 Sep 29, 2004

AMLODIPINE BESYLATE

TABLET; ORAL

AMLODIPINE BESYLATE

AMNEAL PHARMS NY

EQ 2.5MG BASE

A078477 001 Jan 16, 2008

EQ 5MG BASE

A078477 002 Jan 16, 2008

EQ 10MG BASE

A078477 003 Jan 16, 2008

GEDEON RICHTER USA

EQ 2.5MG BASE

A077333 001 Jul 17, 2007

EQ 5MG BASE

A077333 002 Jul 17, 2007

EQ 10MG BASE

A077333 003 Jul 17, 2007

GENPHARM

EQ 2.5MG BASE

A077362 001 Jul 09, 2007

EQ 5MG BASE

A077362 002 Jul 09, 2007

EQ 10MG BASE

A077362 003 Jul 09, 2007

MYLAN PHARMS INC

EQ 2.5MG BASE

A078224 001 Feb 27, 2008

EQ 5MG BASE

A078224 002 Feb 27, 2008

EQ 10MG BASE

A078224 003 Feb 27, 2008

PURACAP PHARM

EQ 2.5MG BASE

A078131 001 Sep 04, 2007

EQ 5MG BASE

A078131 002 Sep 04, 2007

EQ 10MG BASE

A078131 003 Sep 04, 2007

SOVEREIGN PHARMS

EQ 2.5MG BASE

A204900 001 Jul 23, 2015

EQ 5MG BASE

A204900 002 Jul 23, 2015

EQ 10MG BASE

A204900 003 Jul 23, 2015

SUN PHARM INDUSTRIES

EQ 2.5MG BASE

A078081 001 Jan 31, 2008

EQ 5MG BASE

A078081 002 Jan 31, 2008

EQ 10MG BASE

A078081 003 Jan 31, 2008

SUNSHINE LAKE

EQ 2.5MG BASE

A206524 001 May 04, 2018

EQ 5MG BASE

A206524 002 May 04, 2018

EQ 10MG BASE

A206524 003 May 04, 2018

SYNTHON PHARMS

EQ 2.5MG BASE

A077080 001 Jun 27, 2007

EQ 5MG BASE

A077080 002 Jun 27, 2007

EQ 10MG BASE

A077080 003 Jun 27, 2007

YAOPHARMA CO LTD

EQ 2.5MG BASE

A076859 001 Sep 10, 2007

EQ 5MG BASE

A076859 002 Sep 10, 2007

EQ 10MG BASE

A076859 003 Sep 10, 2007

TABLET, ORALLY DISINTEGRATING; ORAL

AMLODIPINE BESYLATE

SYNTHON PHARMS

EQ 2.5MG BASE

N022026 001 Sep 27, 2007

EQ 5MG BASE

N022026 002 Sep 27, 2007

EQ 10MG BASE

N022026 003 Sep 27, 2007

AMLODIPINE MALEATE

TABLET; ORAL

AMVAZ

DR REDDYS LABS INC

2.5MG

N021435 001 Oct 31, 2003

5MG

N021435 002 Oct 31, 2003

10MG

N021435 003 Oct 31, 2003

AMMONIA N-13

INJECTABLE; INTRAVENOUS

AMMONIA N 13

CENTRAL RADIOPHARM

3.75-260mCi/ML

A204539 001 Jun 23, 2015

UNIV TX MD ANDERSON

30mCi-300mCi/8ML (3.75-37.5mCi/ML)

A203933 001 Jun 27, 2014

AMMONIUM CHLORIDE

INJECTABLE; INJECTION

AMMONIUM CHLORIDE

ABBOTT

5MEQ/ML

A083130 001

GD SEARLE LLC

3MEQ/ML

A086205 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMMONIUM CHLORIDE

INJECTABLE; INJECTION

AMMONIUM CHLORIDE 0.9% IN NORMAL SALINE
MCGAW 900MG/100ML

N006580 001

AMMONIUM CHLORIDE 2.14%
B BRAUN 40MEQ/100ML

A085734 001

AMMONIUM LACTATE

CREAM; TOPICAL

LAC-HYDRIN
SUN PHARM INDS INC EQ 12% BASE **

N020508 001 Aug 29, 1996

LOTION; TOPICAL

LAC-HYDRIN
+ SUN PHARM INDS INC EQ 12% BASE **

N019155 001 Apr 24, 1985

AMODIAQUINE HYDROCHLORIDE

TABLET; ORAL

CAMOQUIN HYDROCHLORIDE
PARKE DAVIS EQ 200MG BASE

N006441 001

AMOXAPINE

TABLET; ORAL

AMOXAPINE
UPSHER SMITH LABS 25MG A072943 001 Jun 28, 1991
50MG A072944 001 Jun 28, 1991
100MG A072878 001 Jun 28, 1991
150MG A072879 001 Jun 28, 1991
WATSON PHARMS TEVA 25MG A072418 001 May 11, 1989
50MG A072419 001 May 11, 1989
100MG A072420 001 May 11, 1989
150MG A072421 001 May 11, 1989

ASENDIN

LEDERLE 25MG N018021 001
50MG N018021 002
100MG N018021 003
150MG N018021 004AMOXICILLIN

CAPSULE; ORAL

AMOXICILLIN
LABS ATRAL 250MG A062528 001 Aug 07, 1985
500MG A062528 002 Aug 07, 1985
MYLAN 250MG A062067 001
500MG A062067 002
SUN PHARM INDS LTD 250MG A065016 001 Apr 08, 1999
500MG A065016 002 Apr 08, 1999
TEVA 250MG A062853 001 Dec 22, 1987
250MG A063030 001 Feb 28, 1989
500MG A062854 001 Dec 22, 1987
500MG A063031 001 Feb 28, 1989

AMOXIL

+ GLAXOSMITHKLINE 250MG ** N050459 001
+ 500MG ** N050459 002

TRIMOX

APOTHECON 250MG A061885 001
250MG A062098 001
250MG A062152 001
250MG A063099 001 Mar 20, 1992
500MG A061885 002
500MG A062098 002
500MG A062152 002
500MG A063099 002 Mar 20, 1992

UTIMOX

PARKE DAVIS 250MG A062107 001
500MG A062107 002

WYMOX

WYETH AYERST 250MG A062120 001
500MG A062120 002

FOR SUSPENSION; ORAL

AMOXICILLIN
AM ANTIBIOTICS 125MG/5ML A062059 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMOXICILLIN

FOR SUSPENSION;ORAL

AMOXICILLIN

| | | | | | |
|-------------------------------|--------------|--|---------|-----|--------------|
| | 250MG/5ML | | A062059 | 002 | |
| MYLAN | 125MG/5ML | | A062090 | 001 | |
| | 250MG/5ML | | A062090 | 002 | |
| SUN PHARM INDS LTD | 200MG/5ML | | A065113 | 001 | Nov 29, 2002 |
| | 400MG/5ML | | A065113 | 002 | Nov 29, 2002 |
| TEVA | 125MG/5ML | | A062946 | 001 | Nov 01, 1988 |
| | 250MG/5ML | | A063001 | 001 | Jan 06, 1989 |
| AMOXIL | | | | | |
| + GLAXOSMITHKLINE | 50MG/ML ** | | N050460 | 005 | |
| + | 125MG/5ML ** | | N050460 | 001 | |
| + | 250MG/5ML ** | | N050460 | 002 | |
| + NEOPHARMA | 200MG/5ML ** | | N050760 | 001 | Apr 15, 1999 |
| + | 400MG/5ML ** | | N050760 | 002 | Apr 15, 1999 |
| LAROTID | | | | | |
| + GLAXOSMITHKLINE | 50MG/ML ** | | N050460 | 006 | |
| POLYMOX | | | | | |
| APOTHECON | 125MG/5ML | | A061851 | 001 | |
| | 125MG/5ML | | A062323 | 001 | |
| | 250MG/5ML | | A061851 | 002 | |
| | 250MG/5ML | | A062323 | 002 | |
| TRIMOX | | | | | |
| APOTHECON | 50MG/ML | | A061886 | 001 | |
| | 125MG/5ML | | A061886 | 002 | |
| | 125MG/5ML | | A062099 | 001 | |
| | 125MG/5ML | | A062154 | 001 | |
| | 125MG/5ML | | A062885 | 001 | Mar 08, 1988 |
| | 250MG/5ML | | A061886 | 003 | |
| | 250MG/5ML | | A062099 | 002 | |
| | 250MG/5ML | | A062154 | 002 | |
| | 250MG/5ML | | A062885 | 002 | Mar 08, 1988 |
| UTIMOX | | | | | |
| PARKE DAVIS | 125MG/5ML | | A062127 | 001 | |
| | 250MG/5ML | | A062127 | 002 | |
| WYMOX | | | | | |
| WYETH AYERST | 125MG/5ML | | A062131 | 001 | |
| | 250MG/5ML | | A062131 | 002 | |
| TABLET;ORAL | | | | | |
| AMOXICILLIN | | | | | |
| DAVA PHARMS INC | 875MG | | A065344 | 001 | Jan 15, 2009 |
| SUN PHARM INDS LTD | 500MG | | A065059 | 001 | Nov 24, 2000 |
| | 875MG | | A065059 | 002 | Nov 24, 2000 |
| AMOXIL | | | | | |
| + NEOPHARMA | 500MG ** | | N050754 | 002 | Jul 10, 1998 |
| + | 875MG ** | | N050754 | 001 | Jul 10, 1998 |
| TABLET, CHEWABLE;ORAL | | | | | |
| AMOXICILLIN | | | | | |
| APOTHECON | 125MG | | A064131 | 001 | May 06, 1996 |
| | 250MG | | A064131 | 002 | May 06, 1996 |
| DAVA PHARMS INC | 125MG | | A064139 | 001 | Jan 29, 1996 |
| | 250MG | | A064139 | 002 | Jan 29, 1996 |
| SUN PHARM INDS LTD | 125MG | | A065021 | 001 | Dec 23, 1999 |
| | 200MG | | A065060 | 001 | Nov 29, 2000 |
| | 250MG | | A065021 | 002 | Dec 23, 1999 |
| | 400MG | | A065060 | 002 | Nov 29, 2000 |
| TEVA | 125MG | | A064031 | 001 | Dec 19, 1996 |
| | 250MG | | A064031 | 002 | Dec 19, 1996 |
| AMOXIL | | | | | |
| + NEOPHARMA | 125MG ** | | N050542 | 002 | |
| | 200MG | | N050761 | 001 | Apr 15, 1999 |
| + | 250MG ** | | N050542 | 001 | |
| | 400MG | | N050761 | 002 | Apr 15, 1999 |
| TABLET, EXTENDED RELEASE;ORAL | | | | | |
| MOXATAG | | | | | |
| + PRAGMA | 775MG | | N050813 | 001 | Jan 23, 2008 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMOXICILLIN

TABLET, FOR SUSPENSION;ORAL

AMOXICILLIN

| | | | |
|----------------------|-------|-------------|--------------|
| AUROBINDO PHARMA LTD | 200MG | A065324 001 | Jan 17, 2007 |
| | 400MG | A065324 002 | Jan 17, 2007 |
| DISPERMOX | | | |
| RANBAXY LABS LTD | 200MG | A065080 002 | Aug 11, 2003 |
| | 400MG | A065080 001 | Aug 11, 2003 |
| | 600MG | A065159 001 | Dec 04, 2003 |

AMOXICILLIN; CLARITHROMYCIN; LANSOPRAZOLE

CAPSULE, CAPSULE, DELAYED REL PELLETS, TABLET;ORAL

LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN

| | | | |
|---------------------|------------------------------------------|-------------|--------------|
| TEVA PHARMS USA | 500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,30MG | A200218 001 | Aug 30, 2013 |
| PREVPAC | | | |
| + TAKEDA PHARMS USA | 500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,30MG | N050757 001 | Dec 02, 1997 |

AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

| | | | |
|--------------------|------------------------------|-------------|--------------|
| SUN PHARM INDS LTD | 200MG/5ML;EQ 28.5MG BASE/5ML | A065132 001 | Mar 19, 2003 |
| | 400MG/5ML;EQ 57MG BASE/5ML | A065132 002 | Mar 19, 2003 |
| | 600MG/5ML;EQ 42.9MG BASE/5ML | A065207 002 | Jan 30, 2007 |
| AUGMENTIN '200' | | | |
| + NEOPHARMA | 200MG/5ML;EQ 28.5MG BASE/5ML | N050725 001 | May 31, 1996 |
| AUGMENTIN '400' | | | |
| + NEOPHARMA | 400MG/5ML;EQ 57MG BASE/5ML | N050725 002 | May 31, 1996 |
| AUGMENTIN ES-600 | | | |
| + NEOPHARMA | 600MG/5ML;EQ 42.9MG BASE/5ML | N050755 001 | Jun 22, 2001 |

TABLET;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

| | | | |
|--------------------|------------------------|-------------|--------------|
| APOTEX INC | 250MG;EQ 125MG BASE | A065333 001 | Feb 24, 2009 |
| | 500MG;EQ 125MG BASE | A065333 002 | Feb 24, 2009 |
| | 875MG;EQ 125MG BASE | A065317 003 | Oct 20, 2008 |
| SUN PHARM INDS LTD | 500MG;EQ 125MG BASE | A065109 001 | Nov 04, 2002 |
| | 875MG;EQ 125MG BASE | A065102 001 | Sep 17, 2002 |
| AUGMENTIN '250' | | | |
| + NEOPHARMA | 250MG;EQ 125MG BASE ** | N050564 001 | Aug 06, 1984 |
| AUGMENTIN '500' | | | |
| + NEOPHARMA | 500MG;EQ 125MG BASE ** | N050564 002 | Aug 06, 1984 |

TABLET, CHEWABLE;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

| | | | |
|--------------------|--------------------------|-------------|--------------|
| SANDOZ | 200MG;EQ 28.5MG BASE | A065065 001 | Apr 18, 2002 |
| | 400MG;EQ 57MG BASE | A065065 002 | Apr 18, 2002 |
| SUN PHARM INDS LTD | 200MG;EQ 28.5MG BASE | A065161 001 | Dec 03, 2003 |
| | 400MG;EQ 57MG BASE | A065161 002 | Dec 03, 2003 |
| AUGMENTIN '125' | | | |
| + NEOPHARMA | 125MG;EQ 31.25MG BASE ** | N050597 001 | Jul 22, 1985 |
| AUGMENTIN '200' | | | |
| + NEOPHARMA | 200MG;EQ 28.5MG BASE | N050726 001 | May 31, 1996 |
| AUGMENTIN '250' | | | |
| + NEOPHARMA | 250MG;EQ 62.5MG BASE ** | N050597 002 | Jul 22, 1985 |
| AUGMENTIN '400' | | | |
| + NEOPHARMA | 400MG;EQ 57MG BASE | N050726 002 | May 31, 1996 |

AMPHETAMINE ADIPATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE ADIPATE; DEXTROAMPHETAMINE SULFATE

CAPSULE;ORAL

DELCOBESE

| | | |
|------|--------------------------------|-------------|
| TEVA | 1.25MG;1.25MG;1.25MG;1.25MG ** | A083564 001 |
| | 2.5MG;2.5MG;2.5MG;2.5MG ** | A083564 002 |
| | 3.75MG;3.75MG;3.75MG;3.75MG ** | A083564 003 |
| | 5MG;5MG;5MG;5MG ** | A083564 004 |

TABLET;ORAL

DELCOBESE

| | | |
|------|-----------------------------|-------------|
| TEVA | 1.25MG;1.25MG;1.25MG;1.25MG | A083563 004 |
| | 2.5MG;2.5MG;2.5MG;2.5MG | A083563 003 |
| | 3.75MG;3.75MG;3.75MG;3.75MG | A083563 002 |
| | 5MG;5MG;5MG;5MG | A083563 001 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

| | | | | | |
|------------------------------------------------------------------------|------------------------------------|---------|---------|--------------|--------------|
| ADDERALL 10 | | | | | |
| + TEVA WOMENS | 2.5MG;2.5MG;2.5MG;2.5MG ** | | N011522 | 007 | Feb 13, 1996 |
| ADDERALL 12.5 | | | | | |
| + TEVA WOMENS | 3.125MG;3.125MG;3.125MG;3.125MG ** | | N011522 | 012 | Aug 31, 2000 |
| ADDERALL 15 | | | | | |
| + TEVA WOMENS | 3.75MG;3.75MG;3.75MG;3.75MG ** | | N011522 | 013 | Aug 31, 2000 |
| ADDERALL 20 | | | | | |
| + TEVA WOMENS | 5MG;5MG;5MG;5MG ** | | N011522 | 008 | Feb 13, 1996 |
| ADDERALL 30 | | | | | |
| + TEVA WOMENS | 7.5MG;7.5MG;7.5MG;7.5MG ** | | N011522 | 010 | May 12, 1997 |
| ADDERALL 5 | | | | | |
| + TEVA WOMENS | 1.25MG;1.25MG;1.25MG;1.25MG ** | | N011522 | 009 | May 12, 1997 |
| ADDERALL 7.5 | | | | | |
| + TEVA WOMENS | 1.875MG;1.875MG;1.875MG;1.875MG ** | | N011522 | 011 | Aug 31, 2000 |
| DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE | | | | | |
| ACTAVIS ELIZABETH | 1.25MG;1.25MG;1.25MG;1.25MG | A040456 | 001 | May 06, 2003 | |
| | 2.5MG;2.5MG;2.5MG;2.5MG | A040456 | 002 | May 06, 2003 | |
| | 5MG;5MG;5MG;5MG | A040456 | 003 | May 06, 2003 | |
| | 7.5MG;7.5MG;7.5MG;7.5MG | A040456 | 004 | May 06, 2003 | |
| TEVA PHARMS | 1.25MG;1.25MG;1.25MG;1.25MG | A040472 | 001 | Sep 30, 2003 | |
| | 2.5MG;2.5MG;2.5MG;2.5MG | A040472 | 002 | Sep 30, 2003 | |
| | 5MG;5MG;5MG;5MG | A040472 | 003 | Sep 30, 2003 | |
| | 7.5MG;7.5MG;7.5MG;7.5MG | A040472 | 004 | Sep 30, 2003 | |

AMPHETAMINE RESIN COMPLEX; DEXTROAMPHETAMINE RESIN COMPLEX

CAPSULE, EXTENDED RELEASE; ORAL

| | | | | | |
|------------------|-------------------------------|--|---------|-----|--|
| BIPHETAMINE 12.5 | | | | | |
| UCB INC | EQ 6.25MG BASE;EQ 6.25MG BASE | | N010093 | 007 | |
| BIPHETAMINE 20 | | | | | |
| UCB INC | EQ 10MG BASE;EQ 10MG BASE | | N010093 | 003 | |
| BIPHETAMINE 7.5 | | | | | |
| UCB INC | EQ 3.75MG BASE;EQ 3.75MG BASE | | N010093 | 009 | |

AMPHETAMINE SULFATE

TABLET; ORAL

| | | | | | |
|---------------------|------|---------|-----|--------------|--|
| AMPHETAMINE SULFATE | | | | | |
| LANNETT | 5MG | A083901 | 001 | Aug 31, 1984 | |
| | 10MG | A083901 | 002 | Aug 31, 1984 | |

AMPHOTERICIN B

CREAM; TOPICAL

| | | | | | |
|-----------|----|--|---------|-----|--|
| FUNGIZONE | | | | | |
| APOTHECON | 3% | | N050314 | 001 | |

INJECTABLE; INJECTION

| | | | | | |
|--------------------------------------|------------|---------|-----|--------------|--|
| AMPHOTERICIN B | | | | | |
| ABBOTT | 50MG/VIAL | A064141 | 001 | Dec 23, 1996 | |
| ABRAXIS PHARM | 50MG/VIAL | A062728 | 001 | Apr 13, 1987 | |
| TEVA PARENTERAL | 50MG/VIAL | A064062 | 001 | Mar 31, 1995 | |
| FUNGIZONE | | | | | |
| APOTHECON | 50MG/VIAL | A060517 | 001 | | |
| INJECTABLE, LIPID COMPLEX; INJECTION | | | | | |
| AMPHOTEC | | | | | |
| ALKOPHARMA USA | 50MG/VIAL | N050729 | 001 | Nov 22, 1996 | |
| | 100MG/VIAL | N050729 | 002 | Nov 22, 1996 | |

LOTION; TOPICAL

| | | | | | |
|-----------|----|--|---------|-----|--|
| FUNGIZONE | | | | | |
| APOTHECON | 3% | | A060570 | 001 | |

OINTMENT; TOPICAL

| | | | | | |
|-----------|----|--|---------|-----|--|
| FUNGIZONE | | | | | |
| APOTHECON | 3% | | N050313 | 001 | |

SUSPENSION; ORAL

| | | | | | |
|----------------------|----------|--|---------|-----|--|
| FUNGIZONE | | | | | |
| BRISTOL MYERS SQUIBB | 100MG/ML | | N050341 | 003 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPICILLIN SODIUM

INJECTABLE; INJECTION

AMPICILLIN SODIUM

| | | | |
|----------------------|-----------------------|-------------|--------------|
| ACS DOBFAR SPA | EQ 500MG BASE/VIAL | A090884 001 | Apr 03, 2013 |
| | EQ 1GM BASE/VIAL | A090884 002 | Apr 03, 2013 |
| | EQ 2GM BASE/VIAL | A090884 003 | Apr 03, 2013 |
| APOTHECON | EQ 125MG BASE/VIAL | A062860 001 | Feb 05, 1988 |
| | EQ 250MG BASE/VIAL | A062860 002 | Feb 05, 1988 |
| | EQ 500MG BASE/VIAL | A062860 003 | Feb 05, 1988 |
| | EQ 1GM BASE/VIAL | A062860 004 | Feb 05, 1988 |
| | EQ 2GM BASE/VIAL | A062860 005 | Feb 05, 1988 |
| AUROBINDO PHARMA | EQ 125MG BASE/VIAL | A065499 001 | Aug 17, 2010 |
| CONSOLIDATED PHARM | EQ 125MG BASE/VIAL | A061936 005 | |
| | EQ 250MG BASE/VIAL | A061936 001 | |
| | EQ 500MG BASE/VIAL | A061936 002 | |
| | EQ 1GM BASE/VIAL | A061936 003 | |
| | EQ 2GM BASE/VIAL | A061936 004 | |
| HANFORD GC | EQ 125MG BASE/VIAL | A062772 005 | Apr 15, 1993 |
| | EQ 500MG BASE/VIAL | A062772 008 | Apr 15, 1993 |
| | EQ 1GM BASE/VIAL | A062772 002 | Apr 15, 1993 |
| | EQ 2GM BASE/VIAL | A062772 004 | Apr 15, 1993 |
| INTL MEDICATION | EQ 1GM BASE/VIAL | A062634 002 | Jan 09, 1987 |
| | EQ 2GM BASE/VIAL | A062634 003 | Jan 09, 1987 |
| ISTITUTO BIO ITA SPA | EQ 125MG BASE/VIAL | A062797 001 | Jul 12, 1993 |
| LILLY | EQ 500MG BASE/VIAL | A062565 001 | Apr 04, 1985 |
| | EQ 1GM BASE/VIAL | A062565 002 | Apr 04, 1985 |
| | EQ 2GM BASE/VIAL | A062565 003 | Jun 24, 1986 |
| WATSON LABS INC | EQ 125MG BASE/VIAL | A062816 001 | Oct 24, 1988 |
| | EQ 250MG BASE/VIAL | A062816 002 | Oct 24, 1988 |
| | EQ 500MG BASE/VIAL | A062816 003 | Oct 24, 1988 |
| | EQ 1GM BASE/VIAL | A062816 004 | Oct 24, 1988 |
| | EQ 2GM BASE/VIAL | A062816 005 | Oct 24, 1988 |
| | EQ 10GM BASE/VIAL | A062994 001 | Sep 15, 1988 |
| WEST-WARD PHARMS INT | EQ 125MG BASE/VIAL | A062692 001 | Jun 24, 1986 |
| | EQ 250MG BASE/VIAL | A062692 002 | Jun 24, 1986 |
| | EQ 500MG BASE/VIAL | A062692 003 | Jun 24, 1986 |
| | EQ 1GM BASE/VIAL | A062692 004 | Jun 24, 1986 |
| | EQ 2GM BASE/VIAL | A062692 005 | Jun 24, 1986 |
| | EQ 10GM BASE/VIAL | A062692 006 | Jun 24, 1986 |
| OMNIPEN-N | | | |
| WYETH AYERST | EQ 125MG BASE/VIAL | A060626 001 | |
| | EQ 125MG BASE/VIAL | A062718 001 | Dec 16, 1986 |
| | EQ 250MG BASE/VIAL | A060626 002 | |
| | EQ 250MG BASE/VIAL | A062718 002 | Dec 16, 1986 |
| | EQ 500MG BASE/VIAL | A060626 003 | |
| | EQ 500MG BASE/VIAL | A062718 003 | Dec 16, 1986 |
| | EQ 1GM BASE/VIAL | A060626 004 | |
| | EQ 1GM BASE/VIAL | A062718 004 | Dec 16, 1986 |
| | EQ 2GM BASE/VIAL | A060626 005 | |
| | EQ 2GM BASE/VIAL | A062718 005 | Dec 16, 1986 |
| PENBRITIN-S | | | |
| + WYETH AYERST | EQ 125MG BASE/VIAL ** | N050072 001 | |
| + | EQ 250MG BASE/VIAL ** | N050072 002 | |
| + | EQ 500MG BASE/VIAL ** | N050072 003 | |
| + | EQ 1GM BASE/VIAL ** | N050072 004 | |
| + | EQ 2GM BASE/VIAL ** | N050072 005 | |
| + | EQ 4GM BASE/VIAL ** | N050072 006 | |
| POLYCILLIN-N | | | |
| BRISTOL | EQ 125MG BASE/VIAL ** | N050309 001 | |
| | EQ 250MG BASE/VIAL ** | N050309 002 | |
| | EQ 500MG BASE/VIAL ** | N050309 003 | |
| | EQ 1GM BASE/VIAL ** | N050309 004 | |
| | EQ 2GM BASE/VIAL ** | N050309 005 | |
| TOTACILLIN-N | | | |
| GLAXOSMITHKLINE | EQ 125MG BASE/VIAL | A060677 001 | |
| | EQ 250MG BASE/VIAL | A060677 002 | |
| | EQ 500MG BASE/VIAL | A060677 003 | |
| | EQ 1GM BASE/VIAL | A060677 004 | |
| | EQ 1GM BASE/VIAL | A062727 001 | Dec 19, 1986 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPICILLIN SODIUM

INJECTABLE; INJECTION

TOTACILLIN-N

| | | | |
|-------------------|---------|-----|--------------|
| EQ 2GM BASE/VIAL | A060677 | 005 | |
| EQ 2GM BASE/VIAL | A062727 | 002 | Dec 19, 1986 |
| EQ 10GM BASE/VIAL | A060677 | 006 | |

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

UNASYN

| | | | | |
|--------|---------------------------------------|---------|-----|--------------|
| PFIZER | EQ 500MG BASE/VIAL;EQ 250MG BASE/VIAL | N050608 | 003 | Dec 31, 1986 |
|--------|---------------------------------------|---------|-----|--------------|

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMCILL

| | | | | |
|-------------|---------------|---------|-----|--|
| PARKE DAVIS | EQ 250MG BASE | A062041 | 001 | |
| | EQ 500MG BASE | A062041 | 002 | |

AMPICILLIN TRIHYDRATE

| | | | | |
|-----------------|---------------|---------|-----|--|
| AM ANTI-BIOTICS | EQ 250MG BASE | A061602 | 001 | |
|-----------------|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 500MG BASE | A061602 | 002 | |
|--|---------------|---------|-----|--|

| | | | | |
|----------------------|---------------|---------|-----|--|
| IVAX SUB TEVA PHARMS | EQ 250MG BASE | A060765 | 001 | |
|----------------------|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 500MG BASE | A060765 | 002 | |
|--|---------------|---------|-----|--|

| | | | | |
|---------|---------------|---------|-----|--|
| LEDERLE | EQ 250MG BASE | A062208 | 001 | |
|---------|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 500MG BASE | A062208 | 002 | |
|--|---------------|---------|-----|--|

| | | | | |
|-------|---------------|---------|-----|--|
| MYLAN | EQ 250MG BASE | A061755 | 001 | |
|-------|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 500MG BASE | A061755 | 002 | |
|--|---------------|---------|-----|--|

| | | | | |
|---------------|---------------|---------|-----|--|
| PUREPAC PHARM | EQ 250MG BASE | A061853 | 001 | |
|---------------|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 500MG BASE | A061853 | 002 | |
|--|---------------|---------|-----|--|

| | | | | |
|------|---------------|---------|-----|--|
| TEVA | EQ 250MG BASE | A061502 | 001 | |
|------|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 500MG BASE | A061502 | 002 | |
|--|---------------|---------|-----|--|

| | | | | |
|----------|---------------|---------|-----|--|
| VITARINE | EQ 250MG BASE | A061387 | 001 | |
|----------|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 500MG BASE | A061387 | 003 | |
|--|---------------|---------|-----|--|

OMNIPEN (AMPICILLIN)

| | | | | |
|--------------|-------|---------|-----|--|
| WYETH AYERST | 250MG | A060624 | 001 | |
|--------------|-------|---------|-----|--|

| | | | | |
|--|-------|---------|-----|--|
| | 500MG | A060624 | 002 | |
|--|-------|---------|-----|--|

PENBRITIN

| | | | | |
|--------------|---------------|---------|-----|--|
| WYETH AYERST | EQ 250MG BASE | A060908 | 001 | |
|--------------|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 500MG BASE | A060908 | 002 | |
|--|---------------|---------|-----|--|

PFIZERPEN-A

| | | | | |
|--------|---------------|---------|-----|--|
| PFIZER | EQ 250MG BASE | A062050 | 001 | |
|--------|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 500MG BASE | A062050 | 002 | |
|--|---------------|---------|-----|--|

POLYCILLIN

| | | | | |
|---------|---------------|---------|-----|--|
| BRISTOL | EQ 250MG BASE | N050310 | 001 | |
|---------|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 500MG BASE | N050310 | 002 | |
|--|---------------|---------|-----|--|

PRINCIPEN

| | | | | |
|-----------|---------------|---------|-----|--------------|
| APOTHECON | EQ 250MG BASE | A062888 | 001 | Mar 04, 1988 |
|-----------|---------------|---------|-----|--------------|

| | | | | |
|--|---------------|---------|-----|--------------|
| | EQ 500MG BASE | A062888 | 002 | Mar 04, 1988 |
|--|---------------|---------|-----|--------------|

| | | | | |
|----------------------|---------------|---------|-----|--|
| BRISTOL MYERS SQUIBB | EQ 250MG BASE | A061392 | 001 | |
|----------------------|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 500MG BASE | A061392 | 002 | |
|--|---------------|---------|-----|--|

PRINCIPEN '250'

| | | | | |
|-----------|---------------|---------|-----|--|
| APOTHECON | EQ 250MG BASE | A062157 | 002 | |
|-----------|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 250MG BASE | N050056 | 001 | |
|--|---------------|---------|-----|--|

PRINCIPEN '500'

| | | | | |
|-----------|---------------|---------|-----|--|
| APOTHECON | EQ 500MG BASE | A062157 | 001 | |
|-----------|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 500MG BASE | N050056 | 002 | |
|--|---------------|---------|-----|--|

TOTACILLIN

| | | | | |
|-----------------|---------------|---------|-----|--|
| GLAXOSMITHKLINE | EQ 250MG BASE | A060060 | 001 | |
|-----------------|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 250MG BASE | A062212 | 001 | |
|--|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 500MG BASE | A060060 | 002 | |
|--|---------------|---------|-----|--|

| | | | | |
|--|---------------|---------|-----|--|
| | EQ 500MG BASE | A062212 | 002 | |
|--|---------------|---------|-----|--|

FOR SUSPENSION; ORAL

AMCILL

| | | | | |
|-------------|-------------------|---------|-----|--|
| PARKE DAVIS | EQ 125MG BASE/5ML | A062030 | 001 | |
|-------------|-------------------|---------|-----|--|

| | | | | |
|--|-------------------|---------|-----|--|
| | EQ 250MG BASE/5ML | A062030 | 002 | |
|--|-------------------|---------|-----|--|

AMPICILLIN TRIHYDRATE

| | | | | |
|-----------------|-------------------|---------|-----|--|
| AM ANTI-BIOTICS | EQ 125MG BASE/5ML | A061601 | 001 | |
|-----------------|-------------------|---------|-----|--|

| | | | | |
|--|-------------------|---------|-----|--|
| | EQ 250MG BASE/5ML | A061601 | 002 | |
|--|-------------------|---------|-----|--|

| | | | | |
|-------|-------------------|---------|-----|--|
| MYLAN | EQ 125MG BASE/5ML | A061829 | 002 | |
|-------|-------------------|---------|-----|--|

| | | | | |
|--|-------------------|---------|-----|--|
| | EQ 250MG BASE/5ML | A061829 | 001 | |
|--|-------------------|---------|-----|--|

| | | | | |
|---------------|-------------------|---------|-----|--|
| PUREPAC PHARM | EQ 125MG BASE/5ML | A061980 | 001 | |
|---------------|-------------------|---------|-----|--|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AMPICILLIN/AMPICILLIN TRIHYDRATE

FOR SUSPENSION;ORAL

AMPICILLIN TRIHYDRATE

| | | |
|----------------------|-------------------|-------------|
| | EQ 250MG BASE/5ML | A061980 002 |
| TEVA | EQ 125MG BASE/5ML | A061370 001 |
| | EQ 250MG BASE/5ML | A061370 002 |
| OMNIPEN (AMPICILLIN) | | |
| WYETH AYERST | 100MG/ML | A060625 001 |
| | 125MG/5ML | A060625 002 |
| | 250MG/5ML | A060625 003 |
| | 500MG/5ML | A060625 004 |
| PENBRITIN | | |
| WYETH AYERST | EQ 100MG BASE/ML | N050019 001 |
| | EQ 125MG BASE/5ML | N050019 002 |
| | EQ 250MG BASE/5ML | N050019 003 |
| PFIZERPEN-A | | |
| PFIZER | EQ 125MG BASE/5ML | A062049 001 |
| | EQ 250MG BASE/5ML | A062049 002 |
| POLYCILLIN | | |
| APOTHECON | EQ 125MG BASE/5ML | A062297 001 |
| | EQ 250MG BASE/5ML | A062297 002 |
| BRISTOL | EQ 100MG BASE/ML | N050308 004 |
| | EQ 125MG BASE/5ML | N050308 001 |
| | EQ 250MG BASE/5ML | N050308 002 |
| | EQ 500MG BASE/5ML | N050308 003 |
| PRINCIPEN | | |
| APOTHECON | EQ 100MG BASE/ML | A061394 001 |
| | EQ 125MG BASE/5ML | A061394 002 |
| | EQ 250MG BASE/5ML | A061394 003 |
| PRINCIPEN '125' | | |
| APOTHECON | EQ 125MG BASE/5ML | A060127 002 |
| | EQ 125MG BASE/5ML | A062151 001 |
| PRINCIPEN '250' | | |
| APOTHECON | EQ 250MG BASE/5ML | A060127 001 |
| | EQ 250MG BASE/5ML | A062151 002 |
| TOTACILLIN | | |
| GLAXOSMITHKLINE | EQ 125MG BASE/5ML | A060666 001 |
| | EQ 125MG BASE/5ML | A062223 001 |
| | EQ 250MG BASE/5ML | A060666 002 |
| | EQ 250MG BASE/5ML | A062223 002 |

TABLET, CHEWABLE;ORAL

POLYCILLIN

BRISTOL EQ 125MG BASE N050093 001

AMPICILLIN/AMPICILLIN TRIHYDRATE; PROBENECID

CAPSULE;ORAL

PRINCIPEN W/ PROBENECID

APOTHECON EQ 389MG BASE;111MG A062150 001
EQ 389MG BASE;111MG N050488 001

FOR SUSPENSION;ORAL

POLYCILLIN-PRB

APOTHECON EQ 3.5GM BASE/BOT;1GM/BOT A061898 001
BRISTOL EQ 3.5GM BASE/BOT;1GM/BOT N050457 001

PROBAMPACIN

G AND W LABS INC EQ 3.5GM BASE/BOT;1GM/BOT A061741 001

AMPRENAVIR

CAPSULE;ORAL

AGENERASE

GLAXOSMITHKLINE 50MG N021007 001 Apr 15, 1999
150MG N021007 002 Apr 15, 1999

SOLUTION;ORAL

AGENERASE

+ GLAXOSMITHKLINE 15MG/ML ** N021039 001 Apr 15, 1999

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ANAGRELIDE HYDROCHLORIDE

CAPSULE; ORAL

AGRYLIN

+ SHIRE LLC

EQ 1MG BASE **

N020333 002 Mar 14, 1997

ANAGRELIDE HYDROCHLORIDE

MYLAN PHARMS INC

EQ 0.5MG BASE

A076811 001 Apr 18, 2005

EQ 0.5MG BASE

A077613 001 Jun 27, 2006

EQ 1MG BASE

A076811 002 Apr 18, 2005

EQ 1MG BASE

A077613 002 Jun 27, 2006

ROXANE

EQ 0.5MG BASE

A076489 001 Apr 18, 2005

EQ 1MG BASE

A076489 002 Apr 18, 2005

UPSHER SMITH LABS

EQ 0.5MG BASE

A076683 001 Apr 18, 2005

EQ 1MG BASE

A076683 002 Apr 18, 2005

WATSON LABS

EQ 0.5MG BASE

A076417 001 Apr 18, 2005

EQ 1MG BASE

A076417 002 Apr 18, 2005

ANASTROZOLE

TABLET; ORAL

ANASTROZOLE

IMPAX LABS INC

1MG

A091242 001 May 31, 2012

LANNETT CO INC

1MG

A091331 001 Jan 05, 2011

SANDOZ

1MG

A079007 001 Jun 28, 2010

SUN PHARM INDS LTD

1MG

A091177 001 Jul 15, 2011

SYNTHON PHARMS

1MG

A078322 001 Jun 28, 2010

WATSON LABS TEVA

1MG

A078984 001 Jun 28, 2010

ANGIOTENSIN II ACETATE

SOLUTION; INTRAVENOUS

GIAPREZA

+ LA JOLLA PHARMA

EQ 5MG BASE/2ML (EQ 2.5MG BASE/ML)

N209360 002 Dec 21, 2017

ANILERIDINE HYDROCHLORIDE

TABLET; ORAL

LERITINE

MERCK

EQ 25MG BASE

N010585 002

ANILERIDINE PHOSPHATE

INJECTABLE; INJECTION

LERITINE

MERCK

25MG/ML

N010520 003

ANISINDIONE

TABLET; ORAL

MIRADON

SCHERING

50MG

N010909 003

ANISOTROPINE METHYLBROMIDE

TABLET; ORAL

ANISOTROPINE METHYLBROMIDE

WATSON LABS

50MG

A086046 001

VALPIN 50

ENDO PHARMS

50MG

N013428 001

ANTAZOLINE PHOSPHATE; NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

VASOCON-A

NOVARTIS

0.5%; 0.05%

N018746 002 Jul 11, 1994

APOMORPHINE HYDROCHLORIDE

INJECTABLE; SUBCUTANEOUS

APOKYN

US WORLDMEDS

20MG/2ML (10MG/ML)

N021264 001 Apr 20, 2004

APROTININ

INJECTABLE; INJECTION

TRASYLOL

BAYER HLTHCARE

10,000KIU/ML

N020304 001 Dec 29, 1993

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ARBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

GENESA

GENSIA AUTOMEDICS 0.05MG/ML N020420 001 Sep 12, 1997

ARDEPARIN SODIUM

INJECTABLE; INJECTION

NORMIFLO

+ PHARMACIA AND UPJOHN 5,000 UNITS/0.5ML ** N020227 002 May 23, 1997

+ 10,000 UNITS/0.5ML ** N020227 001 May 23, 1997

ARGATROBAN

SOLUTION; INTRAVENOUS

ARGATROBAN IN DEXTROSE

SANDOZ

125MG/125ML (1MG/ML) N201743 001 May 09, 2011

ARIPIPIRAZOLE

INJECTABLE; INTRAMUSCULAR

ABILIFY

OTSUKA 9.75MG/1.3ML (7.5MG/ML) N021866 001 Sep 20, 2006

SOLUTION; ORAL

ABILIFY

+ OTSUKA 1MG/ML ** N021713 001 Dec 10, 2004

TABLET; ORAL

ARIPIPIRAZOLE

MYLAN PHARMS INC 2MG A206240 001 Sep 19, 2018

5MG A206240 002 Sep 19, 2018

10MG A206240 003 Sep 19, 2018

15MG A206240 004 Sep 19, 2018

20MG A206240 005 Sep 19, 2018

30MG A206240 006 Sep 19, 2018

TABLET, ORALLY DISINTEGRATING; ORAL

ABILIFY

+ OTSUKA 10MG ** N021729 002 Jun 07, 2006

+ 15MG ** N021729 003 Jun 07, 2006

+ 20MG ** N021729 004 Jun 07, 2006

+ 30MG ** N021729 005 Jun 07, 2006

ARMODAFINIL

TABLET; ORAL

ARMODAFINIL

WATSON LABS INC 100MG A200156 002 Aug 29, 2012

200MG A200156 004 Aug 29, 2012

NUVIGIL

+ CEPHALON 100MG ** N021875 002 Mar 26, 2009

ARSENIC TRIOXIDE

INJECTABLE; INJECTION

TRISENOX

+ CEPHALON 1MG/ML N021248 001 Sep 25, 2000

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; PALMITATE; VITAMIN E

INJECTABLE; INJECTION

BEROCCA PN

ROCHE 50MG/ML; 0.03MG/ML; 0.0025MG/ML; 7.5MG/ML; 100 N006071 003 Oct 10, 1985

IU/ML; 0.2MG/ML; 20MG/ML; 2MG/ML; 1.8MG/ML;

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.C. 9+3

ABRAXIS PHARM 10MG/ML; 0.006MG/ML; 0.5MCG/ML; 1.5MG/ML; 2 N018440 002 Aug 08, 1985

0

IU/ML; 0.04MG/ML; 4MG/ML; 0.4MG/ML; 0.36MG/

M.V.I.-12 ADULT

HOSPIRA 10MG/ML; 0.006MG/ML; 0.5MCG/ML; 1.5MG/ML; 2 N008809 004 Aug 08, 1985

0

+ 20MG/ML; 0.006MG/ML; 0.05MCG/ML; 1.5MG/ML; N008809 006 Sep 09, 2004

0.0005MG/ML; 0.06MG/ML; 4MG/ML; 0.6MG/ML;

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.I.-12 ADULT

0.36MG/ML; 0.6MG/ML; 0.1MG/ML; 1MG/ML

MVC PLUS

WATSON LABS

10MG/ML; 0.006MG/ML; 0.5MCG/ML; 1.5MG/ML; 2

N018439 002 Aug 08, 1985

0

IU/ML; 0.04MG/ML; 4MG/ML; 0.4MG/ML; 0.36MG/

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.I.-12 ADULT

HOSPIRA

20MG/ML; 0.006MG/ML; 0.5MCG/ML; 1.5MG/ML; 2

N008809 005 Apr 22, 2004

0

IU/ML; 0.6MG/ML; 4MG/ML; 0.4MG/ML; 0.36MG/M

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.I.-12 LYOPHILIZED

TELIGENT PHARMA INC

100MG/VIAL; 0.06MG/VIAL; 0.005MG/VIAL; 15M

N018933 002 Aug 08, 1985

G/VIAL; 5MCG/VIAL; 0.4MG/VIAL; 40MG/VIAL; 4

MG/VIAL; 3.6MG/VIAL; 3MG/VIAL; 1MG/VIAL; 10

MG/VIAL

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PANTOTHENIC ACID; PHYTONADIONE; PYRIDOXINE; RIBOFLAVIN; THIAMINE; VITAMIN A PALMITATE; VITAMIN E

INJECTABLE; INJECTION

VITAPED

HOSPIRA

N/A, 80MG/VIAL; N/A, 0.02MG/VIAL; N/A, 0.001

N020176 001 Dec 29, 1993

MG/VIAL; 400

IU/10ML, N/A; N/A, 0.14MG/VIAL; N/A, 17MG/VI

AL; N/A, 5MG/VIAL; 0.2MG/10ML, N/A; N/A, 1MG/

VIAL; N/A, 1.4MG/VIAL; N/A, 1.2MG/VIAL; EQ

2, 300 UNITS BASE/10ML, N/A; 7 IU/10ML, N/A

ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE

FOR SOLUTION; ORAL

PEG-3350, SODIUM SULFATE, SODIUM CHLORIDE, POTASSIUM CHLORIDE, SODIUM ASCORBATE AND ASCORBIC

NOVEL LABS INC

4.7GM; 100GM; 1.015GM; 5.9GM; 2.691GM; 7.5GM

A090145 001 Jan 25, 2012

ASPIRIN

TABLET; ORAL

BAYER EXTRA STRENGTH ASPIRIN FOR MIGRAINE PAIN

BAYER

500MG

N021317 001 Oct 18, 2001

TABLET, EXTENDED RELEASE; ORAL

8-HOUR BAYER

BAYER

650MG

N016030 001

MEASURIN

BAYER

650MG

N016030 002

ASPIRIN; BUTALBITAL

TABLET; ORAL

AXOTAL

SAVAGE LABS

650MG; 50MG

A088305 001 Oct 13, 1983

ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

NOSTRUM LABS INC

325MG; 50MG; 40MG

A078149 001 Jun 13, 2007

WATSON LABS

325MG; 50MG; 40MG

A086231 002 Feb 12, 1985

TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

ACTAVIS ELIZABETH

325MG; 50MG; 40MG

A086710 002 Aug 23, 1983

FOSUN PHARMA

325MG; 50MG; 40MG

A086398 002 Apr 06, 1984

HALSEY

325MG; 50MG; 40MG

A089448 001 Dec 01, 1986

IVAX PHARMS

325MG; 50MG; 40MG

A085441 002 Oct 31, 1984

PURACAP PHARM

325MG; 50MG; 40MG

A087048 002 Dec 09, 1983

QUANTUM PHARMICS

325MG; 50MG; 40MG

A088972 001 Jun 18, 1985

WATSON LABS

325MG; 50MG; 40MG

A086237 002 Mar 23, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASPIRIN; BUTALBITAL; CAFFEINE

TABLET;ORAL

FIORINAL

+ ALLERGAN SALES LLC 325MG;50MG;40MG ** N017534 003 Apr 16, 1986

LANORINAL

LANNETT 325MG;50MG;40MG A086986 002 Oct 18, 1985

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE;ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

VINTAGE PHARMS LLC 325MG;50MG;40MG;30MG A075351 001 Mar 05, 1999

WATSON LABS 325MG;50MG;40MG;30MG A074359 001 Aug 31, 1995

ASPIRIN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE;ORAL

SYNALGOS-DC

+ SUN PHARM INDUSTRIES 356.4MG;30MG;16MG N011483 004 Sep 06, 1983

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET;ORAL

INVAGESIC

SANDOZ 385MG;30MG;25MG A074817 001 Nov 27, 1996

INVAGESIC FORTE

SANDOZ 770MG;60MG;50MG A074817 002 Nov 27, 1996

NORGESIC

+ MEDICIS 385MG;30MG;25MG ** N013416 003 Oct 27, 1982

NORGESIC FORTE

+ MEDICIS 770MG;60MG;50MG ** N013416 004 Oct 27, 1982

ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE

STEVENS J 385MG;30MG;25MG A074988 001 Apr 30, 1999

770MG;60MG;50MG A074988 002 Apr 30, 1999

ORPHENGESIC

GALT PHARMS 385MG;30MG;25MG A075141 001 May 29, 1998

ORPHENGESIC FORTE

GALT PHARMS 770MG;60MG;50MG A075141 002 May 29, 1998

ASPIRIN; CAFFEINE; PROPOXYPHENE HYDROCHLORIDE

CAPSULE;ORAL

COMPOUND 65

ALRA 389MG;32.4MG;65MG A084553 002 Aug 17, 1983

DARVON COMPOUND

XANODYNE PHARM 389MG;32.4MG;32MG N010996 006 Mar 08, 1983

DARVON COMPOUND-65

XANODYNE PHARM 389MG;32.4MG;65MG N010996 007 Mar 08, 1983

PROPOXYPHENE COMPOUND 65

IVAX SUB TEVA PHARMS 389MG;32.4MG;65MG A083077 002 Dec 07, 1984

SANDOZ 389MG;32.4MG;65MG A080044 002 Sep 16, 1983

TEVA 389MG;32.4MG;65MG A089025 001 Mar 29, 1985

PROPOXYPHENE COMPOUND-65

SANDOZ 389MG;32.4MG;65MG A083101 002 Jun 24, 1985

PROPOXYPHENE HYDROCHLORIDE W/ ASPIRIN AND CAFFEINE

WATSON LABS 389MG;32.4MG;65MG A085732 002 Sep 03, 1984

ASPIRIN; CARISOPRODOL

TABLET;ORAL

CARISOPRODOL AND ASPIRIN

OXFORD PHARMS 325MG;200MG A040252 001 Dec 10, 1997

CARISOPRODOL COMPOUND

WATSON LABS 325MG;200MG A088809 001 Oct 03, 1985

SOMA COMPOUND

MEDA PHARMS 325MG;200MG ** N012365 005 Jul 11, 1983

ASPIRIN; CARISOPRODOL; CODEINE PHOSPHATE

TABLET;ORAL

CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE

OXFORD PHARMS 325MG;200MG;16MG A040283 001 Dec 29, 1998

SOMA COMPOUND W/ CODEINE

MEDA PHARMS 325MG;200MG;16MG ** N012366 002 Jul 11, 1983

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASPIRIN; HYDROCODONE BITARTRATE

TABLET; ORAL

AZDONE

SCHWARZ PHARMA 500MG; 5MG ** A089420 001 Jan 25, 1988

VICOPRIN

ABBOTT 500MG; 5MG A086333 001 Sep 14, 1983

ASPIRIN; MEPROBAMATE

TABLET; ORAL

EQUAGESIC

SUN PHARM INDUSTRIES 325MG; 200MG N011702 003 Dec 29, 1983

MEPRO-ASPIRIN

SANDOZ 325MG; 200MG A089127 001 Mar 02, 1987

MEPROBAMATE AND ASPIRIN

PAR PHARM 325MG; 200MG A089126 001 Aug 19, 1986

MICRAININ

MEDPOINTE PHARM HLC 325MG; 200MG A084978 001

Q-GESIC

QUANTUM PHARMICS 325MG; 200MG A088740 001 Jun 01, 1984

ASPIRIN; METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL AND ASPIRIN

IVAX SUB TEVA PHARMS 325MG; 400MG A087211 001 Dec 22, 1982

MCNEIL 325MG; 400MG A089193 001 Feb 12, 1986

PAR PHARM 325MG; 400MG A089657 001 Nov 04, 1988

ROBAXISAL

ROBINS AH 325MG; 400MG N012281 001

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET; ORAL

CODOXY

HALSEY 325MG; 4.5MG; 0.38MG A087464 001 Jul 01, 1982

OXYCODONE AND ASPIRIN

SUN PHARM INDUSTRIES 325MG; 4.5MG; 0.38MG A040260 001 Jul 17, 1998

325MG; 4.5MG; 0.38MG A087794 001 May 26, 1982

WATSON LABS 325MG; 4.5MG; 0.38MG A040255 001 Feb 27, 1998

OXYCODONE AND ASPIRIN (HALF-STRENGTH)

ROXANE 325MG; 2.25MG; 0.19MG A087742 001 Jun 04, 1982

PERCODAN

ENDO PHARMS 325MG; 4.5MG; 0.38MG ** N007337 006

PERCODAN-DEMI

ENDO PHARMS 325MG; 2.25MG; 0.19MG ** N007337 005

ROXIPRIN

ROXANE 325MG; 4.5MG; 0.38MG A087743 001 Jun 04, 1982

ASPIRIN; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

TALWIN COMPOUND

+ SANOFI AVENTIS US 325MG; EQ 12.5MG BASE ** N016891 001

ASPIRIN; PRAVASTATIN SODIUM

TABLET, TABLET; ORAL

PRAVIGARD PAC (COPACKAGED)

BRISTOL MYERS SQUIBB 325MG, N/A; N/A, 80MG N021387 006 Jun 24, 2003

TABLET, TABLET, TABLET; ORAL

PRAVIGARD PAC (COPACKAGED)

BRISTOL MYERS SQUIBB 81MG, N/A; N/A, 20MG N021387 001 Jun 24, 2003

81MG, N/A; N/A, 40MG N021387 002 Jun 24, 2003

81MG, N/A; N/A, 80MG N021387 003 Jun 24, 2003

325MG, N/A; N/A, 20MG N021387 004 Jun 24, 2003

325MG, N/A; N/A, 40MG N021387 005 Jun 24, 2003

ASPIRIN; PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

DARVON W/ ASA

XANODYNE PHARM 325MG; 65MG N010996 005

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ASPIRIN; PROPOXYPHENE NAPSYLATE

CAPSULE; ORAL

DARVON-N W/ ASA

AAIPHARMA LLC

325MG;100MG

N016829 001

TABLET; ORAL

DARVON-N W/ ASA

AAIPHARMA LLC

325MG;100MG

N016863 001

ATAZANAVIR SULFATE

CAPSULE; ORAL

REYATAZ

+ BRISTOL MYERS SQUIBB EQ 100MG BASE **

N021567 001 Jun 20, 2003

ATENOLOL

INJECTABLE; INJECTION

TENORMIN

+ ASTRAZENECA

0.5MG/ML **

N019058 001 Sep 13, 1989

TABLET; ORAL

ATENOLOL

ABLE

25MG

A076907 001 Jul 30, 2004

50MG

A076907 002 Jul 30, 2004

100MG

A076907 003 Jul 30, 2004

APOTHECON

50MG

A073317 001 Mar 20, 1992

100MG

A073318 001 Mar 20, 1992

DAVA PHARMS INC

25MG

A074099 001 Apr 28, 1992

MYLAN

25MG

A074126 003 Aug 26, 1998

50MG

A074126 001 Mar 23, 1994

100MG

A074126 002 Mar 23, 1994

NORTHSTAR HLTHCARE

25MG

A078254 001 Sep 25, 2009

50MG

A078254 002 Sep 25, 2009

100MG

A078254 003 Sep 25, 2009

NOSTRUM LABS

50MG

A074127 001 Feb 21, 1995

100MG

A074127 002 Feb 21, 1995

PLIVA

25MG

A074101 001 Jul 17, 1997

50MG

A074101 002 Jul 17, 1997

100MG

A074101 003 Jul 17, 1997

SANDOZ

25MG

A074265 001 Feb 28, 1994

50MG

A074265 002 Feb 28, 1994

100MG

A074265 003 Feb 28, 1994

SCS

50MG

A073676 001 Oct 30, 1992

100MG

A073676 002 Oct 30, 1992

TEVA

50MG

A073315 001 May 28, 1993

100MG

A073316 001 May 28, 1993

TEVA PHARMS

50MG

A074120 001 Feb 24, 1995

100MG

A074120 002 Feb 24, 1995

WATSON LABS

50MG

A073352 001 Dec 27, 1991

WATSON LABS TEVA

100MG

A073353 001 Dec 27, 1991

ATENOLOL; CHLORTHALIDONE

TABLET; ORAL

ATENOLOL AND CHLORTHALIDONE

NOSTRUM LABS

50MG;25MG

A074404 001 May 14, 1998

100MG;25MG

A074404 002 May 14, 1998

PLIVA

50MG;25MG

A074107 001 Sep 24, 1997

100MG;25MG

A074107 002 Sep 24, 1997

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

ATOMOXETINE HYDROCHLORIDE

ZYDUS PHARMS USA INC

18MG

A079017 001 Sep 17, 2010

25MG

A079017 002 Sep 17, 2010

40MG

A079017 003 Sep 17, 2010

60MG

A079017 004 Sep 17, 2010

80MG

A079017 005 Sep 17, 2010

100MG

A079017 006 Sep 17, 2010

STRATTERA

LILLY

5MG

N021411 001 Nov 26, 2002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

TEVA PHARMS

EQ 10MG BASE

A078773 001 May 29, 2012

EQ 20MG BASE

A078773 002 May 29, 2012

EQ 40MG BASE

A078773 003 May 29, 2012

EQ 80MG BASE

A078773 004 May 29, 2012

ATORVASTATIN CALCIUM; EZETIMIBE

TABLET; ORAL

LIPTRUZET

+ MERCK SHARP DOHME

EQ 10MG BASE;10MG **

N200153 001 May 03, 2013

+

EQ 20MG BASE;10MG **

N200153 002 May 03, 2013

+

EQ 40MG BASE;10MG **

N200153 003 May 03, 2013

+

EQ 80MG BASE;10MG **

N200153 004 May 03, 2013

ATOVAQUONE

TABLET; ORAL

MEPRON

+ GLAXOSMITHKLINE LLC

250MG **

N020259 001 Nov 25, 1992

ATRACURIUM BESYLATE

INJECTABLE; INJECTION

ATRACURIUM BESYLATE

BAXTER HLTHCARE

10MG/ML

A074824 001 Sep 30, 1997

BAXTER HLTHCARE CORP

10MG/ML

A074753 001 Jan 23, 1997

HOSPIRA

10MG/ML

A074632 001 Dec 23, 1996

10MG/ML

A074740 001 Mar 28, 1997

TEVA PARENTERAL

10MG/ML

A074784 001 Jun 11, 1997

WATSON PHARMS TEVA

10MG/ML

A074945 001 Jul 28, 1998

ATRACURIUM BESYLATE PRESERVATIVE FREE

BAXTER HLTHCARE

10MG/ML

A074825 001 Sep 30, 1997

BAXTER HLTHCARE CORP

10MG/ML

A074768 001 Jan 23, 1997

HOSPIRA

10MG/ML

A074633 001 Dec 23, 1996

10MG/ML

A074639 001 Mar 25, 1997

10MG/ML

A074741 001 Mar 28, 1997

WATSON LABS INC

10MG/ML

A074944 001 Jul 28, 1998

TRACRIUM

+ HOSPIRA

10MG/ML **

N018831 002 Jun 20, 1985

TRACRIUM PRESERVATIVE FREE

+ HOSPIRA

10MG/ML **

N018831 001 Nov 23, 1983

ATROPINE

INJECTABLE; INJECTION

ATROPINE

ABBVIE

EQ 2MG SULFATE/0.7ML

A071295 001 Jan 30, 1987

SOLUTION; INTRAMUSCULAR

ATROPINE (AUTOINJECTOR)

RAFA LABS LTD

EQ 2MG SULFATE/0.7ML (EQ 2MG
SULFATE/0.7ML)

N212319 001 Jul 09, 2018

ATROPINE SULFATE

AEROSOL, METERED; INHALATION

ATROPINE SULFATE

US ARMY

EQ 0.36MG BASE/INH

N020056 001 Sep 19, 1990

SOLUTION; INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS, ENDOTRACHEAL

ATROPINE SULFATE ANSYR PLASTIC SYRINGE

+ HOSPIRA

0.5MG/5ML (0.1MG/ML)

N021146 001 Jul 09, 2001

ATROPINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET; ORAL

MOTOFEN HALF-STRENGTH

SEBELA IRELAND LTD

0.025MG;0.5MG

N017744 001

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENOXYLATE HYDROCHLORIDE W/ ATROPINE SULFATE

SCHERER RP

0.025MG;2.5MG

A086440 001

SOLUTION; ORAL

COLONAIID

MEDPOINTE PHARM HLC

0.025MG/5ML;2.5MG/5ML

A085735 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

SOLUTION; ORAL

LOMANATE

ALPHARMA US PHARMS 0.025MG/5ML; 2.5MG/5ML A085746 001

LOMOTIL

GD SEARLE LLC 0.025MG/5ML; 2.5MG/5ML N012699 001

TABLET; ORAL

COLONAIID

MEDPOINTE PHARM HLC 0.025MG; 2.5MG A085737 001

DI-ATRO

MD PHARM 0.025MG; 2.5MG A085266 001

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

ABLE 0.025MG; 2.5MG A040395 001 Nov 27, 2000

ASCOT 0.025MG; 2.5MG A087934 001 Jul 19, 1983

FOSUN PHARMA 0.025MG; 2.5MG A086173 001

HEATHER 0.025MG; 2.5MG A086798 001

HIKMA PHARMS 0.025MG; 2.5MG A087765 001 Mar 15, 1982

INWOOD LABS 0.025MG; 2.5MG A085509 001

KV PHARM 0.025MG; 2.5MG A085659 001

LEDERLE 0.025MG; 2.5MG A086950 001

PARKE DAVIS 0.025MG; 2.5MG A087131 001

PVT FORM 0.025MG; 2.5MG A085766 001

R AND S PHARMA 0.025MG; 2.5MG A085035 001

ROXANE 0.025MG; 2.5MG A086057 001

SUN PHARM INDUSTRIES 0.025MG; 2.5MG A085506 001

USL PHARMA 0.025MG; 2.5MG A087842 001 Mar 29, 1982

VALEANT PHARM INTL 0.025MG; 2.5MG A087195 001 Feb 16, 1982

WATSON LABS 0.025MG; 2.5MG A085876 001

LO-TROL

VANGARD 0.025MG; 2.5MG A088009 001 Mar 25, 1983

LOGEN

SUPERPHARM 0.025MG; 2.5MG A088962 001 May 10, 1985

LONOX

FOSUN PHARMA 0.025MG; 2.5MG A085311 002

LOW-QUEL

HALSEY 0.025MG; 2.5MG A085211 001

ATROPINE SULFATE; EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

ENLON-PLUS

MYLAN INSTITUTIONAL 0.14MG/ML; 10MG/ML N019677 001 Nov 06, 1991

+ 0.14MG/ML; 10MG/ML N019678 001 Nov 06, 1991

ATROPINE SULFATE; MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

ATROPINE AND DEMEROL

ABBVIE 0.4MG/ML; 50MG/ML A087853 001 Nov 26, 1982

0.4MG/ML; 75MG/ML A087847 001 Nov 26, 1982

0.4MG/ML; 100MG/ML A087848 001 Nov 26, 1982

MEPERIDINE AND ATROPINE SULFATE

WYETH AYERST 0.4MG/ML; 50MG/ML A085121 001

0.4MG/ML; 75MG/ML A085121 002

0.4MG/ML; 100MG/ML A085121 003

ATROPINE; PRALIDOXIME CHLORIDE

INJECTABLE; INTRAMUSCULAR

ATNAA

US ARMY 2.1MG/0.7ML; 600MG/2ML N021175 001 Jan 17, 2002

AVOBENZONE; OCTINOXATE; OXYBENZONE

LOTION; TOPICAL

SHADE UVAGUARD

+ BAYER HEALTHCARE LLC 3%; 7.5%; 3% N020045 001 Dec 07, 1992

AZATADINE MALEATE

TABLET; ORAL

OPTIMINE

SCHERING 1MG N017601 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AZATADINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

TRINALIN

SCHERING

1MG;120MG

N018506 001 Mar 23, 1982

AZATHIOPRINE

TABLET;ORAL

IMURAN

+

SEBELA IRELAND LTD

25MG **

N016324 002

AZATHIOPRINE SODIUM

INJECTABLE;INJECTION

IMURAN

+

CASPER PHARMA LLC

EQ 100MG BASE/VIAL **

N017391 001

AZELASTINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

OPTIVAR

+

MYLAN SPECIALITY LP

0.05% **

N021127 001 May 22, 2000

SPRAY, METERED;NASAL

ASTEPRO

MYLAN SPECIALITY LP

EQ 0.125MG BASE/SPRAY

N022203 001 Oct 15, 2008

AZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE

SPRAY, METERED;NASAL

AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE

APOTEX INC

EQ 0.125MG BASE/SPRAY;0.05MG/SPRAY

A207712 001 Apr 28, 2017

AZITHROMYCIN

CAPSULE;ORAL

ZITHROMAX

+

PFIZER

EQ 250MG BASE **

N050670 001 Nov 01, 1991

FOR SUSPENSION;ORAL

AZITHROMYCIN

SANDOZ

EQ 100MG BASE/5ML

A065297 001 Sep 18, 2006

EQ 200MG BASE/5ML

A065297 002 Sep 18, 2006

FOR SUSPENSION, EXTENDED RELEASE;ORAL

ZMAX

+

PF PRISM CV

EQ 2GM BASE/BOT

N050797 001 Jun 10, 2005

INJECTABLE;INJECTION

AZITHROMYCIN

CSPC OUYI PHARM CO

EQ 500MG BASE/VIAL

A065265 001 Jan 18, 2007

TEVA PARENTERAL

EQ 500MG BASE/VIAL

N050809 001 Dec 19, 2006

EQ 2.5GM BASE/VIAL

N050809 002 Dec 19, 2006

TABLET;ORAL

AZITHROMYCIN

APOTEX CORP

EQ 250MG BASE

A065507 001 Jul 13, 2011

EQ 500MG BASE

A065509 001 Jul 13, 2011

EQ 600MG BASE

A065508 001 Jul 13, 2011

MYLAN

EQ 250MG BASE

A065365 001 May 30, 2007

EQ 500MG BASE

A065366 001 May 30, 2007

AZITHROMYCIN DIHYDRATE; TROVAFLOXACIN MESYLATE

FOR SUSPENSION, TABLET;ORAL

TROVAN/ZITHROMAX COMPLIANCE PAK

PFIZER

EQ 1GM BASE,N/A;N/A,EQ 100MG BASE

N050762 001 Dec 18, 1998

AZLOCILLIN SODIUM

INJECTABLE;INJECTION

AZLIN

BAYER PHARMS

EQ 2GM BASE/VIAL

A062388 001 Sep 08, 1982

EQ 2GM BASE/VIAL

A062417 001 Oct 12, 1982

EQ 2GM BASE/VIAL

N050562 001 Sep 03, 1982

EQ 3GM BASE/VIAL

A062388 002 Sep 08, 1982

EQ 3GM BASE/VIAL

A062417 002 Oct 12, 1982

EQ 3GM BASE/VIAL

N050562 002 Sep 03, 1982

EQ 4GM BASE/VIAL

A062388 003 Sep 08, 1982

EQ 4GM BASE/VIAL

A062417 003 Oct 12, 1982

EQ 4GM BASE/VIAL

N050562 003 Sep 03, 1982

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

AZTREONAM

INJECTABLE; INJECTION

AZACTAM

BRISTOL MYERS SQUIBB 500MG/VIAL

N050580 001 Dec 31, 1986

AZACTAM IN PLASTIC CONTAINER

BRISTOL MYERS SQUIBB 10MG/ML

N050632 003 May 24, 1989

AZTREONAM

WEST-WARD PHARMS INT 1GM/VIAL

A065286 001 Mar 23, 2011

2GM/VIAL

A065286 002 Mar 23, 2011

BACAMPICILLIN HYDROCHLORIDE

FOR SUSPENSION; ORAL

SPECTROBID

PFIZER 125MG/5ML

N050556 001 Mar 23, 1982

TABLET; ORAL

SPECTROBID

PFIZER 400MG

N050520 001

800MG

N050520 002 Sep 12, 1983

BACITRACIN

INJECTABLE; INJECTION

BACITRACIN

MYLAN ASI 50,000 UNITS/VIAL

A090211 001 May 11, 2010

PFIZER 50,000 UNITS/VIAL

A060282 001

PHARMACIA AND UPJOHN 10,000 UNITS/VIAL

A060733 001

OINTMENT; OPHTHALMIC

BACIGUENT

PHARMACIA AND UPJOHN 500 UNITS/GM

A060734 001

BACITRACIN

LILLY 500 UNITS/GM

A060687 001

PHARMADERM 500 UNITS/GM

A062158 001

PHARMAFAIR 500 UNITS/GM

A062453 001 Mar 28, 1984

OINTMENT; TOPICAL

BACITRACIN

COMBE 500 UNITS/GM

A062799 001 May 14, 1987

NASKA 500 UNITS/GM

A062857 001 Nov 13, 1987

POWDER; FOR RX COMPOUNDING

BACI-RX

X GEN PHARMS 5,000,000 UNITS/BOT

A061580 001

BACITRACIN

APOTHEKERNES 5,000,000 UNITS/BOT

A061699 001

PADDOCK LLC 5,000,000 UNITS/BOT

A062456 001 Jul 27, 1983

BACITRACIN ZINC

POWDER; FOR RX COMPOUNDING

ZIBA-RX

X GEN PHARMS 500,000 UNITS/BOT

A061737 001

BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

CORTISPORIN

+ CASPER PHARMA LLC 400 UNITS/GM; 1%; EQ 3.5MG BASE/GM; 10,000 UNITS/GM **

N050416 002

ZINC BACITRACIN, NEOMYCIN SULFATE, POLYMYXIN B SULFATE & HYDROCORTISONE

PHARMAFAIR 400 UNITS/GM; 1%; EQ 3.5MG BASE/GM; 10,000 UNITS/GM

A062389 001 Jul 02, 1982

OINTMENT; TOPICAL

NEOMYCIN & POLYMYXIN B SULFATES & BACITRACIN ZINC & HYDROCORTISONE

PHARMAFAIR 400 UNITS/GM; 1%; EQ 3.5MG BASE/GM; 5,000 UNITS/GM

A062381 001 Sep 06, 1985

BACITRACIN ZINC; LIDOCAINE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; TOPICAL

LANABIOTIC

COMBE 400 UNITS/GM; 40MG/GM; EQ 5MG BASE/GM; 5,000 UNITS/GM

A062499 001 Jun 03, 1985

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BACITRACIN ZINC; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

BACITRACIN ZINC-NEOMYCIN SULFATE-POLYMYXIN B SULFATE

| | | | |
|------------|-----------------------------------------------|-------------|--------------|
| PHARMAFAIR | 400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM | A062386 001 | Sep 09, 1982 |
|------------|-----------------------------------------------|-------------|--------------|

BACITRACIN-NEOMYCIN-POLYMYXIN

| | | | |
|------------|----------------------------------------------|-------------|--|
| PHARMADERM | 400 UNITS/GM;EQ 3.5MG BASE/GM;5,000 UNITS/GM | A062167 001 | |
|------------|----------------------------------------------|-------------|--|

NEO-POLYICIN

| | | | |
|-----------|-----------------------------------------------|-------------|--|
| DOW PHARM | 500 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM | A060647 001 | |
|-----------|-----------------------------------------------|-------------|--|

OINTMENT;TOPICAL

BACITRACIN ZINC-NEOMYCIN SULFATE-POLYMYXIN B SULFATE

| | | | |
|-------|----------------------------------------------|-------------|--------------|
| NASKA | 400 UNITS/GM;EQ 3.5MG BASE/GM;5,000 UNITS/GM | A062833 001 | Nov 09, 1987 |
|-------|----------------------------------------------|-------------|--------------|

BACITRACIN ZINC; POLYMYXIN B SULFATE

AEROSOL;TOPICAL

POLYSPORIN

| | | | |
|-----------------|------------------------------------|-------------|--------------|
| GLAXOSMITHKLINE | 10,000 UNITS/GM;2,000,000 UNITS/GM | N050167 002 | Mar 01, 1985 |
|-----------------|------------------------------------|-------------|--------------|

OINTMENT;OPHTHALMIC

OCUMYCIN

| | | | |
|------------|------------------------------|-------------|--------------|
| PHARMAFAIR | 500 UNITS/GM;10,000 UNITS/GM | A062430 001 | Apr 08, 1983 |
|------------|------------------------------|-------------|--------------|

POLYSPORIN

| | | | |
|----------------|---------------------------------|-------------|--|
| MONARCH PHARMS | 500 UNITS/GM;10,000 UNITS/GM ** | A061229 001 | |
|----------------|---------------------------------|-------------|--|

OINTMENT;TOPICAL

BACITRACIN ZINC-POLYMYXIN B SULFATE

| | | | |
|-------|------------------------------|-------------|--------------|
| NASKA | 500 UNITS/GM;10,000 UNITS/GM | A062849 001 | Nov 13, 1987 |
|-------|------------------------------|-------------|--------------|

BACITRACIN; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE

| | | | |
|--------|--------------------------------------------------|-------------|--|
| ALTANA | 400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM | A060731 002 | |
|--------|--------------------------------------------------|-------------|--|

BACITRACIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

MYCITRACIN

| | | | |
|----------------------|-----------------------------------------------|-------------|--|
| PHARMACIA AND UPJOHN | 500 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM | A061048 001 | |
|----------------------|-----------------------------------------------|-------------|--|

BACITRACIN; POLYMYXIN B SULFATE

DISC;TOPICAL

LANABIOTIC

| | | | |
|-------|-----------------------------|-------------|--------------|
| COMBE | 500 UNITS/GM;5,000 UNITS/GM | N050598 001 | Sep 22, 1986 |
|-------|-----------------------------|-------------|--------------|

BACLOFEN

TABLET;ORAL

BACLOFEN

| | | | |
|-------|------|-------------|--------------|
| MYLAN | 10MG | A077181 001 | Jul 29, 2005 |
|-------|------|-------------|--------------|

| | | | |
|------|------|-------------|--------------|
| TEVA | 10MG | A073043 001 | Feb 27, 1992 |
|------|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 20MG | A073044 001 | Feb 27, 1992 |
|--|------|-------------|--------------|

| | | | |
|------------|------|-------------|--------------|
| USL PHARMA | 10MG | A071260 001 | May 06, 1988 |
|------------|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 20MG | A071261 001 | May 06, 1988 |
|--|------|-------------|--------------|

| | | | |
|-------------|------|-------------|--------------|
| WATSON LABS | 10MG | A072824 001 | Sep 18, 1991 |
|-------------|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 10MG | A073092 001 | Jan 28, 1994 |
|--|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 10MG | A074698 001 | Aug 20, 1996 |
|--|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 20MG | A072825 001 | Sep 18, 1991 |
|--|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 20MG | A073093 001 | Jan 28, 1994 |
|--|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 20MG | A074698 002 | Aug 20, 1996 |
|--|------|-------------|--------------|

Lioresal

| | | | |
|------------|---------|-------------|--|
| + NOVARTIS | 10MG ** | N017851 001 | |
|------------|---------|-------------|--|

| | | | |
|---|---------|-------------|--------------|
| + | 20MG ** | N017851 003 | Jan 20, 1982 |
|---|---------|-------------|--------------|

TABLET, ORALLY DISINTEGRATING;ORAL

KEMSTRO

| | | | |
|---------|------|-------------|--------------|
| UCB INC | 10MG | N021589 001 | Oct 30, 2003 |
|---------|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 20MG | N021589 002 | Oct 30, 2003 |
|--|------|-------------|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BARIUM SULFATE

FOR SUSPENSION;ORAL

E-Z-CAT DRY

+ BRACCO 40% (9GM/POUCH) N208036 003 Jan 03, 2017

BECLOMETHASONE DIPROPIONATE

AEROSOL, METERED;INHALATION

BECLOVENT

GLAXOSMITHKLINE 0.042MG/INH N018153 001

QVAR 40

+ TEVA BRANDED PHARM 0.04MG/INH N020911 002 Sep 15, 2000

QVAR 80

+ TEVA BRANDED PHARM 0.08MG/INH N020911 001 Sep 15, 2000

VANCERIL

SCHERING 0.042MG/INH N017573 001

VANCERIL DOUBLE STRENGTH

SCHERING 0.084MG/INH N020486 001 Dec 24, 1996

AEROSOL, METERED;NASAL

BECONASE

GLAXOSMITHKLINE 0.042MG/INH N018584 001

VANCENASE

SCHERING 0.042MG/INH N018521 001

BECLOMETHASONE DIPROPIONATE MONOHYDRATE

SPRAY, METERED;NASAL

VANCENASE AQ

SCHERING EQ 0.042MG DIPROP/SPRAY N019589 001 Dec 23, 1987

EQ 0.084MG DIPROP/SPRAY N020469 001 Jun 26, 1996

BENZAEPRIIL HYDROCHLORIDE

TABLET;ORAL

BENZAEPRIIL HYDROCHLORIDE

ACTAVIS LABS FL INC 5MG A076267 001 Feb 11, 2004

10MG A076267 002 Feb 11, 2004

20MG A076267 003 Feb 11, 2004

40MG A076267 004 Feb 11, 2004

GENPHARM 5MG A076476 001 Feb 11, 2004

10MG A076476 002 Feb 11, 2004

20MG A076476 003 Feb 11, 2004

40MG A076476 004 Feb 11, 2004

BENZAEPRIIL HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

BENZAEPRIIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

ACTAVIS LABS FL INC 5MG;6.25MG A076342 001 Feb 11, 2004

10MG;12.5MG A076342 002 Feb 11, 2004

20MG;12.5MG A076342 003 Feb 11, 2004

20MG;25MG A076342 004 Feb 11, 2004

IVAX SUB TEVA PHARMS 5MG;6.25MG A076348 001 Feb 11, 2004

10MG;12.5MG A076348 002 Feb 11, 2004

20MG;12.5MG A076348 003 Feb 11, 2004

20MG;25MG A076348 004 Feb 11, 2004

MYLAN PHARMS INC 5MG;6.25MG A076612 001 Feb 11, 2004

10MG;12.5MG A076612 002 Feb 11, 2004

20MG;12.5MG A076612 003 Feb 11, 2004

20MG;25MG A076612 004 Feb 11, 2004

SUN PHARM INDS LTD 5MG;6.25MG A077483 001 Sep 08, 2005

10MG;12.5MG A077483 002 Sep 08, 2005

20MG;12.5MG A077483 003 Sep 08, 2005

20MG;25MG A077483 004 Sep 08, 2005

LOTENSIN HCT

+ US PHARMS HOLDINGS I 5MG;6.25MG ** N020033 001 May 19, 1992

BENDAMUSTINE HYDROCHLORIDE

SOLUTION;IV (INFUSION)

TREANDA

+ CEPHALON 45MG/0.5ML (90MG/ML) N022249 003 Sep 13, 2013

+ 180MG/2ML (90MG/ML) N022249 004 Sep 13, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BENDROFLUMETHIAZIDE

TABLET; ORAL

NATURETIN-10

APOTHECON

10MG

N012164 003

NATURETIN-2.5

APOTHECON

2.5MG

N012164 001

NATURETIN-5

APOTHECON

5MG

N012164 002

BENDROFLUMETHIAZIDE; NADOLOL

TABLET; ORAL

NADOLOL AND BENDROFLUMETHIAZIDE

MYLAN

5MG; 40MG

A078688 001 Feb 15, 2008

5MG; 80MG

A078688 002 Feb 15, 2008

BENOXINATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

BENOXINATE HYDROCHLORIDE

SOLA BARNES HIND

0.4%

A084149 001

BENTIROMIDE

SOLUTION; ORAL

CHYMEX

SAVAGE LABS

500MG/7.5ML

N018366 001 Dec 29, 1983

BENZONATATE

CAPSULE; ORAL

BENZONATATE

NESHER PHARMS

100MG

A040795 001 Oct 31, 2007

200MG

A040795 002 Oct 31, 2007

TESSALON

+ PFIZER

200MG **

N011210 003 Jun 25, 1999

BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

BENZACLIN

VALEANT BERMUDA

5%; EQ 1% BASE

N050756 002 Apr 20, 2007

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL

BENZPHETAMINE HYDROCHLORIDE

EPIC PHARMA LLC

50MG

A040714 001 Oct 29, 2007

IMPAX LABS

50MG

A040845 001 Nov 18, 2008

TEDOR PHARM

25MG

A040747 002 Nov 20, 2015

50MG

A040747 001 Mar 30, 2007

DIDREX

+ PHARMACIA AND UPJOHN

25MG **

N012427 003

+

50MG **

N012427 002

BENZQUINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

EMETE-CON

PFIZER

EQ 50MG BASE/VIAL

N016820 001

SUPPOSITORY; RECTAL

EMETE-CON

ROERIG

EQ 100MG BASE

N016818 006

BENZTHIAZIDE

TABLET; ORAL

AQUATAG

SOLVAY

25MG

N016001 001

50MG

N016001 002

BENZTHIAZIDE

PVT FORM

50MG

A083206 001

EXNA

AH ROBINS INC

50MG

N012489 001

FOVANE

PFIZER

50MG

N012128 002

URESE

PFIZER

25MG

N012128 003

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BENZTROPINE MESYLATE

TABLET; ORAL

BENZTROPINE MESYLATE

| | | | |
|------------------|----------|-------------|--------------|
| CHARTWELL RX | 1MG | A081265 002 | Jan 23, 1992 |
| | 2MG | A081265 001 | Jan 23, 1992 |
| LANNETT CO INC | 0.5MG ** | A088877 001 | Apr 11, 1985 |
| | 1MG ** | A088894 001 | Apr 11, 1985 |
| | 2MG ** | A088895 001 | Apr 11, 1985 |
| OXFORD PHARMS | 2MG | A040706 001 | Feb 14, 2008 |
| QUANTUM PHARMICS | 0.5MG | A088514 001 | Jan 31, 1984 |
| | 1MG | A088510 001 | Jan 31, 1984 |
| | 2MG | A088511 001 | Jan 31, 1984 |
| USL PHARMA | 0.5MG | A089211 001 | Jun 14, 1988 |
| | 1MG | A089212 001 | Jun 14, 1988 |
| | 2MG | A089213 001 | Jun 14, 1988 |
| COGENTIN | | | |
| + MERCK | 0.5MG ** | N009193 004 | |
| + | 1MG ** | N009193 003 | |
| + | 2MG ** | N009193 002 | |

BENZYL BENZOATE

EMULSION; TOPICAL

BENZYL BENZOATE

| | | | |
|---------|-----|-------------|--|
| LANNETT | 50% | A084535 001 | |
|---------|-----|-------------|--|

BEPRIDIL HYDROCHLORIDE

TABLET; ORAL

BEPADIN

| | | | |
|---------------------|-------|-------------|--------------|
| MEDPOINTE PHARM HLC | 200MG | N019001 001 | Dec 28, 1990 |
| | 300MG | N019001 002 | Dec 28, 1990 |
| | 400MG | N019001 003 | Dec 28, 1990 |
| VASCOR | | | |
| JOHNSON AND JOHNSON | 200MG | N019002 001 | Dec 28, 1990 |
| | 300MG | N019002 002 | Dec 28, 1990 |
| | 400MG | N019002 003 | Dec 28, 1990 |

BETA CAROTENE

CAPSULE; ORAL

SOLATENE

| | | | |
|-------|------|-------------|--|
| ROCHE | 30MG | N017589 001 | |
|-------|------|-------------|--|

BETAMETHASONE

CREAM; TOPICAL

CELESTONE

| | | | |
|----------|------|-------------|--|
| SCHERING | 0.2% | N014762 001 | |
|----------|------|-------------|--|

SYRUP; ORAL

CELESTONE

| | | | |
|-------------------|-----------|-------------|--|
| MERCK SHARP DOHME | 0.6MG/5ML | N014215 002 | |
|-------------------|-----------|-------------|--|

TABLET; ORAL

CELESTONE

| | | | |
|----------|-------|-------------|--|
| SCHERING | 0.6MG | N012657 003 | |
|----------|-------|-------------|--|

BETAMETHASONE BENZOATE

CREAM; TOPICAL

UTICORT

| | | | |
|-------------|--------|-------------|--|
| PARKE DAVIS | 0.025% | N016998 002 | |
|-------------|--------|-------------|--|

GEL; TOPICAL

UTICORT

| | | | |
|-------------|--------|-------------|--|
| PARKE DAVIS | 0.025% | N017244 001 | |
|-------------|--------|-------------|--|

LOTION; TOPICAL

UTICORT

| | | | |
|-------------|--------|-------------|--|
| PARKE DAVIS | 0.025% | N017528 001 | |
|-------------|--------|-------------|--|

OINTMENT; TOPICAL

UTICORT

| | | | |
|-------------|--------|-------------|--|
| PARKE DAVIS | 0.025% | N018089 001 | |
|-------------|--------|-------------|--|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BETAMETHASONE DIPROPIONATE

CREAM;TOPICAL

ALPHATREX

SAVAGE LABS EQ 0.05% BASE N019138 001 Jun 26, 1984

BETAMETHASONE DIPROPIONATE

PERRIGO NEW YORK EQ 0.05% BASE A072536 001 Jan 31, 1990

EQ 0.05% BASE A074579 001 Nov 26, 1997

PHARMADERM EQ 0.05% BASE N019136 001 Jun 26, 1984

TARO EQ 0.05% BASE A071143 001 Jun 17, 1987

TEVA EQ 0.05% BASE A071476 001 Aug 10, 1987

DIPROSONE

SCHERING EQ 0.05% BASE N017536 001

CREAM, AUGMENTED;TOPICAL

DIPROLENE

SCHERING EQ 0.05% BASE N019408 001 Jan 31, 1986

DISC;TOPICAL

DIPROSONE

SCHERING EQ 0.1% BASE N017829 001

GEL, AUGMENTED;TOPICAL

DIPROLENE

SCHERING EQ 0.05% BASE N019408 002 Nov 22, 1991

LOTION;TOPICAL

ALPHATREX

SAVAGE LABS EQ 0.05% BASE A070273 001 Aug 12, 1985

BETAMETHASONE DIPROPIONATE

ALPHARMA US PHARMS EQ 0.05% BASE A071085 001 Feb 03, 1987

G AND W LABS INC EQ 0.05% BASE A071882 001 Jun 06, 1988

PHARMADERM EQ 0.05% BASE A070274 001 Aug 12, 1985

TARO EQ 0.05% BASE A072276 001 Aug 24, 1988

EQ 0.05% BASE A074272 001 Sep 30, 1994

DIPROSONE

+ SCHERING EQ 0.05% BASE ** N017781 001

LOTION, AUGMENTED;TOPICAL

DIPROLENE

+ MERCK SHARP DOHME EQ 0.05% BASE N019716 001 Aug 01, 1988

OINTMENT;TOPICAL

ALPHATREX

SAVAGE LABS EQ 0.05% BASE N019143 001 Sep 04, 1984

BETAMETHASONE DIPROPIONATE

PERRIGO NEW YORK EQ 0.05% BASE A072526 001 Jan 31, 1990

PHARMADERM EQ 0.05% BASE N019140 001 Sep 04, 1984

TEVA EQ 0.05% BASE A071477 001 Aug 10, 1987

DIPROSONE

SCHERING EQ 0.05% BASE N017691 001

BETAMETHASONE SODIUM PHOSPHATE

INJECTABLE;INJECTION

BETAMETHASONE SODIUM PHOSPHATE

WATSON LABS EQ 3MG BASE/ML A085738 001

CELESTONE

+ SCHERING EQ 3MG BASE/ML ** N017561 001

BETAMETHASONE VALERATE

CREAM;TOPICAL

BETADERM

ROACO EQ 0.1% BASE N018839 001 Jun 30, 1983

BETAMETHASONE VALERATE

PERRIGO NEW YORK EQ 0.1% BASE A070053 001 Jun 10, 1986

PHARMADERM EQ 0.1% BASE N018860 002 Aug 31, 1983

PHARMAFAIR EQ 0.1% BASE A070485 001 May 29, 1987

TARO EQ 0.1% BASE A070062 001 May 14, 1985

BETATREX

SAVAGE LABS EQ 0.1% BASE N018862 001 Aug 31, 1983

VALISONE

SCHERING EQ 0.01% BASE N016322 002

EQ 0.1% BASE N016322 001

LOTION;TOPICAL

BETA-VAL

G AND W LABS INC EQ 0.1% BASE A070072 001 Jun 27, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BETAMETHASONE VALERATE

LOTION; TOPICAL

BETAMETHASONE VALERATE

| | | | |
|-------------|--------------|-------------|--------------|
| PHARMADERM | EQ 0.1% BASE | N018870 001 | Aug 31, 1983 |
| PHARMAFAIR | EQ 0.1% BASE | A070484 001 | May 29, 1987 |
| TEVA PHARMS | EQ 0.1% BASE | A071883 001 | Apr 22, 1988 |

BETATREX

| | | | |
|-------------|--------------|-------------|--------------|
| SAVAGE LABS | EQ 0.1% BASE | N018867 001 | Aug 31, 1983 |
|-------------|--------------|-------------|--------------|

VALISONE

| | | | |
|----------|--------------|-------------|--|
| SCHERING | EQ 0.1% BASE | N016932 001 | |
|----------|--------------|-------------|--|

OINTMENT; TOPICAL

BETAMETHASONE VALERATE

| | | | |
|------------------|--------------|-------------|--------------|
| PERRIGO NEW YORK | EQ 0.1% BASE | A071478 001 | Dec 23, 1987 |
| PHARMADERM | EQ 0.1% BASE | N018864 001 | Aug 31, 1983 |
| PHARMAFAIR | EQ 0.1% BASE | A070486 001 | May 29, 1987 |

BETATREX

| | | | |
|-------------|--------------|-------------|--------------|
| SAVAGE LABS | EQ 0.1% BASE | N018863 001 | Aug 31, 1983 |
|-------------|--------------|-------------|--------------|

VALISONE

| | | | |
|----------|--------------|-------------|--|
| SCHERING | EQ 0.1% BASE | N016740 001 | |
|----------|--------------|-------------|--|

BETAXOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

BETAXOLOL HYDROCHLORIDE

| | | | |
|------------|--------------|-------------|--------------|
| APOTEX INC | EQ 0.5% BASE | A075446 001 | Sep 28, 2000 |
|------------|--------------|-------------|--------------|

TABLET; ORAL

KERLONE

| | | | |
|-------------------|------|-------------|--------------|
| SANOFI AVENTIS US | 10MG | N019507 001 | Oct 27, 1989 |
| | 20MG | N019507 002 | Oct 27, 1989 |

BETAXOLOL HYDROCHLORIDE; CHLORTHALIDONE

TABLET; ORAL

KERLEDEX

| | | | |
|-------------------|-------------|-------------|--------------|
| SANOFI AVENTIS US | 5MG;12.5MG | N019807 001 | Oct 30, 1992 |
| | 10MG;12.5MG | N019807 002 | Oct 30, 1992 |

BETAXOLOL HYDROCHLORIDE; Pilocarpine Hydrochloride

SUSPENSION/DROPS; OPHTHALMIC

BETOPTIC PILO

| | | | |
|-------|---------------------|-------------|--------------|
| ALCON | EQ 0.25% BASE;1.75% | N020619 001 | Apr 17, 1997 |
|-------|---------------------|-------------|--------------|

BETAZOLE HYDROCHLORIDE

INJECTABLE; INJECTION

HISTALOG

| | | | |
|-------|---------|-------------|--|
| LILLY | 50MG/ML | N009344 001 | |
|-------|---------|-------------|--|

BETHANECHOL CHLORIDE

INJECTABLE; INJECTION

URECHOLINE

+ ODYSSEY PHARMS

| | | |
|-----------|-------------|--|
| 5MG/ML ** | N006536 001 | |
|-----------|-------------|--|

TABLET; ORAL

BETHANECHOL CHLORIDE

| | | | |
|----------------------|------|-------------|--------------|
| ABLE | 5MG | A040492 001 | Jul 27, 2004 |
| | 10MG | A040483 001 | Jul 27, 2004 |
| | 25MG | A040485 001 | Jul 27, 2004 |
| | 50MG | A040509 001 | Jul 27, 2004 |
| ACTAVIS ELIZABETH | 5MG | A040552 001 | Oct 28, 2004 |
| | 10MG | A040553 001 | Oct 28, 2004 |
| | 25MG | A040554 001 | Oct 28, 2004 |
| | 50MG | A040551 001 | Oct 28, 2004 |
| ASCOT | 10MG | A088288 001 | Jun 08, 1983 |
| | 25MG | A088289 001 | Jun 08, 1983 |
| IMPAX LABS | 5MG | A040721 001 | Nov 01, 2006 |
| | 10MG | A040721 002 | Nov 01, 2006 |
| | 25MG | A040721 003 | Nov 01, 2016 |
| | 50MG | A040721 004 | Nov 01, 2006 |
| IVAX SUB TEVA PHARMS | 25MG | A084689 001 | |
| LANNETT | 5MG | A084702 001 | |
| | 10MG | A084712 001 | |
| | 25MG | A084074 001 | |
| SANDOZ | 5MG | A084353 001 | |
| | 10MG | A084378 001 | |
| | 10MG | A084379 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BETHANECHOL CHLORIDE

TABLET; ORAL

BETHANECHOL CHLORIDE

| | | | | |
|--------------------|---------|---------|-----|--------------|
| | 25MG | A084383 | 001 | |
| | 25MG | A084384 | 001 | |
| SUN PHARM INDS INC | 5MG | A040897 | 001 | Apr 22, 2009 |
| | 10MG | A040897 | 002 | Apr 22, 2009 |
| | 25MG | A040897 | 003 | Apr 22, 2009 |
| | 50MG | A040897 | 004 | Apr 22, 2009 |
| WATSON LABS | 5MG | A084402 | 001 | |
| | 5MG | A085230 | 002 | |
| | 5MG | A085841 | 001 | |
| | 10MG | A084408 | 001 | |
| | 10MG | A085228 | 001 | |
| | 10MG | A085842 | 001 | |
| | 25MG | A084441 | 001 | |
| | 25MG | A085229 | 001 | |
| | 25MG | A085839 | 001 | |
| | 50MG | A087397 | 001 | |
| | 50MG | A087444 | 001 | |
| MYOTONACHOL | | | | |
| GLENWOOD | 5MG | A084188 | 001 | |
| | 10MG | A084188 | 003 | |
| | 25MG | A084188 | 004 | |
| URECHOLINE | | | | |
| + ODYSSEY PHARMS | 5MG ** | N006536 | 003 | |
| + | 10MG ** | N006536 | 002 | |
| + | 25MG ** | N006536 | 004 | |
| + | 50MG ** | N006536 | 005 | |

BETHANIDINE SULFATE

TABLET; ORAL

TENATHAN

| | | | | |
|-----------|------|---------|-----|--|
| ROBINS AH | 10MG | N017675 | 001 | |
| | 25MG | N017675 | 002 | |

BICALUTAMIDE

TABLET; ORAL

BICALUTAMIDE

| | | | | |
|----------------|------|---------|-----|--------------|
| KUDCO IRELAND | 50MG | A077995 | 001 | Jul 06, 2009 |
| ROXANE | 50MG | A078285 | 001 | Mar 24, 2011 |
| SYNTHON PHARMS | 50MG | A077973 | 001 | Jul 06, 2009 |

BIMATOPROST

SOLUTION/DROPS; OPHTHALMIC

LUMIGAN

| | | | | |
|------------|----------|---------|-----|--------------|
| + ALLERGAN | 0.03% ** | N021275 | 001 | Mar 16, 2001 |
|------------|----------|---------|-----|--------------|

BIPERIDEN HYDROCHLORIDE

TABLET; ORAL

AKINETON

| | | | | |
|--------|-----|---------|-----|--|
| ABBVIE | 2MG | N012003 | 001 | |
|--------|-----|---------|-----|--|

BIPERIDEN LACTATE

INJECTABLE; INJECTION

AKINETON

| | | | | |
|--------|--------|---------|-----|--|
| ABBVIE | 5MG/ML | N012418 | 002 | |
|--------|--------|---------|-----|--|

BISACODYL; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION, TABLET, DELAYED RELEASE; ORAL

HALFLYTELY

| | | | | |
|--------------------------------------------------------------------------------------|---------------------------------------------------------------|---------|-----|--------------|
| + BRAINTREE | 5MG, N/A; N/A, 210GM; N/A, 0.74GM; N/A, 2.86GM; N/A, 5.6GM ** | N021551 | 003 | Jul 16, 2010 |
| PEG-3350, SODIUM CHLORIDE, SODIUM BICARBONATE, POTASSIUM CHLORIDE AND NOVEL LABS INC | 5MG, N/A; N/A, 210GM; N/A, 0.74GM; N/A, 2.86GM; N/A, 5.6GM | A202217 | 001 | Aug 20, 2014 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BISMUTH SUBSALICYLATE; METRONIDAZOLE; TETRACYCLINE HYDROCHLORIDE

TABLET, CHEWABLE, TABLET, CAPSULE; ORAL

HELIDAC

| | | | | |
|---|-------------------|------------------------------------------------|-------------|--------------|
| + | CASPER PHARMA LLC | 262.4MG,N/A,N/A;N/A,250MG,N/A;N/A,N/A,500MG ** | N050719 001 | Aug 15, 1996 |
|---|-------------------|------------------------------------------------|-------------|--------------|

BISOPROLOL FUMARATE

TABLET; ORAL

BISOPROLOL FUMARATE

| | | | | |
|--|-----------------|------|-------------|--------------|
| | ANDA REPOSITORY | 5MG | A075474 001 | Oct 25, 2002 |
| | | 10MG | A075474 002 | Oct 25, 2002 |

ZEBETA

| | | | | |
|---|-------------|---------|-------------|--------------|
| + | TEVA WOMENS | 5MG ** | N019982 002 | Jul 31, 1992 |
| + | | 10MG ** | N019982 001 | Jul 31, 1992 |

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

| | | | | |
|--|----------------------|--------------|-------------|--------------|
| | ACTAVIS ELIZABETH | 2.5MG;6.25MG | A075672 001 | Sep 25, 2000 |
| | | 5MG;6.25MG | A075672 002 | Sep 25, 2000 |
| | | 10MG;6.25MG | A075672 003 | Sep 25, 2000 |
| | APOTHECON | 2.5MG;6.25MG | A075642 002 | Dec 27, 2000 |
| | | 5MG;6.25MG | A075642 001 | Dec 27, 2000 |
| | | 10MG;6.25MG | A075642 003 | Dec 27, 2000 |
| | IVAX SUB TEVA PHARMS | 2.5MG;6.25MG | A075632 001 | Sep 27, 2000 |
| | | 5MG;6.25MG | A075632 002 | Sep 27, 2000 |
| | | 10MG;6.25MG | A075632 003 | Sep 27, 2000 |
| | SANDOZ | 2.5MG;6.25MG | A075527 001 | Sep 25, 2000 |
| | | 5MG;6.25MG | A075527 003 | Sep 25, 2000 |
| | | 10MG;6.25MG | A075527 002 | Sep 25, 2000 |
| | TEVA | 2.5MG;6.25MG | A075686 001 | Jan 19, 2001 |
| | | 5MG;6.25MG | A075686 002 | Jan 19, 2001 |
| | | 10MG;6.25MG | A075686 003 | Jan 19, 2001 |
| | WATSON LABS TEVA | 2.5MG;6.25MG | A075469 001 | Sep 25, 2000 |
| | | 5MG;6.25MG | A075469 002 | Sep 25, 2000 |
| | | 10MG;6.25MG | A075469 003 | Sep 25, 2000 |

BITOLTEROL MESYLATE

AEROSOL, METERED; INHALATION

TORNALATE

| | | | | |
|--|-------------------|------------|-------------|--------------|
| | SANOFI AVENTIS US | 0.37MG/INH | N018770 001 | Dec 28, 1984 |
|--|-------------------|------------|-------------|--------------|

SOLUTION; INHALATION

TORNALATE

| | | | | |
|--|-------------------|------|-------------|--------------|
| | SANOFI AVENTIS US | 0.2% | N019548 001 | Feb 19, 1992 |
|--|-------------------|------|-------------|--------------|

BLEOMYCIN SULFATE

INJECTABLE; INJECTION

BLENOXANE

| | | | | |
|---|----------------------|--------------------------|-------------|--------------|
| + | BRISTOL MYERS SQUIBB | EQ 15 UNITS BASE/VIAL ** | N050443 001 | |
| + | | EQ 30 UNITS BASE/VIAL ** | N050443 002 | Sep 07, 1995 |

BLEOMYCIN SULFATE

| | | | | |
|--|-----------------|-----------------------|-------------|--------------|
| | PHARMACHEMIE BV | EQ 15 UNITS BASE/VIAL | A065201 001 | Dec 13, 2007 |
| | TEVA PARENTERAL | EQ 15 UNITS BASE/VIAL | A064084 001 | Jun 01, 1996 |
| | | EQ 30 UNITS BASE/VIAL | A064084 002 | Jun 01, 1996 |

BOCEPREVIR

CAPSULE; ORAL

VICTRELIS

| | | | | |
|--|-------------------|-------|-------------|--------------|
| | MERCK SHARP DOHME | 200MG | N202258 001 | May 13, 2011 |
|--|-------------------|-------|-------------|--------------|

BORTEZOMIB

POWDER; INTRAVENOUS, SUBCUTANEOUS

BORTEZOMIB

| | | | | |
|---|-------------|------------|-------------|--------------|
| + | HOSPIRA INC | 2.5MG/VIAL | N209191 001 | Jul 12, 2018 |
|---|-------------|------------|-------------|--------------|

BRETYLIUM TOSYLATE

INJECTABLE; INJECTION

BRETYLIUM TOSYLATE

| | | | | |
|--|---------------|----------|-------------|--------------|
| | ABRAXIS PHARM | 50MG/ML | A070134 001 | Apr 29, 1986 |
| | | 100MG/ML | A071298 001 | Feb 13, 1987 |
| | ASTRAZENECA | 50MG/ML | A071151 001 | Aug 10, 1987 |
| | | 50MG/ML | A071152 001 | Aug 10, 1987 |
| | | 50MG/ML | A071153 001 | Aug 10, 1987 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BRETYLIUM TOSYLATE

INJECTABLE; INJECTION

BRETYLIUM TOSYLATE

| | | | | |
|--------------------------------------------------------|-------------|---------|-----|--------------|
| EUROHLTH INTL SARL | 50MG/ML | A070546 | 001 | May 14, 1986 |
| + HOSPIRA | 50MG/ML ** | N019030 | 001 | Apr 29, 1986 |
| | 50MG/ML | N019033 | 001 | Apr 29, 1986 |
| INTL MEDICATION | 50MG/ML | A070119 | 001 | Apr 29, 1986 |
| LUITPOLD | 50MG/ML | A070891 | 001 | Jul 26, 1988 |
| WEST-WARD PHARMS INT | 50MG/ML | A070545 | 001 | May 14, 1986 |
| BRETYLIUM TOSYLATE IN DEXTROSE 5% | | | | |
| ABBOTT | 200MG/100ML | N019005 | 002 | Apr 29, 1986 |
| | 400MG/100ML | N019005 | 003 | Apr 29, 1986 |
| | 800MG/100ML | N019005 | 001 | Apr 29, 1986 |
| BRETYLIUM TOSYLATE IN DEXTROSE 5% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 100MG/100ML | N019121 | 001 | Apr 29, 1986 |
| | 200MG/100ML | N019121 | 002 | Apr 29, 1986 |
| | 400MG/100ML | N019121 | 003 | Apr 29, 1986 |
| BAXTER HLTHCARE | 200MG/100ML | N019837 | 002 | Apr 12, 1989 |
| | 400MG/100ML | N019837 | 001 | Apr 12, 1989 |
| HOSPIRA INC | 200MG/100ML | N019008 | 002 | Apr 29, 1986 |
| | 400MG/100ML | N019008 | 003 | Apr 29, 1986 |
| | 800MG/100ML | N019008 | 001 | Apr 29, 1986 |
| BRETYLOL | | | | |
| HOSPIRA | 50MG/ML | N017954 | 001 | |

BRIMONIDINE TARTRATE

SOLUTION/DROPS; OPHTHALMIC

ALPHAGAN

| | | | | |
|----------------------|---------|---------|-----|--------------|
| + ALLERGAN | 0.2% ** | N020613 | 001 | Sep 06, 1996 |
| | 0.5% | N020490 | 001 | Mar 13, 1997 |
| BRIMONIDINE TARTRATE | | | | |
| TEVA PARENTERAL | 0.2% | A076372 | 001 | Sep 10, 2004 |

BROMFENAC SODIUM

SOLUTION/DROPS; OPHTHALMIC

BROMDAY

| | | | | |
|-----------------------|------------------|---------|-----|--------------|
| + BAUSCH AND LOMB INC | EQ 0.09% ACID ** | N021664 | 002 | Oct 16, 2010 |
| BROMFENAC SODIUM | | | | |
| AMRING PHARMS | EQ 0.09% ACID | A202030 | 001 | Jan 09, 2013 |
| APOTEX INC | EQ 0.09% ACID | A202435 | 001 | Jun 19, 2014 |
| | EQ 0.09% ACID | A202620 | 001 | Jun 23, 2014 |
| COASTAL PHARMS | EQ 0.09% ACID | A201211 | 001 | May 11, 2011 |
| PADDOCK LLC | EQ 0.09% ACID | A201941 | 001 | Feb 10, 2015 |
| XIBROM | | | | |
| + BAUSCH AND LOMB INC | EQ 0.09% ACID ** | N021664 | 001 | Mar 24, 2005 |

BROMOCRIPTINE MESYLATE

CAPSULE; ORAL

BROMOCRIPTINE MESYLATE

| | | | | |
|-----------|-------------|---------|-----|--------------|
| LEK PHARM | EQ 5MG BASE | A075100 | 001 | Dec 10, 1998 |
|-----------|-------------|---------|-----|--------------|

BROMODIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

AMBODRYL

| | | | | |
|-------------|------|---------|-----|--|
| PARKE DAVIS | 25MG | N007984 | 001 | |
|-------------|------|---------|-----|--|

BROMODIPHENHYDRAMINE HYDROCHLORIDE; CODEINE PHOSPHATE

SYRUP; ORAL

AMBENYL

| | | | | |
|----------------------------------------------------------|----------------------|---------|-----|--------------|
| FOREST LABS | 12.5MG/5ML; 10MG/5ML | N009319 | 006 | Jan 10, 1984 |
| BROMANYL | | | | |
| ALPHARMA US PHARMS | 12.5MG/5ML; 10MG/5ML | A088343 | 001 | Aug 15, 1984 |
| BROMODIPHENHYDRAMINE HYDROCHLORIDE AND CODEINE PHOSPHATE | | | | |
| WOCKHARDT | 12.5MG/5ML; 10MG/5ML | A088626 | 001 | Oct 12, 1984 |

BROMPHENIRAMINE MALEATE

ELIXIR; ORAL

BROMPHENIRAMINE MALEATE

| | | | | |
|--------------------|---------|---------|-----|--------------|
| ALPHARMA US PHARMS | 2MG/5ML | A086936 | 001 | |
| KV PHARM | 2MG/5ML | A085466 | 001 | |
| PHARM ASSOC | 2MG/5ML | A087517 | 001 | |
| USL PHARMA | 2MG/5ML | A087964 | 001 | Jan 25, 1983 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BROMPHENIRAMINE MALEATE

INJECTABLE; INJECTION

BROMPHENIRAMINE MALEATE

WATSON LABS

10MG/ML

A083821 001

100MG/ML

A083820 001

DIMETANE-TEN

WYETH AYERST

10MG/ML

N011418 002

TABLET; ORAL

BROMPHENIRAMINE MALEATE

BARR

4MG

A084468 001

IVAX SUB TEVA PHARMS

4MG

A084351 001

NEWTRON PHARMS

4MG

A086987 001

NEXGEN PHARMA INC

4MG

A086187 001

PAR PHARM

4MG

A087009 001

PIONEER PHARMS

4MG

A088604 001 Jul 13, 1984

UPSHER SMITH LABS

4MG

A083215 001

VITARINE

4MG

A085850 001

WATSON LABS

4MG

A083123 001

4MG

A085769 001

DIMETANE

WYETH CONS

4MG

N010799 003

TABLET, EXTENDED RELEASE; ORAL

DIMETANE

WYETH CONS

8MG

N010799 010 Jun 10, 1983

12MG

N010799 011 Jun 10, 1983

BROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SYRUP; ORAL

BROMANATE DM

ALPHARMA US PHARMS

2MG/5ML; 10MG/5ML; 30MG/5ML

A088722 001 Mar 07, 1985

BROMFED-DM

WOCKHARDT

2MG/5ML; 10MG/5ML; 30MG/5ML

A089681 001 Dec 22, 1988

DIMETANE-DX

+ ROBINS AH

2MG/5ML; 10MG/5ML; 30MG/5ML **

N019279 001 Aug 24, 1984

BROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

EFIDAC 24 PSEUDOEPHEDRINE HYDROCHLORIDE/BROMPHENIRAMINE MALEATE

ALZA

16MG; 240MG

N019672 001 Mar 29, 1996

BUCLIZINE HYDROCHLORIDE

TABLET; ORAL

BUCLADIN-S

STUART PHARMS

50MG

N010911 006

BUDESONIDE

AEROSOL, METERED; NASAL

RHINOCORT

ASTRAZENECA

0.032MG/INH

N020233 001 Feb 14, 1994

POWDER, METERED; INHALATION

PULMICORT

ASTRAZENECA

0.16MG/INH

N020441 002 Jun 24, 1997

0.32MG/INH

N020441 003 Jun 24, 1997

BUMETANIDE

INJECTABLE; INJECTION

BUMETANIDE

ATHENEX INC

0.25MG/ML

A074441 001 Jan 27, 1995

HOSPIRA

0.25MG/ML

A074160 001 Oct 30, 1997

TEVA PARENTERAL

0.25MG/ML

A074613 001 Nov 18, 1997

BUMEX

+ VALIDUS PHARMS

0.25MG/ML **

N018226 001 Feb 28, 1983

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE

HOSPIRA

0.75%

A070587 001 Mar 03, 1987

BUPIVACAINE HYDROCHLORIDE KIT

HOSPIRA

0.075%

N019978 001 Sep 03, 1992

0.114%

N019978 002 Sep 03, 1992

0.23%

N019978 003 Sep 03, 1992

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE

| | | | |
|----------------|-------|-------------|--------------|
| INTL MEDICATED | 0.25% | A076012 001 | Jan 09, 2002 |
| | 0.5% | A076012 002 | Jan 09, 2002 |
| | 0.75% | A076012 003 | Jan 09, 2002 |

INJECTABLE; SPINAL

SENSORCAINE

| | | | |
|--------------------|-------|-------------|--------------|
| FRESENIUS KABI USA | 0.75% | A071202 001 | Apr 15, 1987 |
|--------------------|-------|-------------|--------------|

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

| | | | |
|---------|-------------------|-------------|--------------|
| HOSPIRA | 0.25%; 0.005MG/ML | A071166 001 | Jun 16, 1988 |
| | 0.5%; 0.005MG/ML | A071169 001 | Jun 16, 1988 |
| | 0.75%; 0.005MG/ML | A071171 001 | Jun 16, 1988 |

BUPIVACAINE HYDROCHLORIDE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DUOCAINE

| | | | |
|----------------------|---------------------------------------------|-------------|--------------|
| AMPHASTAR PHARMS INC | EQ 0.375% (37.5MG/10ML); EQ 1% (100MG/10ML) | N021496 001 | May 23, 2003 |
|----------------------|---------------------------------------------|-------------|--------------|

BUPRENORPHINE HYDROCHLORIDE

TABLET; SUBLINGUAL

SUBUTEX

| | | | | |
|---|--------------|----------------|-------------|--------------|
| + | INDIVIOR INC | EQ 2MG BASE ** | N020732 002 | Oct 08, 2002 |
| + | | EQ 8MG BASE ** | N020732 003 | Oct 08, 2002 |

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE

TABLET; SUBLINGUAL

SUBOXONE

| | | | | |
|---|--------------|-------------------------------|-------------|--------------|
| + | INDIVIOR INC | EQ 2MG BASE; EQ 0.5MG BASE ** | N020733 001 | Oct 08, 2002 |
| + | | EQ 8MG BASE; EQ 2MG BASE | N020733 002 | Oct 08, 2002 |

BUPROPION HYDROCHLORIDE

TABLET; ORAL

BUPROPION HYDROCHLORIDE

| | | | |
|--------|-------|-------------|--------------|
| SANDOZ | 75MG | A075613 002 | Oct 10, 2000 |
| | 100MG | A075613 001 | Oct 10, 2000 |
| TEVA | 75MG | A075310 001 | Nov 29, 1999 |
| | 100MG | A075310 002 | Nov 29, 1999 |

WELLBUTRIN

| | | | | |
|---|-----------------|----------|-------------|--------------|
| + | GLAXOSMITHKLINE | 50MG ** | N018644 001 | Dec 30, 1985 |
| + | | 75MG ** | N018644 002 | Dec 30, 1985 |
| + | | 100MG ** | N018644 003 | Dec 30, 1985 |

TABLET, EXTENDED RELEASE; ORAL

BUPROPION HYDROCHLORIDE

| | | | |
|---------------------|-------|-------------|--------------|
| ACTAVIS LABS FL INC | 300MG | A077715 002 | Jun 13, 2007 |
| IMPAX LABS | 300MG | A077415 002 | Dec 15, 2006 |
| SANDOZ | 100MG | A076845 001 | Jul 14, 2005 |
| | 150MG | A076834 001 | Jul 14, 2005 |
| | 150MG | A076845 002 | Jul 14, 2005 |
| WOCKHARDT LTD | 100MG | A201331 001 | Aug 30, 2012 |
| | 150MG | A201331 002 | Aug 30, 2012 |
| | 200MG | A201331 003 | Aug 30, 2012 |

WELLBUTRIN SR

| | | | |
|-----------------|------|-------------|--------------|
| GLAXOSMITHKLINE | 50MG | N020358 001 | Oct 04, 1996 |
|-----------------|------|-------------|--------------|

ZYBAN

| | | | |
|-----------------|-------|-------------|--------------|
| GLAXOSMITHKLINE | 100MG | N020711 002 | May 14, 1997 |
|-----------------|-------|-------------|--------------|

BUSPIRONE HYDROCHLORIDE

CAPSULE; ORAL

BUSPAR

| | | | |
|----------------------|-------|-------------|--------------|
| BRISTOL MYERS SQUIBB | 5MG | N021190 001 | Dec 20, 2000 |
| | 7.5MG | N021190 002 | Dec 20, 2000 |
| | 10MG | N021190 003 | Dec 20, 2000 |
| | 15MG | N021190 004 | Dec 20, 2000 |

TABLET; ORAL

BUSPAR

| | | | | |
|---|----------------------|---------|-------------|--------------|
| + | BRISTOL MYERS SQUIBB | 5MG ** | N018731 001 | Sep 29, 1986 |
| + | | 10MG ** | N018731 002 | Sep 29, 1986 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL

BUSPAR

| | | | |
|---|---------|-------------|--------------|
| + | 15MG ** | N018731 003 | Apr 22, 1996 |
| + | 30MG ** | N018731 004 | Apr 22, 1996 |

BUSPIRONE HYDROCHLORIDE

APOTEX

| | | |
|------|-------------|--------------|
| 5MG | A075521 001 | Apr 05, 2002 |
| 10MG | A075521 002 | Apr 05, 2002 |
| 15MG | A075521 003 | Apr 05, 2002 |

EGIS

| | | |
|------|-------------|--------------|
| 5MG | A075119 001 | Mar 14, 2002 |
| 10MG | A075119 002 | Mar 14, 2002 |
| 15MG | A075119 003 | Jan 23, 2003 |

FOSUN PHARMA

| | | |
|------|-------------|--------------|
| 5MG | A075413 001 | Mar 19, 2002 |
| 10MG | A075413 002 | Mar 19, 2002 |
| 15MG | A075413 003 | Mar 19, 2002 |

IVAX SUB TEVA PHARMS

| | | |
|---------|-------------|--------------|
| 5MG ** | A075385 001 | Mar 01, 2002 |
| 10MG ** | A075385 002 | Mar 01, 2002 |
| 15MG ** | A075385 003 | Mar 01, 2002 |

MYLAN

| | | |
|------|-------------|--------------|
| 5MG | A075467 001 | Feb 28, 2002 |
| 10MG | A075467 003 | Feb 28, 2002 |
| 15MG | A075467 004 | Feb 28, 2002 |

NESHER PHARMS

| | | |
|------|-------------|--------------|
| 5MG | A075572 001 | Feb 27, 2002 |
| 10MG | A075572 002 | Feb 27, 2002 |
| 15MG | A075572 003 | Feb 27, 2002 |

OXFORD PHARMS

| | | |
|------|-------------|--------------|
| 5MG | A075388 001 | May 09, 2002 |
| 10MG | A075388 002 | May 09, 2002 |
| 15MG | A075388 003 | May 09, 2002 |

BUTABARBITAL SODIUM

CAPSULE; ORAL

BUTICAPS

MEDPOINTE PHARM HLC

| | |
|-------|-------------|
| 15MG | A085381 001 |
| 30MG | A085381 002 |
| 50MG | A085381 003 |
| 100MG | A085381 004 |

ELIXIR; ORAL

BUTABARB

ALPHARMA US PHARMS

| | |
|----------|-------------|
| 30MG/5ML | A085873 001 |
|----------|-------------|

BUTABARBITAL SODIUM

WOCKHARDT

| | |
|----------|-------------|
| 30MG/5ML | A085383 001 |
|----------|-------------|

BUTALAN

LANNETT

| | |
|------------|-------------|
| 33.3MG/5ML | A085880 001 |
|------------|-------------|

BUTISOL SODIUM

MEDA PHARMS

| | |
|----------|-------------|
| 30MG/5ML | A085380 001 |
|----------|-------------|

SARISOL

HALSEY

| | |
|----------|-------------|
| 30MG/5ML | A084723 001 |
|----------|-------------|

TABLET; ORAL

BUTABARBITAL

BUNDY

| | |
|------|-------------|
| 30MG | A085550 001 |
|------|-------------|

BUTABARBITAL SODIUM

SANDOZ

| | | |
|--------|-------------|--------------|
| 15MG | A084292 003 | Feb 09, 1982 |
| 15MG | A085938 001 | |
| 30MG | A084272 002 | |
| 30MG | A085934 001 | |
| SOLVAY | 16.2MG | A083606 001 |
| | 32.4MG | A083898 001 |
| | 48.6MG | A083897 001 |
| | 97.2MG | A083896 001 |

TEVA

| | | |
|------|-------------|--------------|
| 15MG | A088632 001 | May 18, 1985 |
| 30MG | A088631 001 | May 01, 1985 |

WATSON LABS

| | |
|------|-------------|
| 15MG | A085764 001 |
| 30MG | A085772 001 |

WHITEWORTH TOWN PLSN

| | |
|------|-------------|
| 15MG | A083325 002 |
| 30MG | A083337 001 |

BUTISOL SODIUM

MYLAN SPECIALITY LP

| | |
|-------|-------------|
| 15MG | N000793 002 |
| 50MG | N000793 003 |
| 100MG | N000793 005 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

BUTABARBITAL SODIUM

TABLET; ORAL

| | | | | |
|----------------------|--------|--|---------|-----|
| SARISOL NO. 1 | | | | |
| HALSEY | 15MG | | A084719 | 001 |
| SARISOL NO. 2 | | | | |
| HALSEY | 30MG | | A084719 | 002 |
| SODIUM BUTABARBITAL | | | | |
| HIKMA PHARMS | 15MG | | A085418 | 001 |
| | 30MG | | A085432 | 001 |
| IVAX SUB TEVA PHARMS | 15MG | | A083484 | 001 |
| | 30MG | | A084040 | 001 |
| LANNETT | 15MG | | A085849 | 001 |
| | 30MG | | A085866 | 001 |
| | 100MG | | A085881 | 001 |
| MARSHALL PHARMA | 16.2MG | | A083524 | 001 |
| | 32.4MG | | A083858 | 001 |

BUTENAFINE HYDROCHLORIDE

CREAM; TOPICAL

MENTAX-TC

| | | | | | |
|-------|----|--|---------|-----|--------------|
| MYLAN | 1% | | N021408 | 001 | Oct 17, 2002 |
|-------|----|--|---------|-----|--------------|

BUTOCONAZOLE NITRATE

CREAM; VAGINAL

BUTOCONAZOLE NITRATE

| | | | | | |
|---------------------|----|--|---------|-----|--------------|
| PERRIGO PHARMA INTL | 2% | | N019881 | 001 | Feb 07, 1997 |
|---------------------|----|--|---------|-----|--------------|

FEMSTAT

| | | | | | |
|------------|----|--|---------|-----|--------------|
| ROCHE PALO | 2% | | N019215 | 001 | Nov 25, 1985 |
|------------|----|--|---------|-----|--------------|

FEMSTAT 3

| | | | | | |
|---------|----|--|---------|-----|--------------|
| + BAYER | 2% | | N020421 | 001 | Dec 21, 1995 |
|---------|----|--|---------|-----|--------------|

SUPPOSITORY; VAGINAL

FEMSTAT

| | | | | | |
|------------|-------|--|---------|-----|--------------|
| ROCHE PALO | 100MG | | N019359 | 001 | Nov 25, 1985 |
|------------|-------|--|---------|-----|--------------|

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

BUTORPHANOL TARTRATE

| | | | | | |
|----------------------|--------|--|---------|-----|--------------|
| BAXTER HLTHCARE CORP | 2MG/ML | | A075697 | 001 | Oct 23, 2001 |
|----------------------|--------|--|---------|-----|--------------|

| | | | | | |
|--------------------|--------|--|---------|-----|--------------|
| HIKMA FARMACEUTICA | 2MG/ML | | A078247 | 001 | Apr 29, 2009 |
|--------------------|--------|--|---------|-----|--------------|

| | | | | | |
|---------|--------|--|---------|-----|--------------|
| HOSPIRA | 1MG/ML | | A075342 | 001 | Nov 04, 1999 |
|---------|--------|--|---------|-----|--------------|

| | | | | | |
|--|--------|--|---------|-----|--------------|
| | 1MG/ML | | A075559 | 001 | Mar 20, 2000 |
|--|--------|--|---------|-----|--------------|

| | | | | | |
|--|--------|--|---------|-----|--------------|
| | 2MG/ML | | A075342 | 002 | Nov 04, 1999 |
|--|--------|--|---------|-----|--------------|

| | | | | | |
|--|--------|--|---------|-----|--------------|
| | 2MG/ML | | A075559 | 002 | Mar 20, 2000 |
|--|--------|--|---------|-----|--------------|

BUTORPHANOL TARTRATE PRESERVATIVE FREE

| | | | | | |
|----------------------|--------|--|---------|-----|--------------|
| BAXTER HLTHCARE CORP | 1MG/ML | | A075695 | 001 | Oct 23, 2001 |
|----------------------|--------|--|---------|-----|--------------|

| | | | | | |
|--|--------|--|---------|-----|--------------|
| | 2MG/ML | | A075695 | 002 | Oct 23, 2001 |
|--|--------|--|---------|-----|--------------|

| | | | | | |
|---------|--------|--|---------|-----|--------------|
| HOSPIRA | 1MG/ML | | A074620 | 001 | Jan 22, 1997 |
|---------|--------|--|---------|-----|--------------|

| | | | | | |
|--|--------|--|---------|-----|--------------|
| | 1MG/ML | | A075170 | 001 | Sep 28, 1998 |
|--|--------|--|---------|-----|--------------|

| | | | | | |
|--|--------|--|---------|-----|--------------|
| | 2MG/ML | | A074620 | 002 | Jan 22, 1997 |
|--|--------|--|---------|-----|--------------|

| | | | | | |
|--|--------|--|---------|-----|--------------|
| | 2MG/ML | | A075170 | 002 | Sep 28, 1998 |
|--|--------|--|---------|-----|--------------|

STADOL

| | | | | | |
|-------------|--------|----|---------|-----|--|
| + APOTHECON | 2MG/ML | ** | N017857 | 004 | |
|-------------|--------|----|---------|-----|--|

STADOL PRESERVATIVE FREE

| | | | | | |
|-------------|--------|----|---------|-----|--|
| + APOTHECON | 1MG/ML | ** | N017857 | 001 | |
|-------------|--------|----|---------|-----|--|

| | | | | | |
|-------------|--------|----|---------|-----|--|
| + APOTHECON | 2MG/ML | ** | N017857 | 002 | |
|-------------|--------|----|---------|-----|--|

SPRAY, METERED; NASAL

STADOL

| | | | | | |
|----------------------|-----------|----|---------|-----|--------------|
| BRISTOL MYERS SQUIBB | 1MG/SPRAY | ** | N019890 | 001 | Dec 12, 1991 |
|----------------------|-----------|----|---------|-----|--------------|

CABERGOLINE

TABLET; ORAL

CABERGOLINE

| | | | | | |
|-------------|-------|--|---------|-----|--------------|
| APOTEX CORP | 0.5MG | | A201503 | 001 | Mar 08, 2013 |
|-------------|-------|--|---------|-----|--------------|

| | | | | | |
|----------------|-------|--|---------|-----|--------------|
| IMPAX LABS INC | 0.5MG | | A077843 | 001 | Jul 03, 2007 |
|----------------|-------|--|---------|-----|--------------|

DOSTINEX

| | | | | | |
|------------------------|-------|----|---------|-----|--------------|
| + PHARMACIA AND UPJOHN | 0.5MG | ** | N020664 | 001 | Dec 23, 1996 |
|------------------------|-------|----|---------|-----|--------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CAFFEINE; ERGOTAMINE TARTRATE

SUPPOSITORY;RECTAL

CAFERGOT

+ NOVARTIS

100MG;2MG **

N009000 002

TABLET;ORAL

CAFERGOT

NOVARTIS

100MG;1MG

N006620 001

WIGRAINE

ORGANON USA INC

100MG;1MG

A086562 001

CALCIFEDIOL

CAPSULE;ORAL

CALDEROL

ORGANON USA INC

0.02MG

N018312 001

0.05MG

N018312 002

CALCIPOTRIENE

OINTMENT;TOPICAL

DOVONEX

+ LEO PHARMA AS

0.005% **

N020273 001 Dec 29, 1993

SOLUTION;TOPICAL

DOVONEX

+ LEO PHARM

0.005% **

N020611 001 Mar 03, 1997

CALCITONIN HUMAN

INJECTABLE;INJECTION

CIBACALCIN

NOVARTIS

0.5MG/VIAL

N018470 001 Oct 31, 1986

CALCITONIN SALMON

INJECTABLE;INJECTION

CALCIMAR

SANOFI AVENTIS US

200 IU/ML

N017769 001

400 IU/VIAL

N017497 001

CALCITONIN-SALMON

IGI LABS INC

200 IU/ML

A073690 001 Apr 14, 1995

MIACALCIN

MYLAN IRELAND LTD

100 IU/ML

N017808 001 Jul 03, 1986

SPRAY, METERED;NASAL

MIACALCIN

+ MYLAN IRELAND LTD

200 IU/SPRAY

N020313 002 Aug 17, 1995

CALCITONIN SALMON RECOMBINANT

SPRAY, METERED;NASAL

FORTICAL

UPSHER SMITH LABS

200 IU/SPRAY **

N021406 001 Aug 12, 2005

CALCITRIOL

INJECTABLE;INJECTION

CALCIJEX

+ ABBVIE

0.001MG/ML **

N018874 001 Sep 25, 1986

+

0.002MG/ML **

N018874 002 Sep 25, 1986

CALCITRIOL

AKORN

0.002MG/ML

A078066 002 Jan 29, 2008

FRESENIUS KABI USA

0.001MG/ML

A075836 001 Dec 31, 2002

0.002MG/ML

A075836 002 Dec 31, 2002

FRESENIUS MEDCL

0.001MG/ML

A075766 001 Feb 20, 2003

0.002MG/ML

A075766 002 Feb 20, 2003

HOSPIRA

0.001MG/ML

A075816 001 Jan 16, 2004

0.002MG/ML

A075816 002 Jan 16, 2004

LUITPOLD

0.001MG/ML

A075746 001 Sep 26, 2003

0.002MG/ML

A075746 002 Sep 26, 2003

ROCKWELL MEDCL

0.001MG/ML

A076206 001 Sep 17, 2003

SAGENT PHARMS

0.001MG/ML

A077102 001 Feb 08, 2006

TEVA PARENTERAL

0.001MG/ML

A075823 001 Mar 31, 2003

0.002MG/ML

A075823 002 Mar 31, 2003

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM ACETATE

CAPSULE; ORAL

PHOSLO

| | | | |
|-----------------|---------|-------------|--------------|
| FRESENIUS MEDCL | 333.5MG | N021160 001 | Apr 02, 2001 |
| | 667MG | N021160 002 | Apr 02, 2001 |

TABLET; ORAL

CALCIUM ACETATE

| | | | |
|----------------------|-------|-------------|--------------|
| WEST-WARD PHARMS INT | 667MG | A077693 001 | Jan 30, 2008 |
|----------------------|-------|-------------|--------------|

ELIPHOS

| | | | |
|---------------|-------|-------------|--------------|
| CYPRESS PHARM | 667MG | A078502 001 | Nov 25, 2008 |
|---------------|-------|-------------|--------------|

PHOSLO

| | | | |
|-------------------|----------|-------------|--------------|
| + FRESENIUS MEDCL | 667MG ** | N019976 001 | Dec 10, 1990 |
|-------------------|----------|-------------|--------------|

CALCIUM CARBONATE; RISEDRONATE SODIUM

TABLET, TABLET; ORAL

ACTONEL WITH CALCIUM (COPACKAGED)

| | | | |
|-------------------|----------------------------------|-------------|--------------|
| + WARNER CHILCOTT | EQ 500MG BASE, N/A; N/A, 35MG ** | N021823 001 | Aug 12, 2005 |
|-------------------|----------------------------------|-------------|--------------|

CALCIUM CHLORIDE; DEXTROSE; GLUTATHIONE DISULFIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

METHOTREXATE SODIUM

| | | | |
|-------|----------------------------------------------------------------------------------------|-------------|--------------|
| AKORN | 0.154MG/ML; 0.92MG/ML; 0.184MG/ML; 0.2MG/ML; 0.38MG/ML; 2.1MG/ML; 7.14MG/ML; 0.42MG/ML | N020079 001 | Feb 26, 1999 |
|-------|----------------------------------------------------------------------------------------|-------------|--------------|

CALCIUM CHLORIDE; DEXTROSE; LACTIC ACID; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PRISMASOL B22GK 2/0 IN PLASTIC CONTAINER

| | | | |
|------------------------|-------------------------------------------------------------------------------------------------------------|-------------|--------------|
| + BAXTER HLTHCARE CORP | N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.157GM/1000ML; 2.21GM/1000ML; 7.07GM/1000ML (5000ML) | N021703 010 | Oct 10, 2008 |
|------------------------|-------------------------------------------------------------------------------------------------------------|-------------|--------------|

PRISMASOL B22GK 2/2.5 IN PLASTIC CONTAINER

| | | | |
|----------------------|----------------------------------------------------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE CORP | 3.68GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.157GM/1000ML; 2.21GM/1000ML; 7.07GM/1000ML (5000ML) | N021703 012 | Oct 10, 2008 |
|----------------------|----------------------------------------------------------------------------------------------------------------|-------------|--------------|

PRISMASOL B22GK 4/2.5 IN PLASTIC CONTAINER

| | | | |
|------------------------|----------------------------------------------------------------------------------------------------------------|-------------|--------------|
| + BAXTER HLTHCARE CORP | 3.68GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 2.21GM/1000ML; 7.07GM/1000ML (5000ML) | N021703 013 | Oct 10, 2008 |
|------------------------|----------------------------------------------------------------------------------------------------------------|-------------|--------------|

PRISMASOL BGK 4/0 IN PLASTIC CONTAINER

| | | | |
|----------------------|-------------------------------------------------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE CORP | N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML) | N021703 005 | Oct 25, 2006 |
|----------------------|-------------------------------------------------------------------------------------------------------------|-------------|--------------|

PRISMASOL BGK 4/3.5 IN PLASTIC CONTAINER

| | | | |
|----------------------|----------------------------------------------------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE CORP | 5.15GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 2.03GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML) | N021703 008 | Oct 25, 2006 |
|----------------------|----------------------------------------------------------------------------------------------------------------|-------------|--------------|

PRISMASOL BK 0/0 IN PLASTIC CONTAINER

| | | | |
|----------------------|--------------------------------------------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE CORP | N/A/1000ML; N/A/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; N/A/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML) | N021703 007 | Oct 25, 2006 |
|----------------------|--------------------------------------------------------------------------------------------------------|-------------|--------------|

PRISMASOL BK 0/3.5 IN PLASTIC CONTAINER

| | | | |
|------------------------|-----------------------------------------------------------------------------------------------------------|-------------|--------------|
| + BAXTER HLTHCARE CORP | 5.15GM/1000ML; N/A/1000ML; 5.4GM/1000ML; 2.03GM/1000ML; N/A/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML) | N021703 001 | Oct 25, 2006 |
|------------------------|-----------------------------------------------------------------------------------------------------------|-------------|--------------|

PRISMASOL BK 4/2.5 IN PLASTIC CONTAINER

| | | | |
|----------------------|---------------------------------------------------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE CORP | 3.68GM/1000ML; N/A/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML; 6.46GM/1000ML (5000ML) | N021703 009 | Oct 25, 2006 |
|----------------------|---------------------------------------------------------------------------------------------------------------|-------------|--------------|

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; OXIGLUTATHIONE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

NAVSTEL

| | | | |
|------------------|----------------------------------------------------------------------------------------|-------------|--------------|
| ALCON PHARMS LTD | 0.154MG/ML; 0.92MG/ML; 0.2MG/ML; 0.184MG/ML; 0.38MG/ML; 2.1MG/ML; 7.14MG/ML; 0.42MG/ML | N022193 001 | Jul 24, 2008 |
|------------------|----------------------------------------------------------------------------------------|-------------|--------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | | |
|-----------------------------------------------|-------------------------------------------------------------------------|-------------|--------------|
| ISOLYTE R IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 37MG/100ML; 5GM/100ML; 31MG/100ML; 120MG/100ML; 330MG/100ML; 88MG/100ML | N019864 001 | Jun 10, 1993 |
| ISOLYTE R W/ DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 37MG/100ML; 5GM/100ML; 31MG/100ML; 120MG/100ML; 330MG/100ML; 88MG/100ML | N018271 001 | |

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

INJECTABLE; INJECTION

| | | | |
|-----------------------------------------------|-------------------------------------------------------------------------------------|-------------|--------------|
| ISOLYTE E IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 35MG/100ML; 5GM/100ML; 30MG/100ML; 74MG/100ML; 640MG/100ML; 500MG/100ML; 74MG/100ML | N019867 001 | Dec 20, 1993 |
| ISOLYTE E W/ DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 35MG/100ML; 5GM/100ML; 30MG/100ML; 74MG/100ML; 640MG/100ML; 500MG/100ML; 74MG/100ML | N018269 002 | Jan 17, 1983 |

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

| | | | |
|----------------------------------------------------|--------------------------------------------------------------------------------------|-------------|--|
| PLASMA-LYTE M AND DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| + BAXTER HLTHCARE | 37MG/100ML; 5GM/100ML; 30MG/100ML; 119MG/100ML; 161MG/100ML; 94MG/100ML; 138MG/100ML | N017390 001 | |

**

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

SOLUTION; INTRAPERITONEAL

| | | | |
|----------------------------------------------------------|----------------------------------------------------------------|-------------|--------------|
| DIALYTE CONCENTRATE W/ DEXTROSE 30% IN PLASTIC CONTAINER | | | |
| B BRAUN | 510MG/100ML; 30GM/100ML; 200MG/100ML; 9.2GM/100ML; 9.6GM/100ML | N018807 001 | Aug 26, 1983 |
| | 510MG/100ML; 30GM/100ML; 200MG/100ML; 9.4GM/100ML; 11GM/100ML | N018807 003 | Aug 26, 1983 |
| DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER | | | |
| B BRAUN | 510MG/100ML; 50GM/100ML; 200MG/100ML; 9.2GM/100ML; 9.6GM/100ML | N018807 002 | Aug 26, 1983 |
| | 510MG/100ML; 50GM/100ML; 200MG/100ML; 9.4GM/100ML; 11GM/100ML | N018807 004 | Aug 26, 1983 |
| DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 29MG/100ML; 2.5GM/100ML; 15MG/100ML; 610MG/100ML; 560MG/100ML | N018460 006 | Jan 29, 1986 |
| DIALYTE W/ DEXTROSE 1.5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 29MG/100ML; 1.5GM/100ML; 15MG/100ML; 610MG/100ML; 560MG/100ML | N018460 001 | |
| DIALYTE W/ DEXTROSE 4.25% IN PLASTIC CONTAINER | | | |
| B BRAUN | 29MG/100ML; 4.25GM/100ML; 15MG/100ML; 610MG/100ML; 560MG/100ML | N018460 003 | |

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

| | | | |
|-------------------------------------------------|--------------------------------------------------------------------|-------------|--------------|
| DELFLX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER | | | |
| FRESENIUS MEDCL | 25.7MG/100ML; 1.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML | N018379 002 | |
| DELFLX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER | | | |
| FRESENIUS MEDCL | 25.7MG/100ML; 2.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML | N018379 003 | |
| DELFLX W/ DEXTROSE 3.5% IN PLASTIC CONTAINER | | | |
| FRESENIUS MEDCL | 25.7MG/100ML; 3.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML | N018379 007 | Jun 24, 1988 |
| DELFLX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER | | | |
| FRESENIUS MEDCL | 25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML | N018379 001 | |
| DELFLX-LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER | | | |
| FRESENIUS MEDCL | 25.7MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML | N018379 004 | Jul 07, 1982 |
| DELFLX-LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER | | | |
| FRESENIUS MEDCL | 25.7MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML | N018379 005 | Jul 07, 1982 |
| DELFLX-LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER | | | |
| FRESENIUS MEDCL | 25.7MG/100ML; 3.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML | N018379 008 | Jun 24, 1988 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

| | | | |
|-------------------------------------------------------|--------------------------------------------------------------------|-------------|--------------|
| DELFLEX-LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER | | | |
| FRESENIUS MEDCL | 25.7MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML | N018379 006 | Jul 07, 1982 |
| DIALYTE LM/ DEXTROSE 1.5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 26MG/100ML; 1.5GM/100ML; 5MG/100ML; 530MG/100ML; 450MG/100ML | N018460 007 | Jan 29, 1986 |
| | 26MG/100ML; 1.5GM/100ML; 15MG/100ML; 560MG/100ML; 390MG/100ML | N018460 002 | |
| DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 26MG/100ML; 2.5GM/100ML; 5MG/100ML; 530MG/100ML; 450MG/100ML | N018460 005 | Nov 02, 1983 |
| | 26MG/100ML; 5GM/100ML; 5MG/100ML; 530MG/100ML; 450MG/100ML | N018460 008 | Jan 29, 1986 |
| DIALYTE LM/ DEXTROSE 4.25% IN PLASTIC CONTAINER | | | |
| B BRAUN | 26MG/100ML; 4.25GM/100ML; 5MG/100ML; 530MG/100ML; 450MG/100ML | N018460 009 | Jan 29, 1986 |
| | 26MG/100ML; 4.25GM/100ML; 15MG/100ML; 560MG/100ML; 390MG/100ML | N018460 004 | |
| DIANEAL 137 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 25.7MG/100ML; 1.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML | N017512 001 | |
| DIANEAL 137 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 25.7MG/100ML; 2.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML | N017512 003 | |
| DIANEAL 137 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML | N017512 002 | |
| DIANEAL PD-1 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 25.7MG/100ML; 1.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML | N017512 007 | Jul 09, 1984 |
| DIANEAL PD-1 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 25.7MG/100ML; 2.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML | N017512 008 | Jul 09, 1984 |
| DIANEAL PD-1 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 25.7MG/100ML; 3.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML | N017512 010 | Nov 18, 1985 |
| DIANEAL PD-1 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML | N017512 009 | Jul 09, 1984 |
| DIANEAL PD-2 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 25.7MG/100ML; 3.5GM/100ML; 5.08MG/100ML; 38MG/100ML; 448MG/100ML | N017512 011 | Nov 18, 1985 |
| INPERSOL-LC/LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER | | | |
| FRESENIUS | 18.4MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 38MG/100ML; 448MG/100ML | A020374 001 | Jun 13, 1994 |
| INPERSOL-LC/LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER | | | |
| FRESENIUS | 18.4MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 38MG/100ML; 448MG/100ML | A020374 002 | Jun 13, 1994 |
| INPERSOL-LC/LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER | | | |
| FRESENIUS | 18.4MG/100ML; 3.5GM/100ML; 5.08MG/100ML; 38MG/100ML; 448MG/100ML | A020374 003 | Jun 13, 1994 |
| INPERSOL-LC/LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER | | | |
| FRESENIUS | 18.4MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML | A020374 004 | Jun 13, 1994 |

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | | |
|-------------------------------------------------------|-------------------------------------------------------------|-------------|--|
| DEXTROSE 5% IN ACETATED RINGER'S IN PLASTIC CONTAINER | | | |
| B BRAUN | 20MG/100ML; 5GM/100ML; 30MG/100ML; 380MG/100ML; 600MG/100ML | N018258 001 | |

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | | |
|-----------------------------------------------|------------------------------------------------|-------------|--------------|
| DEXTROSE 5% AND RINGER'S IN PLASTIC CONTAINER | | | |
| HOSPIRA | 33MG/100ML; 5GM/100ML; 30MG/100ML; 860MG/100ML | N018254 001 | |
| DEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER | | | |
| B BRAUN | 33MG/100ML; 5GM/100ML; 30MG/100ML; 860MG/100ML | N018256 001 | |
| | 33MG/100ML; 5GM/100ML; 30MG/100ML; 860MG/100ML | N020000 001 | Apr 17, 1992 |
| BAXTER HLTHCARE | 33MG/100ML; 5GM/100ML; 30MG/100ML; 860MG/100ML | N016695 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 4% IN MODIFIED LACTATED RINGER'S IN PLASTIC CONTAINER

| | | | |
|---------|----------------------------------------------------------|-------------|--------------|
| B BRAUN | 4MG/100ML; 4GM/100ML; 6MG/100ML; 120MG/100ML; 62MG/100ML | N019634 002 | Feb 24, 1988 |
|---------|----------------------------------------------------------|-------------|--------------|

DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER

| | | | |
|---------|-------------------------------------------------------------|-------------|--|
| B BRAUN | 20MG/100ML; 5GM/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML | N017510 001 | |
|---------|-------------------------------------------------------------|-------------|--|

| | | | |
|-------|-------------------------------------------------------------|-------------|--|
| MILES | 20MG/100ML; 5GM/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML | N018499 001 | |
|-------|-------------------------------------------------------------|-------------|--|

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

| | | | |
|-------------------|--------------------------------------------------------------|-------------|--------------|
| + ICU MEDICAL INC | 20MG/100ML; 5GM/100ML; 104MG/100ML; 600MG/100ML; 310MG/100ML | N019685 005 | Oct 17, 1988 |
|-------------------|--------------------------------------------------------------|-------------|--------------|

| | | | |
|-------------------|--------------------------------------------------------------|-------------|--------------|
| + ICU MEDICAL INC | 20MG/100ML; 5GM/100ML; 179MG/100ML; 600MG/100ML; 310MG/100ML | N019685 006 | Oct 17, 1988 |
|-------------------|--------------------------------------------------------------|-------------|--------------|

POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

| | | | |
|-------------------|--------------------------------------------------------------|-------------|--------------|
| + ICU MEDICAL INC | 20MG/100ML; 5GM/100ML; 254MG/100ML; 600MG/100ML; 310MG/100ML | N019685 007 | Oct 17, 1988 |
|-------------------|--------------------------------------------------------------|-------------|--------------|

POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

| | | | |
|-------------------|--------------------------------------------------------------|-------------|--------------|
| + ICU MEDICAL INC | 20MG/100ML; 5GM/100ML; 328MG/100ML; 600MG/100ML; 310MG/100ML | N019685 008 | Oct 17, 1988 |
|-------------------|--------------------------------------------------------------|-------------|--------------|

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

| | | | |
|-------------------|--------------------------------------------------------------|-------------|--------------|
| + ICU MEDICAL INC | 20MG/100ML; 5GM/100ML; 254MG/100ML; 600MG/100ML; 310MG/100ML | N019685 003 | Oct 17, 1988 |
|-------------------|--------------------------------------------------------------|-------------|--------------|

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

| | | | |
|-------------------|--------------------------------------------------------------|-------------|--------------|
| + ICU MEDICAL INC | 20MG/100ML; 5GM/100ML; 328MG/100ML; 600MG/100ML; 310MG/100ML | N019685 004 | Oct 17, 1988 |
|-------------------|--------------------------------------------------------------|-------------|--------------|

POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

| | | | |
|-------------------|--------------------------------------------------------------|-------------|--------------|
| + ICU MEDICAL INC | 20MG/100ML; 5GM/100ML; 104MG/100ML; 600MG/100ML; 310MG/100ML | N019685 001 | Oct 17, 1988 |
|-------------------|--------------------------------------------------------------|-------------|--------------|

CALCIUM CHLORIDE; DEXTROSE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

INPERSOL-ZM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

| | | | |
|-----------------|-----------------------------------------------------|-------------|--------------|
| FRESENIUS MEDCL | 25.7MG/100ML; 1.5GM/100ML; 538MG/100ML; 448MG/100ML | N019395 001 | Mar 26, 1986 |
|-----------------|-----------------------------------------------------|-------------|--------------|

INPERSOL-ZM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

| | | | |
|-----------------|-----------------------------------------------------|-------------|--------------|
| FRESENIUS MEDCL | 25.7MG/100ML; 2.5GM/100ML; 538MG/100ML; 448MG/100ML | N019395 002 | Mar 26, 1986 |
|-----------------|-----------------------------------------------------|-------------|--------------|

INPERSOL-ZM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

| | | | |
|-----------------|------------------------------------------------------|-------------|--------------|
| FRESENIUS MEDCL | 25.7MG/100ML; 4.25GM/100ML; 538MG/100ML; 448MG/100ML | N019395 003 | Mar 26, 1986 |
|-----------------|------------------------------------------------------|-------------|--------------|

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TPN ELECTROLYTES IN PLASTIC CONTAINER

| | | | |
|--------|------------------------------------------------------|-------------|--------------|
| ABBOTT | 16.5MG/ML; 25.4MG/ML; 74.6MG/ML; 121MG/ML; 16.1MG/ML | N019399 001 | Jun 16, 1986 |
|--------|------------------------------------------------------|-------------|--------------|

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

INJECTABLE; INJECTION

ISOLYTE E IN PLASTIC CONTAINER

| | | | |
|---------|--------------------------------------------------------------------------|-------------|--------------|
| B BRAUN | 35MG/100ML; 30MG/100ML; 74MG/100ML; 640MG/100ML; 500MG/100ML; 74MG/100ML | N018899 001 | Oct 31, 1983 |
|---------|--------------------------------------------------------------------------|-------------|--------------|

| | | | |
|---------|--------------------------------------------------------------------------|-------------|--------------|
| B BRAUN | 35MG/100ML; 30MG/100ML; 74MG/100ML; 640MG/100ML; 500MG/100ML; 74MG/100ML | N019718 001 | Sep 29, 1989 |
|---------|--------------------------------------------------------------------------|-------------|--------------|

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

PLASMA-LYTE R IN PLASTIC CONTAINER

| | | | |
|-----------------|----------------------------------------------------------------------------------|-------------|--|
| BAXTER HLTHCARE | 36.8MG/100ML; 30.5MG/100ML; 74.6MG/100ML; 640MG/100ML; 496MG/100ML; 89.6MG/100ML | N017438 001 | |
|-----------------|----------------------------------------------------------------------------------|-------------|--|

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ACETATED RINGER'S IN PLASTIC CONTAINER

| | | | |
|---------|--------------------------------------------------|-------------|--------------|
| B BRAUN | 20MG/100ML; 30MG/100ML; 380MG/100ML; 600MG/100ML | N018725 001 | Nov 29, 1982 |
|---------|--------------------------------------------------|-------------|--------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

RINGER'S IN PLASTIC CONTAINER

B BRAUN 33MG/100ML; 30MG/100ML; 860MG/100ML N018721 001 Nov 09, 1982

SOLUTION; IRRIGATION

RINGER'S IN PLASTIC CONTAINER

ABBOTT 33MG/100ML; 30MG/100ML; 860MG/100ML N018462 001

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

LACTATED RINGER'S IN PLASTIC CONTAINER

ABBOTT 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG /100ML N019485 001 Oct 24, 1985

B BRAUN 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG /100ML N018023 001

MILES 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG /100ML N018417 001

SOLUTION; IRRIGATION

LACTATED RINGER'S IN PLASTIC CONTAINER

BAXTER HLTHCARE 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG /100ML N019933 001 Aug 29, 1989

CALCIUM GLUCEPTATE

INJECTABLE; INJECTION

CALCIUM GLUCEPTATE

ABBOTT EQ 90MG CALCIUM/5ML A080001 001
EQ 90MG CALCIUM/5ML A083159 001

ABRAXIS PHARM EQ 90MG CALCIUM/5ML A089373 001 Apr 30, 1987

LILLY EQ 90MG CALCIUM/5ML N006470 001

CALCIUM METRIZOATE; MEGLUMINE METRIZOATE; METRIZOATE MAGNESIUM; METRIZOATE SODIUM

INJECTABLE; INJECTION

ISOPAQUE 440

GE HEALTHCARE 0.78MG/ML; 75.9MG/ML; 0.15MG/ML; 16.6MG/ML N016847 001

CALCIUM; MEGLUMINE; METRIZOIC ACID

INJECTABLE; INJECTION

ISOPAQUE 280

GE HEALTHCARE 0.35MG/ML; 140.1MG/ML; 461.8MG/ML N017506 001

CANDESARTAN CILEXETIL

TABLET; ORAL

CANDESARTAN CILEXETIL

APOTEX INC 4MG A202079 001 Jan 10, 2014

8MG A202079 002 Jan 10, 2014

16MG A202079 003 Jan 10, 2014

32MG A202079 004 Jan 10, 2014

CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE

APOTEX INC 16MG; 12.5MG A202884 001 Dec 04, 2012

32MG; 12.5MG A202884 002 Dec 04, 2012

32MG; 25MG A202884 003 Jun 03, 2013

CANDICIDIN

OINTMENT; VAGINAL

VANOVID

SANOFI AVENTIS US 0.6MG/GM A061596 001

TABLET; VAGINAL

VANOVID

SANOFI AVENTIS US 3MG A061613 001

CAPTOPRIL

TABLET; ORAL

CAPOTEN

+ PAR PHARM 12.5MG ** N018343 005 Jan 17, 1985

+ 25MG ** N018343 002

+ 37.5MG ** N018343 006 Sep 17, 1986

+ 50MG ** N018343 001

+ 75MG ** N018343 007 Jun 13, 1995

+ 100MG ** N018343 003

+ 150MG ** N018343 004 Jun 13, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

| | | | |
|------------------|--------|-------------|--------------|
| APOTEX | 12.5MG | A074737 001 | Oct 28, 1998 |
| | 25MG | A074737 002 | Oct 28, 1998 |
| | 50MG | A074737 003 | Oct 28, 1998 |
| | 100MG | A074737 004 | Oct 28, 1998 |
| APOTHECON | 12.5MG | A074472 001 | Mar 31, 1995 |
| | 25MG | A074472 002 | Mar 31, 1995 |
| | 50MG | A074472 003 | Mar 31, 1995 |
| | 100MG | A074472 004 | Mar 31, 1995 |
| BOSCOGEN | 12.5MG | A074677 004 | May 30, 1997 |
| | 25MG | A074677 002 | May 30, 1997 |
| | 50MG | A074677 001 | May 30, 1997 |
| | 100MG | A074677 003 | May 30, 1997 |
| DAVA PHARMS INC | 12.5MG | A074423 001 | Feb 13, 1996 |
| | 25MG | A074423 002 | Feb 13, 1996 |
| | 50MG | A074423 003 | Feb 13, 1996 |
| | 100MG | A074423 004 | Feb 13, 1996 |
| EGIS PHARMS | 12.5MG | A074748 004 | May 29, 1997 |
| | 25MG | A074748 002 | May 29, 1997 |
| | 50MG | A074748 001 | May 29, 1997 |
| | 100MG | A074748 003 | May 29, 1997 |
| G AND W LABS INC | 12.5MG | A074433 001 | Feb 13, 1996 |
| | 12.5MG | A074462 001 | Feb 13, 1996 |
| | 12.5MG | A074483 001 | Feb 13, 1996 |
| | 12.5MG | A074590 004 | Aug 30, 1996 |
| | 25MG | A074433 002 | Feb 13, 1996 |
| | 25MG | A074462 002 | Feb 13, 1996 |
| | 25MG | A074483 002 | Feb 13, 1996 |
| | 25MG | A074590 002 | Aug 30, 1996 |
| | 50MG | A074433 003 | Feb 13, 1996 |
| | 50MG | A074462 003 | Feb 13, 1996 |
| | 50MG | A074483 003 | Feb 13, 1996 |
| | 50MG | A074590 001 | Aug 30, 1996 |
| | 100MG | A074433 004 | Feb 13, 1996 |
| | 100MG | A074462 004 | Feb 13, 1996 |
| | 100MG | A074483 004 | Feb 13, 1996 |
| | 100MG | A074590 003 | Aug 30, 1996 |
| OXFORD PHARMS | 12.5MG | A074418 001 | Feb 13, 1996 |
| | 25MG | A074418 002 | Feb 13, 1996 |
| | 50MG | A074418 003 | Feb 13, 1996 |
| | 100MG | A074418 004 | Feb 13, 1996 |
| PAR PHARM | 12.5MG | A074493 001 | Feb 13, 1996 |
| | 25MG | A074493 002 | Feb 13, 1996 |
| | 50MG | A074493 003 | Feb 13, 1996 |
| | 100MG | A074493 004 | Feb 13, 1996 |
| PUREPAC PHARM | 12.5MG | A074640 001 | Mar 31, 1997 |
| | 25MG | A074640 002 | Mar 31, 1997 |
| | 50MG | A074640 003 | Mar 31, 1997 |
| | 100MG | A074640 004 | Mar 31, 1997 |
| SANDOZ | 12.5MG | A074481 001 | Feb 13, 1996 |
| | 25MG | A074481 002 | Feb 13, 1996 |
| | 50MG | A074481 003 | Feb 13, 1996 |
| | 100MG | A074481 004 | Feb 13, 1996 |
| WATSON LABS | 12.5MG | A074451 001 | Feb 13, 1996 |
| | 12.5MG | A074576 001 | Apr 23, 1996 |
| | 25MG | A074451 002 | Feb 13, 1996 |
| | 25MG | A074576 002 | Apr 23, 1996 |
| | 50MG | A074451 003 | Feb 13, 1996 |
| | 50MG | A074576 003 | Apr 23, 1996 |
| | 100MG | A074451 004 | Feb 13, 1996 |
| | 100MG | A074576 004 | Apr 23, 1996 |
| YAOPHARMA CO LTD | 12.5MG | A074363 001 | Nov 09, 1995 |
| | 12.5MG | A074519 001 | Feb 13, 1996 |
| | 25MG | A074363 002 | Nov 09, 1995 |
| | 25MG | A074519 002 | Feb 13, 1996 |
| | 50MG | A074363 003 | Nov 09, 1995 |
| | 50MG | A074519 003 | Feb 13, 1996 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CAPTOPRILTABLET; ORAL
CAPTOPRIL

| | | | |
|-------|---------|-----|--------------|
| 100MG | A074363 | 004 | Nov 09, 1995 |
| 100MG | A074519 | 004 | Feb 13, 1996 |

CAPTOPRIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CAPOZIDE 25/15

+ APOTHECON

25MG;15MG **

N018709 001 Oct 12, 1984

CAPOZIDE 25/25

+ APOTHECON

25MG;25MG **

N018709 002 Oct 12, 1984

CAPOZIDE 50/15

+ APOTHECON

50MG;15MG **

N018709 004 Oct 12, 1984

CAPOZIDE 50/25

+ APOTHECON

50MG;25MG **

N018709 003 Oct 12, 1984

CAPTOPRIL AND HYDROCHLOROTHIAZIDE

G AND W LABS INC

25MG;15MG

A074827 001 Dec 29, 1997

25MG;25MG

A074827 002 Dec 29, 1997

50MG;15MG

A074827 004 Dec 29, 1997

50MG;25MG

A074827 003 Dec 29, 1997

IVAX SUB TEVA PHARMS

25MG;15MG

A075055 001 Jun 18, 1998

25MG;25MG

A075055 002 Jun 18, 1998

50MG;15MG

A075055 004 Jun 18, 1998

50MG;25MG

A075055 003 Jun 18, 1998

VINTAGE PHARMS LLC

25MG;15MG

A074788 001 Dec 29, 1997

25MG;25MG

A074788 002 Dec 29, 1997

50MG;15MG

A074788 004 Dec 29, 1997

50MG;25MG

A074788 003 Dec 29, 1997

WATSON LABS

50MG;25MG

A074832 001 Dec 29, 1997

CARBACHOL

SOLUTION; INTRAOCULAR

CARBACHOL

PHARMAFAIR

0.01%

A070292 001 May 21, 1986

CARBASTAT

NOVARTIS

0.01%

A073677 001 Apr 28, 1995

CARBAMAZEPINE

SOLUTION; INTRAVENOUS

CARNEXIV

+ LUNDBECK PHARMS LLC

200MG/20ML (10MG/ML)

N206030 001 Oct 07, 2016

SUSPENSION; ORAL

CARBAMAZEPINE

TARO

100MG/5ML

A075875 001 Dec 21, 2000

TABLET; ORAL

CARBAMAZEPINE

ACTAVIS ELIZABETH

200MG

A071696 001 Nov 09, 1987

INWOOD LABS

200MG

A070231 001 Aug 14, 1986

PLIVA

200MG

A071479 001 Jul 24, 1987

USL PHARMA

200MG

A070300 001 May 15, 1986

WARNER CHILCOTT

200MG

A070429 001 Jan 02, 1987

TERIL

TARO

200MG

A076525 001 Sep 26, 2003

TABLET, CHEWABLE; ORAL

CARBAMAZEPINE

JUBILANT CADISTA

100MG

A071940 001 Feb 01, 1988

CARBENICILLIN DISODIUM

INJECTABLE; INJECTION

GEOPEN

ROERIG

EQ 1GM BASE/VIAL

N050306 001

EQ 2GM BASE/VIAL

N050306 004

EQ 5GM BASE/VIAL

N050306 002

EQ 10GM BASE/VIAL

N050306 006

EQ 30GM BASE/VIAL

N050306 007

PYOPEN

GLAXOSMITHKLINE

EQ 1GM BASE/VIAL

N050298 001

EQ 2GM BASE/VIAL

N050298 002

EQ 5GM BASE/VIAL

N050298 003

EQ 10GM BASE/VIAL

N050298 006

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARBENICILLIN DISODIUMINJECTABLE; INJECTION
PYOPEN

EQ 20GM BASE/VIAL N050298 007

CARBENICILLIN INDANYL SODIUMTABLET; ORAL
GEOCILLIN

PFIZER EQ 382MG BASE N050435 001

CARBIDOPA; LEVODOPA

TABLET; ORAL

CARBIDOPA AND LEVODOPA

ANI PHARMS INC 10MG; 100MG A073587 002 Jun 29, 1995

25MG; 100MG A073587 001 Jun 29, 1995

25MG; 250MG A073587 003 Jun 29, 1995

SCS 10MG; 100MG A074080 001 Mar 25, 1994

25MG; 100MG A074080 002 Mar 25, 1994

25MG; 250MG A074080 003 Mar 25, 1994

WATSON LABS 10MG; 100MG A073381 001 Sep 28, 1993

25MG; 100MG A073382 001 Sep 28, 1993

25MG; 250MG A073383 001 Sep 28, 1993

TABLET, EXTENDED RELEASE; ORAL

CARBIDOPA AND LEVODOPA

KV PHARM 50MG; 200MG A076663 001 Jun 24, 2004

TABLET, FOR SUSPENSION; ORAL

CARBILEV

RANBAXY 10MG; 100MG A076643 001 Jun 10, 2005

25MG; 100MG A076643 002 Jun 10, 2005

25MG; 250MG A076643 003 Jun 10, 2005

TABLET, ORALLY DISINTEGRATING; ORAL

CARBIDOPA AND LEVODOPA

IMPAX LABS 10MG; 100MG A090631 001 Jun 08, 2010

25MG; 100MG A090631 002 Jun 08, 2010

25MG; 250MG A090631 003 Jun 08, 2010

PARCOPA

UCB INC 10MG; 100MG ** A076699 001 Aug 27, 2004

25MG; 100MG ** A076699 002 Aug 27, 2004

25MG; 250MG ** A076699 003 Aug 27, 2004

CARBINOXAMINE MALEATE

ELIXIR; ORAL

CLISTIN

+ MCNEIL 4MG/5ML **

N008955 001

SOLUTION; ORAL

CARBINOXAMINE MALEATE

CYPRESS PHARM 4MG/5ML A090418 001 May 04, 2010

TABLET; ORAL

CARBINOXAMINE MALEATE

CYPRESS PHARM 4MG A090417 001 Aug 23, 2010

CLISTIN

+ ORTHO MCNEIL PHARM 4MG ** N008915 001

CARBOPLATIN

INJECTABLE; INJECTION

CARBOPLATIN

CIPLA LTD 50MG/VIAL A077383 001 Jan 27, 2006

150MG/VIAL A077383 002 Jan 27, 2006

450MG/VIAL A077383 003 Jan 27, 2006

FRESENIUS KABI USA 50MG/VIAL A076235 001 Oct 14, 2004

150MG/VIAL A076235 002 Oct 14, 2004

450MG/VIAL A076235 003 Oct 14, 2004

HOSPIRA 50MG/VIAL A076473 001 Oct 27, 2004

150MG/VIAL A076473 002 Oct 27, 2004

450MG/VIAL A076473 003 Oct 27, 2004

MYLAN LABS LTD 50MG/VIAL A091510 001 May 29, 2012

150MG/VIAL A091510 002 May 29, 2012

450MG/VIAL A091510 003 May 29, 2012

PLIVA 50MG/VIAL A076602 001 Nov 16, 2004

150MG/VIAL A076602 002 Nov 16, 2004

450MG/VIAL A076602 003 Nov 16, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARBOPLATIN

INJECTABLE; INJECTION

CARBOPLATIN

| | | | |
|----------------------|------------|-------------|--------------|
| SANDOZ | 50MG/VIAL | A076959 001 | Mar 18, 2005 |
| | 150MG/VIAL | A076959 002 | Mar 18, 2005 |
| | 450MG/VIAL | A076959 003 | Mar 18, 2005 |
| WATSON LABS TEVA | 50MG/VIAL | A076162 001 | Oct 14, 2004 |
| | 150MG/VIAL | A076162 002 | Oct 14, 2004 |
| | 450MG/VIAL | A076162 003 | Oct 14, 2004 |
| WEST-WARD PHARMS INT | 50MG/VIAL | A076099 001 | Oct 14, 2004 |
| | 150MG/VIAL | A076099 002 | Oct 14, 2004 |
| | 450MG/VIAL | A076099 003 | Oct 14, 2004 |

PARAPLATIN

| | | | | |
|---|---------------|---------------|-------------|--------------|
| + | CORDEN PHARMA | 50MG/VIAL ** | N019880 001 | Mar 03, 1989 |
| + | | 150MG/VIAL ** | N019880 002 | Mar 03, 1989 |
| + | | 450MG/VIAL ** | N019880 003 | Mar 03, 1989 |

INJECTABLE; IV (INFUSION)

CARBOPLATIN

| | | | |
|--------------------|----------------------|-------------|--------------|
| ACTAVIS TOTOWA | 50MG/5ML (10MG/ML) | A078732 001 | Feb 06, 2012 |
| | 150MG/15ML (10MG/ML) | A078732 002 | Feb 06, 2012 |
| | 450MG/45ML (10MG/ML) | A078732 003 | Feb 06, 2012 |
| | 600MG/60ML (10MG/ML) | A078732 004 | Feb 06, 2012 |
| FRESENIUS KABI USA | 50MG/5ML (10MG/ML) | A077247 001 | Oct 21, 2004 |
| | 50MG/5ML (10MG/ML) | A077266 001 | Feb 15, 2006 |
| | 150MG/15ML (10MG/ML) | A077247 002 | Oct 21, 2004 |
| MYLAN LABS LTD | 150MG/15ML (10MG/ML) | A077266 002 | Feb 15, 2006 |
| | 1GM/100ML (10MG/ML) | A091478 001 | Nov 23, 2011 |
| PHARMACHEMIE BV | 50MG/5ML (10MG/ML) | A077679 001 | Feb 25, 2009 |
| | 150MG/15ML (10MG/ML) | A077679 002 | Feb 25, 2009 |
| | 450MG/45ML (10MG/ML) | A077679 003 | Feb 25, 2009 |
| TEVA PARENTERAL | 50MG/5ML (10MG/ML) | A077389 001 | Mar 30, 2007 |
| | 150MG/15ML (10MG/ML) | A077389 002 | Mar 30, 2007 |
| | 450MG/45ML (10MG/ML) | A077389 003 | Mar 30, 2007 |

PARAPLATIN

| | | | | |
|---|--------------|-------------------------|-------------|--------------|
| + | CORDENPHARMA | 50MG/5ML (10MG/ML) ** | N020452 001 | Jul 14, 2003 |
| + | | 150MG/15ML (10MG/ML) ** | N020452 002 | Jul 14, 2003 |
| + | | 450MG/45ML (10MG/ML) ** | N020452 003 | Jul 14, 2003 |
| + | | 600MG/60ML (10MG/ML) ** | N020452 004 | Jan 15, 2004 |

CARISOPRODOL

CAPSULE; ORAL

SOMA

| | | | |
|---|---------------------|-------|-------------|
| + | MYLAN SPECIALITY LP | 250MG | N011792 003 |
|---|---------------------|-------|-------------|

TABLET; ORAL

CARISOPRODOL

| | | | |
|--------------------|-------|-------------|--------------|
| ABLE | 350MG | A040421 001 | Jun 21, 2001 |
| EPIC PHARMA LLC | 350MG | A040397 001 | Sep 21, 2000 |
| FOSUN PHARMA | 350MG | A081025 001 | Apr 13, 1989 |
| OXFORD PHARMS | 350MG | A040188 001 | Mar 07, 1997 |
| PIONEER PHARMS | 350MG | A089390 001 | Oct 13, 1988 |
| SANDOZ | 350MG | A089566 001 | Aug 30, 1988 |
| SUN PHARM INDS LTD | 350MG | A040755 001 | Feb 27, 2007 |
| WATSON LABS | 350MG | A040152 001 | Dec 03, 1996 |
| | 350MG | A085433 001 | |
| WATSON LABS TEVA | 350MG | A086179 001 | |

RELA

| | | |
|----------|-------|-------------|
| SCHERING | 350MG | N012155 001 |
|----------|-------|-------------|

CARPHENAZINE MALEATE

CONCENTRATE; ORAL

PROKETAZINE

| | | |
|--------------|---------|-------------|
| WYETH AYERST | 50MG/ML | N014173 001 |
|--------------|---------|-------------|

TABLET; ORAL

PROKETAZINE

| | | |
|--------------|--------|-------------|
| WYETH AYERST | 12.5MG | N012768 001 |
| | 25MG | N012768 002 |
| | 50MG | N012768 004 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CARPROFEN

TABLET;ORAL

RIMADYL

ROCHE

100MG

N018550 002 Dec 31, 1987

150MG

N018550 003 Dec 31, 1987

CARTEOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

CARTEOLOL HYDROCHLORIDE

APOTEX INC

1%

A076097 001 Feb 06, 2002

OCUPRESS

+ NOVARTIS

1% **

N019972 001 May 23, 1990

TABLET;ORAL

CARTROL

ABBVIE

2.5MG

N019204 001 Dec 28, 1988

5MG

N019204 002 Dec 28, 1988

10MG

N019204 003 Dec 28, 1988

CARVEDILOL

TABLET;ORAL

CARVEDILOL

HIKMA

3.125MG

A077887 001 Sep 07, 2007

6.25MG

A077887 002 Sep 07, 2007

12.5MG

A077887 003 Sep 07, 2007

25MG

A077887 004 Sep 07, 2007

PLIVA HRVATSKA DOO

3.125MG

A078240 001 Oct 30, 2007

6.25MG

A078240 002 Oct 30, 2007

12.5MG

A078240 003 Oct 30, 2007

25MG

A078240 004 Oct 30, 2007

WOCKHARDT LTD

3.125MG

A078786 001 Dec 22, 2009

6.25MG

A078786 002 Dec 22, 2009

12.5MG

A078786 003 Dec 22, 2009

25MG

A078786 004 Dec 22, 2009

CEFACLOR

CAPSULE;ORAL

CECLOR

+ LILLY

EQ 250MG BASE **

N050521 001

+

EQ 500MG BASE **

N050521 002

CEFACLOR

CEPH INTL

EQ 250MG BASE

A062205 001

EQ 500MG BASE

A062205 002

DAVA PHARMS INC

EQ 250MG BASE

A064107 001 Apr 27, 1995

EQ 500MG BASE

A064107 002 Apr 27, 1995

IVAX SUB TEVA PHARMS

EQ 250MG BASE

A064061 001 Apr 27, 1995

EQ 500MG BASE

A064061 002 Apr 27, 1995

RANBAXY

EQ 250MG BASE

A064156 001 Aug 28, 1997

EQ 500MG BASE

A064156 002 Aug 28, 1997

TEVA

EQ 250MG BASE

A064081 001 Sep 16, 1996

EQ 250MG BASE

A064145 001 Jun 24, 1996

EQ 500MG BASE

A064081 002 Sep 16, 1996

EQ 500MG BASE

A064145 002 Jun 24, 1996

WATSON LABS INC

EQ 250MG BASE

A064148 001 May 23, 1996

EQ 500MG BASE

A064148 002 May 23, 1996

FOR SUSPENSION;ORAL

CECLOR

+ LILLY

EQ 125MG BASE/5ML **

N050522 001

+

EQ 250MG BASE/5ML **

N050522 002

CEFACLOR

DAVA PHARMS INC

EQ 125MG BASE/5ML

A064114 001 Apr 28, 1995

EQ 187MG BASE/5ML

A064115 001 Apr 28, 1995

EQ 250MG BASE/5ML

A064116 001 Apr 28, 1995

EQ 375MG BASE/5ML

A064110 001 Apr 28, 1995

FACTA FARMA

EQ 125MG BASE/5ML

A062206 001

EQ 187MG BASE/5ML

A062206 003 Apr 20, 1988

EQ 250MG BASE/5ML

A062206 002

EQ 375MG BASE/5ML

A062206 004 Apr 20, 1988

IVAX SUB TEVA PHARMS

EQ 125MG BASE/5ML

A064087 001 Apr 28, 1995

EQ 187MG BASE/5ML

A064086 001 Apr 28, 1995

EQ 250MG BASE/5ML

A064085 001 Apr 28, 1995

EQ 375MG BASE/5ML

A064070 001 Apr 28, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFACTOR

FOR SUSPENSION;ORAL

CEFACTOR

| | | | |
|-----------------|-------------------|-------------|--------------|
| RANBAXY | EQ 125MG BASE/5ML | A064166 001 | Oct 02, 1997 |
| | EQ 187MG BASE/5ML | A064165 001 | Oct 02, 1997 |
| | EQ 250MG BASE/5ML | A064164 001 | Oct 02, 1997 |
| | EQ 375MG BASE/5ML | A064155 001 | Oct 02, 1997 |
| WATSON LABS INC | EQ 125MG BASE/5ML | A064204 001 | Feb 18, 1998 |
| | EQ 187MG BASE/5ML | A064205 001 | Feb 18, 1998 |
| | EQ 250MG BASE/5ML | A064206 001 | Feb 18, 1998 |
| | EQ 375MG BASE/5ML | A064207 001 | Feb 18, 1998 |

TABLET, CHEWABLE;ORAL

RANICLOR

| | | | |
|------------------|---------------|-------------|--------------|
| RANBAXY LABS LTD | EQ 125MG BASE | A065092 001 | Dec 22, 2003 |
| | EQ 187MG BASE | A065092 002 | Dec 22, 2003 |
| | EQ 250MG BASE | A065092 003 | Dec 22, 2003 |
| | EQ 375MG BASE | A065092 004 | Dec 22, 2003 |

TABLET, EXTENDED RELEASE;ORAL

CECLOR CD

| | | | |
|-------|---------------|-------------|--------------|
| LILLY | EQ 375MG BASE | N050673 001 | Jun 28, 1996 |
| | EQ 500MG BASE | N050673 002 | Jun 28, 1996 |

CEFACTOR

| | | | |
|-----------|---------------|-------------|--------------|
| WORLD GEN | EQ 500MG BASE | A065057 001 | Jan 05, 2001 |
|-----------|---------------|-------------|--------------|

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE;ORAL

CEFADROXIL

| | | | |
|----------------------|---------------|-------------|--------------|
| CSPC OUYI PHARM CO | EQ 500MG BASE | A205072 001 | Jul 28, 2017 |
| IVAX SUB TEVA PHARMS | EQ 500MG BASE | A062766 001 | Mar 03, 1987 |
| PUREPAC PHARM | EQ 500MG BASE | A063017 001 | Jan 05, 1989 |
| RANBAXY LABS LTD | EQ 500MG BASE | A065015 001 | Jun 22, 1999 |
| SANDOZ | EQ 500MG BASE | A062291 001 | |
| TEVA | EQ 500MG BASE | A062695 001 | Feb 10, 1989 |

DURICEF

| | | | |
|-----------------|---------------|-------------|--|
| WARNER CHILCOTT | EQ 250MG BASE | N050512 002 | |
|-----------------|---------------|-------------|--|

+

| | | | |
|--|------------------|-------------|--|
| | EQ 500MG BASE ** | N050512 001 | |
|--|------------------|-------------|--|

ULTRACEF

| | | | |
|---------|---------------|-------------|--------------|
| BRISTOL | EQ 500MG BASE | A062378 001 | Mar 16, 1982 |
|---------|---------------|-------------|--------------|

FOR SUSPENSION;ORAL

CEFADROXIL

| | | | |
|--------------------|-------------------|-------------|--------------|
| ANI PHARMS INC | EQ 125MG BASE/5ML | A062698 001 | Mar 01, 1989 |
| | EQ 250MG BASE/5ML | A062698 002 | Mar 01, 1989 |
| | EQ 250MG BASE/5ML | A065278 001 | Jan 20, 2006 |
| | EQ 500MG BASE/5ML | A062698 003 | Mar 01, 1989 |
| | EQ 500MG BASE/5ML | A065278 002 | Jan 20, 2006 |
| APOTHECON | EQ 125MG BASE/5ML | A062334 001 | |
| | EQ 250MG BASE/5ML | A062334 002 | |
| | EQ 500MG BASE/5ML | A062334 003 | |
| SUN PHARM INDS LTD | EQ 125MG BASE/5ML | A065115 001 | Mar 26, 2003 |
| | EQ 250MG BASE/5ML | A065115 002 | Mar 26, 2003 |
| | EQ 500MG BASE/5ML | A065115 003 | Mar 26, 2003 |

DURICEF

+

| | | | |
|-----------------|----------------------|-------------|--|
| WARNER CHILCOTT | EQ 125MG BASE/5ML ** | N050527 002 | |
|-----------------|----------------------|-------------|--|

+

| | | | |
|--|----------------------|-------------|--|
| | EQ 250MG BASE/5ML ** | N050527 003 | |
|--|----------------------|-------------|--|

+

| | | | |
|--|----------------------|-------------|--|
| | EQ 500MG BASE/5ML ** | N050527 001 | |
|--|----------------------|-------------|--|

ULTRACEF

| | | | |
|---------|-------------------|-------------|--------------|
| BRISTOL | EQ 125MG BASE/5ML | A062376 001 | Mar 16, 1982 |
| | EQ 250MG BASE/5ML | A062376 002 | Mar 16, 1982 |
| | EQ 500MG BASE/5ML | A062376 003 | Mar 16, 1982 |

TABLET;ORAL

CEFADROXIL

| | | | |
|---------|-------------|-------------|--------------|
| RANBAXY | EQ 1GM BASE | A065018 001 | Apr 23, 1999 |
|---------|-------------|-------------|--------------|

DURICEF

+

| | | | |
|-----------------|----------------|-------------|--|
| WARNER CHILCOTT | EQ 1GM BASE ** | N050528 001 | |
|-----------------|----------------|-------------|--|

ULTRACEF

| | | | |
|-----------|-------------|-------------|--------------|
| APOTHECON | EQ 1GM BASE | A062390 001 | Jun 10, 1982 |
| BRISTOL | EQ 1GM BASE | A062408 001 | Aug 31, 1982 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFAMANDOLE NAFATE

INJECTABLE; INJECTION

MANDOL

| | | | |
|-------|--------------------|-------------|--------------|
| LILLY | EQ 500MG BASE/VIAL | N050504 001 | |
| | EQ 1GM BASE/VIAL | A062560 001 | Sep 10, 1985 |
| | EQ 1GM BASE/VIAL | N050504 002 | |
| | EQ 2GM BASE/VIAL | A062560 002 | Sep 10, 1985 |
| | EQ 2GM BASE/VIAL | N050504 003 | |
| | EQ 10GM BASE/VIAL | N050504 004 | |

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

ANCEF

| | | | |
|----------------------------------------------------|-----------------|-----------------------|--------------------------|
| + | GLAXOSMITHKLINE | EQ 250MG BASE/VIAL ** | N050461 001 |
| + | | EQ 500MG BASE/VIAL | N050461 002 |
| + | | EQ 1GM BASE/VIAL ** | N050461 003 |
| + | | EQ 5GM BASE/VIAL ** | N050461 004 |
| + | | EQ 10GM BASE/VIAL ** | N050461 005 |
| ANCEF IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | EQ 10MG BASE/ML | N050566 003 Jun 08, 1983 |
| | | EQ 20MG BASE/ML | N050566 004 Jun 08, 1983 |
| ANCEF IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | EQ 10MG BASE/ML | A063002 001 Mar 28, 1991 |
| ANCEF IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | |
| | BAXTER HLTHCARE | EQ 10MG BASE/ML | N050566 001 Jun 08, 1983 |
| | | EQ 20MG BASE/ML | N050566 002 Jun 08, 1983 |

CEFAZOLIN AND DEXTROSE

| | | | |
|---------|--------------------|-------------|--------------|
| B BRAUN | EQ 500MG BASE/VIAL | N050779 001 | Jul 27, 2000 |
|---------|--------------------|-------------|--------------|

CEFAZOLIN SODIUM

| | | | |
|----------------------|-----------------------|-------------|--------------|
| ABRAXIS PHARM | EQ 500MG BASE/VIAL | A062688 002 | Nov 17, 1986 |
| | EQ 1GM BASE/VIAL | A062688 003 | Nov 17, 1986 |
| | EQ 10GM BASE/VIAL | A062688 004 | Nov 17, 1986 |
| | EQ 20GM BASE/VIAL | A062688 005 | Aug 03, 1987 |
| AUROBINDO PHARMA | EQ 500MG BASE/VIAL | A065395 001 | Aug 08, 2008 |
| | EQ 1GM BASE/VIAL | A065395 002 | Aug 08, 2008 |
| BEDFORD | EQ 250MG BASE/VIAL | A062894 001 | Jul 21, 1988 |
| | EQ 500MG BASE/VIAL | A062894 002 | Jul 21, 1988 |
| | EQ 1GM BASE/VIAL | A062894 003 | Jul 21, 1988 |
| | EQ 5GM BASE/VIAL | A062894 004 | Jul 21, 1988 |
| | EQ 10GM BASE/VIAL | A062894 005 | Jul 21, 1988 |
| CEPHAZONE PHARMA | EQ 500MG BASE/VIAL | A065280 001 | Mar 18, 2009 |
| | EQ 1GM BASE/VIAL | A065280 002 | Mar 18, 2009 |
| | EQ 10GM BASE/VIAL | A065295 001 | Mar 18, 2009 |
| | EQ 20GM BASE/VIAL | A065296 001 | Mar 18, 2009 |
| FACTA FARMA | EQ 500MG BASE/VIAL | A063214 001 | Dec 27, 1991 |
| | EQ 1GM BASE/VIAL | A063207 001 | Dec 27, 1991 |
| | EQ 10GM BASE/VIAL | A063209 001 | Dec 27, 1991 |
| | EQ 20GM BASE/VIAL | A063209 002 | Apr 30, 1999 |
| FRESENIUS KABI USA | EQ 500MG BASE/VIAL ** | A064169 001 | Aug 14, 1998 |
| | EQ 1GM BASE/VIAL ** | A064169 002 | Aug 14, 1998 |
| | EQ 10GM BASE/VIAL | A064170 001 | Mar 18, 1998 |
| | EQ 20GM BASE/VIAL | A064170 002 | Mar 18, 1998 |
| GLAXOSMITHKLINE | EQ 1GM BASE/VIAL | A064033 001 | Oct 31, 1993 |
| STERI PHARMA | EQ 500MG BASE/VIAL | A063216 001 | Dec 27, 1991 |
| | EQ 1GM BASE/VIAL | A063208 001 | Dec 27, 1991 |
| TEVA PHARMS | EQ 250MG BASE/VIAL | A063016 001 | Mar 14, 1989 |
| | EQ 500MG BASE/VIAL | A063016 002 | Mar 14, 1989 |
| | EQ 1GM BASE/VIAL | A063016 003 | Mar 14, 1989 |
| | EQ 5GM BASE/VIAL | A063018 001 | Mar 05, 1990 |
| | EQ 10GM BASE/VIAL | A063018 002 | Mar 05, 1990 |
| WATSON LABS INC | EQ 250MG BASE/VIAL | A062988 001 | Dec 29, 1989 |
| | EQ 500MG BASE/VIAL | A062988 002 | Dec 29, 1989 |
| | EQ 1GM BASE/VIAL | A062988 003 | Dec 29, 1989 |
| | EQ 5GM BASE/VIAL | A062989 001 | Dec 29, 1989 |
| | EQ 10GM BASE/VIAL | A062989 002 | Dec 29, 1989 |
| | EQ 20GM BASE/VIAL | A062989 003 | Dec 29, 1989 |
| WEST-WARD PHARMS INT | EQ 250MG BASE/VIAL | A062807 001 | Jan 12, 1988 |
| | EQ 500MG BASE/VIAL | A062807 002 | Jan 12, 1988 |
| | EQ 1GM BASE/VIAL | A062807 003 | Jan 12, 1988 |
| | EQ 5GM BASE/VIAL | A062807 004 | Jan 12, 1988 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

CEFAZOLIN SODIUM

| | | |
|-------------------|-------------|--------------|
| EQ 10GM BASE/VIAL | A062807 005 | Jan 12, 1988 |
| EQ 20GM BASE/VIAL | A062807 006 | Jan 12, 1988 |

KEFZOL

ACS DOBFAR

| | | |
|--------------------|-------------|--------------|
| EQ 250MG BASE/VIAL | A061773 001 | |
| EQ 500MG BASE/VIAL | A061773 002 | |
| EQ 1GM BASE/VIAL | A061773 003 | |
| EQ 10GM BASE/VIAL | A061773 004 | |
| EQ 20GM BASE/VIAL | A061773 005 | Sep 08, 1987 |
| EQ 500MG BASE/VIAL | A062557 001 | Sep 10, 1985 |
| EQ 1GM BASE/VIAL | A062557 002 | Sep 10, 1985 |

LILLY

CEFDINIR

CAPSULE; ORAL

OMNICEF

+ ABBVIE

| | | |
|----------|-------------|--------------|
| 300MG ** | N050739 001 | Dec 04, 1997 |
|----------|-------------|--------------|

FOR SUSPENSION; ORAL

OMNICEF

+ ABBVIE

| | | |
|--------------|-------------|--------------|
| 125MG/5ML ** | N050749 001 | Dec 04, 1997 |
|--------------|-------------|--------------|

+

| | | |
|--------------|-------------|--------------|
| 250MG/5ML ** | N050749 002 | Jul 29, 2004 |
|--------------|-------------|--------------|

CEFDITOREN PIVOXIL

TABLET; ORAL

SPECTRACEF

VANSEN PHARMA

| | | |
|-------|-------------|--------------|
| 200MG | N021222 001 | Aug 29, 2001 |
| 400MG | N021222 002 | Jul 21, 2008 |

CEFEPIME HYDROCHLORIDE

INJECTABLE; INJECTION

CEFEPIME HYDROCHLORIDE

FOSUN PHARMA

| | | |
|--------------------|-------------|--------------|
| EQ 500MG BASE/VIAL | A090291 001 | Dec 21, 2010 |
| EQ 1GM BASE/VIAL | A090291 002 | Dec 21, 2010 |
| EQ 2GM BASE/VIAL | A090291 003 | Dec 21, 2010 |

CEFIXIME

FOR SUSPENSION; ORAL

SUPRAX

+ LEDERLE

| | | |
|--------------|-------------|--------------|
| 100MG/5ML ** | N050622 001 | Apr 28, 1989 |
|--------------|-------------|--------------|

TABLET; ORAL

SUPRAX

+ LEDERLE

| | | |
|----------|-------------|--------------|
| 200MG ** | N050621 001 | Apr 28, 1989 |
|----------|-------------|--------------|

+

| | | |
|----------|-------------|--------------|
| 400MG ** | N050621 002 | Apr 28, 1989 |
|----------|-------------|--------------|

CEFMENOXIME HYDROCHLORIDE

INJECTABLE; INJECTION

CEFMAX

TAP PHARM

| | | |
|--------------------|-------------|--------------|
| EQ 500MG BASE/VIAL | N050571 001 | Dec 30, 1987 |
| EQ 1GM BASE/VIAL | N050571 002 | Dec 30, 1987 |
| EQ 2GM BASE/VIAL | N050571 003 | Dec 30, 1987 |

CEFMETAZOLE SODIUM

INJECTABLE; INJECTION

ZEFAZONE

+ PHARMACIA AND UPJOHN

| | | |
|---------------------|-------------|--------------|
| EQ 1GM BASE/VIAL ** | N050637 001 | Dec 11, 1989 |
|---------------------|-------------|--------------|

+

| | | |
|---------------------|-------------|--------------|
| EQ 2GM BASE/VIAL ** | N050637 002 | Dec 11, 1989 |
|---------------------|-------------|--------------|

ZEFAZONE IN PLASTIC CONTAINER

+ PHARMACIA AND UPJOHN

| | | |
|--------------------|-------------|--------------|
| EQ 20MG BASE/ML ** | N050683 001 | Dec 29, 1992 |
|--------------------|-------------|--------------|

+

| | | |
|--------------------|-------------|--------------|
| EQ 40MG BASE/ML ** | N050683 002 | Dec 29, 1992 |
|--------------------|-------------|--------------|

CEFONICID SODIUM

INJECTABLE; INJECTION

MONOCID

GLAXOSMITHKLINE

| | | |
|--------------------|-------------|--------------|
| EQ 500MG BASE/VIAL | N050579 001 | May 23, 1984 |
| EQ 1GM BASE/VIAL | A063295 001 | Jul 26, 1993 |
| EQ 1GM BASE/VIAL | N050579 002 | May 23, 1984 |
| EQ 2GM BASE/VIAL | N050579 003 | May 23, 1984 |
| EQ 10GM BASE/VIAL | N050579 004 | May 23, 1984 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFOPERAZONE SODIUM

INJECTABLE; INJECTION

CEFOBID

| | | | |
|--------|-------------------|-------------|--------------|
| PFIZER | EQ 1GM BASE/VIAL | A063333 001 | Mar 31, 1995 |
| | EQ 1GM BASE/VIAL | N050551 001 | Nov 18, 1982 |
| | EQ 2GM BASE/VIAL | A063333 002 | Mar 31, 1995 |
| | EQ 2GM BASE/VIAL | N050551 002 | Nov 18, 1982 |
| | EQ 10GM BASE/VIAL | N050551 003 | Mar 05, 1990 |

CEFOBID IN PLASTIC CONTAINER

| | | | |
|--------|-----------------|-------------|--------------|
| PFIZER | EQ 20MG BASE/ML | N050613 002 | Jul 31, 1987 |
| | EQ 40MG BASE/ML | N050613 001 | Jul 23, 1986 |

CEFORANIDE

INJECTABLE; INJECTION

PRECEF

| | | | |
|-----------|------------|-------------|--------------|
| APOTHECON | 500MG/VIAL | A062579 001 | Nov 26, 1984 |
| | 1GM/VIAL | A062579 002 | Nov 26, 1984 |
| | 2GM/VIAL | A062579 003 | Nov 26, 1984 |
| | 10GM/VIAL | A062579 004 | Nov 26, 1984 |
| | 20GM/VIAL | A062579 005 | Nov 26, 1984 |
| BRISTOL | 500MG/VIAL | N050554 001 | May 24, 1984 |
| | 1GM/VIAL | N050554 002 | May 24, 1984 |
| | 2GM/VIAL | N050554 003 | May 24, 1984 |
| | 10GM/VIAL | N050554 004 | May 24, 1984 |
| | 20GM/VIAL | N050554 005 | May 24, 1984 |

CEFOTAXIME SODIUM

INJECTABLE; INJECTION

CEFOTAXIME

| | | | |
|---------------------------------------------------|--------------------|-------------|--------------|
| FRESENIUS KABI USA | EQ 500MG BASE/VIAL | A064200 001 | Mar 24, 2000 |
| | EQ 1GM BASE/VIAL | A064200 002 | Mar 24, 2000 |
| | EQ 2GM BASE/VIAL | A064200 003 | Mar 24, 2000 |
| | EQ 10GM BASE/VIAL | A064201 001 | Mar 24, 2000 |
| | EQ 20GM BASE/VIAL | A064201 002 | Mar 24, 2000 |
| CEFOTAXIME AND DEXTROSE 2.4% IN PLASTIC CONTAINER | | | |
| B BRAUN | EQ 2GM BASE | N050792 001 | Jul 29, 2004 |
| CEFOTAXIME AND DEXTROSE 3.9% IN PLASTIC CONTAINER | | | |
| B BRAUN | EQ 1GM BASE | N050792 002 | Jul 29, 2004 |
| CEFOTAXIME SODIUM | | | |
| AUROBINDO PHARMA | EQ 500MG BASE/VIAL | A065517 001 | Nov 06, 2009 |
| | EQ 1GM BASE/VIAL | A065517 002 | Nov 06, 2009 |
| | EQ 2GM BASE/VIAL | A065517 003 | Nov 06, 2009 |
| AUROBINDO PHARMA LTD | EQ 10GM BASE/VIAL | A065516 001 | Nov 06, 2009 |
| CEPHAZONE PHARMA | EQ 10GM BASE/VIAL | A065348 001 | Jan 25, 2010 |
| LUPIN | EQ 500MG BASE/VIAL | A065124 001 | Sep 24, 2003 |
| | EQ 1GM BASE/VIAL | A065124 002 | Sep 24, 2003 |
| | EQ 2GM BASE/VIAL | A065124 003 | Sep 24, 2003 |

CLAFORAN

| | | | |
|-------------------------------------------------------|--------------------|-------------|--------------|
| SANOFI AVENTIS US | EQ 1GM BASE/VIAL | A062659 001 | Jan 13, 1987 |
| | EQ 2GM BASE/VIAL | A062659 002 | Jan 13, 1987 |
| + US PHARM HOLDINGS | EQ 500MG BASE/VIAL | N050547 001 | |
| + | EQ 1GM BASE/VIAL | N050547 002 | |
| + | EQ 2GM BASE/VIAL | N050547 003 | |
| + | EQ 10GM BASE/VIAL | N050547 004 | Dec 29, 1983 |
| CLAFORAN IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| US PHARM HOLDINGS | EQ 20MG BASE/ML | N050596 002 | May 20, 1985 |
| | EQ 40MG BASE/ML | N050596 004 | May 20, 1985 |
| CLAFORAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | |
| US PHARM HOLDINGS | EQ 20MG BASE/ML | N050596 001 | May 20, 1985 |
| | EQ 40MG BASE/ML | N050596 003 | May 20, 1985 |

CEFOTETAN DISODIUM

INJECTABLE; INJECTION

CEFOTAN

| | | | |
|------------------------------|-------------------|-------------|--------------|
| TELIGENT | EQ 10GM BASE/VIAL | N050588 003 | Apr 25, 1988 |
| TELIGENT PHARMA INC | EQ 1GM BASE/VIAL | A063293 001 | Apr 29, 1993 |
| | EQ 2GM BASE/VIAL | A063293 002 | Apr 29, 1993 |
| CEFOTAN IN PLASTIC CONTAINER | | | |
| TELIGENT | EQ 20MG BASE/ML | N050694 002 | Jul 30, 1993 |
| | EQ 40MG BASE/ML | N050694 001 | Jul 30, 1993 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFOTIAM HYDROCHLORIDE

INJECTABLE; INJECTION

CERADON

| | | | | |
|--------|------------------|---------|-----|--------------|
| TAKEDA | EQ 1GM BASE/VIAL | N050601 | 001 | Dec 30, 1988 |
|--------|------------------|---------|-----|--------------|

CEFOXITIN SODIUM

INJECTABLE; INJECTION

CEFOXITIN

| | | | | |
|--------------------|---------------------|---------|-----|--------------|
| ACS DOBFAR SPA | EQ 1GM BASE/VIAL | A065467 | 001 | Aug 31, 2011 |
| | EQ 2GM BASE/VIAL | A065467 | 002 | Aug 31, 2011 |
| | EQ 10GM BASE/VIAL | A065464 | 001 | Aug 31, 2011 |
| FRESENIUS KABI USA | EQ 1GM BASE/VIAL ** | A065012 | 001 | Jul 03, 2000 |
| | EQ 2GM BASE/VIAL ** | A065012 | 002 | Jul 03, 2000 |
| | EQ 10GM BASE/VIAL | A065011 | 001 | Jul 03, 2000 |

MEFOXIN

| | | | | |
|------------------------------------------------------|----------------------|---------|-----|--------------|
| MYLAN INSTITUTIONAL | EQ 1GM BASE/VIAL | A062757 | 001 | Jan 08, 1987 |
| + | EQ 1GM BASE/VIAL ** | N050517 | 001 | |
| | EQ 2GM BASE/VIAL | A062757 | 002 | Jan 08, 1987 |
| + | EQ 2GM BASE/VIAL ** | N050517 | 002 | |
| + | EQ 10GM BASE/VIAL ** | N050517 | 003 | |
| MEFOXIN IN DEXTROSE 5% IN PLASTIC CONTAINER | | | | |
| + | EQ 20MG BASE/ML ** | N050581 | 003 | Sep 20, 1984 |
| + | EQ 40MG BASE/ML ** | N050581 | 004 | Sep 20, 1984 |
| MEFOXIN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | | |
| + | EQ 20MG BASE/ML ** | N050581 | 002 | Sep 20, 1984 |
| + | EQ 40MG BASE/ML ** | N050581 | 001 | Sep 20, 1984 |

CEFPYRAMIDE SODIUM

INJECTABLE; INJECTION

CEFPYRAMIDE SODIUM

| | | | | |
|--------------|-------------------|---------|-----|--------------|
| WYETH AYERST | EQ 1GM BASE/VIAL | N050633 | 002 | Jan 31, 1989 |
| | EQ 2GM BASE/VIAL | N050633 | 003 | Jan 31, 1989 |
| | EQ 10GM BASE/VIAL | N050633 | 005 | Jan 31, 1989 |

CEFPODOXIME PROXETIL

FOR SUSPENSION; ORAL

BANAN

| | | | | |
|--------|-------------------|---------|-----|--------------|
| SANKYO | EQ 50MG BASE/5ML | N050688 | 002 | Aug 07, 1992 |
| | EQ 100MG BASE/5ML | N050688 | 001 | Aug 07, 1992 |

CEFPODOXIME PROXETIL

| | | | | |
|--------------------|-------------------|---------|-----|--------------|
| SUN PHARM INDS LTD | EQ 50MG BASE/5ML | A065082 | 001 | May 31, 2002 |
| | EQ 100MG BASE/5ML | A065082 | 002 | May 31, 2002 |

VANTIN

| | | | | | |
|---|----------------------|----------------------|---------|-----|--------------|
| + | PHARMACIA AND UPJOHN | EQ 50MG BASE/5ML ** | N050675 | 001 | Aug 07, 1992 |
| + | | EQ 100MG BASE/5ML ** | N050675 | 002 | Aug 07, 1992 |

TABLET; ORAL

BANAN

| | | | | |
|--------|---------------|---------|-----|--------------|
| SANKYO | EQ 100MG BASE | N050687 | 001 | Aug 07, 1992 |
| | EQ 200MG BASE | N050687 | 002 | Aug 07, 1992 |

CEFPODOXIME PROXETIL

| | | | | |
|--------------------|---------------|---------|-----|--------------|
| SUN PHARM INDS LTD | EQ 100MG BASE | A065083 | 001 | Aug 20, 2003 |
| | EQ 200MG BASE | A065083 | 002 | Aug 20, 2003 |

VANTIN

| | | | | | |
|---|----------------------|------------------|---------|-----|--------------|
| + | PHARMACIA AND UPJOHN | EQ 100MG BASE ** | N050674 | 001 | Aug 07, 1992 |
| + | | EQ 200MG BASE ** | N050674 | 002 | Aug 07, 1992 |

CEFPROZIL

FOR SUSPENSION; ORAL

CEFPROZIL

| | | | | |
|------------------|-----------|---------|-----|--------------|
| RANBAXY LABS LTD | 125MG/5ML | A065202 | 001 | Jun 30, 2006 |
| | 250MG/5ML | A065202 | 002 | Jun 30, 2006 |

CEFZIL

| | | | | | |
|---|---------------|-----------|---------|-----|--------------|
| + | CORDEN PHARMA | 125MG/5ML | N050665 | 001 | Dec 23, 1991 |
| + | | 250MG/5ML | N050665 | 002 | Dec 23, 1991 |

TABLET; ORAL

CEFPROZIL

| | | | | |
|------------------|-------|---------|-----|--------------|
| RANBAXY LABS LTD | 250MG | A065198 | 001 | Dec 13, 2006 |
| | 500MG | A065198 | 002 | Dec 13, 2006 |

CEFZIL

| | | | | | |
|---|---------------|-------|---------|-----|--------------|
| + | CORDEN PHARMA | 250MG | N050664 | 001 | Dec 23, 1991 |
| + | | 500MG | N050664 | 002 | Dec 23, 1991 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFTAZIDIME

INJECTABLE; INJECTION

CEFTAZIDIME

| | | | | |
|----------------------|------------|---------|-----|--------------|
| ACS DOBFAR | 500MG/VIAL | A062640 | 001 | Nov 20, 1985 |
| AUROBINDO PHARMA LTD | 500MG/VIAL | A065481 | 001 | May 28, 2010 |
| | 1GM/VIAL | A065481 | 002 | May 28, 2010 |
| | 2GM/VIAL | A065481 | 003 | May 28, 2010 |
| | 6GM/VIAL | A065482 | 001 | May 28, 2010 |

CEPTAZ

| | | | | |
|-----------------|------------|---------|-----|--------------|
| GLAXOSMITHKLINE | 500MG/VIAL | N050646 | 001 | Sep 27, 1990 |
| | 1GM/VIAL | N050646 | 002 | Sep 27, 1990 |
| | 2GM/VIAL | N050646 | 003 | Sep 27, 1990 |
| | 10GM/VIAL | N050646 | 004 | Sep 27, 1990 |

PENTACEF

| | | | | |
|-----------------|-----------|---------|-----|--------------|
| GLAXOSMITHKLINE | 1GM/VIAL | A063322 | 001 | Nov 07, 1995 |
| | 1GM/VIAL | A064006 | 001 | Mar 31, 1992 |
| | 2GM/VIAL | A063322 | 002 | Nov 07, 1995 |
| | 2GM/VIAL | A064006 | 002 | Mar 31, 1992 |
| | 6GM/VIAL | A064008 | 001 | Mar 31, 1992 |
| | 10GM/VIAL | A064008 | 002 | Mar 31, 1992 |

TAZIDIME

| | | | | |
|-------|----------|---------|-----|--------------|
| LILLY | 1GM/VIAL | A062655 | 001 | Nov 20, 1985 |
| | 2GM/VIAL | A062655 | 002 | Nov 20, 1985 |

TAZIDIME IN PLASTIC CONTAINER

| | | | | |
|-------|----------|---------|-----|--------------|
| LILLY | 1GM/VIAL | A062739 | 001 | Jul 10, 1986 |
| | 2GM/VIAL | A062739 | 002 | Jul 10, 1986 |

CEFTAZIDIME SODIUM

INJECTABLE; INJECTION

CEFTAZIDIME SODIUM IN PLASTIC CONTAINER

| | | | | |
|-----------------|-----------------|---------|-----|--------------|
| BAXTER HLTHCARE | EQ 10MG BASE/ML | A063221 | 001 | Apr 29, 1993 |
| | EQ 20MG BASE/ML | A063221 | 002 | Apr 29, 1993 |
| | EQ 40MG BASE/ML | A063221 | 003 | Apr 29, 1993 |

FORTAZ IN PLASTIC CONTAINER

| | | | | |
|----------|-----------------|---------|-----|--------------|
| TELIGENT | EQ 10MG BASE/ML | N050634 | 001 | Apr 28, 1989 |
| + | EQ 20MG BASE/ML | N050634 | 002 | Apr 28, 1989 |
| + | EQ 40MG BASE/ML | N050634 | 003 | Apr 28, 1989 |

CEFTIBUTEN DIHYDRATE

CAPSULE; ORAL

CEDAX

| | | | | |
|-----------|---------------|---------|-----|--------------|
| SI PHARMS | EQ 400MG BASE | N050685 | 002 | Dec 20, 1995 |
|-----------|---------------|---------|-----|--------------|

FOR SUSPENSION; ORAL

CEDAX

| | | | | | |
|---|-----------|----------------------|---------|-----|--------------|
| + | SI PHARMS | EQ 90MG BASE/5ML ** | N050686 | 001 | Dec 20, 1995 |
| + | | EQ 180MG BASE/5ML ** | N050686 | 002 | Dec 20, 1995 |

CEFTIZOXIME SODIUM

INJECTABLE; INJECTION

CEFIZOX

| | | | | |
|----------|--------------------|---------|-----|--------------|
| ASTELLAS | EQ 500MG BASE/VIAL | N050560 | 001 | Sep 15, 1983 |
| | EQ 1GM BASE/VIAL | A063294 | 002 | Mar 31, 1994 |
| | EQ 1GM BASE/VIAL | N050560 | 002 | Sep 15, 1983 |
| | EQ 2GM BASE/VIAL | A063294 | 003 | Mar 31, 1994 |
| | EQ 2GM BASE/VIAL | N050560 | 003 | Sep 15, 1983 |
| | EQ 10GM BASE/VIAL | N050560 | 005 | Mar 19, 1993 |

CEFIZOX IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|----------|-----------------|---------|-----|--------------|
| ASTELLAS | EQ 20MG BASE/ML | N050589 | 001 | Oct 03, 1984 |
| | EQ 40MG BASE/ML | N050589 | 002 | Oct 03, 1984 |

CEFIZOX IN PLASTIC CONTAINER

| | | | | |
|----------|-----------------|---------|-----|--------------|
| ASTELLAS | EQ 20MG BASE/ML | N050589 | 003 | Apr 13, 1995 |
| | EQ 40MG BASE/ML | N050589 | 004 | Apr 13, 1995 |

CEFTRIAZONE SODIUM

INJECTABLE; INJECTION

CEFTRIAZONE

| | | | | |
|----------------------|-------------------|---------|-----|--------------|
| AGILA SPECLTS | EQ 10GM BASE/VIAL | A091068 | 001 | Jan 07, 2013 |
| AUROBINDO PHARMA LTD | EQ 10GM BASE/VIAL | A065504 | 001 | Jul 31, 2008 |
| BEDFORD | EQ 10GM BASE/VIAL | A065475 | 001 | Aug 18, 2008 |
| FACTA FARMA | EQ 10GM BASE/VIAL | A065269 | 001 | Feb 28, 2007 |
| FRESENIUS KABI USA | EQ 10GM BASE/VIAL | A065252 | 001 | Feb 15, 2006 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFTRIAXONE SODIUM

INJECTABLE; INJECTION

CEFTRIAXONE

| | | | |
|-------------------------------------------|--------------------------------------|-------------|--------------|
| HOSPIRA INC | EQ 1GM BASE/VIAL | A065231 001 | Aug 02, 2005 |
| | EQ 1GM BASE/VIAL | A202563 001 | Aug 20, 2012 |
| | EQ 2GM BASE/VIAL | A065231 002 | Aug 02, 2005 |
| | EQ 2GM BASE/VIAL | A202563 002 | Aug 20, 2012 |
| LUPIN | EQ 10GM BASE/VIAL | A065263 001 | Sep 12, 2006 |
| TEVA | EQ 10GM BASE/VIAL | A065274 001 | May 01, 2006 |
| ROCEPHIN | | | |
| HOFFMANN LA ROCHE | EQ 250MG BASE/VIAL | A063239 001 | Aug 13, 1993 |
| | EQ 500MG BASE/VIAL | A062654 001 | Apr 30, 1987 |
| | EQ 500MG BASE/VIAL | A063239 002 | Aug 13, 1993 |
| | EQ 1GM BASE/VIAL | A062654 002 | Apr 30, 1987 |
| | EQ 1GM BASE/VIAL | A063239 003 | Aug 13, 1993 |
| | EQ 2GM BASE/VIAL | A062654 003 | Apr 30, 1987 |
| + | EQ 10GM BASE/VIAL | N050585 005 | Dec 21, 1984 |
| ROCHE | EQ 250MG BASE/VIAL | A062510 001 | Mar 12, 1985 |
| | EQ 500MG BASE/VIAL | A062510 002 | Mar 12, 1985 |
| | EQ 1GM BASE/VIAL | A062510 003 | Mar 12, 1985 |
| ROCEPHIN W/ DEXTROSE IN PLASTIC CONTAINER | | | |
| + | HOFFMANN LA ROCHE EQ 10MG BASE/ML ** | N050624 001 | Feb 11, 1987 |
| + | EQ 20MG BASE/ML ** | N050624 002 | Feb 11, 1987 |
| + | EQ 40MG BASE/ML ** | N050624 003 | Feb 11, 1987 |

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFTRIAXONE

| | | | |
|----------------------|--------------------------------------|-------------|--------------|
| AUROBINDO PHARMA LTD | EQ 250MG BASE/VIAL | A065505 001 | Jul 31, 2008 |
| | EQ 500MG BASE/VIAL | A065505 002 | Jul 31, 2008 |
| | EQ 1GM BASE/VIAL | A065505 003 | Jul 31, 2008 |
| | EQ 2GM BASE/VIAL | A065505 004 | Jul 31, 2008 |
| BEDFORD | EQ 250MG BASE/VIAL | A065465 001 | Aug 18, 2008 |
| | EQ 500MG BASE/VIAL | A065465 002 | Aug 18, 2008 |
| | EQ 1GM BASE/VIAL | A065465 003 | Aug 18, 2008 |
| | EQ 2GM BASE/VIAL | A065465 004 | Aug 18, 2008 |
| CEPHAZONE PHARMA | EQ 250MG BASE/VIAL | A065294 001 | Mar 26, 2007 |
| | EQ 500MG BASE/VIAL | A065294 002 | Mar 26, 2007 |
| | EQ 1GM BASE/VIAL | A065294 003 | Mar 26, 2007 |
| | EQ 2GM BASE/VIAL | A065294 004 | Mar 26, 2007 |
| FACTA FARMA | EQ 1GM BASE/VIAL | A065268 001 | Feb 28, 2007 |
| | EQ 2GM BASE/VIAL | A065268 002 | Feb 28, 2007 |
| FRESENIUS KABI USA | EQ 250MG BASE/VIAL | A065245 001 | Feb 15, 2006 |
| | EQ 500MG BASE/VIAL | A065245 002 | Feb 15, 2006 |
| | EQ 1GM BASE/VIAL | A065245 003 | Feb 15, 2006 |
| | EQ 2GM BASE/VIAL | A065245 004 | Feb 15, 2006 |
| TEVA | EQ 1GM BASE/VIAL | A065262 001 | Jun 29, 2006 |
| | EQ 2GM BASE/VIAL | A065262 002 | Jun 29, 2006 |
| TEVA PHARMS USA | EQ 250MG BASE/VIAL | A065227 001 | Mar 15, 2007 |
| | EQ 500MG BASE/VIAL | A065227 002 | Mar 15, 2007 |
| | EQ 1GM BASE/VIAL | A065227 003 | Mar 15, 2007 |
| | EQ 2GM BASE/VIAL | A065227 004 | Mar 15, 2007 |
| ROCEPHIN | | | |
| + | HOFFMANN LA ROCHE EQ 250MG BASE/VIAL | N050585 001 | Dec 21, 1984 |
| + | EQ 500MG BASE/VIAL | N050585 002 | Dec 21, 1984 |
| + | EQ 1GM BASE/VIAL | N050585 003 | Dec 21, 1984 |
| + | EQ 2GM BASE/VIAL | N050585 004 | Dec 21, 1984 |

CEFTRIAXONE SODIUM; LIDOCAINE

INJECTABLE; INJECTION

ROCEPHIN KIT

| | | | |
|-------------------|----------------------------------|-------------|--------------|
| HOFFMANN LA ROCHE | EQ 500MG BASE/VIAL, N/A; N/A, 1% | N050585 007 | May 08, 1996 |
| | EQ 1GM BASE/VIAL, N/A; N/A, 1% | N050585 006 | May 08, 1996 |

CEFUROXIME AXETIL

FOR SUSPENSION; ORAL

CEFTIN

| | | | |
|---|-----------------------------------|-------------|--------------|
| + | GLAXOSMITHKLINE EQ 125MG BASE/5ML | N050672 001 | Jun 30, 1994 |
| + | EQ 250MG BASE/5ML | N050672 002 | Apr 29, 1997 |

CEFUROXIME AXETIL

| | | | |
|--------------------|-------------------|-------------|--------------|
| SUN PHARM INDS LTD | EQ 125MG BASE/5ML | A065323 001 | Feb 05, 2008 |
| | EQ 250MG BASE/5ML | A065323 002 | Feb 05, 2008 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEFUROXIME AXETIL

TABLET; ORAL

CEFTIN

| | | | | |
|---|-----------------|---------------|-------------|--------------|
| + | GLAXOSMITHKLINE | EQ 125MG BASE | N050605 001 | Dec 28, 1987 |
| + | | EQ 250MG BASE | N050605 002 | Dec 28, 1987 |
| + | | EQ 500MG BASE | N050605 003 | Dec 28, 1987 |

CEFUROXIME AXETIL

| | | | |
|--------------------|---------------|-------------|--------------|
| FOSUN PHARMA | EQ 250MG BASE | A065126 001 | Oct 28, 2003 |
| | EQ 500MG BASE | A065126 002 | Oct 28, 2003 |
| RANBAXY LABS LTD | EQ 125MG BASE | A065043 003 | Feb 15, 2002 |
| | EQ 250MG BASE | A065043 002 | Feb 15, 2002 |
| | EQ 500MG BASE | A065043 001 | Feb 15, 2002 |
| SUN PHARM INDS LTD | EQ 125MG BASE | A065118 001 | Apr 25, 2003 |
| | EQ 250MG BASE | A065118 002 | Apr 25, 2003 |
| | EQ 500MG BASE | A065118 003 | Apr 25, 2003 |

CEFUROXIME SODIUM

INJECTABLE; INJECTION

CEFUROXIME SODIUM

| | | | |
|--------------------|--------------------|-------------|--------------|
| FRESENIUS KABI USA | EQ 1.5GM BASE/VIAL | A065001 002 | May 30, 2001 |
| | EQ 7.5GM BASE/VIAL | A065002 001 | Sep 28, 1998 |
| TEVA PHARMS | EQ 7.5GM BASE/VIAL | A064191 001 | Apr 16, 1998 |
| WATSON LABS INC | EQ 1.5GM BASE/VIAL | A064035 002 | Feb 26, 1993 |
| | EQ 7.5GM BASE/VIAL | A064036 001 | Feb 26, 1993 |

CEFUROXIME SODIUM IN PLASTIC CONTAINER

| | | | |
|--------------|--------------------|-------------|--------------|
| SAMSON MEDCL | EQ 75GM BASE/VIAL | A065251 001 | Dec 30, 2009 |
| | EQ 225GM BASE/VIAL | A065251 002 | Dec 30, 2009 |

KEFUROX

| | | | |
|------------|--------------------|-------------|--------------|
| ACS DOBFAR | EQ 1.5GM BASE/VIAL | A062591 002 | Jan 10, 1986 |
| | EQ 7.5GM BASE/VIAL | A062591 003 | Dec 17, 1987 |
| LILLY | EQ 1.5GM BASE/VIAL | A062592 002 | Jan 10, 1986 |

KEFUROX IN PLASTIC CONTAINER

| | | | |
|-------|--------------------|-------------|--------------|
| LILLY | EQ 1.5GM BASE/VIAL | A062590 002 | Jan 10, 1986 |
|-------|--------------------|-------------|--------------|

ZINACEF IN PLASTIC CONTAINER

| | | | |
|----------|-----------------|-------------|--------------|
| TELIGENT | EQ 15MG BASE/ML | N050643 001 | Apr 28, 1989 |
| + | EQ 30MG BASE/ML | N050643 002 | Apr 28, 1989 |

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFUROXIME SODIUM

| | | | |
|--------------------|--------------------|-------------|--------------|
| FRESENIUS KABI USA | EQ 750MG BASE/VIAL | A065001 001 | May 30, 2001 |
| TEVA PHARMS | EQ 750MG BASE/VIAL | A064192 002 | Apr 16, 1998 |
| | EQ 1.5GM BASE/VIAL | A064192 001 | Apr 16, 1998 |
| WATSON LABS INC | EQ 750MG BASE/VIAL | A064035 001 | Feb 26, 1993 |

KEFUROX

| | | | |
|------------|--------------------|-------------|--------------|
| ACS DOBFAR | EQ 750MG BASE/VIAL | A062591 001 | Jan 10, 1986 |
|------------|--------------------|-------------|--------------|

INJECTABLE; INTRAVENOUS

KEFUROX

| | | | |
|-------|--------------------|-------------|--------------|
| LILLY | EQ 750MG BASE/VIAL | A062592 001 | Jan 10, 1986 |
|-------|--------------------|-------------|--------------|

KEFUROX IN PLASTIC CONTAINER

| | | | |
|-------|--------------------|-------------|--------------|
| LILLY | EQ 750MG BASE/VIAL | A062590 001 | Jan 10, 1986 |
|-------|--------------------|-------------|--------------|

CELLULOSE SODIUM PHOSPHATE

POWDER; ORAL

CALCIBIND

| | | | |
|----------------|--------------|-------------|--------------|
| MISSION PHARMA | 2.5GM/PACKET | N018757 002 | Dec 28, 1982 |
| | 300GM/BOT | N018757 003 | Oct 16, 1984 |

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

| | | | |
|----------------------|---------------|-------------|--------------|
| APOTHECON | EQ 250MG BASE | A062973 001 | Nov 08, 1988 |
| | EQ 250MG BASE | A063063 001 | Sep 29, 1989 |
| | EQ 250MG BASE | A063186 001 | Dec 30, 1994 |
| | EQ 500MG BASE | A062974 001 | Nov 23, 1988 |
| | EQ 500MG BASE | A063063 002 | Sep 29, 1989 |
| | EQ 500MG BASE | A063186 002 | Dec 30, 1994 |
| BARR | EQ 250MG BASE | A062773 001 | Jun 26, 1987 |
| | EQ 500MG BASE | A062775 001 | Apr 22, 1987 |
| FACTA FARMA | EQ 250MG BASE | A062118 001 | |
| | EQ 500MG BASE | A062118 002 | |
| IVAX SUB TEVA PHARMS | EQ 250MG BASE | A061969 001 | |
| | EQ 500MG BASE | A061969 002 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

| | | | |
|--------------------|---------------|-------------|--------------|
| PUREPAC PHARM | EQ 250MG BASE | A062809 001 | Apr 22, 1987 |
| | EQ 500MG BASE | A062809 002 | Apr 22, 1987 |
| STEVENS J | EQ 250MG BASE | A062870 001 | Mar 17, 1988 |
| | EQ 500MG BASE | A062869 001 | Mar 17, 1988 |
| SUN PHARM INDS LTD | EQ 250MG BASE | A065007 001 | Sep 16, 1999 |
| | EQ 500MG BASE | A065007 002 | Sep 16, 1999 |
| TEVA | EQ 250MG BASE | A062760 001 | Apr 24, 1987 |
| | EQ 250MG BASE | A062821 001 | Feb 05, 1988 |
| | EQ 500MG BASE | A062761 001 | Apr 24, 1987 |
| | EQ 500MG BASE | A062823 001 | Feb 05, 1988 |
| YOSHITOMI | EQ 250MG BASE | A062872 001 | Jun 20, 1988 |
| | EQ 500MG BASE | A062871 001 | Jul 05, 1988 |

KEFLEX

+ PRAGMA

FOR SUSPENSION; ORAL

CEPHALEXIN

| | | | |
|--------------------|----------------------|-------------|--------------|
| APOTHECON | EQ 125MG BASE/5ML | A062986 001 | Apr 18, 1991 |
| | EQ 250MG BASE/5ML | A062987 001 | Jul 25, 1989 |
| BARR | EQ 125MG BASE/5ML | A062778 001 | Aug 06, 1987 |
| | EQ 250MG BASE/5ML | A062777 001 | Aug 06, 1987 |
| FACTA FARMA | EQ 100MG BASE/ML ** | A062117 001 | |
| | EQ 125MG BASE/5ML ** | A062117 002 | |
| | EQ 250MG BASE/5ML ** | A062117 003 | |
| HIKMA PHARMS | EQ 125MG BASE/5ML | A065444 001 | Aug 28, 2009 |
| | EQ 250MG BASE/5ML | A065444 002 | Aug 28, 2009 |
| SUN PHARM INDS LTD | EQ 125MG BASE/5ML | A065081 001 | Jul 27, 2001 |
| | EQ 250MG BASE/5ML | A065081 002 | Jul 27, 2001 |
| TEVA | EQ 125MG BASE/5ML | A062767 001 | Jun 16, 1987 |
| | EQ 125MG BASE/5ML | A062873 001 | May 23, 1988 |
| | EQ 250MG BASE/5ML | A062768 001 | Jun 16, 1987 |
| | EQ 250MG BASE/5ML | A062867 001 | Apr 15, 1988 |
| VITARINE | EQ 125MG BASE/5ML | A062779 001 | Dec 22, 1987 |
| | EQ 250MG BASE/5ML | A062781 001 | Dec 22, 1987 |

KEFLEX

+ PRAGMA

+

+

TABLET; ORAL

CEPHALEXIN

| | | | |
|----------|---------------|-------------|--------------|
| BARR | EQ 250MG BASE | A062826 001 | Aug 17, 1987 |
| | EQ 500MG BASE | A062827 001 | Aug 17, 1987 |
| VITARINE | EQ 250MG BASE | A062863 001 | Aug 11, 1988 |
| | EQ 500MG BASE | A062863 002 | Aug 11, 1988 |
| | EQ 1GM BASE | A062863 003 | Aug 11, 1988 |

KEFLET

LILLY

| | | |
|---------------|-------------|--------------|
| EQ 250MG BASE | A062745 001 | Dec 01, 1986 |
| EQ 250MG BASE | N050440 003 | Feb 26, 1987 |
| EQ 500MG BASE | A062745 002 | Dec 01, 1986 |
| EQ 500MG BASE | N050440 001 | |
| EQ 1GM BASE | N050440 002 | |

TABLET, FOR SUSPENSION; ORAL

PANIXINE DISPERDOSE

RANBAXY LABS LTD

| | | |
|---------------|-------------|--------------|
| EQ 125MG BASE | A065100 002 | Sep 11, 2003 |
| EQ 250MG BASE | A065100 001 | Sep 11, 2003 |

CEPHALEXIN HYDROCHLORIDE

TABLET; ORAL

KEFTAB

LILLY

| | | |
|---------------|-------------|--------------|
| EQ 250MG BASE | N050614 001 | Oct 29, 1987 |
| EQ 333MG BASE | N050614 003 | May 16, 1988 |
| EQ 500MG BASE | N050614 002 | Oct 29, 1987 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEPHALOGLYCIN

CAPSULE; ORAL

KAFOCIN

LILLY

250MG

N050219 001

CEPHALOTHIN SODIUM

INJECTABLE; INJECTION

CEPHALOTHIN

INTL MEDICATION

EQ 500MG BASE/VIAL

A062426 001 May 03, 1985

EQ 1GM BASE/VIAL

A062426 002 May 03, 1985

EQ 2GM BASE/VIAL

A062426 003 May 03, 1985

EQ 4GM BASE/VIAL

A062426 004 May 03, 1985

CEPHALOTHIN SODIUM

ABBOTT

EQ 1GM BASE/VIAL

A062547 001 Sep 11, 1985

EQ 1GM BASE/VIAL

A062548 001 Sep 11, 1985

EQ 2GM BASE/VIAL

A062547 002 Sep 11, 1985

EQ 2GM BASE/VIAL

A062548 002 Sep 11, 1985

ABRAXIS PHARM

EQ 1GM BASE/VIAL

A062666 002 Jun 10, 1987

EQ 2GM BASE/VIAL

A062666 001 Jun 10, 1987

BRISTOL

EQ 1GM BASE/VIAL

A062464 001 May 07, 1984

EQ 2GM BASE/VIAL

A062464 002 May 07, 1984

EQ 4GM BASE/VIAL

A062464 003 May 07, 1984

CEPHALOTHIN SODIUM W/ DEXTROSE IN PLASTIC CONTAINER

BAXTER HLTHCARE

EQ 20MG BASE/ML

A062422 003 Jan 31, 1984

EQ 20MG BASE/ML

A062422 005 Jul 16, 1991

EQ 20MG BASE/ML

A062730 001 Mar 05, 1987

EQ 40MG BASE/ML

A062422 004 Jan 31, 1984

EQ 40MG BASE/ML

A062422 006 Jul 16, 1991

EQ 40MG BASE/ML

A062730 002 Mar 05, 1987

CEPHALOTHIN SODIUM W/ SODIUM CHLORIDE IN PLASTIC CONTAINER

BAXTER HLTHCARE

EQ 20MG BASE/ML

A062422 001 Jan 31, 1984

EQ 40MG BASE/ML

A062422 002 Jan 31, 1984

KEFLIN

LILLY

EQ 1GM BASE/VIAL

N050482 001

EQ 2GM BASE/VIAL

N050482 002

EQ 4GM BASE/VIAL

N050482 003

EQ 20GM BASE/VIAL

N050482 007

KEFLIN IN PLASTIC CONTAINER

LILLY

EQ 1GM BASE/VIAL

A062549 001 Sep 10, 1985

EQ 2GM BASE/VIAL

A062549 002 Sep 10, 1985

SEFFIN

GLAXOSMITHKLINE

EQ 1GM BASE/VIAL

A062435 001 Nov 15, 1983

EQ 2GM BASE/VIAL

A062435 002 Nov 15, 1983

EQ 10GM BASE/VIAL

A062435 003 Nov 15, 1983

CEPHAPIRIN SODIUM

INJECTABLE; INJECTION

CEFADYL

APOTHECON

EQ 500MG BASE/VIAL

A062961 001 Sep 20, 1988

EQ 500MG BASE/VIAL

N050446 005

EQ 1GM BASE/VIAL

A061769 001

EQ 1GM BASE/VIAL

A062724 001 Dec 23, 1986

EQ 1GM BASE/VIAL

A062961 002 Sep 20, 1988

EQ 1GM BASE/VIAL

N050446 001

EQ 2GM BASE/VIAL

A061769 002

EQ 2GM BASE/VIAL

A062724 002 Dec 23, 1986

EQ 2GM BASE/VIAL

A062961 003 Sep 20, 1988

EQ 2GM BASE/VIAL

N050446 002

EQ 4GM BASE/VIAL

A061769 003

EQ 4GM BASE/VIAL

A062961 004 Sep 20, 1988

EQ 4GM BASE/VIAL

N050446 003

EQ 20GM BASE/VIAL

N050446 004

CEPHAPIRIN SODIUM

ABRAXIS PHARM

EQ 500MG BASE/VIAL

A062723 001 Nov 17, 1986

EQ 1GM BASE/VIAL

A062723 002 Nov 17, 1986

EQ 2GM BASE/VIAL

A062723 003 Nov 17, 1986

EQ 4GM BASE/VIAL

A062723 004 Nov 17, 1986

EQ 20GM BASE/VIAL

A062723 005 Nov 17, 1986

WEST-WARD PHARMS INT

EQ 500MG BASE/VIAL

A062720 001 Jul 02, 1987

EQ 1GM BASE/VIAL

A062720 002 Jul 02, 1987

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CEPHAPIRIN SODIUM

INJECTABLE; INJECTION

CEPHAPIRIN SODIUM

| | | |
|-------------------|-------------|--------------|
| EQ 2GM BASE/VIAL | A062720 003 | Jul 02, 1987 |
| EQ 20GM BASE/VIAL | A062720 004 | Jul 02, 1987 |

CEPHRADINE

CAPSULE; ORAL

ANSPOR

| | | |
|-----------------|-------|-------------|
| GLAXOSMITHKLINE | 250MG | A061859 001 |
| | 500MG | A061859 002 |

CEPHRADINE

| | | | |
|----------------------|-------|-------------|--------------|
| BARR | 250MG | A062850 001 | Apr 22, 1988 |
| | 500MG | A062851 001 | Apr 22, 1988 |
| IVAX SUB TEVA PHARMS | 250MG | A062762 001 | Mar 06, 1987 |
| | 500MG | A062762 002 | Mar 06, 1987 |
| TEVA | 250MG | A062683 001 | Jan 09, 1987 |
| | 500MG | A062683 002 | Jan 09, 1987 |
| VITARINE | 250MG | A062813 001 | Feb 25, 1988 |
| | 500MG | A062813 002 | Feb 25, 1988 |

VELOSEF

| | | |
|-----------|-------|-------------|
| APOTHECON | 250MG | A061764 001 |
| | 500MG | A061764 002 |

VELOSEF '250'

| | | |
|--------|-------|-------------|
| ERSANA | 250MG | N050548 001 |
|--------|-------|-------------|

VELOSEF '500'

| | | |
|--------|-------|-------------|
| ERSANA | 500MG | N050548 002 |
|--------|-------|-------------|

FOR SUSPENSION; ORAL

ANSPOR

| | | |
|-----------------|-----------|-------------|
| GLAXOSMITHKLINE | 125MG/5ML | A061866 001 |
| | 250MG/5ML | A061866 002 |

CEPHRADINE

| | | | |
|------|-----------|-------------|--------------|
| BARR | 125MG/5ML | A062858 001 | May 19, 1988 |
| | 250MG/5ML | A062859 001 | May 19, 1988 |
| TEVA | 125MG/5ML | A062693 001 | Jan 09, 1987 |
| | 250MG/5ML | A062693 002 | Jan 09, 1987 |

VELOSEF '125'

| | | |
|-----------|-----------|-------------|
| APOTHECON | 125MG/5ML | A061763 001 |
|-----------|-----------|-------------|

VELOSEF '250'

| | | |
|-----------|-----------|-------------|
| APOTHECON | 250MG/5ML | A061763 002 |
|-----------|-----------|-------------|

INJECTABLE; INJECTION

VELOSEF

| | | |
|-----------|------------|-------------|
| APOTHECON | 250MG/VIAL | A061976 001 |
| | 500MG/VIAL | A061976 002 |
| | 1GM/VIAL | A061976 004 |
| | 2GM/VIAL | A061976 003 |
| | 4GM/VIAL | A061976 005 |

TABLET; ORAL

VELOSEF

| | | |
|----------------------|-----|-------------|
| BRISTOL MYERS SQUIBB | 1GM | N050530 001 |
|----------------------|-----|-------------|

CERIVASTATIN SODIUM

TABLET; ORAL

BAYCOL

| | | | |
|--------------|--------|-------------|--------------|
| BAYER PHARMS | 0.05MG | N020740 001 | Jun 26, 1997 |
| | 0.1MG | N020740 002 | Jun 26, 1997 |
| | 0.2MG | N020740 003 | Jun 26, 1997 |
| | 0.3MG | N020740 004 | Jun 26, 1997 |
| | 0.4MG | N020740 005 | May 24, 1999 |
| | 0.8MG | N020740 006 | Jul 24, 2000 |

CERULETIDE DIETHYLAMINE

INJECTABLE; INJECTION

TYMTRAN

| | | |
|----------------------|-----------|-------------|
| PHARMACIA AND UPJOHN | 0.02MG/ML | N018296 001 |
|----------------------|-----------|-------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CETIRIZINE HYDROCHLORIDE

SYRUP;ORAL

CETIRIZINE HYDROCHLORIDE

| | | | | |
|----------------------|---------|---------|-----|--------------|
| ACTAVIS MID ATLANTIC | 5MG/5ML | A078617 | 001 | Feb 02, 2010 |
| APOTEX INC | 5MG/5ML | A078412 | 001 | Jun 18, 2008 |
| AUROBINDO PHARMA LTD | 5MG/5ML | A090751 | 001 | Dec 16, 2009 |
| RANBAXY LABS LTD | 5MG/5ML | A077472 | 001 | Jun 18, 2008 |
| WOCKHARDT | 5MG/5ML | A078757 | 001 | Aug 28, 2009 |

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

| | | | | |
|----------------------|---------|---------|-----|--------------|
| ACTAVIS MID ATLANTIC | 5MG/5ML | A090378 | 002 | May 09, 2008 |
| APOTEX INC | 5MG/5ML | A090188 | 002 | Apr 22, 2008 |
| CYPRESS PHARM | 5MG/5ML | A090300 | 001 | Oct 10, 2008 |
| RANBAXY LABS LTD | 5MG/5ML | A090183 | 002 | Apr 24, 2008 |

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

| | | | | |
|----------------------|---------|---------|-----|--------------|
| ACTAVIS MID ATLANTIC | 5MG/5ML | A090378 | 001 | May 09, 2008 |
| APOTEX INC | 5MG/5ML | A090188 | 001 | Apr 22, 2008 |
| CYPRESS PHARM | 5MG/5ML | A090300 | 002 | Oct 10, 2008 |
| RANBAXY LABS LTD | 5MG/5ML | A090183 | 001 | Apr 24, 2008 |

ZYRTEC

| | | | | |
|----------------------|------------|---------|-----|--------------|
| J AND J CONSUMER INC | 5MG/5ML ** | N020346 | 001 | Sep 27, 1996 |
|----------------------|------------|---------|-----|--------------|

TABLET;ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

| | | | | |
|-------------------|------|---------|-----|--------------|
| ACTAVIS ELIZABETH | 5MG | A078615 | 003 | Dec 28, 2007 |
| | 10MG | A078615 | 004 | Dec 28, 2007 |

ZYRTEC ALLERGY

| | | | | |
|------------------------|-----|---------|-----|--------------|
| + J AND J CONSUMER INC | 5MG | N019835 | 003 | Nov 16, 2007 |
|------------------------|-----|---------|-----|--------------|

ZYRTEC HIVES RELIEF

| | | | | |
|------------------------|------|---------|-----|--------------|
| + J AND J CONSUMER INC | 5MG | N019835 | 005 | Nov 16, 2007 |
| + J AND J CONSUMER INC | 10MG | N019835 | 006 | Nov 16, 2007 |

TABLET, CHEWABLE;ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

| | | | | |
|--------------------|------|---------|-----|--------------|
| SUN PHARM INDS INC | 5MG | A077631 | 004 | Jan 11, 2008 |
| | 10MG | A077631 | 003 | Jan 11, 2008 |

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

| | | | | |
|--------------------|------|---------|-----|--------------|
| SUN PHARM INDS INC | 5MG | A077631 | 001 | Jan 11, 2008 |
| | 10MG | A077631 | 002 | Jan 11, 2008 |

CHILDREN'S ZYRTEC ALLERGY

| | | | | |
|------------------------|---------|---------|-----|--------------|
| + J AND J CONSUMER INC | 5MG ** | N021621 | 003 | Nov 16, 2007 |
| + J AND J CONSUMER INC | 10MG ** | N021621 | 004 | Nov 16, 2007 |

CHILDREN'S ZYRTEC HIVES RELIEF

| | | | | |
|------------------------|---------|---------|-----|--------------|
| + J AND J CONSUMER INC | 5MG ** | N021621 | 005 | Nov 16, 2007 |
| + J AND J CONSUMER INC | 10MG ** | N021621 | 006 | Nov 16, 2007 |

CETRORELIX

INJECTABLE;INJECTION

CETROTIDE

| | | | | |
|----------------|----------------|---------|-----|--------------|
| EMD SERONO INC | EQ 3MG BASE/ML | N021197 | 002 | Aug 11, 2000 |
|----------------|----------------|---------|-----|--------------|

CETYL ALCOHOL; COLFOSCERIL PALMITATE; TYLOXAPOL

FOR SUSPENSION;INTRATRACHEAL

EXOSURF NEONATAL

| | | | | |
|-----------------|-------------------------------|---------|-----|--------------|
| GLAXOSMITHKLINE | 12MG/VIAL;108MG/VIAL;8MG/VIAL | N020044 | 001 | Aug 02, 1990 |
|-----------------|-------------------------------|---------|-----|--------------|

CEVIMELINE HYDROCHLORIDE

CAPSULE;ORAL

CEVIMELINE HYDROCHLORIDE

| | | | | |
|------------|------|---------|-----|--------------|
| APOTEX INC | 30MG | A091260 | 001 | Aug 25, 2011 |
|------------|------|---------|-----|--------------|

CHENODIOL

TABLET;ORAL

CHENIX

| | | | | |
|-----------------------|----------|---------|-----|--------------|
| + LEADIANT BIOSCI INC | 250MG ** | N018513 | 002 | Jul 28, 1983 |
|-----------------------|----------|---------|-----|--------------|

CHLOPHEDIANOL HYDROCHLORIDE

SYRUP;ORAL

ULO

| | | | | |
|----|----------|---------|-----|--|
| 3M | 25MG/5ML | N012126 | 001 | |
|----|----------|---------|-----|--|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORAMPHENICOL

CREAM; TOPICAL

CHLOROMYCETIN

PARKE DAVIS 1% N050183 001

FOR SOLUTION; OPHTHALMIC

CHLOROMYCETIN

PARKEDALE 25MG/VIAL N050143 001

INJECTABLE; INJECTION

CHLOROMYCETIN

PARKE DAVIS 250MG/ML N050153 001

OINTMENT; OPHTHALMIC

CHLORAMPHENICOL

ALTANA 1% A060133 001

CHLOROFAIR

PHARMAFAIR 1% A062439 001 Apr 21, 1983

CHLOROMYCETIN

PARKEDALE 1% N050156 001

CHLOROPTIC S.O.P.

ALLERGAN 1% A061187 001

ECONOCHLOR

ALCON 1% A061648 001

SOLUTION/DROPS; OPHTHALMIC

CHLORAMPHENICOL

AKORN 0.5% A062042 001

ALCON 0.5% A062628 001 Sep 25, 1985

CHLOROFAIR

PHARMAFAIR 0.5% A062437 001 Apr 14, 1983

CHLOROPTIC

ALLERGAN 0.5% N050091 001

ECONOCHLOR

ALCON 0.5% A061645 001

OPHTHOCHLOR

PARKEDALE 0.5% A061220 001

OPTOMYCIN

OPTOPICS 0.5% A062171 001 Mar 31, 1982

SOLUTION/DROPS; OTIC

CHLOROMYCETIN

PARKEDALE 0.5% N050205 001

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE; INJECTION

CHLORAMPHENICOL

ELKINS SINN EQ 1GM BASE/VIAL A062406 001 Nov 09, 1982

CHLORAMPHENICOL SODIUM SUCCINATE

GRUPPO LEPETIT EQ 1GM BASE/VIAL A062278 001

CHLOROMYCETIN

+ PARKEDALE EQ 1GM BASE/VIAL N050155 001

MYCHEL-S

ANGUS EQ 1GM BASE/VIAL A060132 001

CHLORAMPHENICOL; DESOXYRIBONUCLEASE; FIBRINOLYSIN

OINTMENT; TOPICAL

ELASE-CHLOROMYCETIN

PARKE DAVIS 10MG/GM; 666 UNITS/GM; 1 UNITS/GM N050294 001

CHLORAMPHENICOL; HYDROCORTISONE ACETATE

FOR SUSPENSION; OPHTHALMIC

CHLOROMYCETIN HYDROCORTISONE

PARKEDALE 12.5MG/VIAL; 25MG/VIAL N050202 001

CHLORAMPHENICOL; HYDROCORTISONE ACETATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

OPHTHOCORT

PARKEDALE 10MG/GM; 5MG/GM; 10,000 UNITS/GM N050201 002

CHLORAMPHENICOL; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

CHLOROMYCIN

PARKE DAVIS 1%; 10,000 UNITS/GM N050203 002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORAMPHENICOL; PREDNISOLONE

OINTMENT;OPHTHALMIC

CHLOROPTIC-P S.O.P.

ALLERGAN

1%;0.5%

A061188 001

CHLORDIAZEPOXIDE

CAPSULE, EXTENDED RELEASE;ORAL

LIBRELEASE

VALEANT PHARM INTL

30MG

N017813 001 Sep 12, 1983

TABLET;ORAL

LIBRITABS

VALEANT PHARM INTL

5MG

A085482 001

10MG

A085481 001

25MG

A085488 001

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE;ORAL

A-POXIDE

ABBOTT

5MG

A085447 001

5MG

A085517 001

10MG

A085447 002

10MG

A085518 001

25MG

A085447 003

25MG

A085513 001

CHLORDIAZACHEL

RACHELLE

5MG

A085086 001

10MG

A084639 001

25MG

A085087 001

CHLORDIAZEPOXIDE HYDROCHLORIDE

ASCOT

5MG

A087525 001 Jan 07, 1982

10MG

A087524 001 Jan 07, 1982

25MG

A087512 001 Jan 07, 1982

FERRANTE

5MG

A085118 001

10MG

A085119 001

25MG

A085120 001

HALSEY

5MG

A085340 001

10MG

A085339 001

25MG

A084685 001

IMPAX LABS

5MG

A086213 001

10MG

A085113 001

25MG

A086212 001

IVAX SUB TEVA PHARMS

5MG

A083741 001

10MG

A083742 001

25MG

A083570 001

LEDERLE

5MG

A086892 001

5MG

A087234 001

10MG

A086876 001

10MG

A087037 001

25MG

A086893 001

25MG

A087231 001

MAST MM

10MG

A086217 001

MYLAN

5MG

A084886 001

10MG

A084601 001

25MG

A084887 001

PARKE DAVIS

5MG

A085163 001

10MG

A084598 001

25MG

A085164 001

PIONEER PHARMS

10MG

A089533 001 Jul 15, 1988

25MG

A089558 001 Jul 15, 1988

PUREPAC PHARM

5MG

A085155 001

10MG

A084939 002

25MG

A085144 001

ROXANE

5MG

A084706 001

10MG

A084700 001

25MG

A084705 001

SUPERPHARM

5MG

A088987 001 Apr 25, 1985

10MG

A088986 001 Apr 25, 1985

25MG

A088988 001 Apr 25, 1985

TEVA

5MG

A088705 001 Jan 18, 1985

10MG

A088706 001 Jan 18, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE

| | | | | |
|----------------------|---------|---------|-----|--------------|
| | 25MG | A086494 | 001 | |
| | 25MG | A088707 | 001 | Jan 18, 1985 |
| UPSHER SMITH LABS | 5MG | A084678 | 001 | |
| | 10MG | A084041 | 001 | |
| | 25MG | A084679 | 002 | |
| UPSHER-SMITH LABS | 5MG | A084919 | 001 | |
| | 10MG | A084920 | 001 | |
| | 25MG | A084823 | 001 | |
| USL PHARMA | 5MG | A084644 | 001 | |
| | 10MG | A084623 | 001 | |
| | 25MG | A084645 | 001 | |
| VANGARD | 5MG | A088129 | 001 | Mar 28, 1983 |
| | 10MG | A088010 | 001 | Mar 28, 1983 |
| | 25MG | A088130 | 001 | Mar 28, 1983 |
| WATSON LABS | 5MG | A086383 | 001 | |
| | 10MG | A086294 | 001 | |
| | 25MG | A086382 | 001 | |
| WEST WARD | 5MG | A085014 | 001 | |
| | 10MG | A085000 | 001 | |
| | 25MG | A085294 | 001 | |
| LIBRIUM | | | | |
| + VALEANT PHARM INTL | 5MG ** | N012249 | 002 | |
| + | 10MG ** | N012249 | 001 | |
| + | 25MG ** | N012249 | 003 | |
| LYGEN | | | | |
| ALRA | 5MG | A085107 | 001 | |
| | 10MG | A085009 | 001 | |
| | 25MG | A085108 | 001 | |

INJECTABLE; INJECTION

LIBRIUM

VALEANT PHARMS LLC 100MG/AMP N012301 001

CHLORDIAZEPOXIDE; ESTROGENS, ESTERIFIED

TABLET; ORAL

| | | | | |
|--------------|-------------|---------|-----|--|
| MENRIUM 10-4 | | | | |
| ROCHE | 10MG; 0.4MG | N014740 | 006 | |
| MENRIUM 5-2 | | | | |
| ROCHE | 5MG; 0.2MG | N014740 | 002 | |
| MENRIUM 5-4 | | | | |
| ROCHE | 5MG; 0.4MG | N014740 | 004 | |

CHLORHEXIDINE GLUCONATE

SOLUTION; DENTAL

CHLORHEXIDINE GLUCONATE

BAJAJ 0.12% A075561 001 Nov 14, 2000

SOLUTION; TOPICAL

DYNA-HEX

BAJAJ 0.75% N020111 001 Sep 11, 1997

EXIDINE

XTTRIUM 2.5% N019421 001 Dec 17, 1985

MICRODERM

J AND J 4% A072255 001 Apr 15, 1991

PREVACARE R

J AND J 0.5% A072292 001 Jan 28, 1992

STERI-STAT

MATRIX MEDCL 4% A070104 001 Jul 24, 1986

SPONGE; TOPICAL

CHLORHEXIDINE GLUCONATE

KENDALL IL 4% N019490 001 Mar 27, 1987

E-Z SCRUB

BECTON DICKINSON 4% A073416 001 Mar 14, 2000

HIBICLENS

+ MOLNLYCKE HLTH 4% ** N018423 001

MICRODERM

J AND J 4% A072295 001 Feb 28, 1991

PHARMASEAL SCRUB CARE

CAREFUSION 2200 4% N019793 001 Dec 02, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORMERODRIN HG-197

INJECTABLE; INJECTION

CHLORMERODRIN HG 197

BRACCO

0.6-1.4mCi/ML

N017269 001

CHLORMEZANONE

TABLET; ORAL

TRANCOPAL

SANOFI AVENTIS US

100MG

N011467 003

200MG

N011467 005

CHLOROPROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

NESACAINE-MPF

FRESENIUS KABI USA

2%

N009435 003

3%

N009435 004

CHLOROQUINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARALEN HYDROCHLORIDE

SANOFI AVENTIS US

EQ 40MG BASE/ML

N006002 002

CHLOROQUINE PHOSPHATE

TABLET; ORAL

ARALEN

+ SANOFI AVENTIS US

EQ 300MG BASE

N006002 001

CHLOROQUINE PHOSPHATE

IMPAX LABS

EQ 150MG BASE

A080880 001

EQ 300MG BASE

A040516 001 Aug 29, 2003

MD PHARM

EQ 150MG BASE

A087228 001

PUREPAC PHARM

EQ 150MG BASE

A080886 001

TEVA

EQ 150MG BASE

A087504 001 Jan 13, 1982

WATSON LABS

EQ 150MG BASE

A087979 001 Dec 21, 1982

EQ 300MG BASE

A088030 001 Dec 21, 1982

CHLOROQUINE PHOSPHATE; PRIMAQUINE PHOSPHATE

TABLET; ORAL

ARALEN PHOSPHATE W/ PRIMAQUINE PHOSPHATE

SANOFI AVENTIS US

EQ 300MG BASE; EQ 45MG BASE

N014860 002

CHLOROTHIAZIDE

TABLET; ORAL

CHLOROTHIAZIDE

ABC HOLDING

250MG

A085569 001

HIKMA INTL PHARMS

250MG

A086028 001 Jul 14, 1982

500MG

A087736 001 Jul 14, 1982

LEDERLE

250MG

A086940 001

500MG

A086938 001

SANDOZ

250MG

A085485 001

WATSON LABS

250MG

A085165 001

250MG

A085173 001

250MG

A086795 001 Aug 15, 1983

500MG

A084026 001 Sep 01, 1982

500MG

A086796 001 Aug 15, 1983

DIURIL

+ OAK PHARMS AKORN

250MG **

N011145 004

+

500MG **

N011145 002

CHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

ALDOCLOR-150

MERCK

150MG; 250MG

N016016 001

ALDOCLOR-250

MERCK

250MG; 250MG

N016016 002

METHYLDOPA AND CHLOROTHIAZIDE

PAR PHARM

150MG; 250MG

A070783 001 Nov 06, 1987

250MG; 250MG

A070654 001 Nov 06, 1987

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CHLOROTHIAZIDE AND RESERPINE

| | | | | |
|--------------|---------------|---------|-----|--------------|
| HIKMA PHARMS | 250MG;0.125MG | A088557 | 001 | Dec 22, 1983 |
| | 500MG;0.125MG | A088365 | 001 | Dec 22, 1983 |

CHLOROTHIAZIDE W/ RESERPINE

| | | | | |
|-------------|---------------|---------|-----|--------------|
| WATSON LABS | 250MG;0.125MG | A084853 | 001 | |
| | 500MG;0.125MG | A088151 | 001 | Jun 09, 1983 |

CHLOROTHIAZIDE-RESERPINE

| | | | | |
|-------|---------------|---------|-----|--------------|
| MYLAN | 250MG;0.125MG | A087744 | 001 | May 06, 1982 |
| | 500MG;0.125MG | A087745 | 001 | May 06, 1982 |

DIUPRES-250

| | | | | |
|-------|---------------|---------|-----|--------------|
| MERCK | 250MG;0.125MG | N011635 | 003 | Aug 26, 1987 |
|-------|---------------|---------|-----|--------------|

DIUPRES-500

| | | | | |
|-------|---------------|---------|-----|--------------|
| MERCK | 500MG;0.125MG | N011635 | 006 | Aug 26, 1987 |
|-------|---------------|---------|-----|--------------|

CHLOROTRIANISENE

CAPSULE; ORAL

CHLOROTRIANISENE

| | | | | |
|-------------------|------|---------|-----|--|
| BANNER PHARMACAPS | 12MG | A084652 | 001 | |
|-------------------|------|---------|-----|--|

TACE

| | | | | |
|-------------------|------|---------|-----|--|
| SANOFI AVENTIS US | 12MG | N008102 | 004 | |
| | 25MG | N011444 | 001 | |
| | 72MG | N016235 | 001 | |

CHLOROXINE

SHAMPOO; TOPICAL

CAPITROL

| | | | | |
|-----------------|----|---------|-----|--|
| WESTWOOD SQUIBB | 2% | N017594 | 001 | |
|-----------------|----|---------|-----|--|

CHLORPHENESIN CARBAMATE

TABLET; ORAL

MAOLATE

| | | | | |
|------------|-------|---------|-----|--|
| PAMLAB LLC | 400MG | N014217 | 002 | |
|------------|-------|---------|-----|--|

CHLORPHENIRAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

CHLORPHENIRAMINE MALEATE

| | | | | |
|---------------------|------|---------|-----|--------------|
| AUROLIFE PHARMA LLC | 12MG | A070797 | 001 | Aug 12, 1988 |
|---------------------|------|---------|-----|--------------|

TELDRIN

| | | | | |
|-----------------|------|---------|-----|--|
| GLAXOSMITHKLINE | 8MG | N017369 | 001 | |
| | 12MG | N017369 | 002 | |

INJECTABLE; INJECTION

CHLOR-TRIMETON

| | | | | |
|-----------------|----------|---------|-----|--|
| SCHERING PLOUGH | 10MG/ML | N008826 | 001 | |
| | 100MG/ML | N008794 | 001 | |

CHLORPHENIRAMINE MALEATE

| | | | | |
|-------------|----------|---------|-----|--|
| BEL MAR | 10MG/ML | A080821 | 001 | |
| ELKINS SINN | 10MG/ML | A080797 | 001 | |
| WATSON LABS | 10MG/ML | A083593 | 001 | |
| | 10MG/ML | A086096 | 001 | |
| | 100MG/ML | A086095 | 001 | |

PYRIDAMAL 100

| | | | | |
|---------|----------|---------|-----|--|
| BEL MAR | 100MG/ML | A083733 | 001 | |
|---------|----------|---------|-----|--|

SYRUP; ORAL

CHLOR-TRIMETON

| | | | | |
|----------|---------|---------|-----|--|
| SCHERING | 2MG/5ML | N006921 | 006 | |
|----------|---------|---------|-----|--|

CHLORPHENIRAMINE MALEATE

| | | | | |
|-------------|---------|---------|-----|--------------|
| PHARM ASSOC | 2MG/5ML | A087520 | 001 | Feb 10, 1982 |
|-------------|---------|---------|-----|--------------|

TABLET; ORAL

ANTAGONATE

| | | | | |
|--------------|-----|---------|-----|--|
| BAYER PHARMS | 4MG | A083381 | 001 | |
|--------------|-----|---------|-----|--|

CHLOR-TRIMETON

| | | | | |
|----------|-----|---------|-----|--|
| SCHERING | 4MG | N006921 | 002 | |
|----------|-----|---------|-----|--|

CHLORPHENIRAMINE MALEATE

| | | | | |
|----------------------|-----|---------|-----|--|
| ANABOLIC | 4MG | A083078 | 001 | |
| AUROLIFE PHARMA LLC | 4MG | A080961 | 001 | |
| BELL PHARMA | 4MG | A083062 | 001 | |
| ELKINS SINN | 4MG | A080938 | 001 | |
| IMPAX LABS | 4MG | A080809 | 001 | |
| IVAX SUB TEVA PHARMS | 4MG | A080779 | 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPHENIRAMINE MALEATE

TABLET; ORAL

CHLORPHENIRAMINE MALEATE

| | | | | |
|----------------------|-----|---------|-----|--------------|
| KV PHARM | 4MG | A087164 | 001 | |
| LEDERLE | 4MG | A086941 | 001 | |
| NEWTRON PHARMS | 4MG | A086519 | 001 | |
| PANRAY | 4MG | A083243 | 001 | |
| PHARMAVITE | 4MG | A085104 | 001 | |
| PHARMERAL | 4MG | A083753 | 001 | |
| PIONEER PHARMS | 4MG | A088556 | 001 | Jul 13, 1984 |
| PUREPAC PHARM | 4MG | A086306 | 001 | |
| PVT FORM | 4MG | A080786 | 001 | |
| ROXANE | 4MG | A080626 | 001 | |
| SUN PHARM INDUSTRIES | 4MG | A080700 | 001 | |
| VITARINE | 4MG | A085837 | 001 | |
| WATSON LABS | 4MG | A080696 | 001 | |
| | 4MG | A080791 | 001 | |
| | 4MG | A085139 | 001 | |
| WEST WARD | 4MG | A083787 | 001 | |

KLOOROMIN

| | | | | |
|--------|-----|---------|-----|--|
| HALSEY | 4MG | A083629 | 001 | |
|--------|-----|---------|-----|--|

PHENETRON

| | | | | |
|---------|-----|---------|-----|--|
| LANNETT | 4MG | A080846 | 001 | |
|---------|-----|---------|-----|--|

TABLET, EXTENDED RELEASE; ORAL

CHLOR-TRIMETON

| | | | | |
|----------------------|-----|---------|-----|--|
| BAYER HEALTHCARE LLC | 8MG | N007638 | 001 | |
|----------------------|-----|---------|-----|--|

EFIDAC 24 CHLORPHENIRAMINE MALEATE

| | | | | |
|------|------|---------|-----|--------------|
| ALZA | 16MG | N019746 | 002 | Nov 18, 1994 |
|------|------|---------|-----|--------------|

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND CHLORPHENIRAMINE MALEATE

| | | | | |
|-----------------|------------------|---------|-----|--------------|
| TRIS PHARMA INC | 4MG/5ML; 5MG/5ML | A206438 | 001 | Jan 27, 2015 |
|-----------------|------------------|---------|-----|--------------|

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

| | | | | |
|---------------|----------------------------|---------|-----|--------------|
| BIO-PHARM INC | 4MG/5ML; 5MG/5ML; 60MG/5ML | A206660 | 001 | May 15, 2017 |
|---------------|----------------------------|---------|-----|--------------|

| | | | | |
|-----------------|----------------------------|---------|-----|--------------|
| TRIS PHARMA INC | 4MG/5ML; 5MG/5ML; 60MG/5ML | A203838 | 001 | Nov 26, 2014 |
|-----------------|----------------------------|---------|-----|--------------|

CHLORPHENIRAMINE MALEATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

COLD CAPSULE IV

| | | | | |
|-----------|------------|---------|-----|--------------|
| GRAHAM DM | 12MG; 75MG | N018793 | 001 | Apr 25, 1985 |
|-----------|------------|---------|-----|--------------|

COLD CAPSULE V

| | | | | |
|-----------|-----------|---------|-----|--------------|
| GRAHAM DM | 8MG; 75MG | N018794 | 001 | Apr 23, 1985 |
|-----------|-----------|---------|-----|--------------|

TABLET, EXTENDED RELEASE; ORAL

TRIAMINIC-12

| | | | | |
|----------|------------|---------|-----|--|
| NOVARTIS | 12MG; 75MG | N018115 | 001 | |
|----------|------------|---------|-----|--|

CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CODIMAL-L.A. 12

| | | | | |
|----------------|-------------|---------|-----|--------------|
| SCHWARZ PHARMA | 12MG; 120MG | N018935 | 001 | Apr 15, 1985 |
|----------------|-------------|---------|-----|--------------|

ISOCOLOR

| | | | | |
|--------|------------|---------|-----|--------------|
| FISONS | 8MG; 120MG | N018747 | 001 | Mar 06, 1986 |
|--------|------------|---------|-----|--------------|

PSEUDOEPHEDRINE HYDROCHLORIDE AND CHLORPHENIRAMINE MALEATE

| | | | | |
|-------------|------------|---------|-----|--------------|
| CENT PHARMS | 8MG; 120MG | N019428 | 001 | Aug 02, 1988 |
|-------------|------------|---------|-----|--------------|

| | | | | |
|-----------|------------|---------|-----|--------------|
| GRAHAM DM | 8MG; 120MG | N018844 | 001 | Mar 20, 1985 |
|-----------|------------|---------|-----|--------------|

| | | | | |
|--|-------------|---------|-----|--------------|
| | 12MG; 120MG | N018843 | 001 | Mar 18, 1985 |
|--|-------------|---------|-----|--------------|

| | | | | |
|----------|-------------|---------|-----|--------------|
| KV PHARM | 12MG; 120MG | A071455 | 001 | Mar 01, 1989 |
|----------|-------------|---------|-----|--------------|

CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

CHLOR-TRIMETON

| | | | | |
|------------------------|------------|---------|-----|--|
| + BAYER HEALTHCARE LLC | 8MG; 120MG | N018397 | 001 | |
|------------------------|------------|---------|-----|--|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX

SUSPENSION, EXTENDED RELEASE;ORAL

CODEPREX

LANNETT CO INC EQ 4MG MALEATE/5ML;EQ 20MG BASE/5ML N021369 001 Jun 21, 2004

PENNTUSS

FISONS EQ 4MG MALEATE/5ML;EQ 10MG BASE/5ML N018928 001 Aug 14, 1985

CHLORPHENIRAMINE POLISTIREX; HYDROCODONE POLISTIREX

SUSPENSION, EXTENDED RELEASE;ORAL

TUSSIONEX PENNKINETIC

+ UCB INC EQ 8MG MALEATE/5ML;EQ 10MG BITARTRATE/5ML N019111 001 Dec 31, 1987

CHLORPHENTERMINE HYDROCHLORIDE

TABLET;ORAL

PRE-SATE

PARKE DAVIS EQ 65MG BASE N014696 001

CHLORPROMAZINE

SUPPOSITORY;RECTAL

THORAZINE

+ GLAXOSMITHKLINE 25MG ** N009149 024

+ 100MG ** N009149 033

CHLORPROMAZINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

THORAZINE

GLAXOSMITHKLINE 30MG N011120 016

75MG N011120 017

150MG N011120 018

200MG N011120 019

300MG N011120 020

CONCENTRATE;ORAL

CHLORPROMAZINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC 100MG/ML A086863 001

PHARM ASSOC 30MG/ML A040231 001 Dec 30, 1999

100MG/ML A040224 001 Jan 26, 1999

WOCKHARDT 30MG/ML A087032 001 Jul 08, 1982

100MG/ML A087053 001

CHLORPROMAZINE HYDROCHLORIDE INTENSOL

CYCLE PHARMS LTD 30MG/ML A088157 001 Apr 27, 1983

100MG/ML A088158 001 Apr 27, 1983

SONAZINE

FOSUN PHARMA 30MG/ML A080983 004

100MG/ML A080983 005

THORAZINE

+ GLAXOSMITHKLINE 30MG/ML ** N009149 032

+ 100MG/ML ** N009149 043

INJECTABLE;INJECTION

CHLORPROMAZINE HYDROCHLORIDE

ABRAXIS PHARM 25MG/ML A084911 001

MARSAM PHARMS LLC 25MG/ML A089563 001 Apr 15, 1988

WATSON LABS 25MG/ML A080365 001

25MG/ML A085591 001

WYETH AYERST 25MG/ML A080370 001

THORAZINE

+ GLAXOSMITHKLINE 25MG/ML ** N009149 011

SYRUP;ORAL

CHLORPROMAZINE HYDROCHLORIDE

ALPHARMA US PHARMS 10MG/5ML A086712 001

SONAZINE

FOSUN PHARMA 10MG/5ML A083040 001

THORAZINE

+ GLAXOSMITHKLINE 10MG/5ML ** N009149 022

TABLET;ORAL

CHLORPROMAZINE HYDROCHLORIDE

ABBOTT 10MG A084414 001

25MG A084415 001

50MG A084411 001

100MG A084412 001

200MG A084413 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPROMAZINE HYDROCHLORIDE

TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

| | | | | |
|----------------------|----------|---------|-----|--------------|
| CYCLE PHARMS LTD | 10MG | A085331 | 001 | |
| | 25MG | A085331 | 002 | |
| | 50MG | A085331 | 003 | |
| | 100MG | A085331 | 004 | |
| | 200MG | A085331 | 005 | |
| IVAX SUB TEVA PHARMS | 10MG | A083549 | 001 | |
| | 25MG | A083549 | 002 | |
| | 50MG | A083549 | 003 | |
| | 100MG | A083574 | 001 | |
| | 200MG | A083575 | 001 | |
| KV PHARM | 10MG | A085750 | 002 | Jan 04, 1982 |
| | 25MG | A085751 | 001 | |
| | 50MG | A085484 | 001 | |
| | 100MG | A085752 | 001 | |
| | 200MG | A085748 | 002 | Jan 04, 1982 |
| LEDERLE | 10MG | A084803 | 001 | |
| | 25MG | A084801 | 001 | |
| | 50MG | A084800 | 001 | |
| | 100MG | A084789 | 001 | |
| | 200MG | A084802 | 001 | |
| PUREPAC PHARM | 10MG | A080403 | 004 | |
| | 25MG | A080403 | 001 | |
| | 50MG | A080403 | 002 | |
| | 100MG | A080403 | 003 | |
| | 200MG | A080403 | 005 | |
| PVT FORM | 25MG | A080340 | 001 | |
| | 50MG | A080340 | 002 | |
| | 200MG | A080340 | 003 | |
| SANDOZ | 10MG ** | A080439 | 001 | |
| | 25MG ** | A080439 | 002 | |
| | 50MG ** | A080439 | 003 | |
| | 100MG ** | A080439 | 004 | |
| | 200MG ** | A080439 | 005 | |
| VANGARD | 10MG | A088038 | 001 | Aug 16, 1982 |
| | 25MG | A087645 | 001 | |
| | 50MG | A087646 | 001 | |
| WATSON LABS | 10MG | A085959 | 001 | |
| | 25MG | A085956 | 001 | |
| | 50MG | A085960 | 001 | |
| | 100MG | A085957 | 001 | |
| | 200MG | A085958 | 001 | |
| WEST WARD | 10MG | A087783 | 001 | Sep 16, 1982 |
| | 25MG | A087865 | 001 | Sep 16, 1982 |
| | 50MG | A087878 | 001 | Sep 15, 1982 |
| | 100MG | A087884 | 001 | Sep 15, 1982 |
| | 200MG | A087880 | 001 | Sep 16, 1982 |
| PROMAPAR | | | | |
| PARKE DAVIS | 10MG | A086886 | 001 | |
| | 25MG | A084423 | 001 | |
| | 50MG | A086887 | 001 | |
| | 100MG | A086888 | 001 | |
| | 200MG | A086885 | 001 | |
| THORAZINE | | | | |
| GLAXOSMITHKLINE | 10MG ** | N009149 | 002 | |
| | 25MG ** | N009149 | 007 | |
| | 50MG ** | N009149 | 013 | |
| | 100MG ** | N009149 | 018 | |
| | 200MG ** | N009149 | 020 | |

CHLORPROPAMIDE

TABLET; ORAL

CHLORPROPAMIDE

| | | | | |
|----------------|-------|---------|-----|--------------|
| ANI PHARMS INC | 100MG | A088768 | 001 | Oct 11, 1984 |
| | 100MG | A088812 | 001 | Oct 19, 1984 |
| | 100MG | A088840 | 001 | Oct 25, 1984 |
| | 100MG | A088918 | 001 | Oct 16, 1984 |
| | 100MG | A088921 | 001 | Apr 12, 1985 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORPROPAMIDE

TABLET; ORAL

CHLORPROPAMIDE

| | | | |
|---------------------|-------|-------------|--------------|
| | 100MG | A089446 001 | Nov 17, 1986 |
| | 250MG | A087353 001 | |
| | 250MG | A088813 001 | Oct 19, 1984 |
| | 250MG | A088919 001 | Oct 16, 1984 |
| | 250MG | A088922 001 | Apr 12, 1985 |
| | 250MG | A089447 001 | Nov 17, 1986 |
| AUROLIFE PHARMA LLC | 100MG | A088725 001 | Aug 31, 1984 |
| | 250MG | A088726 001 | Aug 31, 1984 |
| DAVA PHARMS INC | 100MG | A089561 001 | Sep 04, 1987 |
| | 250MG | A089562 001 | Sep 04, 1987 |
| HALSEY | 100MG | A089321 001 | Jan 16, 1986 |
| | 250MG | A088662 001 | Jan 09, 1986 |
| PAR PHARM | 100MG | A088175 001 | Feb 27, 1984 |
| | 250MG | A088176 001 | Feb 27, 1984 |
| SANDOZ | 250MG | A084669 001 | |
| SUPERPHARM | 100MG | A088694 001 | Sep 17, 1984 |
| | 250MG | A088695 001 | Sep 17, 1984 |
| USL PHARMA | 100MG | A088708 001 | Aug 30, 1984 |
| | 250MG | A088709 001 | Aug 30, 1984 |
| WATSON LABS | 100MG | A086865 001 | Sep 24, 1984 |
| | 100MG | A088608 001 | Apr 12, 1984 |
| | 250MG | A086866 001 | |
| | 250MG | A088568 001 | Apr 12, 1984 |
| WATSON LABS TEVA | 100MG | A088852 001 | Sep 26, 1984 |
| | 250MG | A088826 001 | Sep 26, 1984 |
| DIABINESE | | | |
| + PFIZER | 100MG | N011641 003 | |
| + | 250MG | N011641 006 | |
| GLUCAMIDE | | | |
| ANI PHARMS INC | 250MG | A088641 001 | Oct 11, 1984 |

CHLORPROTHIXENE

CONCENTRATE; ORAL

TARACTAN

ROCHE

100MG/5ML

N016149 002

INJECTABLE; INJECTION

TARACTAN

ROCHE

12.5MG/ML

N012487 001

TABLET; ORAL

TARACTAN

ROCHE

10MG

N012486 005

25MG

N012486 004

50MG

N012486 003

100MG

N012486 001

CHLORTETRACYCLINE HYDROCHLORIDE

OINTMENT; OPHTHALMIC

AUREOMYCIN

LEDERLE

1%

N050404 001

CHLORTHALIDONE

TABLET; ORAL

CHLORTHALIDONE

ABBOTT

25MG

A087364 001

50MG

A087384 001

ASCOT

25MG

A087698 001 Oct 20, 1982

50MG

A087699 001 Oct 20, 1982

BARR LABS INC

25MG

A088902 001 Sep 19, 1985

50MG

A088903 001 Sep 19, 1985

DAVA PHARMS INC

25MG

A087451 001

50MG

A087450 001

G AND W LABS INC

50MG

A088651 001 May 30, 1985

IVAX PHARMS

25MG

A087555 001

25MG

A088164 001 Jan 09, 1984

50MG

A087176 001

50MG

A087947 001 Feb 27, 1984

KV PHARM

25MG

A087311 001

50MG

A087312 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORTHALIDONE

TABLET; ORAL

CHLORTHALIDONE

| | | | | |
|---------------------|---------|---------|-----|--------------|
| MUTUAL PHARM | 25MG | A087292 | 001 | |
| | 25MG | A089738 | 001 | Sep 19, 1988 |
| | 50MG | A087293 | 001 | |
| | 50MG | A089739 | 001 | Sep 19, 1988 |
| PIONEER PHARMS | 50MG | A089591 | 001 | Jul 21, 1988 |
| PUREPAC PHARM | 25MG | A088139 | 001 | Jul 16, 1986 |
| | 50MG | A088140 | 001 | Aug 11, 1983 |
| SANDOZ | 25MG | A087380 | 001 | |
| | 50MG | A087118 | 001 | |
| | 50MG | A087381 | 001 | |
| SUPERPHARM | 25MG | A087473 | 001 | Feb 09, 1983 |
| | 50MG | A087247 | 001 | Feb 09, 1983 |
| USL PHARMA | 25MG | A089051 | 001 | Jun 01, 1987 |
| | 50MG | A089052 | 001 | Jun 01, 1987 |
| VANGARD | 25MG | A088012 | 001 | Jul 14, 1982 |
| | 50MG | A088073 | 001 | Mar 25, 1983 |
| WARNER CHILCOTT | 25MG | A087515 | 001 | Jan 24, 1983 |
| | 50MG | A087516 | 001 | Feb 09, 1983 |
| WATSON LABS | 25MG | A087050 | 001 | |
| | 25MG | A087100 | 001 | |
| | 25MG | A087296 | 001 | |
| | 25MG | A087706 | 001 | |
| | 50MG | A087029 | 001 | |
| | 50MG | A087082 | 001 | |
| | 50MG | A087521 | 001 | |
| | 50MG | A087689 | 001 | |
| HYGROTON | | | | |
| + SANOFI AVENTIS US | 25MG ** | N012283 | 004 | |
| + | 50MG ** | N012283 | 003 | |
| THALITONE | | | | |
| + CASPER PHARMA LLC | 15MG ** | N019574 | 001 | Dec 20, 1988 |
| | 25MG | N019574 | 002 | Feb 12, 1992 |
| MONARCH PHARMS | 25MG | A088051 | 001 | Nov 12, 1982 |

CHLORTHALIDONE; CLONIDINE HYDROCHLORIDE

TABLET; ORAL

CLONIDINE HYDROCHLORIDE AND CHLORTHALIDONE

| | | | | |
|------------------------|----------------|---------|-----|--------------|
| PAR PHARM | 15MG; 0.1MG | A071179 | 001 | Dec 16, 1987 |
| | 15MG; 0.2MG | A071178 | 001 | Dec 16, 1987 |
| | 15MG; 0.3MG | A071142 | 001 | Dec 16, 1987 |
| CLORPRES | | | | |
| MYLAN | 15MG; 0.1MG | A071325 | 003 | Feb 09, 1987 |
| | 15MG; 0.2MG | A071325 | 002 | Feb 09, 1987 |
| | 15MG; 0.3MG | A071325 | 001 | Feb 09, 1987 |
| COMBIPRES | | | | |
| + BOEHRINGER INGELHEIM | 15MG; 0.1MG ** | N017503 | 001 | |
| + | 15MG; 0.2MG ** | N017503 | 002 | |
| + | 15MG; 0.3MG ** | N017503 | 003 | Apr 10, 1984 |

CHLORTHALIDONE; METOPROLOL TARTRATE

CAPSULE; ORAL

LOPRESSIDONE

| | | | | |
|----------|-------------|---------|-----|--------------|
| NOVARTIS | 25MG; 100MG | N019451 | 001 | Dec 31, 1987 |
| | 25MG; 200MG | N019451 | 002 | Dec 31, 1987 |

CHLORTHALIDONE; RESERPINE

TABLET; ORAL

DEMI-REGROTON

| | | | | |
|-------------------|---------------|---------|-----|--|
| SANOFI AVENTIS US | 25MG; 0.125MG | N015103 | 002 | |
| REGROTON | | | | |
| SANOFI AVENTIS US | 50MG; 0.25MG | N015103 | 001 | |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE

| | | | |
|----------------------|----------|-------------|--------------|
| ACTAVIS ELIZABETH | 250MG | A088928 001 | May 08, 1987 |
| | 500MG | A040113 001 | Sep 29, 1995 |
| AUROLIFE PHARMA LLC | 250MG | A089852 001 | May 04, 1988 |
| | 500MG | A089853 001 | May 04, 1988 |
| BARR | 500MG | A089895 001 | May 04, 1988 |
| OHM LABS | 250MG | A081298 001 | Dec 29, 1993 |
| | 500MG | A081299 001 | Dec 29, 1993 |
| PAR PHARM | 250MG | A087981 001 | Sep 20, 1983 |
| PIONEER PHARMS | 250MG | A089592 001 | Jan 06, 1989 |
| | 500MG | A089948 001 | Jan 06, 1989 |
| SUN PHARM INDUSTRIES | 500MG | A089970 001 | Sep 27, 1990 |
| WATSON LABS | 250MG | A086901 001 | |
| | 250MG | A086948 001 | Aug 09, 1982 |
| | 500MG | A040137 001 | Aug 09, 1996 |
| | 500MG | A081019 001 | Jul 29, 1991 |
| | 500MG | A081040 001 | Aug 22, 1989 |
| PARAFLEX | | | |
| + ORTHO MCNEIL PHARM | 250MG ** | N011300 003 | |
| PARAFON FORTE DSC | | | |
| + JANSSEN R AND D | 500MG ** | N011529 002 | Jun 15, 1987 |
| STRIFON FORTE DSC | | | |
| FERNDALE LABS | 500MG | A081008 001 | Dec 23, 1988 |

CHOLESTYRAMINE

BAR, CHEWABLE; ORAL

CHOLYBAR

| | | | |
|-------------|------------------|-------------|--------------|
| PARKE DAVIS | EQ 4GM RESIN/BAR | A071621 001 | May 26, 1988 |
| | EQ 4GM RESIN/BAR | A071739 001 | May 26, 1988 |

POWDER; ORAL

CHOLESTYRAMINE

| | | | |
|----------------------|--------------------------|-------------|--------------|
| IVAX SUB TEVA PHARMS | EQ 4GM RESIN/PACKET | A074771 001 | Jul 09, 1997 |
| | EQ 4GM RESIN/SCOOPFUL | A074771 002 | Jul 09, 1997 |
| TEVA | EQ 4GM RESIN/PACKET | A074347 001 | May 28, 1998 |
| | EQ 4GM RESIN/SCOOPFUL | A074347 002 | May 28, 1998 |
| CHOLESTYRAMINE LIGHT | | | |
| TEVA | EQ 4GM RESIN/PACKET | A074348 001 | May 28, 1998 |
| | EQ 4GM RESIN/SCOOPFUL | A074348 002 | May 28, 1998 |
| TEVA PHARMS | EQ 4GM RESIN/PACKET | A074555 001 | Sep 30, 1998 |
| | EQ 4GM RESIN/SCOOPFUL | A074555 002 | Sep 30, 1998 |
| LOCHOLEST | | | |
| SANDOZ | EQ 4GM RESIN/PACKET | A074561 001 | Aug 15, 1996 |
| | EQ 4GM RESIN/SCOOPFUL | A074561 002 | Aug 15, 1996 |
| LOCHOLEST LIGHT | | | |
| SANDOZ | EQ 4GM RESIN/PACKET | A074562 001 | Aug 15, 1996 |
| | EQ 4GM RESIN/SCOOPFUL | A074562 002 | Aug 15, 1996 |
| QUESTRAN | | | |
| + BRISTOL MYERS | EQ 4GM RESIN/PACKET ** | N016640 001 | |
| + BRISTOL MYERS | EQ 4GM RESIN/SCOOPFUL ** | N016640 003 | |
| QUESTRAN LIGHT | | | |
| + BRISTOL MYERS | EQ 4GM RESIN/PACKET ** | N019669 001 | Dec 05, 1988 |
| + BRISTOL MYERS | EQ 4GM RESIN/SCOOPFUL ** | N019669 003 | Dec 05, 1988 |

TABLET; ORAL

QUESTRAN

| | | | |
|-----------|----------------|-------------|--------------|
| APOTHECON | EQ 800MG RESIN | A073403 002 | Dec 27, 1999 |
| | EQ 1GM RESIN | A073403 001 | Apr 28, 1994 |

CHORIOGONADOTROPIN ALFA

INJECTABLE; INJECTION

OVIDREL

| | | | |
|------------|-------------|-------------|--------------|
| EMD SERONO | 0.25MG/VIAL | N021149 001 | Sep 20, 2000 |
|------------|-------------|-------------|--------------|

CHROMIC CHLORIDE

INJECTABLE; INJECTION

CHROMIC CHLORIDE

| | | | |
|---------------|------------------------|-------------|--------------|
| ABRAXIS PHARM | EQ 0.004MG CHROMIUM/ML | N019271 001 | May 05, 1987 |
|---------------|------------------------|-------------|--------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CHROMIC PHOSPHATE P-32

INJECTABLE; INJECTION

PHOSPHOCOL P32

MALLINKRODT NUCLEAR 5mCi/ML

N017084 001

CHYMOPAPAIN

INJECTABLE; INJECTION

CHYMODIACTIN

CHART MEDCL 4,000 UNITS/VIAL

N018663 002 Aug 21, 1984

+ 10,000 UNITS/VIAL **

N018663 001 Nov 10, 1982

DISCASE

ABBOTT 12,500 UNITS/VIAL

N018625 001 Jan 18, 1984

CHYMOTRYPSIN

FOR SOLUTION; OPHTHALMIC

ALPHA CHYMAR

SOLA BARNES HIND 750 UNITS/VIAL

N011837 001

CATARASE

CIBA 300 UNITS/VIAL

N016938 001

NOVARTIS 150 UNITS/VIAL

N018121 001

ZOLYSE

ALCON 750 UNITS/VIAL

N011903 001

CICLOPIROX

SOLUTION; TOPICAL

CICLOPIROX

MYLAN PHARMS INC 8%

A078567 001 Sep 18, 2007

TEVA PHARMS 8%

A078079 001 Sep 18, 2007

CIDOFOVIR

INJECTABLE; INJECTION

VISTIDE

+ GILEAD SCIENCES INC EQ 75MG BASE/ML **

N020638 001 Jun 26, 1996

CILASTATIN SODIUM; IMIPENEM

INJECTABLE; INJECTION

PRIMAXIN

MERCK EQ 250MG BASE/VIAL; 250MG/VIAL

A062756 001 Jan 08, 1987

EQ 500MG BASE/VIAL; 500MG/VIAL

A062756 002 Jan 08, 1987

POWDER; INTRAMUSCULAR

PRIMAXIN

MERCK EQ 500MG BASE/VIAL; 500MG/VIAL

N050630 001 Dec 14, 1990

EQ 750MG BASE/VIAL; 750MG/VIAL

N050630 002 Dec 14, 1990

POWDER; INTRAVENOUS

IMIPENEM AND CILASTATIN

HOSPIRA INC EQ 250MG BASE/VIAL; 250MG/VIAL

A090825 001 Nov 16, 2011

PRIMAXIN

+ MERCK EQ 250MG BASE/VIAL; 250MG/VIAL

N050587 001 Nov 26, 1985

CILOSTAZOL

TABLET; ORAL

CILOSTAZOL

ACTAVIS ELIZABETH 100MG

A077028 002 Nov 26, 2004

EPIC PHARMA LLC 50MG

A077150 001 Mar 11, 2005

100MG

A077022 001 Nov 23, 2004

IVAX SUB TEVA PHARMS 100MG

A077020 002 Mar 01, 2005

MYLAN 50MG

A077323 002 Apr 20, 2006

100MG

A077323 001 Apr 20, 2006

MYLAN PHARMS INC 50MG

A077019 001 Nov 23, 2004

100MG

A077019 002 Nov 23, 2004

PLIVA HRVATSKA DOO 50MG

A077898 001 Oct 29, 2007

100MG

A077898 002 Oct 29, 2007

PLETAL

+ OTSUKA 50MG **

N020863 001 Jan 15, 1999

+ 100MG **

N020863 002 Jan 15, 1999

CIMETIDINE

SUSPENSION; ORAL

TAGAMET HB 200

GLAXOSMITHKLINE 200MG/20ML

N020951 001 Jul 09, 1999

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CIMETIDINE

TABLET; ORAL

CIMETIDINE

| | | | | | |
|----------------------|----------|--|---------|-----|--------------|
| CHARTWELL MOLECULES | 200MG | | A074329 | 002 | May 17, 1994 |
| | 300MG | | A074329 | 003 | May 17, 1994 |
| | 400MG | | A074329 | 004 | May 17, 1994 |
| | 800MG | | A074329 | 001 | May 17, 1994 |
| CONTRACT PHARMACAL | 200MG | | A074961 | 001 | Jun 19, 1998 |
| | 200MG | | A074963 | 001 | Jun 19, 1998 |
| CYCLE PHARMS LTD | 300MG | | A074361 | 001 | Dec 23, 1994 |
| | 400MG | | A074361 | 002 | Dec 23, 1994 |
| | 800MG | | A074371 | 001 | Dec 23, 1994 |
| DAVA PHARMS INC | 300MG | | A074340 | 001 | Jun 23, 1995 |
| | 400MG | | A074340 | 002 | Jun 23, 1995 |
| | 800MG | | A074339 | 001 | Jun 23, 1995 |
| IVAX SUB TEVA PHARMS | 200MG | | A074401 | 001 | May 30, 1995 |
| | 200MG | | A074424 | 001 | Jul 28, 1995 |
| | 300MG | | A074401 | 002 | May 30, 1995 |
| | 300MG | | A074424 | 002 | Jul 28, 1995 |
| | 400MG | | A074401 | 003 | May 30, 1995 |
| | 400MG | | A074424 | 003 | Jul 28, 1995 |
| | 800MG | | A074402 | 001 | May 30, 1995 |
| | 800MG | | A074424 | 004 | Jul 28, 1995 |
| PERRIGO | 100MG | | A074972 | 001 | Jun 19, 1998 |
| PLIVA | 200MG | | A074568 | 001 | Feb 27, 1997 |
| | 300MG | | A074568 | 002 | Feb 27, 1997 |
| | 400MG | | A074568 | 003 | Feb 27, 1997 |
| SANDOZ INC | 100MG | | A075122 | 001 | Jun 19, 1998 |
| | 200MG | | A074250 | 001 | Jun 29, 1995 |
| | 200MG | | A075122 | 002 | Jun 19, 1998 |
| | 300MG | | A074250 | 002 | Jun 29, 1995 |
| | 400MG | | A074250 | 003 | Jun 29, 1995 |
| | 800MG | | A074250 | 004 | Jun 29, 1995 |
| TEVA | 200MG | | A074365 | 001 | Feb 28, 1995 |
| | 300MG | | A074365 | 002 | Feb 28, 1995 |
| | 400MG | | A074365 | 003 | Feb 28, 1995 |
| | 800MG | | A074365 | 004 | Feb 28, 1995 |
| UPSHER SMITH LABS | 200MG | | A074506 | 001 | Jan 24, 1996 |
| | 300MG | | A074506 | 002 | Jan 24, 1996 |
| | 400MG | | A074506 | 003 | Jan 24, 1996 |
| | 800MG | | A074506 | 004 | Jan 24, 1996 |
| WATSON LABS INC | 200MG | | A074349 | 001 | Aug 30, 1996 |
| | 300MG | | A074349 | 002 | Aug 30, 1996 |
| | 400MG | | A074349 | 003 | Aug 30, 1996 |
| | 800MG | | A074316 | 001 | Feb 28, 1996 |
| WATSON LABS TEVA | 200MG | | A075425 | 001 | Jul 29, 1999 |
| YAOPHARMA CO LTD | 200MG | | A074100 | 001 | Jan 31, 1995 |
| | 300MG | | A074100 | 002 | Jan 31, 1995 |
| | 400MG | | A074100 | 003 | Jan 31, 1995 |
| | 800MG | | A074100 | 004 | Jan 31, 1995 |
| TAGAMET | | | | | |
| GLAXOSMITHKLINE | 200MG ** | | N017920 | 002 | |
| | 300MG ** | | N017920 | 003 | |
| | 400MG ** | | N017920 | 004 | Dec 14, 1983 |
| | 800MG ** | | N017920 | 005 | Apr 30, 1986 |
| TAGAMET HB | | | | | |
| + MEDTECH PRODUCTS | 100MG ** | | N020238 | 001 | Jun 19, 1995 |

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

| | | | | | |
|--------------------|-------------------|--|---------|-----|--------------|
| HOSPIRA | EQ 300MG BASE/2ML | | A074296 | 001 | Mar 28, 1997 |
| | EQ 300MG BASE/2ML | | A074344 | 001 | Jan 31, 1995 |
| | EQ 300MG BASE/2ML | | A074345 | 001 | Jan 31, 1995 |
| | EQ 300MG BASE/2ML | | A074412 | 001 | Mar 28, 1997 |
| | EQ 300MG BASE/2ML | | A074422 | 001 | Jan 31, 1995 |
| LUITPOLD | EQ 300MG BASE/2ML | | A074353 | 001 | Dec 20, 1994 |
| TEVA PARENTERAL | EQ 300MG BASE/2ML | | A074252 | 001 | Nov 26, 1997 |
| VINTAGE PHARMS LLC | EQ 300MG BASE/2ML | | A074005 | 001 | Aug 31, 1994 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | |
|---------|---------------------|---------|-----|--------------|
| HOSPIRA | EQ 6MG BASE/ML | A074269 | 001 | Dec 27, 1994 |
| | EQ 90MG BASE/100ML | A074468 | 005 | Dec 29, 1994 |
| | EQ 120MG BASE/100ML | A074468 | 006 | Dec 29, 1994 |
| | EQ 180MG BASE/100ML | A074468 | 003 | Dec 29, 1994 |
| | EQ 240MG BASE/100ML | A074468 | 004 | Dec 29, 1994 |
| | EQ 360MG BASE/100ML | A074468 | 001 | Dec 29, 1994 |
| | EQ 480MG BASE/100ML | A074468 | 002 | Dec 29, 1994 |

TAGAMET

| | | | | |
|-----------------|----------------------|---------|-----|--|
| GLAXOSMITHKLINE | EQ 300MG BASE/2ML ** | N017939 | 002 | |
|-----------------|----------------------|---------|-----|--|

TAGAMET HYDROCHLORIDE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | |
|-------------------|-------------------|---------|-----|--------------|
| + GLAXOSMITHKLINE | EQ 6MG BASE/ML ** | N019434 | 001 | Oct 31, 1985 |
|-------------------|-------------------|---------|-----|--------------|

SOLUTION; ORAL

CIMETIDINE HYDROCHLORIDE

| | | | | |
|------------------|-------------------|---------|-----|--------------|
| ANI PHARMS INC | EQ 300MG BASE/5ML | A074859 | 001 | Jul 09, 1998 |
| | EQ 300MG BASE/5ML | A075110 | 001 | Jun 18, 1998 |
| APOTEX INC | EQ 300MG BASE/5ML | A075560 | 001 | Mar 15, 2000 |
| CYCLE PHARMS LTD | EQ 300MG BASE/5ML | A074541 | 001 | Aug 05, 1997 |
| G AND W LABS INC | EQ 300MG BASE/5ML | A074176 | 001 | Jun 01, 1994 |
| LANNETT CO INC | EQ 300MG BASE/5ML | A074251 | 001 | Dec 22, 1994 |

TAGAMET

| | | | | |
|-----------------|----------------------|---------|-----|--|
| GLAXOSMITHKLINE | EQ 300MG BASE/5ML ** | N017924 | 001 | |
|-----------------|----------------------|---------|-----|--|

CINOXACIN

CAPSULE; ORAL

CINOBAC

| | | | | |
|-------|-------|---------|-----|--|
| LILLY | 250MG | N018067 | 001 | |
| | 500MG | N018067 | 002 | |

CINOXACIN

| | | | | |
|------|-------|---------|-----|--------------|
| TEVA | 250MG | A073005 | 001 | Feb 28, 1992 |
| | 500MG | A073006 | 001 | Feb 28, 1992 |

CIPROFLOXACIN

INJECTABLE; INJECTION

CIPRO

| | | | | |
|------------------|---------------------------|---------|-----|--------------|
| + BAYER HLTHCARE | 400MG/40ML (10MG/ML) ** | N019847 | 001 | Dec 26, 1990 |
| + | 200MG/20ML (10MG/ML) ** | N019847 | 002 | Dec 26, 1990 |
| | 1200MG/120ML (10MG/ML) ** | N019847 | 003 | Dec 26, 1990 |

CIPRO IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|------------------|----------------|---------|-----|--------------|
| + BAYER HLTHCARE | 200MG/100ML ** | N019857 | 001 | Dec 26, 1990 |
| + | 400MG/200ML ** | N019857 | 002 | Dec 26, 1990 |

CIPRO IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | | |
|--------------|-------------|---------|-----|--------------|
| BAYER PHARMS | 200MG/100ML | N019858 | 001 | Dec 26, 1990 |
|--------------|-------------|---------|-----|--------------|

CIPROFLOXACIN

| | | | | |
|--------------------|------------------------|---------|-----|--------------|
| BEDFORD LABS | 200MG/20ML (10MG/ML) | A076992 | 001 | Aug 28, 2006 |
| | 400MG/40ML (10MG/ML) | A076992 | 002 | Aug 28, 2006 |
| | 1200MG/120ML (10MG/ML) | A076993 | 001 | Aug 28, 2006 |
| FRESENIUS KABI USA | 200MG/20ML (10MG/ML) | A076484 | 001 | Aug 28, 2006 |
| | 400MG/40ML (10MG/ML) | A076484 | 002 | Aug 28, 2006 |
| TEVA PHARMS USA | 200MG/20ML (10MG/ML) | A077782 | 001 | Aug 28, 2006 |
| | 400MG/40ML (10MG/ML) | A077782 | 002 | Aug 28, 2006 |

CIPROFLOXACIN IN DEXTROSE 5%

| | | | | |
|--------------------|-------------|---------|-----|--------------|
| HIKMA FARMACEUTICA | 200MG/100ML | A076757 | 001 | Apr 21, 2008 |
|--------------------|-------------|---------|-----|--------------|

CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|-----------------|-------------|---------|-----|--------------|
| BAXTER HLTHCARE | 200MG/100ML | A077888 | 001 | Mar 18, 2008 |
| | 400MG/200ML | A077888 | 002 | Mar 18, 2008 |
| BEDFORD | 200MG/100ML | A078114 | 001 | Mar 18, 2008 |
| | 400MG/200ML | A078114 | 002 | Mar 18, 2008 |
| TEVA PHARMS | 200MG/100ML | A077138 | 001 | Mar 18, 2008 |
| | 400MG/200ML | A077138 | 002 | Mar 18, 2008 |

CIPROFLOXACIN HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CIPROFLOXACIN HYDROCHLORIDE

| | | | | |
|---------------|--------------|---------|-----|--------------|
| AMRING PHARMS | EQ 0.3% BASE | A078598 | 001 | Jan 16, 2008 |
| APOTEX INC | EQ 0.3% BASE | A075928 | 001 | Jun 09, 2004 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CIPROFLOXACIN HYDROCHLORIDE

TABLET; ORAL

CIPRO

| | | | | |
|---|----------------|---------------|-------------|--------------|
| + | BAYER HLTHCARE | EQ 100MG BASE | N019537 001 | Apr 08, 1996 |
| + | | EQ 750MG BASE | N019537 004 | Oct 22, 1987 |

CIPROFLOXACIN HYDROCHLORIDE

| | | | |
|--------------------|---------------|-------------|--------------|
| ANI PHARMS INC | EQ 100MG BASE | A075939 001 | Mar 03, 2005 |
| | EQ 250MG BASE | A075939 002 | Jun 09, 2004 |
| | EQ 500MG BASE | A075939 003 | Jun 09, 2004 |
| | EQ 750MG BASE | A075939 004 | Jun 09, 2004 |
| BARR | EQ 250MG BASE | A074124 001 | Jun 09, 2004 |
| | EQ 500MG BASE | A074124 002 | Jun 09, 2004 |
| | EQ 750MG BASE | A074124 003 | Jun 09, 2004 |
| FOSUN PHARMA | EQ 250MG BASE | A076593 002 | Jun 09, 2004 |
| | EQ 500MG BASE | A076593 003 | Jun 09, 2004 |
| | EQ 750MG BASE | A076593 004 | Jun 09, 2004 |
| MYLAN | EQ 100MG BASE | A075817 001 | Jun 25, 2007 |
| | EQ 250MG BASE | A075685 002 | Jun 09, 2004 |
| | EQ 250MG BASE | A075817 002 | Jun 09, 2004 |
| | EQ 500MG BASE | A075685 003 | Jun 09, 2004 |
| | EQ 750MG BASE | A075685 001 | Jun 09, 2004 |
| | EQ 750MG BASE | A075817 004 | Jun 09, 2004 |
| NOSTRUM LABS | EQ 250MG BASE | A076138 001 | Jun 09, 2004 |
| | EQ 500MG BASE | A076138 002 | Jun 09, 2004 |
| | EQ 750MG BASE | A076138 003 | Jun 09, 2004 |
| PLIVA | EQ 100MG BASE | A076426 001 | Jun 15, 2005 |
| | EQ 250MG BASE | A076426 002 | Jun 15, 2005 |
| | EQ 500MG BASE | A076426 003 | Jun 15, 2005 |
| | EQ 750MG BASE | A076426 004 | Jun 15, 2005 |
| SUN PHARM INDS LTD | EQ 250MG BASE | A075747 001 | Jun 09, 2004 |
| | EQ 500MG BASE | A075747 002 | Jun 09, 2004 |
| | EQ 750MG BASE | A075747 003 | Jun 09, 2004 |
| TEVA | EQ 250MG BASE | A076136 001 | Jun 09, 2004 |
| | EQ 500MG BASE | A076136 002 | Jun 09, 2004 |
| | EQ 750MG BASE | A076136 003 | Jun 09, 2004 |

TABLET, EXTENDED RELEASE; ORAL

PROQUIN XR

| | | | |
|-------------|---------------|-------------|--------------|
| DEPOMED INC | EQ 500MG BASE | N021744 001 | May 19, 2005 |
|-------------|---------------|-------------|--------------|

CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

CIPRO XR

| | | | |
|----------------|----------------------------|-------------|--------------|
| BAYER HLTHCARE | 212.6MG;EQ 287.5MG BASE ** | N021473 001 | Dec 13, 2002 |
| | 425.2MG;EQ 574.9MG BASE ** | N021473 002 | Aug 28, 2003 |

CIPROFLOXACIN EXTENDED RELEASE

| | | | |
|---------------------|-------------------------|-------------|--------------|
| ACTAVIS LABS FL INC | 212.6MG;EQ 287.5MG BASE | A077417 001 | Nov 30, 2010 |
| | 425.2MG;EQ 574.9MG BASE | A077809 001 | Nov 30, 2010 |
| DR REDDYS LABS LTD | 212.6MG;EQ 287.5MG BASE | A077701 002 | Oct 31, 2007 |
| FOSUN PHARMA | 212.6MG;EQ 287.5MG BASE | A078712 001 | Dec 11, 2007 |

CISAPRIDE MONOHYDRATE

SUSPENSION; ORAL

PROPULSID

| | | | |
|----------------|----------------|-------------|--------------|
| JANSSEN PHARMS | EQ 1MG BASE/ML | N020398 001 | Sep 15, 1995 |
|----------------|----------------|-------------|--------------|

TABLET; ORAL

PROPULSID

| | | | |
|----------------|--------------|-------------|--------------|
| JANSSEN PHARMS | EQ 10MG BASE | N020210 001 | Jul 29, 1993 |
| | EQ 20MG BASE | N020210 002 | Dec 23, 1993 |

TABLET, ORALLY DISINTEGRATING; ORAL

PROPULSID QUICKSOLV

| | | | |
|----------------|--------------|-------------|--------------|
| JANSSEN PHARMA | EQ 20MG BASE | N020767 001 | Nov 07, 1997 |
|----------------|--------------|-------------|--------------|

CISATRACURIUM BESYLATE

INJECTABLE; INJECTION

CISATRACURIUM BESYLATE

| | | | |
|-------------|-----------------|-------------|--------------|
| HOSPIRA INC | EQ 2MG BASE/ML | A203236 001 | Mar 30, 2018 |
| | EQ 2MG BASE/ML | A203238 001 | Mar 30, 2018 |
| | EQ 10MG BASE/ML | A203236 002 | Mar 30, 2018 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CISPLATIN

INJECTABLE; INJECTION

CISPLATIN

| | | | |
|-------------------|-----------|-------------|--------------|
| BEDFORD | 10MG/VIAL | A074713 001 | Nov 14, 2000 |
| | 50MG/VIAL | A074713 002 | Nov 14, 2000 |
| TEVA PHARMS USA | 1MG/ML | A074814 001 | May 16, 2000 |
| PLATINOL | | | |
| + HQ SPCLT PHARMA | 10MG/VIAL | N018057 001 | |
| + | 50MG/VIAL | N018057 002 | |
| PLATINOL-AQ | | | |
| + HQ SPCLT PHARMA | 0.5MG/ML | N018057 003 | Jul 18, 1984 |

CITALOPRAM HYDROBROMIDE

CAPSULE; ORAL

CITALOPRAM HYDROBROMIDE

| | | | |
|------------------|--------------|-------------|--------------|
| MYLAN PHARMS INC | EQ 10MG BASE | A077668 001 | Feb 28, 2007 |
| | EQ 20MG BASE | A077668 002 | Feb 28, 2007 |
| | EQ 40MG BASE | A077668 003 | Feb 28, 2007 |

SOLUTION; ORAL

CELEXA

| | | | |
|-------------------------|---------------------|-------------|--------------|
| + FOREST LABS | EQ 10MG BASE/5ML ** | N021046 001 | Dec 22, 1999 |
| CITALOPRAM HYDROBROMIDE | | | |
| APOTEX INC | EQ 10MG BASE/5ML | A077601 001 | Nov 15, 2005 |

TABLET; ORAL

CELEXA

| | | | |
|-------------------------|--------------|-------------|--------------|
| ALLERGAN SALES LLC | EQ 60MG BASE | N020822 004 | Jul 17, 1998 |
| CITALOPRAM HYDROBROMIDE | | | |
| ACTAVIS ELIZABETH | EQ 10MG BASE | A077033 001 | Oct 28, 2004 |
| | EQ 20MG BASE | A077033 002 | Oct 28, 2004 |
| | EQ 40MG BASE | A077033 003 | Oct 28, 2004 |
| EPIC PHARMA LLC | EQ 10MG BASE | A077036 001 | Oct 28, 2004 |
| | EQ 20MG BASE | A077036 002 | Oct 28, 2004 |
| | EQ 40MG BASE | A077036 003 | Oct 28, 2004 |
| FOSUN PHARMA | EQ 10MG BASE | A077035 001 | Oct 28, 2004 |
| | EQ 10MG BASE | A077040 001 | Aug 17, 2005 |
| | EQ 20MG BASE | A077035 002 | Oct 28, 2004 |
| | EQ 20MG BASE | A077040 002 | Aug 17, 2005 |
| | EQ 40MG BASE | A077035 003 | Oct 28, 2004 |
| | EQ 40MG BASE | A077040 003 | Aug 17, 2005 |
| MYLAN | EQ 10MG BASE | A077039 001 | Feb 03, 2005 |
| | EQ 20MG BASE | A077039 002 | Feb 03, 2005 |
| | EQ 40MG BASE | A077039 003 | Feb 03, 2005 |
| MYLAN PHARMS INC | EQ 10MG BASE | A077037 001 | Nov 05, 2004 |
| | EQ 20MG BASE | A077037 002 | Nov 05, 2004 |
| | EQ 40MG BASE | A077037 003 | Nov 05, 2004 |
| NATCO PHARMA LTD | EQ 20MG BASE | A077141 002 | Apr 10, 2008 |
| | EQ 40MG BASE | A077141 001 | Apr 10, 2008 |
| ROXANE | EQ 10MG BASE | A077041 001 | Nov 23, 2004 |
| | EQ 20MG BASE | A077041 002 | Nov 23, 2004 |
| | EQ 40MG BASE | A077041 003 | Nov 23, 2004 |
| SUN PHARM INDUSTRIES | EQ 10MG BASE | A077052 001 | Jul 03, 2006 |
| | EQ 20MG BASE | A077052 002 | Jul 03, 2006 |
| | EQ 40MG BASE | A077052 003 | Jul 03, 2006 |
| TARO | EQ 10MG BASE | A077278 001 | Mar 22, 2006 |
| | EQ 20MG BASE | A077278 002 | Mar 22, 2006 |
| | EQ 40MG BASE | A077278 003 | Mar 22, 2006 |
| TEVA PHARMS | EQ 10MG BASE | A077213 001 | Mar 31, 2006 |
| | EQ 20MG BASE | A077213 002 | Mar 31, 2006 |
| | EQ 40MG BASE | A077213 003 | Mar 31, 2006 |
| WATSON LABS | EQ 10MG BASE | A077034 001 | Jun 30, 2005 |
| | EQ 20MG BASE | A077034 002 | Jun 30, 2005 |
| | EQ 40MG BASE | A077034 003 | Jun 30, 2005 |

TABLET, ORALLY DISINTEGRATING; ORAL

CITALOPRAM HYDROBROMIDE

| | | | |
|-------------------|--------------|-------------|--------------|
| BIOVAIL LABS INTL | EQ 10MG BASE | N021763 001 | Dec 20, 2005 |
| | EQ 20MG BASE | N021763 002 | Dec 20, 2005 |
| | EQ 40MG BASE | N021763 003 | Dec 20, 2005 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CITRIC ACID; MAGNESIUM OXIDE; SODIUM CARBONATE

SOLUTION;IRRIGATION

IRRIGATING SOLUTION G IN PLASTIC CONTAINER

BAXTER HLTHCARE 3.24GM/100ML;380MG/100ML;430MG/100ML N018519 001 Jun 22, 1982

UROLOGIC G IN PLASTIC CONTAINER

HOSPIRA 3.24GM/100ML;380MG/100ML;430MG/100ML N018904 001 May 27, 1983

CLADRIBINE

INJECTABLE;INJECTION

LEUSTATIN

+ JANSSEN PHARMS 1MG/ML ** N020229 001 Feb 26, 1993

CLARITHROMYCIN

FOR SUSPENSION;ORAL

BIAXIN

+ ABBVIE 125MG/5ML N050698 001 Dec 23, 1993

187MG/5ML N050698 003 Sep 30, 1998

+ 250MG/5ML N050698 002 Dec 23, 1993

CLARITHROMYCIN

SUN PHARM INDS LTD 125MG/5ML A065382 001 Aug 30, 2007

250MG/5ML A065382 002 Aug 30, 2007

TABLET;ORAL

BIAXIN

+ ABBVIE 250MG ** N050662 001 Oct 31, 1991

+ 500MG ** N050662 002 Oct 31, 1991

CLARITHROMYCIN

IVAX SUB TEVA PHARMS 250MG A065137 001 May 31, 2005

500MG A065137 002 May 31, 2005

MYLAN 250MG A065195 001 Mar 11, 2005

500MG A065195 002 Mar 11, 2005

SUN PHARM INDS LTD 250MG A065174 001 Sep 24, 2004

500MG A065174 002 Sep 24, 2004

TABLET, EXTENDED RELEASE;ORAL

BIAXIN XL

+ ABBVIE 500MG ** N050775 001 Mar 03, 2000

CLARITHROMYCIN

ANI PHARMS INC 500MG A065250 001 Aug 25, 2005

RANBAXY 1GM A065210 001 Jan 26, 2005

CLAVULANATE POTASSIUM; TICARCILLIN DISODIUM

INJECTABLE;INJECTION

TIMENTIN

GLAXOSMITHKLINE EQ 100MG BASE/VIAL;EQ 3GM BASE/VIAL A062691 001 Dec 19, 1986

EQ 100MG BASE/VIAL;EQ 3GM BASE/VIAL N050590 001 Apr 01, 1985

EQ 200MG BASE/VIAL;EQ 3GM BASE/VIAL N050590 002 Apr 01, 1985

EQ 1GM BASE/VIAL;EQ 30GM BASE/VIAL N050590 003 Aug 18, 1987

TIMENTIN IN PLASTIC CONTAINER

GLAXOSMITHKLINE EQ 100MG BASE/100ML;EQ 3GM BASE/100ML N050658 001 Dec 15, 1989

CLEMASTINE FUMARATE

SYRUP;ORAL

CLEMASTINE FUMARATE

ACTAVIS MID ATLANTIC EQ 0.5MG BASE/5ML A074075 001 Oct 31, 1993

APOTEX INC EQ 0.5MG BASE/5ML A075703 001 Nov 27, 2000

LANNETT CO INC EQ 0.5MG BASE/5ML A074884 001 Dec 17, 1997

TEVA PHARMS EQ 0.5MG BASE/5ML A073095 001 Apr 21, 1992

WOCKHARDT BIO AG EQ 0.5MG BASE/5ML A074863 001 Mar 13, 1998

TAVIST

+ NOVARTIS EQ 0.5MG BASE/5ML ** N018675 001 Jun 28, 1985

TABLET;ORAL

CLEMASTINE FUMARATE

ANI PHARMS INC 1.34MG A073282 001 Jan 31, 1992

1.34MG A073282 002 Dec 03, 1992

PLD ACQUISITIONS LLC 1.34MG A073458 001 Oct 31, 1993

SANDOZ 2.68MG A073459 001 Oct 31, 1993

TAVIST

+ NOVARTIS 2.68MG N017661 001

TAVIST-1

+ GLAXOSMITHKLINE CONS 1.34MG N020925 001 Aug 21, 1992

NOVARTIS 1.34MG N017661 002

1.34MG N017661 003 Aug 21, 1992

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLEVIDIPINE

EMULSION; INTRAVENOUS

CLEVIPREX

+ CHIESI USA INC 125MG/250ML (0.5MG/ML) N022156 003 Nov 08, 2013

CLIDINIUM BROMIDE

CAPSULE; ORAL

QUARZAN

ROCHE 2.5MG N010355 001
5MG N010355 002CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL

CLEOCIN

PHARMACIA AND UPJOHN EQ 75MG BASE A061809 001
EQ 150MG BASE A061809 002

CLINDAMYCIN HYDROCHLORIDE

MYLAN PHARMS INC EQ 75MG BASE A091225 001 May 31, 2011
EQ 150MG BASE A091225 002 May 31, 2011
EQ 300MG BASE A091225 003 May 31, 2011
TEVA EQ 75MG BASE A063027 001 Sep 20, 1989
WATSON LABS EQ 75MG BASE A063082 001 Jul 31, 1991CLINDAMYCIN PALMITATE HYDROCHLORIDE

FOR SOLUTION; ORAL

CLEOCIN

PHARMACIA AND UPJOHN EQ 75MG BASE/5ML ** A061827 001

CLINDAMYCIN PHOSPHATE

CREAM; VAGINAL

CLEOCIN

PHARMACIA AND UPJOHN EQ 2% BASE N050680 001 Aug 11, 1992

INJECTABLE; INJECTION

CLEOCIN PHOSPHATE

PHARMACIA AND UPJOHN EQ 150MG BASE/ML A061839 001

CLINDAMYCIN PHOSPHATE

ABRAXIS PHARM EQ 150MG BASE/ML A062747 001 Jun 03, 1988
BEDFORD EQ 150MG BASE/ML A063163 001 Jun 30, 1994
BRISTOL MYERS SQUIBB EQ 150MG BASE/ML A062908 001 Feb 01, 1989
IGI LABS INC EQ 150MG BASE/ML A062928 001 Feb 13, 1989
LOCH EQ 150MG BASE/ML A062905 001 May 09, 1988
MARSAM PHARMS LLC EQ 150MG BASE/ML A062913 001 Oct 20, 1988
SOLOPAK EQ 150MG BASE/ML A062819 001 Mar 15, 1988
EQ 150MG BASE/ML A062852 001 Mar 17, 1988
TEVA PARENTERAL EQ 150MG BASE/ML A063041 001 Dec 29, 1989
EQ 150MG BASE/ML A063282 001 May 29, 1992
WATSON LABS EQ 150MG BASE/ML A062900 001 Jun 08, 1988
EQ 150MG BASE/ML A063079 001 Mar 05, 1990
WEST-WARD PHARMS INT EQ 150MG BASE/ML A062806 001 Oct 15, 1987
EQ 150MG BASE/ML A062953 001 Apr 21, 1988
EQ 150MG BASE/ML A063068 001 Aug 28, 1989

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%

ABRAXIS PHARM EQ 12MG BASE/ML N050636 001 Dec 22, 1989

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER

ABBOTT LABS EQ 6MG BASE/ML A065027 001 Jun 29, 2001
EQ 12MG BASE/ML A065027 002 Jun 29, 2001
EQ 18MG BASE/ML A065027 003 Jun 29, 2001
BAXTER HLTHCARE EQ 6MG BASE/ML N050648 001 Dec 29, 1989
EQ 12MG BASE/ML N050648 002 Dec 29, 1989
EQ 900MG BASE/100ML N050648 003 Dec 29, 1989

SOLUTION; TOPICAL

CLEOCIN T

PHARMACIA AND UPJOHN EQ 1% BASE A062363 001 Feb 08, 1982

CLINDAMYCIN PHOSPHATE

BOCA PHARMA LLC EQ 1% BASE A062944 001 Jan 11, 1989
NOVAST LABS EQ 1% BASE A064108 001 Sep 27, 1996
VINTAGE PHARMS EQ 1% BASE A062930 001 Jun 28, 1989
WOCKHARDT BIO AG EQ 1% BASE A063304 001 Jul 15, 1997

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLIOQUINOL; NYSTATINOINTMENT; TOPICAL
NYSTAFORM

BAYER PHARMS 10MG/GM; 100,000 UNITS/GM N050235 001

CLOBAZAMTABLET; ORAL
ONFI

LUNDBECK PHARMS LLC 5MG N202067 001 Oct 21, 2011

CLOBETASOL PROPIONATE

CREAM; TOPICAL

CLOBETASOL PROPIONATE

TEVA PHARMS USA 0.05% A074087 001 Feb 16, 1994

CLOBETASOL PROPIONATE (EMOLLIENT)

NOVAST LABS 0.05% A075733 001 Aug 22, 2001

TEMOVATE

+ FOUGERA PHARMS 0.05% ** N019322 001 Dec 27, 1985

TEMOVATE E

+ FOUGERA PHARMS 0.05% ** N020340 001 Jun 17, 1994

GEL; TOPICAL

TEMOVATE

+ FOUGERA PHARMS 0.05% ** N020337 001 Apr 29, 1994

OINTMENT; TOPICAL

CLOBETASOL PROPIONATE

ACTAVIS MID ATLANTIC 0.05% A074128 001 Aug 03, 1994

TEMOVATE

+ FOUGERA PHARMS 0.05% ** N019323 001 Dec 27, 1985

SOLUTION; TOPICAL

TEMOVATE

+ FOUGERA PHARMS 0.05% ** N019966 001 Feb 22, 1990

SPRAY; TOPICAL

CLOBETASOL PROPIONATE

APOTEX INC 0.05% A210446 001 Apr 17, 2018

CLOFAZIMINE

CAPSULE; ORAL

LAMPRENE

+ NOVARTIS 50MG N019500 002 Dec 15, 1986

100MG N019500 001 Dec 15, 1986

CLOFIBRATE

CAPSULE; ORAL

ATROMID-S

WYETH AYERST 500MG N016099 002

CLOFIBRATE

BANNER PHARMACAPS 500MG A073396 001 Mar 20, 1992

SANDOZ 500MG A072191 001 May 02, 1988

TEVA 500MG A072600 001 Jul 25, 1991

USL PHARMA 500MG A070531 001 Jun 16, 1986

WATSON LABS 500MG A071603 001 Sep 18, 1987

CLOMIPHENE CITRATE

TABLET; ORAL

CLOMID

+ SANOFI AVENTIS US 50MG N016131 002

MILOPHENE

MILEX 50MG A072196 001 Dec 20, 1988

SEROPHENE

EMD SERONO 50MG N018361 001 Mar 22, 1982

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

CLOMIPRAMINE HYDROCHLORIDE

TEVA 25MG A074849 001 Apr 04, 1997

50MG A074849 002 Apr 04, 1997

75MG A074849 003 Apr 04, 1997

WATSON LABS 25MG A074600 001 Nov 27, 1996

25MG A074751 001 Sep 30, 1998

50MG A074600 002 Nov 27, 1996

50MG A074751 002 Sep 30, 1998

75MG A074600 003 Nov 27, 1996

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE;ORAL

CLOMIPRAMINE HYDROCHLORIDE

75MG

A074751 003 Sep 30, 1998

CLONAZEPAM

TABLET;ORAL

CLONAZEPAM

APOTEX INC

0.5MG

A075468 001 Oct 06, 2000

1MG

A075468 002 Oct 06, 2000

2MG

A075468 003 Oct 06, 2000

MYLAN PHARMS INC

0.5MG

A074940 001 Oct 30, 1997

1MG

A074940 002 Oct 30, 1997

2MG

A074940 003 Oct 30, 1997

SANDOZ

0.5MG

A074925 001 Sep 30, 1997

1MG

A074925 002 Sep 30, 1997

2MG

A074925 003 Sep 30, 1997

TEVA

0.5MG

A074920 001 Aug 04, 1998

1MG

A074920 002 Aug 04, 1998

2MG

A074920 003 Aug 04, 1998

KLONOPIN

ROCHE

0.125MG

N017533 005 Apr 09, 1997

0.25MG

N017533 006 Apr 09, 1997

TABLET, ORALLY DISINTEGRATING;ORAL

KLONOPIN RAPIDLY DISINTEGRATING

+ ROCHE

0.125MG **

N020813 001 Dec 23, 1997

+

0.25MG **

N020813 002 Dec 23, 1997

+

0.5MG **

N020813 003 Dec 23, 1997

+

1MG **

N020813 004 Dec 23, 1997

+

2MG **

N020813 005 Dec 23, 1997

CLONIDINE

SUSPENSION, EXTENDED RELEASE;ORAL

CLONIDINE

TRIS PHARMA INC

EQ 0.09MG BASE/ML

N022499 001 Dec 03, 2009

TABLET, EXTENDED RELEASE;ORAL

CLONIDINE

TRIS PHARMA INC

EQ 0.17MG BASE

N022500 001 Dec 03, 2009

EQ 0.26MG BASE

N022500 002 Dec 03, 2009

CLONIDINE HYDROCHLORIDE

TABLET;ORAL

CLONIDINE HYDROCHLORIDE

AM THERAP

0.1MG

A070881 001 Jul 08, 1986

0.2MG

A070882 001 Jul 08, 1986

0.3MG

A070883 001 Jul 08, 1986

AUROLIFE PHARMA LLC

0.1MG

A070886 002 Aug 31, 1988

0.2MG

A070886 001 Aug 31, 1988

0.3MG

A070886 003 Aug 31, 1988

CHARTWELL MOLECULES

0.1MG

A071785 002 Apr 05, 1988

0.2MG

A071785 003 Apr 05, 1988

0.3MG

A071785 001 Apr 05, 1988

DURAMED PHARMS BARR

0.1MG

A071103 001 Aug 14, 1986

0.2MG

A071102 001 Aug 14, 1986

0.3MG

A071101 001 Aug 14, 1986

INTERPHARM

0.1MG

A071252 001 Oct 01, 1986

0.2MG

A071253 001 Oct 01, 1986

0.3MG

A071254 001 Oct 01, 1986

PAR PHARM

0.1MG

A070461 001 Jul 08, 1986

0.2MG

A070460 001 Jul 08, 1986

0.3MG

A070459 001 Jul 08, 1986

TEVA

0.1MG

A070747 001 Jul 08, 1986

0.2MG

A070702 001 Jul 08, 1986

0.3MG

A070659 001 Jul 08, 1986

WARNER CHILCOTT

0.1MG

A072138 001 Jun 13, 1988

0.2MG

A072139 001 Jun 13, 1988

0.3MG

A072140 001 Jun 13, 1988

WATSON LABS

0.1MG

A070395 001 Mar 23, 1987

0.1MG

A070965 001 Jul 08, 1986

0.2MG

A070396 001 Mar 23, 1987

0.2MG

A070964 001 Jul 08, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLONIDINE HYDROCHLORIDE

TABLET;ORAL

CLONIDINE HYDROCHLORIDE

0.3MG

A070397 001 Mar 23, 1987

0.3MG

A070963 001 Jul 08, 1986

TABLET, EXTENDED RELEASE;ORAL

CLONIDINE HYDROCHLORIDE

ACTAVIS ELIZABETH

0.2MG

A202792 002 May 15, 2015

0.2MG

A203320 002 May 15, 2015

ANCHEN PHARMS

0.1MG

A202983 001 Apr 02, 2014

0.2MG

A202983 002 Apr 02, 2014

0.2MG

A202984 002 Sep 30, 2013

JENLOGA

+ CONCORDIA PHARMS INC

0.1MG **

N022331 001 Sep 30, 2009

+

0.2MG **

N022331 002 May 25, 2010

KAPVAY

+ CONCORDIA PHARMS INC

0.2MG **

N022331 004 Sep 28, 2010

CLOPIDOGREL BISULFATE

TABLET;ORAL

CLOPIDOGREL BISULFATE

ACTAVIS TOTOWA

EQ 75MG BASE

A090307 001 May 28, 2013

CLORAZEPATE DIPOTASSIUM

CAPSULE;ORAL

CLORAZEPATE DIPOTASSIUM

ABLE

3.75MG

A071777 001 Jul 14, 1987

7.5MG

A071778 001 Jul 14, 1987

15MG

A071779 001 Jul 14, 1987

AM THERAP

3.75MG

A071429 001 Jun 23, 1987

7.5MG

A071430 001 Jun 23, 1987

15MG

A071431 001 Jun 23, 1987

AUROLIFE PHARMA LLC

3.75MG

A072112 002 Aug 11, 2017

7.5MG

A072112 003 Aug 11, 2017

15MG

A072112 001 Aug 26, 1988

DAVA PHARMS INC

3.75MG

A071742 001 Dec 14, 1987

7.5MG

A071743 001 Dec 14, 1987

15MG

A071744 001 Dec 14, 1987

GD SEARLE LLC

3.75MG

A071727 001 Dec 18, 1987

7.5MG

A071728 001 Dec 18, 1987

15MG

A071729 001 Dec 18, 1987

MYLAN

3.75MG

A071509 001 Oct 19, 1987

7.5MG

A071510 001 Oct 19, 1987

15MG

A071511 001 Oct 19, 1987

PUREPAC PHARM

3.75MG

A071924 001 Apr 25, 1988

7.5MG

A071925 001 Apr 25, 1988

15MG

A071926 001 Apr 25, 1988

QUANTUM PHARMICS

3.75MG

A071549 001 Sep 12, 1988

7.5MG

A071550 001 Sep 12, 1988

15MG

A071522 001 Sep 12, 1988

USL PHARMA

3.75MG

A071242 001 Jun 23, 1987

7.5MG

A071243 001 Jun 23, 1987

15MG

A071244 001 Jun 23, 1987

WARNER CHILCOTT

3.75MG

A071774 001 Mar 01, 1988

7.5MG

A071775 001 Mar 01, 1988

15MG

A071776 001 Mar 01, 1988

WATSON LABS

3.75MG

A071878 001 Mar 15, 1988

7.5MG

A071879 001 Mar 15, 1988

15MG

A071860 001 Mar 15, 1988

TRANXENE

RECORDATI RARE

3.75MG **

N017105 001

7.5MG **

N017105 002

15MG **

N017105 003

TABLET;ORAL

CLORAZEPATE DIPOTASSIUM

ABLE

3.75MG

A071780 001 Jun 26, 1987

7.5MG

A071781 001 Jun 26, 1987

15MG

A071782 001 Jun 26, 1987

AM THERAP

3.75MG

A071747 001 Jun 23, 1987

7.5MG

A071748 001 Jun 23, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLORAZEPATE DIPOTASSIUM

TABLET;ORAL

CLORAZEPATE DIPOTASSIUM

| | | | | | |
|---------------------|------------|--|---------|-----|--------------|
| | 15MG | | A071749 | 001 | Jun 23, 1987 |
| AUROLIFE PHARMA LLC | 3.75MG | | A072512 | 001 | May 11, 1990 |
| | 7.5MG | | A072513 | 001 | May 11, 1990 |
| | 15MG | | A072514 | 001 | May 11, 1990 |
| LEDERLE | 3.75MG | | A072013 | 001 | Dec 15, 1987 |
| | 7.5MG | | A072014 | 001 | Dec 15, 1987 |
| | 15MG | | A072015 | 001 | Dec 15, 1987 |
| PUREPAC PHARM | 3.75MG | | A072330 | 001 | Aug 08, 1988 |
| | 7.5MG | | A072331 | 001 | Aug 08, 1988 |
| | 15MG | | A072332 | 001 | Aug 08, 1988 |
| QUANTUM PHARMICS | 3.75MG | | A071730 | 001 | Oct 26, 1987 |
| | 7.5MG | | A071731 | 001 | Oct 26, 1987 |
| | 15MG | | A071702 | 001 | Oct 26, 1987 |
| SUN PHARM INDS LTD | 3.75MG | | A076911 | 001 | Sep 29, 2004 |
| | 7.5MG | | A076911 | 002 | Sep 29, 2004 |
| | 15MG | | A076911 | 003 | Sep 29, 2004 |
| WARNER CHILCOTT | 3.75MG | | A071828 | 001 | Mar 03, 1988 |
| | 7.5MG | | A071829 | 001 | Mar 03, 1988 |
| | 15MG | | A071830 | 001 | Mar 03, 1988 |
| WATSON LABS | 3.75MG | | A071852 | 001 | Feb 09, 1988 |
| | 7.5MG | | A071853 | 001 | Feb 09, 1988 |
| | 15MG | | A071854 | 001 | Feb 09, 1988 |
| TRANXENE | | | | | |
| RECORDATI RARE | 3.75MG ** | | N017105 | 006 | |
| + | 15MG ** | | N017105 | 008 | |
| TRANXENE SD | | | | | |
| RECORDATI RARE | 11.25MG ** | | N017105 | 005 | |
| | 22.5MG ** | | N017105 | 004 | |

CLOTRIMAZOLE

CREAM;TOPICAL

LOTRIMIN

SCHERING PLOUGH 1% ** N017619 001

MYCELEX

BAYER HEALTHCARE LLC 1% N018183 001

CREAM;VAGINAL

GYNE-LOTRIMIN

+ BAYER HEALTHCARE LLC 1% ** N018052 002 Nov 30, 1990

GYNE-LOTRIMIN 3

+ BAYER HEALTHCARE LLC 2% N020574 001 Nov 24, 1998

MYCELEX-7

BAYER HEALTHCARE LLC 1% N018230 002 Dec 26, 1991

CREAM, TABLET;TOPICAL, VAGINAL

GYNE-LOTRIMIN 3 COMBINATION PACK

+ BAYER HEALTHCARE LLC 1%,200MG N020526 002 Jul 29, 1996

GYNE-LOTRIMIN COMBINATION PACK

+ BAYER HEALTHCARE LLC 1%,100MG N020289 002 Apr 26, 1993

MYCELEX-7 COMBINATION PACK

BAYER HEALTHCARE LLC 1%,100MG N020389 002 Jun 23, 1994

LOTION;TOPICAL

LOTRIMIN

SCHERING 1% N018813 001 Feb 17, 1984

SOLUTION;TOPICAL

LOTRIMIN

+ SCHERING PLOUGH 1% N017613 001

MYCELEX

+ BAYER HLTHCARE 1% ** N018181 001

TABLET;VAGINAL

GYNE-LOTRIMIN

+ BAYER HEALTHCARE LLC 100MG N017717 002 Nov 30, 1990

GYNE-LOTRIMIN 3

+ BAYER HEALTHCARE LLC 200MG N020525 001 Jul 29, 1996

GYNIX

TEVA PHARMS 100MG A073249 001 Feb 13, 1998

MYCELEX-7

BAYER HEALTHCARE LLC 100MG N018182 002 Dec 26, 1991

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CLOTRIMAZOLE

TABLET;VAGINAL

MYCELEX-G

BAYER PHARMS

500MG

N019069 001 Apr 19, 1985

TROCHE/LOZENGE;ORAL

MYCELEX

+ BAYER HLTHCARE

10MG **

N018713 001 Jun 17, 1983

CLOXACILLIN SODIUM

CAPSULE;ORAL

CLOXACILLIN SODIUM

APOTHECON

EQ 250MG BASE

A061452 001

EQ 500MG BASE

A061452 002

TEVA

EQ 250MG BASE

A062240 001

EQ 500MG BASE

A062240 002

CLOXAPEN

GLAXOSMITHKLINE

EQ 250MG BASE

A061806 001

EQ 250MG BASE

A062233 001

EQ 500MG BASE

A061806 002

EQ 500MG BASE

A062233 002

FOR SOLUTION;ORAL

CLOXACILLIN SODIUM

TEVA

EQ 125MG BASE/5ML

A062268 001

EQ 125MG BASE/5ML

A062978 001 Apr 06, 1989

TEGOPEN

APOTHECON

EQ 125MG BASE/5ML

A061453 001

EQ 125MG BASE/5ML

N050192 001

CLOZAPINE

TABLET;ORAL

CLOZAPINE

MYLAN

12.5MG

A075417 003 Apr 15, 2010

PAR PHARM

25MG

A075162 001 Apr 26, 2005

100MG

A075162 002 Apr 26, 2005

SANDOZ

25MG

A074546 001 Aug 30, 1996

100MG

A074546 002 Aug 30, 1996

ZYDUS PHARMS USA INC

25MG

A209480 001 Dec 06, 2017

50MG

A209480 002 Dec 06, 2017

100MG

A209480 003 Dec 06, 2017

200MG

A209480 004 Dec 06, 2017

TABLET, ORALLY DISINTEGRATING;ORAL

FAZACLO ODT

JAZZ PHARMS III

50MG

N021590 003 Jun 03, 2005

COBALT CHLORIDE CO-57; CYANOCOBALAMIN; CYANOCOBALAMIN CO-57; INTRINSIC FACTOR

N/A;N/A

RUBRATOPE-57 KIT

BRACCO

N/A;N/A;N/A;N/A

N016089 001

COBALT CHLORIDE CO-60; CYANOCOBALAMIN; CYANOCOBALAMIN CO-60; INTRINSIC FACTOR

N/A;N/A

RUBRATOPE-60 KIT

BRACCO

N/A;N/A;N/A;N/A

N016090 001

CODEINE PHOSPHATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PHENERGAN VC W/ CODEINE

+ ANI PHARMS

10MG/5ML;5MG/5ML;6.25MG/5ML **

N008306 005 Apr 02, 1984

PHERAZINE VC W/ CODEINE

HALSEY

10MG/5ML;5MG/5ML;6.25MG/5ML

A088870 001 Mar 02, 1987

PROMETHAZINE VC W/ CODEINE

CENCI

10MG/5ML;5MG/5ML;6.25MG/5ML

A088816 001 Nov 22, 1985

WOCKHARDT

10MG/5ML;5MG/5ML;6.25MG/5ML

A088896 001 Jan 04, 1985

CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PHENERGAN W/ CODEINE

+ ANI PHARMS

10MG/5ML;6.25MG/5ML **

N008306 004 Apr 02, 1984

PHERAZINE W/ CODEINE

HALSEY

10MG/5ML;6.25MG/5ML

A088739 001 Dec 23, 1988

PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE

PHARM ASSOC

10MG/5ML;6.25MG/5ML

A089647 001 Dec 22, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PROMETHAZINE W/ CODEINE

CENCI

10MG/5ML;6.25MG/5ML

A088814 001 Nov 22, 1985

CODEINE PHOSPHATE; PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP;ORAL

ACTIFED W/ CODEINE

GLAXOSMITHKLINE

10MG/5ML;30MG/5ML;1.25MG/5ML

N012575 003 Apr 04, 1984

TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES W/ CODEINE

CENCI

10MG/5ML;30MG/5ML;1.25MG/5ML

A089018 001 Jul 23, 1986

TRIPROLIDINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE AND CODEINE PHOSPHATE

WOCKHARDT

10MG/5ML;30MG/5ML;1.25MG/5ML

A088833 001 Nov 16, 1984

CODEINE SULFATE

SOLUTION;ORAL

CODEINE SULFATE

WEST-WARD PHARMS INT 30MG/5ML

N202245 001 Jun 30, 2011

COLCHICINE; PROBENECID

TABLET;ORAL

COLBENEMID

+ MERCK

0.5MG;500MG **

N012383 001

PROBEN-C

WATSON LABS

0.5MG;500MG

A085552 001

PROBENECID AND COLCHICINE

ANI PHARMS INC

0.5MG;500MG

A083734 001

BEECHAM

0.5MG;500MG

A084321 001

IMPAX LABS

0.5MG;500MG

A083720 002

SANDOZ

0.5MG;500MG

A086130 001

PROBENECID W/ COLCHICINE

LEDERLE

0.5MG;500MG

A086954 001

WATSON LABS

0.5MG;500MG

A083221 001

COLESEVELAM HYDROCHLORIDE

CAPSULE;ORAL

WELCHOL

DAIICHI SANKYO

375MG

N021141 001 May 26, 2000

COLISTIN SULFATE

SUSPENSION;ORAL

COLY-MYCIN S

PARKE DAVIS

EQ 25MG BASE/5ML

N050355 001

CONIVAPTAN HYDROCHLORIDE

INJECTABLE;INTRAVENOUS

VAPRISOL

CUMBERLAND PHARMS

20MG/4ML (5MG/ML)

N021697 001 Dec 29, 2005

COPPER

INTRAUTERINE DEVICE;INTRAUTERINE

CU-7

GD SEARLE LLC

89MG

N017408 001

TATUM-T

GD SEARLE LLC

120MG

N018205 001

CORTICOTROPIN

INJECTABLE;INJECTION

ACTH

PARKEDALE

25 UNITS/VIAL

N008317 002

40 UNITS/VIAL

N008317 004

ACTHAR

SANOFI AVENTIS US

25 UNITS/VIAL

N007504 002

40 UNITS/VIAL

N007504 003

CORTICOTROPIN

ORGANICS LAGRANGE

40 UNITS/ML

N010831 001

80 UNITS/ML

N010831 002

WATSON LABS

40 UNITS/VIAL

A088772 001 Nov 21, 1984

H.P. ACTHAR GEL

MALLINCKRODT ARD

40 UNITS/ML

N008372 006

PURIFIED CORTROPHIN GEL

ANI PHARMS INC

40 UNITS/ML

N008975 001

80 UNITS/ML

N008975 002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CORTICOTROPIN-ZINC HYDROXIDE

INJECTABLE; INJECTION

CORTROPHIN-ZINC

ANI PHARMS INC 40 UNITS/ML N009854 001

CORTISONE ACETATE

INJECTABLE; INJECTION

CORTISONE ACETATE

PHARMACIA AND UPJOHN 25MG/ML N008126 002

WATSON LABS 25MG/ML A083147 003

25MG/ML A085677 001

50MG/ML A083147 004

50MG/ML A085677 002

CORTONE

MERCK 25MG/ML N007110 002

50MG/ML N007110 003

TABLET; ORAL

CORTISONE ACETATE

BARR 25MG A083471 001

ELKINS SINN 25MG A080836 001

EVERYLIFE 25MG A084246 001

HEATHER 25MG A085736 001

IMPAX LABS 25MG N009458 001

INWOOD LABS 25MG A080731 001

IVAX SUB TEVA PHARMS 25MG A080630 001

25MG A083536 001

LANNETT 25MG A080694 001

PANRAY 5MG N008284 002

25MG N008284 001

PHARMACIA AND UPJOHN 5MG N008126 003

10MG N008126 004

25MG N008126 001

PUREPAC PHARM 25MG A080493 001

VITARINE 25MG A080333 001

WATSON LABS 25MG A085884 001

WHITEWORTH TOWN PLSN 25MG A080341 001

CORTONE

+ MERCK 25MG ** N007750 003

COSYNTROPIN

SOLUTION; INTRAVENOUS

COSYNTROPIN

SANDOZ INC 0.25MG/ML (0.25MG/ML) N022028 001 Feb 21, 2008

CROMOLYN SODIUM

AEROSOL, METERED; INHALATION

INTAL

KING PHARMS LLC 0.8MG/INH N018887 001 Dec 05, 1985

CAPSULE; INHALATION

INTAL

+ SANOFI AVENTIS US 20MG ** N016990 001

CAPSULE; ORAL

GASTROCROM

UCB INC 100MG N019188 001 Dec 22, 1989

CONCENTRATE; ORAL

CROMOLYN SODIUM

GENERA PHARMS 100MG/5ML A090954 001 Dec 18, 2009

SOLUTION; INHALATION

CROMOLYN SODIUM

ACTAVIS MID ATLANTIC 10MG/ML A075067 001 Jul 19, 1999

APOTEX INC 10MG/ML A075333 001 Apr 30, 2002

BAUSCH AND LOMB 10MG/ML A075585 001 Dec 21, 2000

FERA PHARMS LLC 10MG/ML A075437 001 Apr 21, 2000

ROXANE 10MG/ML A075175 001 Sep 30, 1999

WATSON LABS 10MG/ML A076469 001 Jun 17, 2005

INTAL

+ KING PHARMS LLC 10MG/ML ** N018596 001 May 28, 1982

SOLUTION/DROPS; OPHTHALMIC

CROLOM

BAUSCH AND LOMB 4% A074443 001 Jan 30, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CROMOLYN SODIUM

SOLUTION/DROPS;OPHTHALMIC

CROMOLYN SODIUM

APOTEX INC

4%

A075615 001 Jan 26, 2001

CROMOPTIC

KING PHARMS

4%

A075088 001 Apr 27, 1999

OPTICROM

+ ALLERGAN

4% **

N018155 001 Oct 03, 1984

SPRAY, METERED;NASAL

CROMOLYN SODIUM

ACTAVIS MID ATLANTIC

5.2MG/SPRAY

A074800 001 Jul 26, 2001

HH AND P

5.2MG/SPRAY

A077976 001 Sep 07, 2007

NASALCROM

+ BLACKSMITH BRANDS

5.2MG/SPRAY **

N020463 001 Jan 03, 1997

CRYPTENAMINE ACETATES

INJECTABLE;INJECTION

UNITENSEN

MEDPOINTE PHARM HLC

260CSR UNIT/ML

N008814 001

CRYPTENAMINE TANNATES

TABLET;ORAL

UNITENSEN

MEDPOINTE PHARM HLC

260CSR UNIT

N009217 001

CUPRIC SULFATE

INJECTABLE;INJECTION

CUPRIC SULFATE

ABRAXIS PHARM

EQ 0.4MG COPPER/ML

N019350 001 May 05, 1987

CYANOCOBALAMIN

GEL, METERED;NASAL

NASCOBAL

PAR PHARM

0.5MG/INH

N019722 001 Nov 05, 1996

INJECTABLE;INJECTION

BERUBIGEN

PHARMACIA AND UPJOHN

1MG/ML

N006798 001

BETALIN 12

LILLY

0.1MG/ML

A080855 001

1MG/ML

A080855 002

COBAVITE

WATSON LABS

0.1MG/ML

A083013 001

1MG/ML

A083064 001

CYANOCOBALAMIN

ABRAXIS PHARM

0.03MG/ML

A080510 003

0.1MG/ML

A080510 001

1MG/ML

A080510 002

AKORN

1MG/ML

A087969 001 Nov 10, 1983

DELL LABS

0.03MG/ML

A080689 001

0.1MG/ML

A080689 002

1MG/ML

A080689 003

FRESENIUS KABI USA

0.1MG/ML

A080557 002

LUITPOLD

0.03MG/ML

A080668 001

LYPHOMED

1MG/ML

A083075 001

MYLAN INSTITUTIONAL

1MG/ML

A040451 001 Sep 23, 2003

SANOFI AVENTIS US

1MG/ML

A080564 001

SOLOPAK

1MG/ML

A087551 001 Feb 29, 1984

WARNER CHILCOTT

1MG/ML

N007085 002

WATSON LABS

0.1MG/ML

A080573 002

0.1MG/ML

A083120 001

1MG/ML

A080573 001

1MG/ML

A083120 002

WYETH AYERST

0.1MG/ML

A080554 001

1MG/ML

A080554 002

DODEX

ACCORD HLTHCARE

1MG/ML

A083022 001

REDISOL

MERCK

1MG/ML

N006668 010

RUBIVITE

BEL MAR

0.03MG/ML

N010791 004

0.05MG/ML

N010791 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CYANOCOBALAMIN

INJECTABLE; INJECTION

RUBIVITE

| | |
|-----------|-------------|
| 0.1MG/ML | N010791 002 |
| 0.12MG/ML | N010791 005 |
| 1MG/ML | N010791 003 |

RUBRAMIN PC

| | | |
|----------------------|----------|-------------|
| BRISTOL MYERS SQUIBB | 0.1MG/ML | N006799 002 |
|----------------------|----------|-------------|

| | | |
|---|-----------|-------------|
| + | 1MG/ML ** | N006799 004 |
|---|-----------|-------------|

| | | | |
|---|-----------|-------------|--------------|
| + | 1MG/ML ** | N006799 010 | Apr 28, 1988 |
|---|-----------|-------------|--------------|

RUVITE

| | | |
|-------------|--------|-------------|
| SAVAGE LABS | 1MG/ML | A080570 002 |
|-------------|--------|-------------|

VI-TWEL

| | | |
|----------------|--------|-------------|
| BAYER HLTHCARE | 1MG/ML | N007012 002 |
|----------------|--------|-------------|

SPRAY, METERED; NASAL

CALOMIST

| | | | |
|-----------|-------------|-------------|--------------|
| PAR PHARM | 25MCG/SPRAY | N022102 001 | Jul 27, 2007 |
|-----------|-------------|-------------|--------------|

TABLET; ORAL

CYANOCOBALAMIN

| | | |
|-----------|-----|-------------|
| WEST WARD | 1MG | A084264 001 |
|-----------|-----|-------------|

CYANOCOBALAMIN CO-57

CAPSULE; ORAL

RUBRATOPE-57

| | | |
|--------|----------|-------------|
| BRACCO | 0.5-1uCi | N016089 002 |
|--------|----------|-------------|

CYANOCOBALAMIN CO-60

CAPSULE; ORAL

RUBRATOPE-60

| | | |
|--------|----------|-------------|
| BRACCO | 0.5-1uCi | N016090 002 |
|--------|----------|-------------|

CYANOCOBALAMIN; CYANOCOBALAMIN CO-57; CYANOCOBALAMIN CO-58

N/A; N/A

DICOPAC KIT

| | | |
|---------------|---------------|-------------|
| GE HEALTHCARE | N/A; N/A; N/A | N017406 001 |
|---------------|---------------|-------------|

CYANOCOBALAMIN; CYANOCOBALAMIN CO-57; INTRINSIC FACTOR

N/A; N/A

CYANOCOBALAMIN CO 57 SCHILLING TEST KIT

| | | |
|--------------|---------------------|-------------|
| MALLINCKRODT | 0.1MG; 0.5uCi; 60MG | N016635 001 |
|--------------|---------------------|-------------|

CYANOCOBALAMIN; TANNIC ACID; ZINC ACETATE

INJECTABLE; INJECTION

DEPINAR

| | | |
|--------------|----------------------------|-------------|
| ARMOUR PHARM | 0.5MG/ML; 2.3MG/ML; 1MG/ML | N011208 001 |
|--------------|----------------------------|-------------|

CYCLACILLIN

FOR SUSPENSION; ORAL

CYCLAPEN-W

| | | |
|--------------|-----------|-------------|
| WYETH AYERST | 125MG/5ML | N050508 001 |
|--------------|-----------|-------------|

| | | |
|--|-----------|-------------|
| | 250MG/5ML | N050508 002 |
|--|-----------|-------------|

| | | |
|--|-----------|-------------|
| | 500MG/5ML | N050508 003 |
|--|-----------|-------------|

TABLET; ORAL

CYCLACILLIN

| | | | |
|------|-------|-------------|--------------|
| TEVA | 250MG | A062895 001 | Aug 04, 1988 |
|------|-------|-------------|--------------|

| | | | |
|--|-------|-------------|--------------|
| | 500MG | A062895 002 | Aug 04, 1988 |
|--|-------|-------------|--------------|

CYCLAPEN-W

| | | |
|--------------|-------|-------------|
| WYETH AYERST | 250MG | N050509 001 |
|--------------|-------|-------------|

| | | |
|--|-------|-------------|
| | 500MG | N050509 002 |
|--|-------|-------------|

CYCLIZINE LACTATE

INJECTABLE; INJECTION

MAREZINE

| | | |
|-----------------|---------|-------------|
| GLAXOSMITHKLINE | 50MG/ML | N009495 001 |
|-----------------|---------|-------------|

CYCLOBENZAPRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

| | | | |
|----------------|------|-------------|--------------|
| TWI PHARMS INC | 15MG | A091281 001 | Jan 31, 2013 |
|----------------|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 30MG | A091281 002 | Jan 31, 2013 |
|--|------|-------------|--------------|

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

| | | | |
|--------|------|-------------|--------------|
| SANDOZ | 10MG | A073683 001 | Feb 26, 1993 |
|--------|------|-------------|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CYCLOBENZAPRINE HYDROCHLORIDE

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

| | | | |
|-------------------|------|-------------|--------------|
| UPSHER SMITH LABS | 5MG | A072854 002 | Feb 03, 2006 |
| | 10MG | A072854 001 | Nov 19, 1991 |
| WATSON LABS | 10MG | A073143 001 | Nov 27, 1991 |
| | 10MG | A074436 001 | Nov 30, 1994 |

FLEXERIL

| | | | |
|-----------------------|---------|-------------|--|
| + JANSSEN RES AND DEV | 5MG ** | N017821 001 | |
| + | 10MG ** | N017821 002 | |

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AK-PENTOLATE

| | | | |
|-------|----|-------------|--|
| AKORN | 1% | A085555 001 | |
|-------|----|-------------|--|

AKPENTOLATE

| | | | |
|-------|----|-------------|--------------|
| AKORN | 2% | A040165 001 | Jan 13, 1997 |
|-------|----|-------------|--------------|

CYCLOPENTOLATE HYDROCHLORIDE

| | | | |
|------------------|----|-------------|--------------|
| ALCON PHARMS LTD | 1% | A089162 001 | Jan 24, 1991 |
| SOLA BARNES HIND | 1% | A084150 001 | |
| | 1% | A084863 001 | |

PENTOLAIR

| | | | |
|------------|------|-------------|--------------|
| PHARMAFAIR | 0.5% | A088643 001 | Feb 09, 1987 |
| | 1% | A088150 001 | Feb 25, 1983 |

CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDE

| | | | |
|-----------------|------------|-------------|--------------|
| BAXTER HLTHCARE | 100MG/VIAL | A088371 001 | Jul 03, 1986 |
| | 200MG/VIAL | A088372 001 | Jul 03, 1986 |
| | 500MG/VIAL | A088373 001 | Jul 03, 1986 |
| | 1GM/VIAL | A088374 001 | Sep 24, 1986 |

CYTOXAN

| | | | |
|-------------------|---------------|-------------|--|
| + BAXTER HLTHCARE | 100MG/VIAL ** | N012142 001 | |
| + | 200MG/VIAL ** | N012142 002 | |

CYTOXAN (LYOPHILIZED)

| | | | |
|-------------------|---------------|-------------|--------------|
| + BAXTER HLTHCARE | 500MG/VIAL | N012142 003 | |
| + | 500MG/VIAL ** | N012142 008 | Jan 04, 1984 |
| + | 1GM/VIAL | N012142 004 | Aug 30, 1982 |
| + | 1GM/VIAL ** | N012142 010 | Sep 24, 1985 |
| + | 2GM/VIAL | N012142 005 | Aug 30, 1982 |
| + | 2GM/VIAL ** | N012142 009 | Dec 10, 1985 |

LYOPHILIZED CYTOXAN

| | | | |
|-------------------|---------------|-------------|--------------|
| + BAXTER HLTHCARE | 100MG/VIAL ** | N012142 006 | Dec 05, 1985 |
| + | 200MG/VIAL ** | N012142 007 | Dec 10, 1985 |

NEOSAR

| | | | |
|-----------------|------------|-------------|--------------|
| BEDFORD | 100MG/VIAL | A087442 001 | Feb 16, 1982 |
| | 200MG/VIAL | A087442 002 | Feb 16, 1982 |
| | 500MG/VIAL | A087442 003 | Feb 16, 1982 |
| | 1GM/VIAL | A087442 004 | Jul 08, 1983 |
| | 2GM/VIAL | A087442 005 | Mar 30, 1989 |
| TEVA PARENTERAL | 100MG/VIAL | A040015 001 | Apr 29, 1993 |
| | 200MG/VIAL | A040015 002 | Apr 29, 1993 |
| | 500MG/VIAL | A040015 003 | Apr 29, 1993 |
| | 1GM/VIAL | A040015 004 | Apr 29, 1993 |
| | 2GM/VIAL | A040015 005 | Apr 29, 1993 |

TABLET; ORAL

CYCLOPHOSPHAMIDE

| | | | |
|--------|------|-------------|--------------|
| ROXANE | 25MG | A040032 001 | Aug 17, 1999 |
| | 50MG | A040032 002 | Aug 17, 1999 |

CYTOXAN

| | | | |
|-------------------|---------|-------------|--|
| + BAXTER HLTHCARE | 25MG ** | N012141 002 | |
| + | 50MG ** | N012141 001 | |

CYCLOSPORINE

CAPSULE; ORAL

GENGRAF

| | | | |
|--------|------|-------------|--------------|
| ABBVIE | 50MG | A065003 002 | May 12, 2000 |
|--------|------|-------------|--------------|

NEORAL

| | | | |
|------------|---------|-------------|--------------|
| + NOVARTIS | 50MG ** | N050715 003 | Jul 14, 1995 |
|------------|---------|-------------|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

CYCLOSPORINE

SOLUTION; ORAL

CYCLOSPORINE

APOTEX INC

100MG/ML

A065167 001 Jan 05, 2005

CYCLOTHIAZIDE

TABLET; ORAL

ANHYDRON

LILLY

2MG

N013157 002

FLUIDIL

PHARMACIA AND UPJOHN

2MG

N018173 001

CYCRIMINE HYDROCHLORIDE

TABLET; ORAL

PAGITANE

LILLY

1.25MG

N008951 001

2.5MG

N008951 002

CYPROHEPTADINE HYDROCHLORIDE

SYRUP; ORAL

CYPROHEPTADINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC 2MG/5ML **

A086833 001

HALSEY

2MG/5ML

A089199 001 Jul 03, 1986

MORTON GROVE

2MG/5ML

A087001 001 Nov 04, 1982

NASKA

2MG/5ML

A089021 001 Dec 21, 1987

PERIACTIN

+ MERCK

2MG/5ML **

N013220 002

TABLET; ORAL

CYPROHEPTADINE HYDROCHLORIDE

AM THERAP

4MG

A088798 001 Feb 15, 1985

ASCOT

4MG

A087685 001 Oct 25, 1982

CHARTWELL RX

4MG

A088212 001 May 26, 1983

DURAMED PHARMS BARR

4MG

A088232 001 Oct 25, 1983

FOSUN PHARMA

4MG

A086808 001

HALSEY

4MG

A089057 001 Jul 03, 1986

KV PHARM

4MG

A086737 001

MD PHARM

4MG

A087566 001 Nov 10, 1982

MYLAN

4MG

A086678 001

PIONEER PHARMS

4MG

A087839 001 Feb 08, 1984

PLIVA

4MG

A088205 001 Jul 26, 1983

SUPERPHARM

4MG

A087405 001

VITARINE

4MG

A087284 001

WATSON LABS

4MG

A085245 001

4MG

A086165 001

4MG

A086580 001

PERIACTIN

+ MERCK

4MG **

N012649 001

CYSTEINE HYDROCHLORIDE

INJECTABLE; INJECTION

CYSTEINE HYDROCHLORIDE

+ HOSPIRA

7.25% **

N019523 001 Oct 22, 1986

CYTARABINE

INJECTABLE; INJECTION

CYTARABINE

+ TEVA PARENTERAL

100MG/VIAL **

N016793 001

+

500MG/VIAL **

N016793 002

+

1GM/VIAL **

N016793 003 Dec 21, 1987

+

2GM/VIAL **

N016793 004 Dec 21, 1987

CYTOSAR-U

TEVA PHARMS USA

100MG/VIAL

A075206 001 Dec 30, 1998

500MG/VIAL

A075206 002 Dec 30, 1998

1GM/VIAL

A075206 004 Dec 30, 1998

2GM/VIAL

A075206 003 Dec 30, 1998

INJECTABLE, LIPOSOMAL; INJECTION

DEPOCYT

+ PACIRA PHARMS INC

10MG/ML

N021041 001 Apr 01, 1999

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DACARBAZINE

INJECTABLE; INJECTION

DACARBAZINE

ABRAXIS PHARM

100MG/VIAL

A070962 001 Aug 28, 1986

200MG/VIAL

A070990 001 Aug 28, 1986

DTIC-DOME

+ BAYER HLTHCARE

100MG/VIAL **

N017575 001

+

200MG/VIAL **

N017575 002

DACTINOMYCIN

INJECTABLE; INJECTION

DACTINOMYCIN

WEST-WARD PHARMS INT 0.5MG/VIAL

A090304 001 Mar 16, 2010

DALFOPRISTIN; QUINUPRISTIN

INJECTABLE; INTRAVENOUS

SYNERCID

KING PHARMS

420MG/VIAL; 180MG/VIAL

N050748 002 Aug 24, 2000

DALTEPARIN SODIUM

INJECTABLE; INJECTION

FRAGMIN

PFIZER INC

7,500 IU/0.75ML

N020287 008 Apr 04, 2002

INJECTABLE; SUBCUTANEOUS

FRAGMIN

PFIZER INC

10,000IU/0.4ML (25,000IU/ML)

N020287 002 May 01, 2007

95,000IU/9.5ML (10,000IU/ML)

N020287 007 Apr 04, 2002

DANAPAROID SODIUM

INJECTABLE; INJECTION

ORGARAN

ASPEN GLOBAL INC

750 UNITS/0.6ML

N020430 001 Dec 24, 1996

DANAZOL

CAPSULE; ORAL

DANAZOL

AM THERAP

200MG

A071569 001 Dec 30, 1987

DANOCRINE

SANOFI AVENTIS US

50MG **

N017557 003

100MG **

N017557 004

200MG **

N017557 002

DAPIPRAZOLE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

DAPIPRAZOLE HYDROCHLORIDE

+ FERA PHARMS

0.5% **

N019849 001 Dec 31, 1990

DAPTOMYCIN

POWDER; IV (INFUSION)

CUBICIN

CUBIST PHARMS LLC

250MG/VIAL

N021572 001 Sep 12, 2003

DARUNAVIR ETHANOLATE

TABLET; ORAL

PREZISTA

+ JANSSEN PRODS

EQ 300MG BASE **

N021976 001 Jun 23, 2006

+

EQ 400MG BASE **

N021976 003 Oct 21, 2008

DASATINIB

TABLET; ORAL

DASATINIB

APOTEX INC

20MG

A202103 001 Jun 10, 2016

50MG

A202103 002 Jun 10, 2016

70MG

A202103 003 Jun 10, 2016

100MG

A202103 004 Jun 10, 2016

DAUNORUBICIN CITRATE

INJECTABLE, LIPOSOMAL; INJECTION

DAUNOXOME

GALEN (UK)

EQ 2MG BASE/ML

N050704 002 Apr 08, 1996

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DAUNORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

CERUBIDINE

SANOFI AVENTIS US EQ 20MG BASE/VIAL A061876 001

WYETH AYERST EQ 20MG BASE/VIAL ** N050484 001

DAUNORUBICIN HYDROCHLORIDE

TEVA PARENTERAL EQ 20MG BASE/VIAL A064212 001 Jun 23, 1998

EQ 50MG BASE/VIAL A064212 002 May 03, 1999

DECAMETHONIUM BROMIDE

INJECTABLE; INJECTION

SYNCURINE

GLAXOSMITHKLINE 1MG/ML N006931 002

DEFEROXAMINE MESYLATE

INJECTABLE; INJECTION

DEFEROXAMINE MESYLATE

WATSON LABS 500MG/VIAL A076806 001 Mar 31, 2006

2GM/VIAL A076806 002 Mar 31, 2006

DEMECARIUM BROMIDE

SOLUTION/DROPS; OPHTHALMIC

HUMORSOL

MERCK 0.125% N011860 002

0.25% N011860 001

DEMECLOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL

DECLOMYCIN

LEDERLE 150MG N050262 001

SYRUP; ORAL

DECLOMYCIN

LEDERLE 75MG/5ML N050257 001

TABLET; ORAL

DECLOMYCIN

COREPHARMA 75MG N050261 001

150MG N050261 002

300MG N050261 003

DEMECLOCYCLINE HYDROCHLORIDE

IMPAX LABS 150MG A065094 001 Mar 22, 2004

300MG A065094 002 Mar 22, 2004

DESERPIDINE

TABLET; ORAL

HARMONYL

ABBVIE 0.1MG N010796 001

0.25MG N010796 002

DESERPIDINE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ORETICYL 25

ABBVIE 0.125MG; 25MG N012148 001

ORETICYL 50

ABBVIE 0.125MG; 50MG N012148 003

ORETICYL FORTE

ABBVIE 0.25MG; 25MG N012148 002

DESERPIDINE; METHYCLOTHIAZIDE

TABLET; ORAL

ENDURONYL

ABBOTT 0.25MG; 5MG N012775 001

ENDURONYL FORTE

ABBOTT 0.5MG; 5MG N012775 002

METHYCLOTHIAZIDE AND DESERPIDINE

WATSON LABS 0.25MG; 5MG A088486 001 Aug 10, 1984

0.5MG; 5MG A088452 001 Aug 10, 1984

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DESIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

PERTOFRANE

| | | | |
|-------------------|------|---------|-----|
| SANOFI AVENTIS US | 25MG | N013621 | 001 |
| | 50MG | N013621 | 002 |

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

ANI PHARMS INC

| | | | |
|-------|---------|-----|--------------|
| 25MG | A071803 | 002 | Dec 08, 1987 |
| 50MG | A071803 | 003 | Dec 08, 1987 |
| 75MG | A071803 | 004 | Dec 08, 1987 |
| 100MG | A071803 | 001 | May 29, 1997 |
| 150MG | A071803 | 005 | May 29, 1997 |

USL PHARMA

| | | | |
|-------|---------|-----|--------------|
| 25MG | A071864 | 001 | Sep 09, 1987 |
| 50MG | A071865 | 001 | Sep 09, 1987 |
| 75MG | A071866 | 001 | Sep 09, 1987 |
| 100MG | A071867 | 001 | Sep 09, 1987 |

DESIRUDIN RECOMBINANT

INJECTABLE; SUBCUTANEOUS

IPRIVASK

| | | | | |
|------------------------|-----------|---------|-----|--------------|
| + VALEANT PHARMS NORTH | 15MG/VIAL | N021271 | 001 | Apr 04, 2003 |
|------------------------|-----------|---------|-----|--------------|

DESLANOSIDE

INJECTABLE; INJECTION

CEDILANID-D

NOVARTIS

| | | |
|----------|---------|-----|
| 0.2MG/ML | N009282 | 002 |
|----------|---------|-----|

DESMOPRESSIN ACETATE

INJECTABLE; INJECTION

DDAVP

| | | | | |
|--------------------|------------|---------|-----|--------------|
| FERRING PHARMS INC | 0.015MG/ML | N018938 | 002 | Apr 25, 1995 |
|--------------------|------------|---------|-----|--------------|

DESMOPRESSIN ACETATE

BEDFORD

| | | | |
|------------|---------|-----|--------------|
| 0.004MG/ML | A074575 | 001 | Feb 18, 2000 |
|------------|---------|-----|--------------|

HOSPIRA

| | | | |
|------------|---------|-----|--------------|
| 0.004MG/ML | A075220 | 001 | Aug 28, 2000 |
|------------|---------|-----|--------------|

TEVA PHARMS USA

| | | | |
|------------|---------|-----|--------------|
| 0.004MG/ML | A074888 | 001 | Oct 15, 1997 |
|------------|---------|-----|--------------|

DESMOPRESSIN ACETATE PRESERVATIVE FREE

BEDFORD

| | | | |
|------------|---------|-----|--------------|
| 0.004MG/ML | A074574 | 001 | Feb 18, 2000 |
|------------|---------|-----|--------------|

SOLUTION; NASAL

CONCENTRAID

FERRING

| | | | |
|-------|---------|-----|--------------|
| 0.01% | N019776 | 001 | Dec 26, 1990 |
|-------|---------|-----|--------------|

SPRAY, METERED; NASAL

DDAVP

| | | | | |
|----------------------|-----------------|---------|-----|--------------|
| + FERRING PHARMS INC | 0.01MG/SPRAY ** | N017922 | 002 | Feb 06, 1989 |
|----------------------|-----------------|---------|-----|--------------|

STIMATE

| | | | | |
|--------------------|--------------|---------|-----|--------------|
| FERRING PHARMS INC | 0.15MG/SPRAY | N020355 | 001 | Mar 07, 1994 |
|--------------------|--------------|---------|-----|--------------|

TABLET; ORAL

DESMOPRESSIN ACETATE

FERRING

| | | | |
|-------|---------|-----|--------------|
| 0.1MG | N021795 | 001 | May 08, 2008 |
|-------|---------|-----|--------------|

| | | | |
|-------|---------|-----|--------------|
| 0.2MG | N021795 | 002 | May 08, 2008 |
|-------|---------|-----|--------------|

DESOGESTREL; ETHINYL ESTRADIOL

TABLET; ORAL-21

DESOGEN

| | | | | |
|-----------------|---------------|---------|-----|--------------|
| ORGANON USA INC | 0.15MG;0.03MG | N020071 | 001 | Dec 10, 1992 |
|-----------------|---------------|---------|-----|--------------|

DESOGESTREL AND ETHINYL ESTRADIOL

DURAMED PHARMS BARR

| | | | |
|---------------|---------|-----|--------------|
| 0.15MG;0.03MG | A075256 | 001 | Aug 12, 1999 |
|---------------|---------|-----|--------------|

ORTHO-CEPT

JANSSEN PHARMS

| | | | |
|---------------|---------|-----|--------------|
| 0.15MG;0.03MG | N020301 | 001 | Dec 14, 1992 |
|---------------|---------|-----|--------------|

TABLET; ORAL-28

MIRCETTE

| | | | | |
|----------------------|-----------------------------|---------|-----|--------------|
| + TEVA BRANDED PHARM | 0.15MG,N/A;0.02MG,0.01MG ** | N020713 | 001 | Apr 22, 1998 |
|----------------------|-----------------------------|---------|-----|--------------|

ORTHO-CEPT

JANSSEN PHARMS

| | | | |
|---------------|---------|-----|--------------|
| 0.15MG;0.03MG | N020301 | 002 | Dec 14, 1992 |
|---------------|---------|-----|--------------|

DESOXIMETASONE

CREAM; TOPICAL

TOPICORT

| | | | |
|-----------------------|----------|---------|-----|
| + TARO PHARM INDS LTD | 0.25% ** | N017856 | 001 |
|-----------------------|----------|---------|-----|

TOPICORT LP

| | | | |
|-----------------------|----------|---------|-----|
| + TARO PHARM INDS LTD | 0.05% ** | N018309 | 001 |
|-----------------------|----------|---------|-----|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DESOXIMETASONE

GEL; TOPICAL

TOPICORT

+ TARO PHARM INDS LTD 0.05% **

N018586 001 Mar 29, 1982

OINTMENT; TOPICAL

DESOXIMETASONE

ALTANA 0.25%

A073440 001 Apr 01, 1998

TOPICORT

+ TARO PHARM INDS LTD 0.25% **

N018763 001 Sep 30, 1983

DESOXYCORTICOSTERONE ACETATE

INJECTABLE; INJECTION

DOCA

ORGANON USA INC 5MG/ML

N001104 001

PELLET; IMPLANTATION

PERCORTEN

NOVARTIS 125MG

N005151 001

DESOXYCORTICOSTERONE PIVALATE

INJECTABLE; INJECTION

PERCORTEN

NOVARTIS 25MG/ML

N008822 001

DESVENLAFAXINE FUMARATE

TABLET, EXTENDED RELEASE; ORAL

DESVENLAFAXINE

+ SUN PHARMA GLOBAL EQ 50MG BASE

N205583 001 Jan 28, 2014

+ EQ 100MG BASE

N205583 002 Jan 28, 2014

TEVA PHARMS USA EQ 50MG BASE

N205208 001 Oct 11, 2013

EQ 100MG BASE

N205208 002 Oct 11, 2013

DEXAMETHASONE

AEROSOL; TOPICAL

AEROSEB-DEX

ALLERGAN HERBERT 0.01% **

A083296 002

DECASPRAY

+ MERCK 0.04% **

N012731 002

ELIXIR; ORAL

DECADRON

MERCK 0.5MG/5ML

N012376 002

DEXAMETHASONE

ALPHARMA US PHARMS 0.5MG/5ML

A088997 001 Oct 10, 1986

HEXADROL

ASPEN GLOBAL INC 0.5MG/5ML

N012674 001

GEL; TOPICAL

DECADERM

MERCK 0.1%

N013538 001

SUSPENSION/DROPS; OPHTHALMIC

DEXAMETHASONE

WATSON LABS 0.1%

A089170 001 May 09, 1989

TABLET; ORAL

DECADRON

+ MERCK 0.25MG **

N011664 004

+ 0.5MG **

N011664 001

+ 0.75MG **

N011664 002

+ 1.5MG **

N011664 003

+ 4MG **

N011664 005

+ 6MG **

N011664 006 Jul 30, 1982

DEXAMETHASONE

ANI PHARMS INC 0.75MG

A080399 001

IMPAX LABS 0.75MG

A085376 001

PAR PHARM 0.25MG

A088149 001 Apr 28, 1983

PHOENIX LABS NY 0.75MG

A083806 001

PVT FORM 0.75MG

A083420 001

ROXANE 0.25MG

A084614 001

SUN PHARM INDUSTRIES 0.25MG

A084013 001

0.25MG

A084764 001

0.5MG

A084084 001

0.5MG

A084766 001

0.75MG

A084081 001

0.75MG

A084765 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXAMETHASONE

TABLET; ORAL

DEXAMETHASONE

| | | |
|----------------------|--------|-------------|
| | 1.5MG | A084086 001 |
| | 1.5MG | A084763 001 |
| UPSHER SMITH | 0.75MG | A087534 001 |
| | 1.5MG | A087533 001 |
| WATSON LABS | 0.25MG | A085455 001 |
| | 0.5MG | A085458 001 |
| | 0.75MG | A080968 001 |
| | 0.75MG | A084457 001 |
| | 0.75MG | A085818 001 |
| | 1.5MG | A085456 001 |
| | 1.5MG | A085840 001 |
| WHITEWORTH TOWN PLSN | 0.75MG | A084327 001 |
| DEXONE 0.5 | | |
| SOLVAY | 0.5MG | A084991 001 |
| DEXONE 0.75 | | |
| SOLVAY | 0.75MG | A084993 001 |
| DEXONE 1.5 | | |
| SOLVAY | 1.5MG | A084990 001 |
| DEXONE 4 | | |
| SOLVAY | 4MG | A084992 001 |
| HEXADROL | | |
| ASPEN GLOBAL INC | 0.5MG | N012675 004 |
| | 0.75MG | N012675 007 |
| | 1.5MG | N012675 009 |
| | 4MG | N012675 010 |

DEXAMETHASONE ACETATE

INJECTABLE; INJECTION

DECADRON-LA

+ MERCK

EQ 8MG BASE/ML **

N016675 001

DEXAMETHASONE ACETATE

WATSON LABS

EQ 8MG BASE/ML

A084315 001

WATSON LABS TEVA

EQ 16MG BASE/ML

A087711 001

May 24, 1982

DEXAMETHASONE SODIUM PHOSPHATE

AEROSOL; NASAL

DEXACORT

UCB INC

EQ 0.1MG PHOSPHATE/INH

N014242 001

AEROSOL, METERED; INHALATION

DEXACORT

UCB INC

EQ 0.1MG PHOSPHATE/INH

N013413 001

CREAM; TOPICAL

DECADRON

MERCK

EQ 0.1% PHOSPHATE

N011983 002

INJECTABLE; INJECTION

DECADRON

+ MERCK

EQ 4MG PHOSPHATE/ML **

N012071 002

+

EQ 24MG PHOSPHATE/ML **

N012071 004

DEXACEN-4

CENT PHARMS

EQ 4MG PHOSPHATE/ML

A084342 001

DEXAMETHASONE

ABRAXIS PHARM

EQ 4MG PHOSPHATE/ML

A088448 001

Jan 25, 1984

FRESENIUS KABI USA

EQ 10MG PHOSPHATE/ML

A088469 001

Jan 25, 1984

DEXAMETHASONE SODIUM PHOSPHATE

AKORN

EQ 4MG PHOSPHATE/ML

A084493 001

BEL MAR

EQ 4MG PHOSPHATE/ML

A084752 001

DELL LABS

EQ 4MG PHOSPHATE/ML

A083161 001

INTL MEDICATION

EQ 20MG PHOSPHATE/ML

A088522 001

Feb 17, 1984

LYPHOMED

EQ 4MG PHOSPHATE/ML

A087065 001

TEVA PARENTERAL

EQ 4MG PHOSPHATE/ML

A081125 001

Aug 31, 1990

EQ 10MG PHOSPHATE/ML

A081126 001

Aug 31, 1990

WATSON LABS

EQ 4MG PHOSPHATE/ML

A083702 001

EQ 4MG PHOSPHATE/ML

A084355 001

EQ 4MG PHOSPHATE/ML

A089169 001

Apr 09, 1986

EQ 10MG PHOSPHATE/ML

A087668 001

Jul 01, 1982

EQ 24MG PHOSPHATE/ML

A085606 001

WYETH AYERST

EQ 4MG PHOSPHATE/ML

A085641 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

HEXADROL

| | | | |
|---|------------------|-------------------------|-------------|
| + | ASPEN GLOBAL INC | EQ 4MG PHOSPHATE/ML ** | N014694 002 |
| + | | EQ 10MG PHOSPHATE/ML ** | N014694 003 |
| | | EQ 20MG PHOSPHATE/ML | N014694 004 |

OINTMENT; OPHTHALMIC

DECADRON

| | | |
|-------|--------------------|-------------|
| MERCK | EQ 0.05% PHOSPHATE | N011977 001 |
|-------|--------------------|-------------|

DEXAIR

| | | |
|------------|--------------------|--------------------------|
| PHARMAFAIR | EQ 0.05% PHOSPHATE | A088071 001 Dec 28, 1982 |
|------------|--------------------|--------------------------|

MAXIDEX

| | | |
|-------|--------------------|-------------|
| ALCON | EQ 0.05% PHOSPHATE | A083342 001 |
|-------|--------------------|-------------|

SOLUTION/DROPS; OPHTHALMIC

DEXAIR

| | | |
|------------|-------------------|--------------------------|
| PHARMAFAIR | EQ 0.1% PHOSPHATE | A088433 001 Dec 15, 1983 |
|------------|-------------------|--------------------------|

DEXAMETHASONE SODIUM PHOSPHATE

| | | |
|------------------|-------------------|-------------|
| SOLA BARNES HIND | EQ 0.1% PHOSPHATE | A084170 001 |
|------------------|-------------------|-------------|

| | | |
|--|-------------------|-------------|
| | EQ 0.1% PHOSPHATE | A084173 001 |
|--|-------------------|-------------|

SOLUTION/DROPS; OPHTHALMIC, OTIC

DECADRON

| | | |
|-------|-------------------|-------------|
| MERCK | EQ 0.1% PHOSPHATE | N011984 001 |
|-------|-------------------|-------------|

SOLUTION/DROPS; OTIC

DEXAMETHASONE SODIUM PHOSPHATE

| | | |
|-------|-------------------|-------------|
| AKORN | EQ 0.1% PHOSPHATE | A084855 001 |
|-------|-------------------|-------------|

DEXAMETHASONE SODIUM PHOSPHATE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DECADRON W/ XYLOCAINE

| | | |
|-------|------------------------------|-------------|
| MERCK | EQ 4MG PHOSPHATE/ML; 10MG/ML | N013334 002 |
|-------|------------------------------|-------------|

DEXAMETHASONE SODIUM PHOSPHATE; NEOMYCIN SULFATE

OINTMENT; OPHTHALMIC

NEODECADRON

| | | |
|-------|--------------------------------------|-------------|
| MERCK | EQ 0.05% PHOSPHATE; EQ 3.5MG BASE/GM | N050324 001 |
|-------|--------------------------------------|-------------|

SOLUTION/DROPS; OPHTHALMIC

NEODECADRON

| | | |
|-------|-------------------------------------|-------------|
| MERCK | EQ 0.1% PHOSPHATE; EQ 3.5MG BASE/ML | N050322 001 |
|-------|-------------------------------------|-------------|

NEOMYCIN SULFATE AND DEXAMETHASONE SODIUM PHOSPHATE

| | | |
|-----------------|-------------------------------------|--------------------------|
| BAUSCH AND LOMB | EQ 0.1% PHOSPHATE; EQ 3.5MG BASE/ML | A064055 001 Oct 30, 1995 |
|-----------------|-------------------------------------|--------------------------|

NEOMYCIN SULFATE-DEXAMETHASONE SODIUM PHOSPHATE

| | | |
|------------------|-------------------------------------|--------------------------|
| ALCON PHARMS LTD | EQ 0.1% PHOSPHATE; EQ 3.5MG BASE/ML | A062714 001 Jul 21, 1986 |
|------------------|-------------------------------------|--------------------------|

| | | |
|------------|-------------------------------------|--------------------------|
| PHARMAFAIR | EQ 0.1% PHOSPHATE; EQ 3.5MG BASE/ML | A062539 001 Jan 10, 1985 |
|------------|-------------------------------------|--------------------------|

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

DEXACIDIN

| | | |
|----------|-----------------------------------------|--------------------------|
| NOVARTIS | 0.1%; EQ 3.5MG BASE/GM; 10,000 UNITS/GM | A062566 001 Feb 22, 1985 |
|----------|-----------------------------------------|--------------------------|

DEXASPORIN

| | | |
|------------|-----------------------------------------|--------------------------|
| PHARMAFAIR | 0.1%; EQ 3.5MG BASE/GM; 10,000 UNITS/GM | A062411 001 May 16, 1983 |
|------------|-----------------------------------------|--------------------------|

SUSPENSION/DROPS; OPHTHALMIC

DEXACIDIN

| | | |
|----------|-----------------------------------------|--------------------------|
| NOVARTIS | 0.1%; EQ 3.5MG BASE/ML; 10,000 UNITS/ML | A062544 001 Oct 29, 1984 |
|----------|-----------------------------------------|--------------------------|

DEXASPORIN

| | | |
|------------|-----------------------------------------|--------------------------|
| PHARMAFAIR | 0.1%; EQ 3.5MG BASE/ML; 10,000 UNITS/ML | A062428 001 May 18, 1983 |
|------------|-----------------------------------------|--------------------------|

NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE

| | | |
|------------------|-----------------------------------------|--------------------------|
| ALCON PHARMS LTD | 0.1%; EQ 3.5MG BASE/ML; 10,000 UNITS/ML | A062721 001 Nov 17, 1986 |
|------------------|-----------------------------------------|--------------------------|

DEXBROMPHENIRAMINE MALEATE

SYRUP; ORAL

DISOMER

| | | |
|----------|---------|-------------|
| SCHERING | 2MG/5ML | N011814 002 |
|----------|---------|-------------|

TABLET; ORAL

DISOMER

| | | |
|----------|-----|-------------|
| SCHERING | 2MG | N011814 001 |
|----------|-----|-------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET; ORAL

DISOPHROL

SCHERING 2MG; 60MG N012394 002

TABLET, EXTENDED RELEASE; ORAL

BROMPHERIL

COPLY PHARM 6MG; 120MG A089116 001 Jan 22, 1987

DEXBROMPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE SULFATE

AVANTHI INC 6MG; 120MG A078648 001 Feb 27, 2013

DISOBROM

SANDOZ 6MG; 120MG A070770 001 Sep 30, 1991

DISOPHROL

SCHERING PLOUGH 6MG; 120MG N013483 004 Sep 13, 1982

DRIXORAL

+ SCHERING PLOUGH 6MG; 120MG ** N013483 003 Sep 13, 1982

RESPORAL

PIONEER PHARMS 6MG; 120MG A089139 001 Jun 16, 1988

DEXCHLORPHENIRAMINE MALEATE

SYRUP; ORAL

POLARAMINE

SCHERING 2MG/5ML A086837 001 Jul 19, 1982

TABLET; ORAL

DEXCHLORPHENIRAMINE MALEATE

ANI PHARMS INC 2MG A088682 001 Jan 17, 1986

POLARAMINE

SCHERING 2MG A086835 001

DEXLANSOPRAZOLE

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE; ORAL

DEXILANT SOLUTAB

+ TAKEDA PHARMS USA 30MG N208056 001 Jan 26, 2016

DEXTROAMPHETAMINE SULFATE

CAPSULE; ORAL

DEXAMPEX

TEVA 15MG A085355 001

CAPSULE, EXTENDED RELEASE; ORAL

DEXTROAMPHETAMINE SULFATE

ABLE 5MG A076814 001 Aug 25, 2004

10MG A076814 002 Aug 25, 2004

15MG A076814 003 Aug 25, 2004

ELIXIR; ORAL

DEXEDRINE

GLAXOSMITHKLINE 5MG/5ML ** A083902 001

TABLET; ORAL

DEXAMPEX

TEVA 5MG A083735 001

10MG A083735 002

DEXEDRINE

GLAXOSMITHKLINE 5MG A084935 001

DEXTROAMPHETAMINE SULFATE

ANI PHARMS INC 5MG A085370 001

EPIC PHARMA LLC 5MG A090652 001 Mar 07, 2014

10MG A090652 002 Mar 07, 2014

HALSEY 10MG A083930 001

LANNETT 5MG A083903 001

10MG A083903 003

15MG A085652 001

MAST MM 5MG A086521 001

NESHER PHARMS 5MG A040365 001 Oct 31, 2002

10MG A040367 001 Oct 31, 2002

PUREPAC PHARM 5MG A084125 001

SANDOZ 10MG A085371 001

VINTAGE PHARMS LLC 5MG A040299 001 May 13, 1999

VITARINE 5MG A084986 001

10MG A085892 001

DEXTROSTAT

SHIRE 5MG ** A084051 001

10MG ** A084051 002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

FERNDEX

FERNDALE LABS

5MG

A084001 001

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PHERAZINE DM

HALSEY

15MG/5ML; 6.25MG/5ML

A088913 001 Mar 02, 1987

PROMETH W/ DEXTROMETHORPHAN

G AND W LABS INC

15MG/5ML; 6.25MG/5ML **

A088762 001 Oct 31, 1984

PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

AMNEAL PHARMS

15MG/5ML; 6.25MG/5ML

A090575 001 Feb 08, 2011

+ ANI PHARMS

15MG/5ML; 6.25MG/5ML **

N011265 002 Apr 02, 1984

TRIS PHARMA INC

15MG/5ML; 6.25MG/5ML

A091687 001 Jun 28, 2012

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 10% IN PLASTIC CONTAINER

B BRAUN

10GM/100ML

N018046 001

MILES

10GM/100ML

N018504 001

DEXTROSE 2.5% IN PLASTIC CONTAINER

B BRAUN

2.5GM/100ML

N018358 001

2.5GM/100ML

N019626 001 Feb 02, 1988

DEXTROSE 20% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE

20GM/100ML

N017521 004

DEXTROSE 30% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE

30GM/100ML

N017521 003

DEXTROSE 38.5% IN PLASTIC CONTAINER

ABBOTT

38.5GM/100ML

N018923 001 Sep 19, 1984

DEXTROSE 40% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE

40GM/100ML

N017521 002

DEXTROSE 5% IN PLASTIC CONTAINER

DHL

5GM/100ML

N019971 001 Sep 28, 1995

+ ICU MEDICAL INC

50MG/ML

N019222 001 Jul 13, 1984

DEXTROSE 50% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE

50GM/100ML

N017521 001

ICU MEDICAL INC

50GM/100ML

N019894 001 Dec 26, 1989

DEXTROSE 60%

B BRAUN

60GM/100ML

N017995 002 Sep 22, 1982

DEXTROSE 60% IN PLASTIC CONTAINER

B BRAUN

60GM/100ML

N017995 001

+ BAXTER HLTHCARE

60GM/100ML

N017521 005 Mar 26, 1982

60GM/100ML

N020047 002 Jul 02, 1991

HOSPIRA

60GM/100ML

N019346 001 Jan 25, 1985

DEXTROSE 7.7% IN PLASTIC CONTAINER

B BRAUN

7.7GM/100ML

N019626 003 Feb 02, 1988

DEXTROSE; MAGNESIUM ACETATE TETRAHYDRATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PLASMA-LYTE 56 AND DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE

5GM/100ML; 32MG/100ML; 128MG/100ML; 234MG/100ML

N017385 001

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE

INJECTABLE; INJECTION

ISOLYTE P W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN

5GM/100ML; 31MG/100ML; 130MG/100ML; 26MG/100ML; 320MG/100ML

N019025 001 Dec 27, 1984

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE; SODIUM LACTATE; SODIUM PHOSPHATE, MONOBASIC ANHYDROUS

INJECTABLE; INJECTION

IONOSOL B AND DEXTROSE 5% IN PLASTIC CONTAINER

HOSPIRA

5GM/100ML; 53MG/100ML; 100MG/100ML; 100MG/100ML; 180MG/100ML; 280MG/100ML; 16MG/100ML

N019515 001 May 08, 1986

L

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | | |
|-----------------------------------------------|-------------------------------------------------------------|-------------|--------------|
| ISOLYTE H IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 30MG/100ML; 97MG/100ML; 220MG/100ML; 140MG/100ML | N019844 001 | Jun 10, 1993 |
| ISOLYTE H W/ DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 30MG/100ML; 97MG/100ML; 220MG/100ML; 140MG/100ML | N018273 001 | |

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

| | | | |
|------------------------------------------------------|--------------------------------------------------------------------------|-------------|--------------|
| ISOLYTE S IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML | N019843 001 | Aug 09, 1993 |
| ISOLYTE S W/ DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML | N018274 001 | |
| PLASMA-LYTE 148 AND DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| + BAXTER HLTHCARE | 5GM/100ML; 30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML | N017451 001 | |

DEXTROSE; POTASSIUM CHLORIDE

INJECTABLE; INJECTION

| | | | |
|---------------------------------------------------------------|------------------------|-------------|--------------|
| POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 37MG/100ML | N019699 001 | Sep 29, 1989 |
| POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 75MG/100ML | N018744 001 | Nov 09, 1982 |
| | 5GM/100ML; 75MG/100ML | N019699 002 | Sep 29, 1989 |
| POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 110MG/100ML | N019699 003 | Sep 29, 1989 |
| POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 150MG/100ML | N018744 002 | Nov 09, 1982 |
| POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 220MG/100ML | N018744 003 | Nov 09, 1982 |
| | 5GM/100ML; 220MG/100ML | N019699 005 | Sep 29, 1989 |
| POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 300MG/100ML | N018744 004 | Nov 09, 1982 |

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM LACTATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, MONOBASIC ANHYDROUS

INJECTABLE; INJECTION

| | | | |
|------------------------------------------------|---------------------------------------------------------------|-------------|--------------|
| IONOSOL T AND DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| HOSPIRA | 5GM/100ML; 111MG/100ML; 256MG/100ML; 146MG/100ML; 207MG/100ML | N019514 001 | May 08, 1986 |

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | | |
|-----------------------------------------------|--------------------------------------------------------------|-------------|--------------|
| ISOLYTE M IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 150MG/100ML; 130MG/100ML; 280MG/100ML; 91MG/100ML | N019870 001 | Jun 10, 1993 |
| ISOLYTE M W/ DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 150MG/100ML; 130MG/100ML; 280MG/100ML; 91MG/100ML | N018270 001 | |

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

| | | | |
|---------------------------------------------------------|---------------------------------------------------------------|-------------|--------------|
| DEXTROSE 5% AND ELECTROLYTE NO. 75 IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 5GM/100ML; 205MG/100ML; 100MG/100ML; 120MG/100ML; 220MG/100ML | N018840 001 | Jun 29, 1983 |

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | | |
|---------------------------------------------------------------------------------------|-------------------------------------|-------------|--------------|
| DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 75MG/100ML; 200MG/100ML | N018268 009 | |
| DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 150MG/100ML; 200MG/100ML | N018268 004 | |
| DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 220MG/100ML; 200MG/100ML | N018268 005 | |
| DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 300MG/100ML; 200MG/100ML | N018268 006 | |
| DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML; 75MG/100ML; 330MG/100ML | N018268 011 | Jan 18, 1986 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | | |
|------------------------------------------------------------------------------------------|------------------------------------|-------------|--------------|
| DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML;150MG/100ML;330MG/100ML | N018268 012 | Jan 18, 1986 |
| DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML;220MG/100ML;330MG/100ML | N018268 013 | Jan 18, 1986 |
| DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.30% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML;300MG/100ML;330MG/100ML | N018268 014 | Jan 18, 1986 |
| DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.075% | | | |
| B BRAUN | 5GM/100ML;75MG/100ML;450MG/100ML | N018268 010 | |
| DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML;150MG/100ML;450MG/100ML | N018268 001 | |
| DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML;220MG/100ML;450MG/100ML | N018268 002 | |
| DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML;300MG/100ML;450MG/100ML | N018268 003 | |
| DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 5GM/100ML;224MG/100ML;450MG/100ML | N018008 003 | |
| DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 5GM/100ML;300MG/100ML;450MG/100ML | N018008 001 | |
| DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER | | | |
| BAXTER HLTHCARE | 5GM/100ML;75MG/100ML;450MG/100ML | N018008 002 | |
| POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | |
| + ICU MEDICAL INC | 5GM/100ML;74.5MG/100ML;900MG/100ML | N019691 002 | Mar 24, 1988 |
| + ICU MEDICAL INC | 5GM/100ML;149MG/100ML;900MG/100ML | N019691 004 | Mar 24, 1988 |
| POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER | | | |
| + ICU MEDICAL INC | 5GM/100ML;224MG/100ML;450MG/100ML | N018362 006 | Mar 28, 1988 |
| POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | |
| + ICU MEDICAL INC | 5GM/100ML;224MG/100ML;900MG/100ML | N019691 006 | Mar 24, 1988 |
| POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER | | | |
| + ICU MEDICAL INC | 5GM/100ML;298MG/100ML;450MG/100ML | N018362 007 | Mar 28, 1988 |
| POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | |
| + ICU MEDICAL INC | 5GM/100ML;298MG/100ML;900MG/100ML | N019691 008 | Mar 24, 1988 |
| POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | |
| + ICU MEDICAL INC | 5GM/100ML;224MG/100ML;900MG/100ML | N019691 007 | Mar 24, 1988 |
| POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER | | | |
| + ICU MEDICAL INC | 5GM/100ML;74.5MG/100ML;450MG/100ML | N018362 008 | Mar 28, 1988 |
| + ICU MEDICAL INC | 5GM/100ML;149MG/100ML;450MG/100ML | N018362 004 | Mar 28, 1988 |
| POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | |
| + ICU MEDICAL INC | 5GM/100ML;74.5MG/100ML;900MG/100ML | N019691 001 | Mar 24, 1988 |
| + ICU MEDICAL INC | 5GM/100ML;149MG/100ML;900MG/100ML | N019691 003 | Mar 24, 1988 |

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | | |
|--------------------------------------------------------------|-------------------------|-------------|--------------|
| DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER | | | |
| B BRAUN | 10GM/100ML;200MG/100ML | N018386 001 | |
| DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER | | | |
| B BRAUN | 10GM/100ML;450MG/100ML | N018229 001 | |
| DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | |
| B BRAUN | 10GM/100ML;900MG/100ML | N018047 001 | |
| BAXTER HLTHCARE | 10GM/100ML;900MG/100ML | N016696 001 | |
| DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER | | | |
| B BRAUN | 2.5GM/100ML;450MG/100ML | N018030 001 | |
| HOSPIRA | 2.5GM/100ML;450MG/100ML | N018096 001 | |
| DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | |
| B BRAUN | 2.5GM/100ML;900MG/100ML | N018376 001 | |
| DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER | | | |
| ABBOTT | 3.3GM/100ML;300MG/100ML | N018055 001 | |
| DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML;110MG/100ML | N018030 005 | |
| DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML;200MG/100ML | N018030 004 | |
| MILES | 5GM/100ML;200MG/100ML | N018399 001 | |
| DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER | | | |
| ABBOTT | 5GM/100ML;225MG/100ML | N019482 001 | Oct 04, 1985 |
| DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER | | | |
| ABBOTT | 5GM/100ML;300MG/100ML | N019486 001 | Oct 04, 1985 |
| MILES | 5GM/100ML;300MG/100ML | N018501 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | | |
|------------------------------------------------------------|-----------------------|---------|------------------|
| DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER | | | |
| B BRAUN | 5GM/100ML;330MG/100ML | N018030 | 003 |
| DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER | | | |
| ABBOTT | 5GM/100ML;450MG/100ML | N019484 | 001 Oct 04, 1985 |
| B BRAUN | 5GM/100ML;450MG/100ML | N018030 | 002 |
| MILES | 5GM/100ML;450MG/100ML | N018400 | 001 |
| DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | |
| ABBOTT | 5GM/100ML;900MG/100ML | N019483 | 001 Oct 04, 1985 |
| B BRAUN | 5GM/100ML;900MG/100ML | N018026 | 001 |
| MILES | 5GM/100ML;900MG/100ML | N018500 | 001 |

DEXTROTHYROXINE SODIUM

TABLET; ORAL

| | | | |
|----------|-----|---------|-----|
| CHOLOXIN | | | |
| ABBVIE | 1MG | N012302 | 005 |
| | 2MG | N012302 | 002 |
| | 4MG | N012302 | 004 |
| | 6MG | N012302 | 006 |

DEZOCINE

INJECTABLE; INJECTION

| | | | |
|-------------|---------|---------|------------------|
| DALGAN | | | |
| ASTRAZENECA | 5MG/ML | N019082 | 001 Dec 29, 1989 |
| | 10MG/ML | N019082 | 002 Dec 29, 1989 |
| | 15MG/ML | N019082 | 003 Dec 29, 1989 |

DIATRIZOATE MEGLUMINE

INJECTABLE; INJECTION

| | | | |
|--------------------------|-----|---------|------------------|
| ANGIOVIST 282 | | | |
| BAYER HLTHCARE | 60% | A087726 | 001 Sep 23, 1982 |
| CARDIOGRAFIN | | | |
| BRACCO | 85% | N011620 | 002 |
| DIATRIZOATE MEGLUMINE | | | |
| BRACCO | 76% | N010040 | 017 |
| HYPaque | | | |
| GE HEALTHCARE | 30% | N016403 | 002 |
| | 60% | N016403 | 001 |
| RENO-60 | | | |
| BRACCO | 60% | N010040 | 016 |
| RENO-DIP | | | |
| BRACCO | 30% | N010040 | 012 |
| UROVIST MEGLUMINE DIU/CT | | | |
| BAYER HLTHCARE | 30% | A087739 | 001 Sep 23, 1982 |
| SOLUTION; URETERAL | | | |
| RENO-30 | | | |
| BRACCO | 30% | N010040 | 021 |
| UROVIST CYSTO | | | |
| BAYER HLTHCARE | 30% | A087729 | 001 Sep 23, 1982 |
| UROVIST CYSTO PEDIATRIC | | | |
| BAYER HLTHCARE | 30% | A087731 | 001 Sep 23, 1982 |
| SOLUTION; URETHRAL | | | |
| HYPaque-CYSTO | | | |
| GE HEALTHCARE | 30% | N016403 | 003 |

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

| | | | |
|-----------------|---------|---------|------------------|
| ANGIOVIST 292 | | | |
| BAYER HLTHCARE | 52%;8% | A087724 | 001 Sep 23, 1982 |
| ANGIOVIST 370 | | | |
| BAYER HLTHCARE | 66%;10% | A087723 | 001 Sep 23, 1982 |
| DIATRIZOATE-60 | | | |
| INTL MEDICATION | 52%;8% | A088166 | 001 Jun 17, 1983 |
| HYPaque-76 | | | |
| GE HEALTHCARE | 66%;10% | A086505 | 001 |
| HYPaque-M, 75% | | | |
| GE HEALTHCARE | 50%;25% | N010220 | 003 |
| HYPaque-M, 90% | | | |
| GE HEALTHCARE | 60%;30% | N010220 | 002 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

| | | | | |
|------------------------|------------------|-------------|---------|------------------|
| MD-60 | | | | |
| | MALLINCKRODT | 52%;8% | A087074 | 001 |
| MD-76 | | | | |
| | MALLINCKRODT | 66%;10% | A087073 | 001 |
| MD-76R | | | | |
| + | LIEBEL-FLARSHEIM | 66%;10% | N019292 | 001 Sep 29, 1989 |
| RENOCAL-76 | | | | |
| | BRACCO | 66%;10% | A089347 | 001 Jun 01, 1988 |
| RENOGRAFIN-60 | | | | |
| | BRACCO | 52%;8% | N010040 | 006 |
| RENOGRAFIN-76 | | | | |
| + | BRACCO | 66%;10% | N010040 | 001 |
| RENOVIST | | | | |
| | BRACCO | 34.3%;35% | N010040 | 020 |
| RENOVIST II | | | | |
| | BRACCO | 28.5%;29.1% | N010040 | 019 |
| SOLUTION; ORAL, RECTAL | | | | |
| GASTROVIST | | | | |
| | BAYER HLTHCARE | 66%;10% | A087728 | 001 Sep 23, 1982 |

DIATRIZOATE MEGLUMINE; IODIPAMIDE MEGLUMINE

SOLUTION; INTRAUTERINE

| | | | | |
|------------|--------|-------------|---------|-----|
| SINOGRAFIN | | | | |
| + | BRACCO | 52.7%;26.8% | N011324 | 002 |

DIATRIZOATE SODIUM

FOR SOLUTION; ORAL, RECTAL

| | | | | |
|------------------------|----------------|------|---------|------------------|
| HYPaque | | | | |
| | GE HEALTHCARE | 100% | N011386 | 001 |
| INJECTABLE; INJECTION | | | | |
| HYPaque | | | | |
| | GE HEALTHCARE | 25% | N009561 | 003 |
| | | 50% | N009561 | 001 |
| MD-50 | | | | |
| | MALLINCKRODT | 50% | A087075 | 001 |
| UROVIST SODIUM 300 | | | | |
| | BAYER HLTHCARE | 50% | A087725 | 001 Sep 23, 1982 |
| SOLUTION; ORAL, RECTAL | | | | |
| HYPaque | | | | |
| | GE HEALTHCARE | 40% | N011386 | 003 |
| SOLUTION; URETERAL | | | | |
| HYPaque SODIUM 20% | | | | |
| | GE HEALTHCARE | 20% | N009561 | 002 |

DIAZEPAM

CAPSULE, EXTENDED RELEASE; ORAL

| | | | | |
|-------------|----------------------|----------------------|---------|------------------|
| VALRELEASE | | | | |
| | ROCHE | 15MG | N018179 | 001 |
| GEL; RECTAL | | | | |
| DIASTAT | | | | |
| + | VALEANT PHARMS NORTH | 5MG/ML (5MG/ML) ** | N020648 | 002 Jul 29, 1997 |
| + | | 10MG/2ML (5MG/ML) ** | N020648 | 003 Jul 29, 1997 |
| + | | 15MG/3ML (5MG/ML) ** | N020648 | 004 Jul 29, 1997 |
| + | | 20MG/4ML (5MG/ML) ** | N020648 | 005 Jul 29, 1997 |

INJECTABLE; INJECTION

| | | | | |
|----------|-------------------|-----------|---------|------------------|
| DIAZEPAM | | | | |
| | ABRAXIS PHARM | 5MG/ML | A070662 | 001 Jun 25, 1986 |
| | HOSPIRA | 5MG/ML | A071584 | 001 Oct 13, 1987 |
| | MARSAM PHARMS LLC | 5MG/ML | A072371 | 001 Jan 29, 1993 |
| | PARENTA PHARMS | 5MG/ML | A076815 | 001 Apr 15, 2004 |
| | US ARMY | 5MG/ML ** | N020124 | 001 Dec 05, 1990 |
| | WARNER CHILCOTT | 5MG/ML | A071613 | 001 Oct 22, 1987 |
| | | 5MG/ML | A071614 | 001 Oct 22, 1987 |
| | WATSON LABS | 5MG/ML | A070296 | 001 Feb 12, 1986 |
| | | 5MG/ML | A070911 | 001 Aug 28, 1986 |
| | | 5MG/ML | A070912 | 001 Aug 28, 1986 |
| | | 5MG/ML | A070930 | 001 Dec 01, 1986 |
| | WATSON LABS INC | 5MG/ML | A072370 | 001 Jan 29, 1993 |
| | | 5MG/ML | A072397 | 001 Jan 29, 1993 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIAZEPAM

INJECTABLE; INJECTION

DIAZEPAM

| | | | | |
|----------------------|--------|---------|-----|--------------|
| WEST-WARD PHARMS INT | 5MG/ML | A070311 | 001 | Dec 16, 1985 |
| | 5MG/ML | A070312 | 001 | Dec 16, 1985 |
| | 5MG/ML | A070313 | 001 | Dec 16, 1985 |
| | 5MG/ML | A071308 | 001 | Jul 17, 1987 |
| | 5MG/ML | A071309 | 001 | Jul 17, 1987 |
| | 5MG/ML | A071310 | 001 | Jul 17, 1987 |

DIZAC

| | | | | |
|----------------------|-----------|---------|-----|--------------|
| PHARMACIA AND UPJOHN | 5MG/ML ** | N019287 | 001 | Jun 18, 1993 |
|----------------------|-----------|---------|-----|--------------|

VALIUM

| | | | | |
|---------|-----------|---------|-----|--|
| + ROCHE | 5MG/ML ** | N016087 | 001 | |
|---------|-----------|---------|-----|--|

TABLET; ORAL

DIAZEPAM

| | | | | |
|----------------------|------|---------|-----|--------------|
| ACTAVIS ELIZABETH | 2MG | A070781 | 001 | Mar 19, 1986 |
| | 5MG | A070706 | 001 | Mar 19, 1986 |
| | 10MG | A070707 | 001 | Mar 19, 1986 |
| DAVA PHARMS INC | 2MG | A070228 | 002 | Sep 26, 1985 |
| | 5MG | A070228 | 003 | Sep 26, 1985 |
| | 10MG | A070228 | 001 | Sep 26, 1985 |
| DURAMED PHARMS BARR | 2MG | A070894 | 001 | Aug 27, 1986 |
| | 5MG | A070895 | 001 | Aug 27, 1986 |
| | 10MG | A070896 | 001 | Aug 27, 1986 |
| FERNDAL LABS | 2MG | A070903 | 001 | Apr 01, 1987 |
| | 5MG | A070904 | 001 | Apr 01, 1987 |
| | 10MG | A070905 | 001 | Apr 01, 1987 |
| HALSEY | 2MG | A070987 | 001 | Aug 15, 1986 |
| | 5MG | A070996 | 001 | Aug 15, 1986 |
| | 10MG | A070956 | 001 | Aug 15, 1986 |
| IVAX SUB TEVA PHARMS | 2MG | A070360 | 001 | Sep 04, 1985 |
| | 5MG | A070361 | 001 | Sep 04, 1985 |
| | 10MG | A070362 | 001 | Sep 04, 1985 |
| MARTEC USA LLC | 10MG | A072402 | 001 | Apr 25, 1989 |
| PIONEER PHARMS | 2MG | A070787 | 001 | Aug 02, 1988 |
| | 5MG | A070788 | 001 | Aug 02, 1988 |
| | 10MG | A070776 | 001 | Aug 02, 1988 |
| ROXANE | 2MG | A070356 | 001 | Jun 17, 1986 |
| | 5MG | A070357 | 001 | Jun 17, 1986 |
| | 10MG | A070358 | 001 | Jun 17, 1986 |
| TEVA PHARMS | 5MG | A070153 | 001 | Nov 01, 1985 |
| UPSHER SMITH LABS | 2MG | A070302 | 001 | Dec 20, 1985 |
| | 5MG | A070303 | 001 | Dec 20, 1985 |
| | 10MG | A070304 | 001 | Dec 20, 1985 |
| VIRTUS PHARMS | 2MG | A070462 | 001 | Feb 25, 1986 |
| | 5MG | A070463 | 001 | Feb 25, 1986 |
| | 10MG | A070464 | 001 | Feb 25, 1986 |
| WARNER CHILCOTT | 2MG | A070209 | 001 | Sep 04, 1985 |
| | 5MG | A070210 | 001 | Sep 04, 1985 |
| | 10MG | A070222 | 001 | Sep 04, 1985 |
| WATSON LABS | 2MG | A070456 | 001 | Nov 01, 1985 |
| | 5MG | A070457 | 001 | Nov 01, 1985 |
| | 10MG | A070458 | 001 | Nov 01, 1985 |

Q-PAM

| | | | | |
|------------------|------|---------|-----|--------------|
| QUANTUM PHARMICS | 2MG | A070423 | 001 | Dec 12, 1985 |
| | 2MG | A072431 | 001 | Apr 29, 1988 |
| | 5MG | A070424 | 001 | Dec 12, 1985 |
| | 5MG | A072432 | 001 | Apr 29, 1988 |
| | 10MG | A070425 | 001 | Dec 12, 1985 |
| | 10MG | A072433 | 001 | Apr 29, 1988 |

DIAZOXIDE

CAPSULE; ORAL

PROGLYCEM

| | | | | |
|--------------------|-------|---------|-----|--|
| TEVA BRANDED PHARM | 50MG | N017425 | 001 | |
| | 100MG | N017425 | 002 | |

INJECTABLE; INJECTION

DIAZOXIDE

| | | | | |
|---------------|---------|---------|-----|--------------|
| ABRAXIS PHARM | 15MG/ML | A071519 | 001 | Aug 26, 1987 |
|---------------|---------|---------|-----|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIAZOXIDE

| | | | |
|-----------------------|---------|--|-------------|
| INJECTABLE; INJECTION | | | |
| HYPERSTAT | | | |
| SCHERING | 15MG/ML | | N016996 001 |

DIBUCAINE HYDROCHLORIDE

| | | | |
|---------------------------|----------|--|-------------|
| INJECTABLE; INJECTION | | | |
| HEAVY SOLUTION NUPERCAINE | | | |
| NOVARTIS | 2.5MG/ML | | N006203 001 |

DICHLORPHENAMIDE

| | | | |
|-------------------|---------|--|-------------|
| TABLET; ORAL | | | |
| DARANIDE | | | |
| + STRONGBRIDGE US | 50MG ** | | N011366 001 |

DICLOFENAC POTASSIUM

| | | | |
|----------------------|---------|-------------|--------------|
| TABLET; ORAL | | | |
| CATAFLAM | | | |
| + NOVARTIS | 25MG ** | N020142 001 | Nov 24, 1993 |
| | 50MG ** | N020142 002 | Nov 24, 1993 |
| DICLOFENAC POTASSIUM | | | |
| SANDOZ | 50MG | A075582 001 | Feb 23, 2001 |
| SUN PHARM INDUSTRIES | 50MG | A075470 001 | Feb 21, 2002 |
| WATSON LABS TEVA | 50MG | A075152 001 | Nov 27, 1998 |

DICLOFENAC SODIUM

| | | | |
|--------------------------------|-----------------------|-------------|--------------|
| SOLUTION; INTRAVENOUS | | | |
| DYLOJECT | | | |
| + JAVELIN PHARMS INC | 37.5MG/ML (37.5MG/ML) | N022396 001 | Dec 23, 2014 |
| SOLUTION; TOPICAL | | | |
| PENNSAID | | | |
| + NUVO PHARMS INC | 1.5% ** | N020947 001 | Nov 04, 2009 |
| SOLUTION/DROPS; OPHTHALMIC | | | |
| DICLOFENAC SODIUM | | | |
| APOTEX INC | 0.1% | A077600 001 | Nov 13, 2008 |
| FALCON PHARMS | 0.1% | N020809 001 | May 04, 1998 |
| TABLET, DELAYED RELEASE; ORAL | | | |
| DICLOFENAC SODIUM | | | |
| ALLIED | 50MG | A074986 001 | Feb 26, 1999 |
| | 75MG | A074986 002 | Feb 26, 1999 |
| PLIVA | 50MG | A074432 002 | Jul 29, 1999 |
| | 75MG | A074432 003 | Jul 29, 1999 |
| ROXANE | 25MG | A074391 001 | Jun 29, 1995 |
| | 50MG | A074391 002 | Jun 29, 1995 |
| | 75MG | A074391 003 | Jun 29, 1995 |
| TEVA | 50MG | A074723 001 | Mar 30, 1999 |
| | 75MG | A074390 001 | Aug 15, 1996 |
| TEVA PHARMS | 25MG | A074459 001 | Jun 25, 1997 |
| | 50MG | A074459 002 | Jun 25, 1997 |
| | 75MG | A074459 003 | Jun 25, 1997 |
| VOLTAREN | | | |
| + NOVARTIS | 25MG ** | N019201 001 | Jul 28, 1988 |
| | 50MG ** | N019201 002 | Jul 28, 1988 |
| | 75MG ** | N019201 003 | Jul 28, 1988 |
| TABLET, EXTENDED RELEASE; ORAL | | | |
| DICLOFENAC SODIUM | | | |
| ACTAVIS ELIZABETH | 100MG | A075910 001 | Jan 07, 2002 |
| VOLTAREN-XR | | | |
| + NOVARTIS | 100MG ** | N020254 001 | Mar 08, 1996 |

DICLOXACILLIN SODIUM

| | | | |
|-----------------|---------------|-------------|--------------|
| CAPSULE; ORAL | | | |
| DYCILL | | | |
| GLAXOSMITHKLINE | EQ 250MG BASE | A060254 002 | |
| | EQ 250MG BASE | A062238 001 | |
| | EQ 500MG BASE | A060254 003 | |
| | EQ 500MG BASE | A062238 002 | |
| PATHOCIL | | | |
| WYETH AYERST | EQ 250MG BASE | N050011 002 | |
| | EQ 500MG BASE | N050011 003 | Mar 28, 1983 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DICLOXACILLIN SODIUM

FOR SUSPENSION;ORAL

DICLOXACILLIN SODIUM

APOTHECON

EQ 62.5MG BASE/5ML

A061455 001

DYNAPEN

APOTHECON

EQ 62.5MG BASE/5ML

N050337 002

PATHOCIL

WYETH AYERST

EQ 62.5MG BASE/5ML

N050092 001

DICUMAROL

CAPSULE;ORAL

DICUMAROL

LILLY

25MG

N005509 003

50MG

N005509 001

TABLET;ORAL

DICUMAROL

ABBVIE

25MG

N005545 003

50MG

N005545 004

100MG

N005545 005

DICYCLOMINE HYDROCHLORIDE

CAPSULE;ORAL

DICYCLOMINE HYDROCHLORIDE

PIONEER PHARMS

10MG

A089361 001 Jan 10, 1989

SUN PHARM INDUSTRIES

10MG

A084505 001 Oct 21, 1986

WATSON LABS

10MG

A083179 001 Feb 12, 1986

INJECTABLE;INJECTION

DICYCLOMINE HYDROCHLORIDE

WATSON LABS

10MG/ML

A080614 001 Feb 11, 1986

SYRUP;ORAL

BENTYL

+ APTALIS PHARMA US

10MG/5ML **

N007961 002 Oct 15, 1984

DICYCLOMINE HYDROCHLORIDE

ALPHARMA US PHARMS

10MG/5ML

A084479 001

TABLET;ORAL

DICYCLOMINE HYDROCHLORIDE

PIONEER PHARMS

20MG

A088585 001 Aug 20, 1986

SUN PHARM INDUSTRIES

20MG

A084600 001 Jul 29, 1985

WATSON LABS

20MG

A084361 001 Feb 06, 1986

DIDANOSINE

CAPSULE, DELAYED REL PELLETS;ORAL

DIDANOSINE

BARR

200MG

A077167 001 Dec 03, 2004

250MG

A077167 002 Dec 03, 2004

400MG

A077167 003 Dec 03, 2004

MYLAN PHARMS INC

125MG

A090788 001 Apr 08, 2010

200MG

A090788 002 Apr 08, 2010

250MG

A090788 003 Apr 08, 2010

400MG

A090788 004 Apr 08, 2010

FOR SOLUTION;ORAL

DIDANOSINE

AUROBINDO PHARMA

10MG/ML

A078112 001 Mar 08, 2007

VIDEX

BRISTOL MYERS SQUIBB

100MG/PACKET

N020155 003 Oct 09, 1991

167MG/PACKET

N020155 004 Oct 09, 1991

250MG/PACKET

N020155 005 Oct 09, 1991

375MG/PACKET

N020155 006 Oct 09, 1991

TABLET, CHEWABLE;ORAL

VIDEX

+ BRISTOL MYERS SQUIBB

25MG **

N020154 002 Oct 09, 1991

+

50MG **

N020154 003 Oct 09, 1991

+

100MG **

N020154 004 Oct 09, 1991

+

150MG **

N020154 005 Oct 09, 1991

+

200MG **

N020154 006 Oct 28, 1999

TABLET, FOR SUSPENSION;ORAL

DIDANOSINE

AUROBINDO

100MG

A077275 001 Aug 14, 2012

150MG

A077275 002 Aug 14, 2012

200MG

A077275 003 Aug 14, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIENESTROL

CREAM;VAGINAL

DIENESTROL

ORTHO MCNEIL PHARM 0.01% N006110 005

DV

SANOFI AVENTIS US 0.01% A083518 001

ESTRAGUARD

SOLVAY 0.01% A084436 001

SUPPOSITORY;VAGINAL

DV

SANOFI AVENTIS US 0.7MG A083517 001

DIETHYLCARBAMAZINE CITRATE

TABLET;ORAL

HETRAZAN

LEDERLE 50MG N006459 001

DIETHYLPROPION HYDROCHLORIDE

TABLET;ORAL

DIETHYLPROPION HYDROCHLORIDE

CHARTWELL RX 25MG A088267 001 Aug 25, 1983

25MG A088268 001 Aug 25, 1983

EPIC PHARMA LLC 25MG A040828 001 Nov 05, 2008

SANDOZ 25MG A085916 001

TEVA 25MG A088642 001 Sep 20, 1984

UCB INC 25MG A085544 001

WATSON LABS 25MG A085741 001

TENUATE

SANOFI AVENTIS US 25MG N017668 001

TEPANIL

3M 25MG N011673 001

TABLET, EXTENDED RELEASE;ORAL

TENUATE

SANOFI AVENTIS US 75MG N017669 001

TEPANIL TEN-TAB

3M 75MG N017956 001

DIETHYLSTILBESTROL

INJECTABLE;INJECTION

STILBESTROL

BRISTOL MYERS SQUIBB 0.2MG/ML N004056 003

0.5MG/ML N004056 004

1MG/ML N004056 005

5MG/ML N004056 006

SUPPOSITORY;VAGINAL

DIETHYLSTILBESTROL

LILLY 0.1MG N004040 001

0.5MG N004040 002

STILBESTROL

BRISTOL MYERS SQUIBB 0.1MG N004056 001

0.5MG N004056 002

TABLET;ORAL

DIETHYLSTILBESTROL

LILLY 0.1MG N004041 002

0.5MG N004041 003

1MG N004041 004

5MG N004041 005

STILBESTROL

TABLICAPS 0.5MG A083004 001

1MG A083002 001

5MG A083006 001

STILBETIN

BRISTOL MYERS SQUIBB 0.1MG N004056 007

0.25MG N004056 017

0.5MG N004056 008

1MG N004056 009

5MG N004056 010

TABLET, DELAYED RELEASE;ORAL

DIETHYLSTILBESTROL

LILLY 0.1MG N004039 002

0.25MG N004039 005

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIETHYLSTILBESTROLTABLET, DELAYED RELEASE;ORAL
DIETHYLSTILBESTROL

| | |
|-------|-------------|
| 0.5MG | N004039 003 |
| 1MG | N004039 004 |
| 5MG | N004039 006 |

STILBESTROL

| | | |
|-----------|-------|-------------|
| TABLICAPS | 0.5MG | A083003 001 |
| | 1MG | A083005 001 |
| | 5MG | A083007 001 |

STILBETIN

| | | |
|----------------------|-------|-------------|
| BRISTOL MYERS SQUIBB | 0.1MG | N004056 011 |
| | 0.5MG | N004056 012 |
| | 1MG | N004056 013 |
| | 5MG | N004056 014 |

DIETHYLSTILBESTROL DIPHOSPHATE

INJECTABLE; INJECTION

STILPHOSTROL

| | | |
|--------------|-----------|-------------|
| BAYER PHARMS | 250MG/5ML | N010010 001 |
|--------------|-----------|-------------|

TABLET; ORAL

STILPHOSTROL

| | | |
|--------------|------|-------------|
| BAYER PHARMS | 50MG | N010010 002 |
|--------------|------|-------------|

DIFLORASONE DIACETATE

CREAM; TOPICAL

DIFLORASONE DIACETATE

| | | | |
|----------------|-------|-------------|--------------|
| FOUGERA PHARMS | 0.05% | A075187 001 | Mar 30, 1998 |
|----------------|-------|-------------|--------------|

FLORONE

| | | | |
|----------------------|----------|-------------|--|
| PHARMACIA AND UPJOHN | 0.05% ** | N017741 001 | |
|----------------------|----------|-------------|--|

FLORONE E

| | | | |
|----------------------|-------|-------------|--------------|
| PHARMACIA AND UPJOHN | 0.05% | N019259 001 | Aug 28, 1985 |
|----------------------|-------|-------------|--------------|

PSORCON

| | | | |
|---------------------|----------|-------------|--------------|
| + TARO PHARMS NORTH | 0.05% ** | N020205 001 | Nov 20, 1992 |
|---------------------|----------|-------------|--------------|

OINTMENT; TOPICAL

PSORCON

| | | | |
|------------------------|-------|-------------|--------------|
| + PHARMACIA AND UPJOHN | 0.05% | N019260 001 | Aug 28, 1985 |
|------------------------|-------|-------------|--------------|

PSORCON E

| | | | |
|----------------------|-------|-------------|--|
| PHARMACIA AND UPJOHN | 0.05% | N017994 001 | |
|----------------------|-------|-------------|--|

DIFLUNISAL

TABLET; ORAL

DIFLUNISAL

| | | | |
|----------------|-------|-------------|--------------|
| ANI PHARMS INC | 500MG | A074604 001 | Jun 10, 1996 |
| PUREPAC PHARM | 250MG | A074285 001 | May 07, 1996 |
| | 500MG | A074285 002 | May 07, 1996 |
| SOCORRO | 250MG | A073562 001 | Nov 27, 1992 |
| | 500MG | A073563 001 | Nov 27, 1992 |
| TEVA | 250MG | A073679 001 | Jul 31, 1992 |
| WATSON LABS | 250MG | A074400 001 | Jul 17, 1997 |
| | 500MG | A074400 002 | Jul 17, 1997 |

DOLOBID

| | | | |
|---------|----------|-------------|--------------|
| + MERCK | 250MG ** | N018445 001 | Apr 19, 1982 |
|---------|----------|-------------|--------------|

| | | | |
|---|----------|-------------|--------------|
| + | 500MG ** | N018445 002 | Apr 19, 1982 |
|---|----------|-------------|--------------|

DIGITOXIN

INJECTABLE; INJECTION

CRYSTODIGIN

| | | | |
|-------|----------|-------------|--|
| LILLY | 0.2MG/ML | A084100 005 | |
|-------|----------|-------------|--|

DIGOXIN

CAPSULE; ORAL

LANOXICAPS

| | | | |
|---------------------|--------|-------------|--------------|
| GLAXOSMITHKLINE LLC | 0.05MG | N018118 002 | Jul 26, 1982 |
| | 0.1MG | N018118 003 | Jul 26, 1982 |
| | 0.15MG | N018118 004 | Sep 24, 1984 |
| | 0.2MG | N018118 001 | Jul 26, 1982 |

INJECTABLE; INJECTION

DIGOXIN

| | | | |
|---------------|-----------|-------------|--------------|
| ABRAXIS PHARM | 0.25MG/ML | A083217 001 | |
| HOSPIRA | 0.25MG/ML | A040093 001 | May 16, 1996 |
| | 0.25MG/ML | A040206 001 | Aug 28, 1998 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIGOXIN

INJECTABLE; INJECTION

DIGOXIN

WYETH AYERST 0.25MG/ML

A084386 001

DIGOXIN PEDIATRIC

HOSPIRA 0.1MG/ML

A040092 001 Apr 25, 1996

TABLET; ORAL

LANOXIN

+ CONCORDIA PHARMS INC 0.1875MG
0.375MG
0.5MG

N020405 003 Sep 30, 1997
N020405 005 Sep 30, 1997
N020405 006 Sep 30, 1997

DIHYDROERGOTAMINE MESYLATE; HEPARIN SODIUM; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

EMBOLEX

NOVARTIS 0.5MG/0.5ML; 2,500
UNITS/0.5ML; 5.33MG/0.5ML
0.5MG/0.7ML; 5,000
UNITS/0.7ML; 7.46MG/0.7ML

N018885 001 Nov 30, 1984
N018885 002 Nov 30, 1984

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CARDIZEM SR

+ BIOVAIL 60MG **
+ 90MG **
+ 120MG **
+ 180MG **

N019471 001 Jan 23, 1989
N019471 002 Jan 23, 1989
N019471 003 Jan 23, 1989
N019471 004 Jan 23, 1989

DILACOR XR

+ ALLERGAN SALES LLC 120MG **
+ 180MG **
+ 240MG **

N020092 001 May 29, 1992
N020092 002 May 29, 1992
N020092 003 May 29, 1992

DILT-CD

APOTEX 120MG
180MG
240MG
300MG

A076151 001 May 20, 2004
A076151 002 May 20, 2004
A076151 003 May 20, 2004
A076151 004 May 20, 2004

DILTIAZEM HYDROCHLORIDE

ACTAVIS LABS FL INC 120MG
180MG
240MG

A074852 001 Oct 10, 1997
A074852 002 Oct 10, 1997
A074852 003 Oct 10, 1997

BIOVAIL 60MG
90MG
120MG
120MG
180MG
240MG
300MG
360MG
420MG

A074845 001 Sep 15, 1999
A074845 002 Sep 15, 1999
A074845 003 Sep 15, 1999
N020939 001 Jan 28, 2000
N020939 002 Jan 28, 2000
N020939 003 Jan 28, 2000
N020939 004 Jan 28, 2000
N020939 005 Sep 14, 2001
N020939 006 Sep 14, 2001

NESHER PHARMS 120MG
180MG
240MG
300MG
360MG
420MG

A076563 002 Sep 12, 2006
A076563 003 Sep 12, 2006
A076563 004 Sep 12, 2006
A076563 005 Sep 12, 2006
A076563 006 Sep 12, 2006

TEVA 60MG
90MG
120MG

A074079 001 Nov 30, 1993
A074079 002 Nov 30, 1993
A074079 003 Nov 30, 1993

INJECTABLE; INJECTION

CARDIZEM

BIOVAIL 100MG/VIAL **
+ BIOVAIL LABS INTL 5MG/ML **
+ 25MG/VIAL **

N020792 001 Sep 05, 1997
N020027 001 Oct 24, 1991
N020027 003 Aug 18, 1995

DILTIAZEM HYDROCHLORIDE

HOSPIRA 5MG/ML
5MG/ML
MYLAN LABS LTD 5MG/ML
TEVA PHARMS USA 5MG/ML

A075004 001 Feb 16, 2000
A075106 001 Apr 29, 1999
A075375 001 Sep 30, 1999
A074894 001 Aug 26, 1997

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DILTIAZEM HYDROCHLORIDE

TABLET; ORAL

DILTIAZEM HYDROCHLORIDE

| | | | |
|----------------------|-------|-------------|--------------|
| APOTHECON | 30MG | A074051 001 | Mar 31, 1993 |
| | 60MG | A074051 002 | Mar 31, 1993 |
| | 90MG | A074051 003 | Mar 31, 1993 |
| | 120MG | A074051 004 | Mar 31, 1993 |
| CHARTWELL MOLECULES | 30MG | A074093 001 | Nov 05, 1992 |
| | 60MG | A074093 002 | Nov 05, 1992 |
| | 90MG | A074093 003 | Nov 05, 1992 |
| | 120MG | A074093 004 | Nov 05, 1992 |
| IVAX SUB TEVA PHARMS | 30MG | A074168 001 | Mar 03, 1995 |
| | 60MG | A074168 002 | Mar 03, 1995 |
| | 90MG | A074168 003 | Mar 03, 1995 |
| | 120MG | A074168 004 | Mar 03, 1995 |
| TEVA | 30MG | A074084 001 | Feb 25, 1994 |
| | 60MG | A074084 002 | Feb 25, 1994 |
| TEVA PHARMS | 30MG | A074067 001 | Nov 05, 1992 |
| | 60MG | A074067 002 | Nov 05, 1992 |
| | 90MG | A074067 003 | Nov 05, 1992 |
| | 120MG | A074067 004 | Nov 05, 1992 |

DILTIAZEM MALATE

TABLET, EXTENDED RELEASE; ORAL

TIAMATE

| | | | |
|-------|------------------------|-------------|--------------|
| MERCK | EQ 120MG HYDROCHLORIDE | N020506 001 | Oct 04, 1996 |
| | EQ 180MG HYDROCHLORIDE | N020506 002 | Oct 04, 1996 |
| | EQ 240MG HYDROCHLORIDE | N020506 003 | Oct 04, 1996 |

DILTIAZEM MALATE; ENALAPRIL MALEATE

TABLET, EXTENDED RELEASE; ORAL

TECZEM

| | | | |
|---------|-----------------------------|-------------|--------------|
| BIOVAIL | EQ 180MG HYDROCHLORIDE; 5MG | N020507 001 | Oct 04, 1996 |
|---------|-----------------------------|-------------|--------------|

DIMENHYDRINATE

INJECTABLE; INJECTION

DIMENHYDRINATE

| | | | |
|------------------|---------|-------------|--|
| BAXTER HLTHCARE | 50MG/ML | A084767 001 | |
| WATSON LABS | 50MG/ML | A083531 001 | |
| WATSON LABS TEVA | 50MG/ML | A080615 001 | |
| WYETH AYERST | 50MG/ML | A084316 001 | |

LIQUID; ORAL

DIMENHYDRINATE

| | | | |
|------|------------|-------------|--|
| ALRA | 12.5MG/4ML | A080715 001 | |
|------|------------|-------------|--|

TABLET; ORAL

DIMENHYDRINATE

| | | | |
|-------------------|------|-------------|--|
| HEATHER | 50MG | A080841 001 | |
| NEXGEN PHARMA INC | 50MG | A085985 001 | |
| WATSON LABS | 50MG | A085166 001 | |

DIMYRISTOYL LECITHIN; PERFLEXANE

INJECTABLE; INTRAVENOUS

IMAGENT

| | | | |
|------------------|---------------------------|-------------|--------------|
| VESSELON SPV LLC | 0.92MG/VIAL; 0.092MG/VIAL | N021191 001 | May 31, 2002 |
|------------------|---------------------------|-------------|--------------|

DINOPROST TROMETHAMINE

INJECTABLE; INJECTION

PROSTIN F2 ALPHA

| | | | |
|----------------------|----------------|-------------|--|
| PHARMACIA AND UPJOHN | EQ 5MG BASE/ML | N017434 001 | |
|----------------------|----------------|-------------|--|

DIPHEMANIL METHYLSULFATE

TABLET; ORAL

PRANTAL

| | | | |
|----------|-------|-------------|--|
| SCHERING | 100MG | N008114 004 | |
|----------|-------|-------------|--|

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

BENADRYL

| | | | |
|-------------|------|-------------|--|
| MCNEIL CONS | 25MG | N005845 007 | |
| | 50MG | N005845 001 | |

DIPHENHYDRAMINE HYDROCHLORIDE

| | | | |
|------|------|-------------|--|
| ALRA | 25MG | A080519 004 | |
| | 50MG | A080519 003 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

| | | | | |
|----------------------|------|---------|-----|--------------|
| ANABOLIC | 50MG | A083275 | 001 | |
| ELKINS SINN | 25MG | A085701 | 001 | |
| | 50MG | A085701 | 002 | |
| FOSUN PHARMA | 25MG | A080832 | 001 | |
| | 25MG | A080845 | 002 | |
| | 50MG | A080832 | 002 | |
| | 50MG | A080845 | 001 | |
| HALSEY | 50MG | A087914 | 001 | Jun 04, 1984 |
| HEATHER | 25MG | A084524 | 001 | |
| | 50MG | A083953 | 001 | |
| HIKMA INTL PHARMS | 50MG | A083567 | 001 | |
| IMPAX LABS | 25MG | A080807 | 001 | |
| | 50MG | A080807 | 002 | |
| IVAX SUB TEVA PHARMS | 25MG | A080762 | 001 | |
| | 50MG | A080762 | 002 | |
| LANNETT | 25MG | A080868 | 001 | |
| | 50MG | A080868 | 002 | |
| LEDERLE | 25MG | A086874 | 001 | |
| | 50MG | A086875 | 001 | |
| LNK | 25MG | A087977 | 001 | Jan 27, 1983 |
| | 50MG | A087978 | 001 | Jan 27, 1983 |
| MK LABS | 25MG | A083087 | 001 | |
| | 50MG | A083087 | 002 | |
| MUTUAL PHARM | 25MG | A084506 | 001 | |
| NEWTRON PHARMS | 25MG | A086543 | 001 | |
| | 50MG | A086544 | 001 | |
| NEXGEN PHARMA INC | 25MG | A083634 | 001 | |
| PERRIGO | 25MG | A083061 | 001 | |
| | 50MG | A083061 | 002 | |
| PIONEER PHARMS | 25MG | A089101 | 001 | Dec 20, 1985 |
| | 50MG | A088880 | 001 | Dec 20, 1985 |
| PUREPAC PHARM | 25MG | A085156 | 001 | |
| | 50MG | A085150 | 001 | |
| PVT FORM | 25MG | A083027 | 001 | |
| | 50MG | A083027 | 002 | |
| ROXANE | 50MG | A080635 | 001 | |
| SUN PHARM INDUSTRIES | 25MG | A089488 | 001 | Jan 02, 1987 |
| | 50MG | A089489 | 001 | Jan 02, 1987 |
| SUPERPHARM | 25MG | A089040 | 001 | May 15, 1985 |
| | 50MG | A089041 | 001 | May 15, 1985 |
| TEVA | 25MG | A085874 | 001 | |
| | 50MG | A085874 | 002 | |
| VALEANT PHARM INTL | 25MG | A080596 | 001 | |
| | 50MG | A080592 | 001 | |
| VANGARD | 25MG | A088034 | 001 | Oct 27, 1982 |
| | 50MG | A087630 | 001 | |
| WATSON LABS | 25MG | A080728 | 001 | |
| | 25MG | A083797 | 001 | |
| | 25MG | A085138 | 001 | |
| | 50MG | A080727 | 001 | |
| | 50MG | A083797 | 002 | |
| | 50MG | A085083 | 001 | |
| WHITEWORTH TOWN PLSN | 25MG | A083441 | 001 | |
| | 50MG | A080800 | 001 | |

ELIXIR; ORAL

BELIX

| | | | | |
|--------|------------|---------|-----|--------------|
| HALSEY | 12.5MG/5ML | A086586 | 001 | Oct 03, 1983 |
|--------|------------|---------|-----|--------------|

BENADRYL

| | | | | |
|-------------|------------|---------|-----|--|
| MCNEIL CONS | 12.5MG/5ML | N005845 | 004 | |
|-------------|------------|---------|-----|--|

DIBENIL

| | | | | |
|-------|------------|---------|-----|--------------|
| CENCI | 12.5MG/5ML | A088304 | 001 | Dec 16, 1983 |
|-------|------------|---------|-----|--------------|

DIPHEN

| | | | | |
|------------|------------|---------|-----|--|
| USL PHARMA | 12.5MG/5ML | A084640 | 001 | |
|------------|------------|---------|-----|--|

DIPHENHYDRAMINE HYDROCHLORIDE

| | | | | |
|-------|------------|---------|-----|--|
| BUNDY | 12.5MG/5ML | A083674 | 001 | |
|-------|------------|---------|-----|--|

| | | | | |
|-------|------------|---------|-----|--------------|
| CENCI | 12.5MG/5ML | A087941 | 001 | Dec 17, 1982 |
|-------|------------|---------|-----|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIPHENHYDRAMINE HYDROCHLORIDE

ELIXIR; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

| | | | | |
|---------------|------------|---------|-----|--------------|
| KV PHARM | 12.5MG/5ML | A085621 | 001 | |
| LANNETT | 12.5MG/5ML | A080939 | 002 | |
| LEDERLE | 12.5MG/5ML | A086937 | 001 | |
| MK LABS | 12.5MG/5ML | A083088 | 002 | |
| NASKA | 12.5MG/5ML | A088680 | 001 | May 31, 1985 |
| PERRIGO | 12.5MG/5ML | A083063 | 001 | |
| PUREPAC PHARM | 12.5MG/5ML | A083237 | 001 | Jan 25, 1982 |
| PVT FORM | 12.5MG/5ML | A085287 | 001 | |
| ROXANE | 12.5MG/5ML | A080643 | 001 | |

HYDRAMINE

| | | | | |
|--------------------|------------|---------|-----|--|
| ALPHARMA US PHARMS | 12.5MG/5ML | A080763 | 002 | |
|--------------------|------------|---------|-----|--|

INJECTABLE; INJECTION

BENADRYL

| | | | | |
|----------------------------|-------------|------------|---------|-----|
| MCNEIL CONS | 10MG/ML | N006146 | 001 | |
| + | 50MG/ML ** | N006146 | 002 | |
| BENADRYL PRESERVATIVE FREE | | | | |
| + | MCNEIL CONS | 50MG/ML ** | N009486 | 001 |

DIPHENHYDRAMINE HYDROCHLORIDE

| | | | | |
|--------------------|---------|---------|-----|--|
| BEL MAR | 10MG/ML | A080822 | 001 | |
| EUROHLTH INTL SARL | 50MG/ML | A083183 | 001 | |
| LYPHOMED | 10MG/ML | A087066 | 001 | |
| WATSON LABS | 10MG/ML | A083533 | 001 | |
| WATSON LABS TEVA | 10MG/ML | A080873 | 001 | |
| | 50MG/ML | A080873 | 002 | |
| WYETH AYERST | 50MG/ML | A080577 | 001 | |

DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE

| | | | | |
|------------------|---------|---------|-----|--|
| ABRAXIS PHARM | 50MG/ML | A080586 | 002 | |
| INTL MEDICATION | 50MG/ML | A084094 | 001 | |
| WATSON LABS TEVA | 50MG/ML | A080873 | 003 | |

SYRUP; ORAL

ANTITUSSIVE

| | | | | |
|-------------------------------|------------|---------|-----|--------------|
| PERRIGO | 12.5MG/5ML | A071292 | 001 | Apr 10, 1987 |
| BELDIN | | | | |
| HALSEY | 12.5MG/5ML | A089179 | 001 | Jun 05, 1986 |
| BENYLIN | | | | |
| PARKE DAVIS | 12.5MG/5ML | N006514 | 004 | |
| DIPHEN | | | | |
| MORTON GROVE | 12.5MG/5ML | A070118 | 001 | Oct 01, 1985 |
| DIPHENHYDRAMINE HYDROCHLORIDE | | | | |
| ALPHARMA US PHARMS | 12.5MG/5ML | A070497 | 001 | Apr 25, 1989 |
| CUMBERLAND SWAN | 12.5MG/5ML | A073611 | 001 | Aug 20, 1992 |
| HI TECH PHARMA | 12.5MG/5ML | A072416 | 001 | Sep 28, 1990 |
| HYDRAMINE | | | | |
| ALPHARMA US PHARMS | 12.5MG/5ML | A070205 | 001 | Jan 28, 1986 |
| SILPHEN | | | | |
| LANNETT CO INC | 12.5MG/5ML | A072646 | 001 | Feb 27, 1992 |
| VICKS FORMULA 44 | | | | |
| WARNER CHILCOTT | 12.5MG/5ML | A070524 | 001 | Jan 14, 1987 |

DIPHENHYDRAMINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

BENYLIN

| | | | | |
|-------------|----------------------|---------|-----|--------------|
| PARKE DAVIS | 12.5MG/5ML; 30MG/5ML | N019014 | 001 | Jun 11, 1985 |
|-------------|----------------------|---------|-----|--------------|

DIPHENIDOL HYDROCHLORIDE

TABLET; ORAL

VONTROL

| | | | | |
|-----------------|--------------|---------|-----|--|
| GLAXOSMITHKLINE | EQ 25MG BASE | N016033 | 001 | |
|-----------------|--------------|---------|-----|--|

DIPHENYLPYRALINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

HISPRIL

| | | | | |
|-----------------|-----|---------|-----|--|
| GLAXOSMITHKLINE | 5MG | N011945 | 001 | |
|-----------------|-----|---------|-----|--|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DIPIVEFRIN HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

AKPRO

AKORN 0.1% A074382 001 Sep 29, 1995

DIPIVEFRIN HYDROCHLORIDE

BAUSCH AND LOMB 0.1% A074188 001 May 19, 1995

FALCON PHARMS 0.1% A073636 001 Jun 30, 1994

PROPINE

ALLERGAN 0.1% N018239 001

DIPYRIDAMOLE

INJECTABLE; INJECTION

DIPYRIDAMOLE

HOSPIRA 5MG/ML A074601 001 Dec 19, 1997

MYLAN LABS LTD 5MG/ML A075769 001 Nov 27, 2002

TEVA PHARMS USA 5MG/ML A074952 001 Nov 26, 1997

IV PERSANTINE

+ BOEHRINGER INGELHEIM 5MG/ML ** N019817 001 Dec 13, 1990

TABLET; ORAL

DIPYRIDAMOLE

ANI PHARMS INC 25MG A086944 002 Apr 16, 1991

50MG A086944 001 Feb 25, 1992

75MG A086944 003 Feb 25, 1992

GLENMARK GENERICS 25MG A088999 001 Feb 05, 1991

50MG A089000 001 Feb 05, 1991

75MG A089001 001 Feb 05, 1991

LANNETT CO INC 25MG A040898 001 Apr 23, 2008

50MG A040898 002 Apr 23, 2008

75MG A040898 003 Apr 23, 2008

OXFORD PHARMS 25MG A040542 001 Apr 21, 2006

50MG A040542 002 Apr 21, 2006

75MG A040542 003 Apr 21, 2006

PUREPAC PHARM 25MG A089425 001 Jul 12, 1990

50MG A089426 001 Jul 12, 1990

75MG A089427 001 Jul 12, 1990

WATSON LABS 50MG A087160 001 Jun 07, 1996

DIRITHROMYCIN

TABLET, DELAYED RELEASE; ORAL

DYNABAC

LILLY RES LABS 250MG N050678 001 Jun 19, 1995

DISOPYRAMIDE PHOSPHATE

CAPSULE; ORAL

DISOPYRAMIDE PHOSPHATE

AUROLIFE PHARMA LLC EQ 100MG BASE A070470 001 Dec 10, 1985

EQ 150MG BASE A070471 001 Dec 10, 1985

INTERPHARM EQ 100MG BASE A071190 001 Jan 15, 1987

EQ 150MG BASE A071191 001 Jan 15, 1987

IVAX SUB TEVA PHARMS EQ 100MG BASE A070186 001 Nov 18, 1985

EQ 150MG BASE A070187 001 Nov 18, 1985

MYLAN EQ 100MG BASE A070138 001 Jun 14, 1985

EQ 150MG BASE A070139 001 Jun 14, 1985

SUN PHARM INDUSTRIES EQ 100MG BASE A070351 001 Dec 17, 1985

EQ 150MG BASE A070352 001 Dec 17, 1985

SUPERPHARM EQ 100MG BASE A070940 001 Feb 09, 1987

EQ 150MG BASE A070941 001 Feb 09, 1987

WATSON LABS EQ 100MG BASE A070240 001 Feb 02, 1986

EQ 150MG BASE A070241 001 Feb 02, 1986

CAPSULE, EXTENDED RELEASE; ORAL

DISOPYRAMIDE PHOSPHATE

NESHER PHARMS EQ 150MG BASE A071200 001 Dec 15, 1987

DISULFIRAM

TABLET; ORAL

ANTABUSE

+ TEVA WOMENS 250MG ** N007883 003

+ 500MG ** N007883 002

DISULFIRAM

PAR PHARM 250MG A088792 001 Aug 14, 1984

500MG A088793 001 Aug 14, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DISULFIRAM

TABLET; ORAL

DISULFIRAM

| | | | | |
|------------------|-------|---------|-----|--------------|
| WATSON LABS | 250MG | A086889 | 001 | |
| | 250MG | A087973 | 001 | Aug 05, 1983 |
| | 500MG | A087974 | 001 | Aug 05, 1983 |
| WATSON LABS TEVA | 500MG | A086890 | 001 | |

DIVALPROEX SODIUM

TABLET, DELAYED RELEASE; ORAL

DEPAKOTE CP

| | | | | |
|--------|---------------|---------|-----|--------------|
| ABBOTT | EQ 250MG BASE | N019794 | 001 | Jul 11, 1990 |
| | EQ 500MG BASE | N019794 | 002 | Jul 11, 1990 |

DIVALPROEX SODIUM

| | | | | |
|-------|------------------------|---------|-----|--------------|
| MYLAN | EQ 125MG VALPROIC ACID | A077254 | 001 | Jul 29, 2008 |
| | EQ 250MG VALPROIC ACID | A077254 | 002 | Jul 29, 2008 |
| | EQ 500MG VALPROIC ACID | A077254 | 003 | Jul 29, 2008 |

TABLET, EXTENDED RELEASE; ORAL

DIVALPROEX SODIUM

| | | | | |
|------------------|------------------------|---------|-----|--------------|
| G AND W LABS INC | EQ 500MG VALPROIC ACID | A078700 | 001 | Aug 03, 2009 |
|------------------|------------------------|---------|-----|--------------|

DOBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOBUTAMINE HYDROCHLORIDE

| | | | | |
|---------------------|----------------------|---------|-----|--------------|
| BAXTER HLTHCARE | EQ 12.5MG BASE/ML | A074381 | 001 | Sep 26, 1996 |
| HOSPIRA | EQ 1.25GM BASE/100ML | A074634 | 001 | Sep 27, 1996 |
| LUITPOLD | EQ 12.5MG BASE/ML | A074545 | 001 | Jun 25, 1998 |
| TELIGENT PHARMA INC | EQ 12.5MG BASE/ML | A074098 | 001 | Feb 21, 1995 |
| TEVA PARENTERAL | EQ 12.5MG BASE/ML | A074206 | 001 | Oct 19, 1993 |
| WATSON LABS | EQ 12.5MG BASE/ML | A074114 | 001 | Nov 30, 1993 |
| WATSON LABS INC | EQ 12.5MG BASE/ML | A074279 | 001 | Feb 18, 1998 |
| | EQ 12.5MG BASE/ML | A074995 | 001 | Mar 31, 1998 |

DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5%

| | | | | |
|---------|---------------------|---------|-----|--------------|
| HOSPIRA | EQ 50MG BASE/100ML | N020269 | 001 | Oct 19, 1993 |
| | EQ 100MG BASE/100ML | N020269 | 002 | Oct 19, 1993 |
| | EQ 200MG BASE/100ML | N020269 | 003 | Oct 19, 1993 |

DOBUTREX

| | | | | |
|---------|-------------------|---------|-----|--|
| + LILLY | EQ 12.5MG BASE/ML | N017820 | 002 | |
|---------|-------------------|---------|-----|--|

DOCETAXEL

INJECTABLE; INJECTION

DOCEFREZ

| | | | | |
|---------------------|-----------|---------|-----|--------------|
| + SUN PHARMA GLOBAL | 20MG/VIAL | N022534 | 001 | May 03, 2011 |
| | 80MG/VIAL | N022534 | 002 | May 03, 2011 |

DOCETAXEL

| | | | | |
|-------------------|----------------------|---------|-----|--------------|
| + ACCORD HLTHCARE | 20MG/0.5ML (40MG/ML) | N201195 | 001 | Jun 08, 2011 |
| | 80MG/2ML (40MG/ML) | N201195 | 002 | Jun 08, 2011 |
| + APOTEX INC | 20MG/0.5ML (40MG/ML) | N022312 | 001 | Jan 11, 2012 |
| | 80MG/2ML (40MG/ML) | N022312 | 002 | Jan 11, 2012 |
| + HOSPIRA INC | 120MG/6ML (20MG/ML) | N022234 | 006 | Jun 23, 2016 |
| + PFIZER LABS | 20MG/2ML (10MG/ML) | N202356 | 001 | Mar 13, 2014 |
| | 80MG/8ML (10MG/ML) | N202356 | 002 | Mar 13, 2014 |
| | 130MG/13ML (10MG/ML) | N202356 | 003 | Mar 13, 2014 |
| | 200MG/20ML (10MG/ML) | N202356 | 004 | Mar 13, 2014 |

TAXOTERE

| | | | | |
|---------------------|------------|---------|-----|--------------|
| + SANOFI AVENTIS US | 40MG/ML ** | N020449 | 001 | May 14, 1996 |
|---------------------|------------|---------|-----|--------------|

DOLASETRON MESYLATE

INJECTABLE; INJECTION

ANZEMET

| | | | | |
|---------------------|--------------------------|---------|-----|--------------|
| + US PHARM HOLDINGS | 12.5MG/0.625ML (20MG/ML) | N020624 | 002 | Sep 11, 1997 |
| | 100MG/5ML (20MG/ML) | N020624 | 001 | Sep 11, 1997 |
| | 500MG/25ML (20MG/ML) | N020624 | 003 | Dec 11, 2001 |

TABLET; ORAL

ANZEMET

| | | | | |
|---------------------|-------|---------|-----|--------------|
| + US PHARM HOLDINGS | 50MG | N020623 | 001 | Sep 11, 1997 |
| | 100MG | N020623 | 002 | Sep 11, 1997 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DONEPEZIL HYDROCHLORIDE

SOLUTION; ORAL

ARICEPT

EISAI INC 5MG/5ML N021719 001 Oct 18, 2004

TABLET; ORAL

DONEPEZIL HYDROCHLORIDE

ACCORD HLTHCARE 5MG A201335 001 Aug 29, 2011

10MG A201335 002 Aug 29, 2011

APOTEX 5MG A078841 001 Jun 02, 2011

10MG A078841 002 Jun 02, 2011

HIKMA PHARMS 5MG A090247 001 May 31, 2011

10MG A090247 002 May 31, 2011

SUN PHARM INDS LTD 5MG A076786 001 Nov 26, 2010

10MG A076786 002 Nov 26, 2010

TABLET, ORALLY DISINTEGRATING; ORAL

ARICEPT ODT

+ EISAI INC 5MG N021720 001 Oct 18, 2004

+ 10MG N021720 002 Oct 18, 2004

DONEPEZIL HYDROCHLORIDE

BARR 5MG A078388 002 Nov 26, 2010

10MG A078388 001 Nov 26, 2010

SUN PHARM INDUSTRIES 5MG A077975 002 Dec 11, 2009

10MG A077975 001 Dec 11, 2009

ZYDUS PHARMS USA INC 5MG A090175 001 May 10, 2011

10MG A090175 002 May 10, 2011

DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MEMANTINE HYDROCHLORIDE AND DONEPEZIL HYDROCHLORIDE

AMNEAL PHARMS 10MG; 14MG A208328 001 Jan 27, 2017

10MG; 28MG A208328 002 Jan 27, 2017

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOPAMINE HYDROCHLORIDE

ABBOTT 40MG/ML A070656 001 Jan 24, 1989

80MG/ML A070657 001 Jan 24, 1989

ABRAXIS PHARM 40MG/ML A070012 001 Jun 12, 1985

40MG/ML A070058 001 Mar 20, 1985

80MG/ML A070013 001 Jun 12, 1985

80MG/ML A070059 001 Mar 20, 1985

160MG/ML A070364 001 Dec 04, 1985

BAXTER HLTHCARE 40MG/ML N018398 001

80MG/ML N018398 002 Mar 22, 1982

HOSPIRA 40MG/ML A074403 001 May 23, 1996

IGI LABS INC 40MG/ML A070087 001 Oct 23, 1985

80MG/ML A070089 001 Oct 23, 1985

80MG/ML A070090 001 Oct 23, 1985

80MG/ML A070091 001 Oct 23, 1985

160MG/ML A070092 001 Oct 23, 1985

160MG/ML A070093 001 Oct 23, 1985

160MG/ML A070094 001 Oct 23, 1985

INTL MEDICATION 40MG/ML N018014 001

LUITPOLD 160MG/ML A070826 001 Feb 11, 1987

LYPHOMED 40MG/ML N018549 001 Mar 11, 1983

SMITH AND NEPHEW 40MG/ML A070011 001 Aug 29, 1985

40MG/ML A070046 001 Aug 29, 1985

80MG/ML A070047 001 Aug 29, 1985

TELIGENT 40MG/ML N018656 001 Jun 28, 1983

TEVA PARENTERAL 40MG/ML A072999 001 Oct 23, 1991

80MG/ML A073000 001 Oct 23, 1991

WARNER CHILCOTT 40MG/ML A070558 001 Sep 20, 1985

40MG/ML N018138 001

80MG/ML A070559 001 Sep 20, 1985

DOPAMINE HYDROCHLORIDE IN DEXTROSE 5%

HOSPIRA 1.6MG/ML N020542 001 Aug 30, 1995

INTROPIN

HOSPIRA 40MG/ML N017395 001

80MG/ML N017395 002

160MG/ML N017395 003

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DORIPENEM

INJECTABLE; INTRAVENOUS

DORIBAX

| | | | | | |
|---|--------------|------------|---------|-----|--------------|
| + | SHIONOGI INC | 250MG/VIAL | N022106 | 002 | Oct 05, 2010 |
| + | | 500MG/VIAL | N022106 | 001 | Oct 12, 2007 |

DORZOLAMIDE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE

| | | | | | |
|--|-----------------|------------|---------|-----|--------------|
| | APOTEX INC | EQ 2% BASE | A078395 | 001 | Oct 28, 2008 |
| | TEVA PHARMS | EQ 2% BASE | A078756 | 001 | Dec 04, 2008 |
| | WATSON LABS INC | EQ 2% BASE | A202053 | 001 | Sep 11, 2014 |
| | ZAMBON SPA | EQ 2% BASE | A091034 | 001 | Dec 04, 2013 |

DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE

| | | | | | |
|--|-----------------|-------------------------|---------|-----|--------------|
| | APOTEX INC | EQ 2% BASE;EQ 0.5% BASE | A078201 | 001 | Oct 28, 2008 |
| | LANNETT CO INC | EQ 2% BASE;EQ 0.5% BASE | A201998 | 001 | Dec 17, 2014 |
| | WATSON LABS INC | EQ 2% BASE;EQ 0.5% BASE | A202054 | 001 | Sep 03, 2014 |
| | ZAMBON SPA | EQ 2% BASE;EQ 0.5% BASE | A091180 | 001 | Dec 04, 2013 |

DOXACURIUM CHLORIDE

INJECTABLE; INJECTION

NUROMAX

| | | | | | |
|--|--------|----------------|---------|-----|--------------|
| | ABBVIE | EQ 1MG BASE/ML | N019946 | 001 | Mar 07, 1991 |
|--|--------|----------------|---------|-----|--------------|

DOXAPRAM HYDROCHLORIDE

INJECTABLE; INJECTION

DOXAPRAM HYDROCHLORIDE

| | | | | | |
|--|-------------|---------|---------|-----|--------------|
| | WATSON LABS | 20MG/ML | A073529 | 001 | Jan 30, 1992 |
|--|-------------|---------|---------|-----|--------------|

DOXAZOSIN MESYLATE

TABLET; ORAL

DOXAZOSIN MESYLATE

| | | | | | |
|--|----------------------|-------------|---------|-----|--------------|
| | ACTAVIS ELIZABETH | EQ 1MG BASE | A075574 | 001 | Oct 18, 2000 |
| | | EQ 2MG BASE | A075574 | 002 | Oct 18, 2000 |
| | | EQ 4MG BASE | A075574 | 003 | Oct 18, 2000 |
| | | EQ 8MG BASE | A075574 | 004 | Oct 18, 2000 |
| | GENPHARM | EQ 1MG BASE | A075466 | 001 | Oct 18, 2000 |
| | | EQ 2MG BASE | A075466 | 002 | Oct 18, 2000 |
| | | EQ 4MG BASE | A075466 | 003 | Oct 18, 2000 |
| | | EQ 8MG BASE | A075466 | 004 | Oct 18, 2000 |
| | IVAX SUB TEVA PHARMS | EQ 1MG BASE | A075453 | 001 | Oct 18, 2000 |
| | | EQ 2MG BASE | A075453 | 002 | Oct 18, 2000 |
| | | EQ 4MG BASE | A075453 | 003 | Oct 18, 2000 |
| | | EQ 8MG BASE | A075453 | 004 | Oct 18, 2000 |
| | NESHER PHARMS | EQ 1MG BASE | A075609 | 001 | Oct 18, 2000 |
| | | EQ 2MG BASE | A075609 | 002 | Oct 18, 2000 |
| | | EQ 4MG BASE | A075609 | 003 | Oct 18, 2000 |
| | | EQ 8MG BASE | A075609 | 004 | Oct 18, 2000 |
| | TEVA | EQ 1MG BASE | A075353 | 001 | Jan 12, 2001 |
| | | EQ 2MG BASE | A075353 | 002 | Jan 12, 2001 |
| | | EQ 4MG BASE | A075353 | 003 | Jan 12, 2001 |
| | | EQ 8MG BASE | A075353 | 004 | Jan 12, 2001 |
| | WATSON LABS INC | EQ 1MG BASE | A075426 | 001 | Oct 18, 2000 |
| | | EQ 2MG BASE | A075426 | 002 | Oct 18, 2000 |
| | | EQ 4MG BASE | A075426 | 003 | Oct 18, 2000 |
| | | EQ 8MG BASE | A075426 | 004 | Oct 18, 2000 |
| | YAOPHARMA CO LTD | EQ 1MG BASE | A075646 | 001 | Oct 18, 2000 |
| | | EQ 2MG BASE | A075646 | 002 | Oct 18, 2000 |
| | | EQ 4MG BASE | A075646 | 003 | Oct 18, 2000 |
| | | EQ 8MG BASE | A075646 | 004 | Oct 18, 2000 |

DOXEPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPIN HYDROCHLORIDE

| | | | | | |
|--|-----------------|---------------|---------|-----|--------------|
| | DAVA PHARMS INC | EQ 10MG BASE | A071685 | 001 | Jan 05, 1988 |
| | | EQ 25MG BASE | A071686 | 001 | Jan 05, 1988 |
| | | EQ 50MG BASE | A071673 | 001 | Jan 05, 1988 |
| | | EQ 75MG BASE | A071674 | 001 | Jan 05, 1988 |
| | | EQ 100MG BASE | A071675 | 001 | Jan 05, 1988 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXEPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPIN HYDROCHLORIDE

| | | | |
|-----------------------|--------------------|-------------|--------------|
| | EQ 150MG BASE | A071676 001 | Jan 05, 1988 |
| NEW RIVER | EQ 10MG BASE | N016987 001 | |
| | EQ 25MG BASE | N016987 002 | |
| | EQ 50MG BASE | N016987 003 | |
| | EQ 75MG BASE | N016987 006 | |
| | EQ 100MG BASE | N016987 004 | |
| | EQ 150MG BASE | N016987 007 | Apr 13, 1987 |
| PAR PHARM | EQ 10MG BASE | A071697 001 | Nov 09, 1987 |
| | EQ 25MG BASE | A071437 001 | Nov 09, 1987 |
| | EQ 75MG BASE | A071608 001 | Nov 09, 1987 |
| PUREPAC PHARM | EQ 10MG BASE | A073054 001 | Dec 28, 1990 |
| | EQ 25MG BASE | A072109 001 | Dec 28, 1990 |
| | EQ 50MG BASE | A073055 001 | Dec 28, 1990 |
| | EQ 75MG BASE | A072386 001 | Sep 08, 1988 |
| | EQ 100MG BASE | A072110 001 | Sep 08, 1988 |
| | EQ 150MG BASE | A072387 001 | Sep 08, 1988 |
| QUANTUM PHARMICS | EQ 10MG BASE | A070972 001 | Sep 29, 1987 |
| | EQ 25MG BASE | A070973 001 | Sep 29, 1987 |
| | EQ 50MG BASE | A070931 001 | Sep 29, 1987 |
| | EQ 75MG BASE | A070932 001 | Sep 29, 1987 |
| | EQ 100MG BASE | A072375 001 | Mar 15, 1989 |
| | EQ 150MG BASE | A072376 001 | Mar 15, 1989 |
| SANDOZ | EQ 10MG BASE | A071487 001 | Mar 02, 1987 |
| | EQ 25MG BASE | A070827 001 | May 15, 1986 |
| | EQ 50MG BASE | A070828 001 | May 15, 1986 |
| | EQ 75MG BASE | A070825 001 | May 15, 1986 |
| | EQ 100MG BASE | A071562 001 | Mar 02, 1987 |
| SUN PHARM INDUSTRIES | EQ 25MG BASE | A071502 001 | Feb 18, 1988 |
| | EQ 50MG BASE | A071653 001 | Feb 18, 1988 |
| | EQ 75MG BASE | A071654 001 | Feb 18, 1988 |
| | EQ 100MG BASE | A071521 001 | Feb 18, 1988 |
| WATSON LABS | EQ 10MG BASE | A070952 001 | Mar 04, 1987 |
| | EQ 10MG BASE | A071485 001 | Apr 30, 1987 |
| | EQ 10MG BASE | A072985 001 | Mar 29, 1991 |
| | EQ 25MG BASE | A070953 001 | May 15, 1986 |
| | EQ 25MG BASE | A071486 001 | Apr 30, 1987 |
| | EQ 25MG BASE | A072986 001 | Mar 29, 1991 |
| | EQ 50MG BASE | A070954 001 | May 15, 1986 |
| | EQ 50MG BASE | A071238 001 | Apr 30, 1987 |
| | EQ 75MG BASE | A071326 001 | Apr 30, 1987 |
| | EQ 75MG BASE | A071763 001 | Feb 09, 1988 |
| | EQ 100MG BASE | A070955 001 | May 15, 1986 |
| | EQ 100MG BASE | A071239 001 | Apr 30, 1987 |
| | EQ 150MG BASE | A071764 001 | Feb 09, 1988 |
| WATSON LABS TEVA | EQ 50MG BASE | A072987 001 | Mar 29, 1991 |
| SINEQUAN | | | |
| + PFIZER | EQ 10MG BASE ** | N016798 003 | |
| + | EQ 25MG BASE ** | N016798 001 | |
| + | EQ 50MG BASE ** | N016798 002 | |
| + | EQ 75MG BASE ** | N016798 006 | |
| + | EQ 100MG BASE ** | N016798 005 | |
| + | EQ 150MG BASE ** | N016798 007 | |
| CONCENTRATE; ORAL | | | |
| DOXEPIN HYDROCHLORIDE | | | |
| PHARM ASSOC | EQ 10MG BASE/ML | A075924 001 | Jan 15, 2004 |
| SINEQUAN | | | |
| + PFIZER | EQ 10MG BASE/ML ** | N017516 001 | |
| TABLET; ORAL | | | |
| DOXEPIN HYDROCHLORIDE | | | |
| ACTAVIS ELIZABETH | EQ 3MG BASE | A201951 001 | Jul 26, 2013 |
| | EQ 6MG BASE | A201951 002 | Jul 26, 2013 |
| MYLAN PHARMS INC | EQ 3MG BASE | A202337 001 | Jan 20, 2016 |
| | EQ 6MG BASE | A202337 002 | Jan 20, 2016 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ADRIAMYCIN PFS

| | | | |
|----------------------|-------------|-------------|--------------|
| PHARMACIA AND UPJOHN | 2MG/ML | A063165 001 | Jan 30, 1991 |
| | 200MG/100ML | A063165 002 | Jan 30, 1991 |

DOXORUBICIN HYDROCHLORIDE

| | | | |
|----------------------|------------|-------------|--------------|
| ALVOGEN INC | 2MG/ML | A065515 001 | Nov 08, 2012 |
| MYLAN LABS LTD | 10MG/VIAL | A200170 001 | Oct 28, 2011 |
| PHARMACIA AND UPJOHN | 10MG/VIAL | N050467 001 | |
| | 20MG/VIAL | N050467 003 | May 20, 1985 |
| | 50MG/VIAL | N050467 002 | |
| | 150MG/VIAL | N050467 004 | Jul 22, 1987 |
| SANDOZ INC | 2MG/ML | A200146 001 | Jul 18, 2012 |

RUBEX

| | | | |
|----------------------|------------|-------------|--------------|
| BRISTOL MYERS SQUIBB | 10MG/VIAL | A062926 001 | Apr 13, 1989 |
| | 50MG/VIAL | A062926 002 | Apr 13, 1989 |
| | 100MG/VIAL | A062926 003 | Apr 13, 1989 |

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

| | | | |
|-------------|---------------|-------------|--------------|
| PAR PHARM | EQ 75MG BASE | A065055 004 | Apr 18, 2005 |
| SANDOZ INC | EQ 50MG BASE | A065032 001 | Jun 30, 2000 |
| | EQ 100MG BASE | A065032 002 | Jun 30, 2000 |
| WATSON LABS | EQ 50MG BASE | A065041 001 | Apr 28, 2000 |
| | EQ 100MG BASE | A065041 002 | Apr 28, 2000 |

FOR SUSPENSION; ORAL

DOXYCHEL

| | | | |
|----------|------------------|-------------|--|
| RACHELLE | EQ 25MG BASE/5ML | A061720 001 | |
|----------|------------------|-------------|--|

TABLET; ORAL

DOXYCYCLINE

| | | | |
|----------------------|---------------|-------------|--------------|
| SANDOZ INC | EQ 50MG BASE | A065353 001 | Nov 27, 2006 |
| | EQ 75MG BASE | A065353 002 | Nov 27, 2006 |
| | EQ 100MG BASE | A065353 003 | Nov 27, 2006 |
| SUN PHARM INDUSTRIES | EQ 50MG BASE | A065471 001 | Apr 17, 2009 |
| | EQ 75MG BASE | A065471 002 | Apr 17, 2009 |
| | EQ 100MG BASE | A065471 003 | Apr 17, 2009 |

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

ACTICLATE CAP

+ AQUA PHARMS

| | | | |
|--|--------------|-------------|--------------|
| | EQ 75MG BASE | N208253 001 | Apr 26, 2016 |
|--|--------------|-------------|--------------|

DOXY-LEMMON

| | | | |
|------|---------------|-------------|--------------|
| TEVA | EQ 50MG BASE | A062497 001 | Aug 23, 1984 |
| | EQ 100MG BASE | A062497 002 | Jun 15, 1984 |

DOXYCYCLINE HYCLATE

| | | | |
|-------------------|---------------|-------------|--------------|
| HALSEY | EQ 50MG BASE | A062119 002 | May 24, 1985 |
| | EQ 100MG BASE | A062119 001 | May 24, 1985 |
| HEATHER | EQ 50MG BASE | A062463 001 | Dec 07, 1983 |
| | EQ 100MG BASE | A062463 002 | Dec 07, 1983 |
| HIKMA INTL PHARMS | EQ 20MG BASE | A065103 001 | May 13, 2005 |
| INTERPHARM | EQ 50MG BASE | A062763 001 | Sep 02, 1988 |
| | EQ 100MG BASE | A062763 002 | Sep 02, 1988 |
| MUTUAL PHARM | EQ 50MG BASE | A062418 001 | Jan 28, 1983 |
| | EQ 100MG BASE | A062418 002 | Jan 28, 1983 |
| PAR PHARM | EQ 50MG BASE | A062434 001 | Oct 19, 1984 |
| | EQ 100MG BASE | A062442 001 | Dec 22, 1983 |
| PVT FORM | EQ 50MG BASE | A062631 001 | Jul 24, 1986 |
| | EQ 100MG BASE | A062631 002 | Jul 24, 1986 |
| RANBAXY | EQ 50MG BASE | A062479 001 | Dec 23, 1983 |
| | EQ 100MG BASE | A062479 002 | Dec 23, 1983 |
| SUPERPHARM | EQ 50MG BASE | A062469 001 | Oct 31, 1984 |
| | EQ 100MG BASE | A062469 002 | Oct 31, 1984 |
| WARNER CHILCOTT | EQ 50MG BASE | A062594 001 | Dec 05, 1985 |
| | EQ 100MG BASE | A062594 002 | Dec 05, 1985 |
| WATSON LABS | EQ 50MG BASE | A061717 001 | |
| | EQ 50MG BASE | A062142 001 | |
| | EQ 100MG BASE | A061717 002 | |
| | EQ 100MG BASE | A062142 002 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXYCYCLINE HYCLATE

| | | | |
|-------------------------------|-----------------------|-------------|--------------|
| CAPSULE;ORAL | | | |
| PERIOSTAT | | | |
| + COLLAGENEX | EQ 20MG BASE ** | N050744 001 | Sep 30, 1998 |
| VIBRAMYCIN | | | |
| + PFIZER | EQ 50MG BASE ** | N050007 001 | |
| CAPSULE, COATED PELLETS;ORAL | | | |
| DOXYCYCLINE HYCLATE | | | |
| PLIVA | EQ 100MG BASE | A063187 001 | Jun 30, 1992 |
| CAPSULE, DELAYED RELEASE;ORAL | | | |
| DORYX | | | |
| + MAYNE PHARMA INTL | EQ 75MG BASE | N050582 002 | Aug 13, 2001 |
| + | EQ 100MG BASE | N050582 001 | Jul 22, 1985 |
| WARNER CHILCOTT | EQ 100MG BASE | A062653 001 | Oct 30, 1985 |
| DOXYCYCLINE HYCLATE | | | |
| MEDICIS | EQ 75MG BASE | A065281 001 | Dec 21, 2005 |
| | EQ 100MG BASE | A065281 002 | Dec 21, 2005 |
| INJECTABLE;INJECTION | | | |
| DOXYCHEL HYCLATE | | | |
| RACHELLE | EQ 100MG BASE/VIAL | A061953 001 | |
| DOXYCYCLINE | | | |
| WEST-WARD PHARMS INT | EQ 100MG BASE/VIAL | A062450 001 | Oct 27, 1983 |
| | EQ 200MG BASE/VIAL | A062450 002 | Oct 27, 1983 |
| | EQ 200MG BASE/VIAL | A062569 002 | Mar 09, 1988 |
| DOXYCYCLINE HYCLATE | | | |
| WEST-WARD PHARMS INT | EQ 100MG BASE/VIAL | A062992 001 | Feb 16, 1989 |
| | EQ 200MG BASE/VIAL | A062992 002 | Feb 16, 1989 |
| VIBRAMYCIN | | | |
| + PFIZER | EQ 100MG BASE/VIAL ** | N050442 002 | |
| + | EQ 200MG BASE/VIAL ** | N050442 001 | |
| TABLET;ORAL | | | |
| DOXY-LEMMON | | | |
| TEVA | EQ 100MG BASE | A062581 001 | Mar 15, 1985 |
| DOXYCYCLINE HYCLATE | | | |
| EPIC PHARMA LLC | EQ 20MG BASE | A065182 001 | May 13, 2005 |
| HEATHER | EQ 100MG BASE | A062462 001 | May 11, 1983 |
| INTERPHARM | EQ 100MG BASE | A062764 001 | Sep 02, 1988 |
| MUTUAL PHARM | EQ 100MG BASE | A062391 001 | Sep 30, 1982 |
| SUPERPHARM | EQ 100MG BASE | A062494 001 | Feb 20, 1985 |
| VINTAGE PHARMS | EQ 100MG BASE | A062538 001 | Apr 07, 1986 |
| WARNER CHILCOTT | EQ 100MG BASE | A062593 001 | Aug 28, 1985 |
| WATSON LABS | EQ 50MG BASE | A062392 001 | Mar 31, 1983 |
| | EQ 100MG BASE | A062392 002 | Mar 31, 1983 |
| LYMEPAK | | | |
| + CHARTWELL PHARMA | EQ 100MG BASE | N209844 001 | Jun 15, 2018 |
| PERIOSTAT | | | |
| + GALDERMA LABS LP | EQ 20MG BASE ** | N050783 001 | Feb 02, 2001 |
| VIBRA-TABS | | | |
| + PFIZER | EQ 100MG BASE ** | N050533 001 | |
| TABLET, DELAYED RELEASE;ORAL | | | |
| DORYX | | | |
| + MAYNE PHARMA | EQ 80MG BASE | N050795 004 | Apr 11, 2013 |
| DORYX MPC | | | |
| + MAYNE PHARMA | EQ 60MG BASE ** | N050795 007 | May 20, 2016 |
| DOXYCYCLINE HYCLATE | | | |
| IMPAX LABS INC | EQ 75MG BASE | A090505 001 | Dec 28, 2010 |
| | EQ 100MG BASE | A090505 002 | Dec 28, 2010 |
| MYLAN | EQ 80MG BASE | A090431 004 | Apr 29, 2016 |
| <u>DOXYLAMINE SUCCINATE</u> | | | |
| CAPSULE;ORAL | | | |
| UNISOM | | | |
| PFIZER | 25MG | N019440 001 | Feb 05, 1986 |
| TABLET;ORAL | | | |
| DECAPRYN | | | |
| SANOFI AVENTIS US | 12.5MG | N006412 015 | |
| | 25MG | N006412 014 | |
| DOXY-SLEEP-AID | | | |
| PAR PHARM | 25MG | A070156 001 | Jul 02, 1987 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DOXYLAMINE SUCCINATE

TABLET; ORAL

DOXYLAMINE SUCCINATE

| | | | | |
|------------------|------|---------|-----|--------------|
| COPLY PHARM | 25MG | A088900 | 002 | Feb 12, 1988 |
| QUANTUM PHARMICS | 25MG | A088603 | 001 | Aug 07, 1984 |

DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

BENDECTIN

| | | | | |
|-------------------|--------------|---------|-----|--|
| SANOFI AVENTIS US | 10MG;10MG ** | N010598 | 002 | |
|-------------------|--------------|---------|-----|--|

DROMOSTANOLONE PROPIONATE

INJECTABLE; INJECTION

DROLBAN

| | | | | |
|-------|---------|---------|-----|--|
| LILLY | 50MG/ML | N012936 | 001 | |
|-------|---------|---------|-----|--|

DRONABINOL

CAPSULE; ORAL

DRONABINOL

| | | | | |
|--------------|-------|---------|-----|--------------|
| INSYS THERAP | 2.5MG | A078501 | 001 | Aug 19, 2011 |
| | 5MG | A078501 | 002 | Aug 19, 2011 |
| | 10MG | A078501 | 003 | Aug 19, 2011 |

DROPERIDOL

INJECTABLE; INJECTION

DROPERIDOL

| | | | | |
|------------------|----------|---------|-----|--------------|
| ABRAXIS PHARM | 2.5MG/ML | A070992 | 001 | Nov 17, 1986 |
| | 2.5MG/ML | A070993 | 001 | Nov 17, 1986 |
| ASTRAZENECA | 2.5MG/ML | A072018 | 001 | Oct 20, 1988 |
| HOSPIRA | 2.5MG/ML | A071645 | 001 | Apr 07, 1988 |
| | 2.5MG/ML | A072272 | 001 | Aug 31, 1995 |
| IGI LABS INC | 2.5MG/ML | A072019 | 001 | Oct 19, 1988 |
| | 2.5MG/ML | A072020 | 001 | Oct 19, 1988 |
| | 2.5MG/ML | A072021 | 001 | Oct 19, 1988 |
| LUITPOLD | 2.5MG/ML | A072335 | 001 | Oct 24, 1988 |
| SMITH AND NEPHEW | 2.5MG/ML | A071750 | 001 | Sep 06, 1988 |
| SOLOPAK | 2.5MG/ML | A071754 | 001 | Sep 06, 1988 |
| | 2.5MG/ML | A071755 | 001 | Sep 06, 1988 |
| WATSON LABS | 2.5MG/ML | A073520 | 001 | Nov 27, 1991 |
| | 2.5MG/ML | A073521 | 001 | Nov 27, 1991 |
| | 2.5MG/ML | A073523 | 001 | Nov 27, 1991 |

DROPERIDOL; FENTANYL CITRATE

INJECTABLE; INJECTION

FENTANYL CITRATE AND DROPERIDOL

| | | | | |
|-------------|----------------------------|---------|-----|--------------|
| ASTRAZENECA | 2.5MG/ML;EQ 0.05MG BASE/ML | A072026 | 001 | Apr 13, 1989 |
| | 2.5MG/ML;EQ 0.05MG BASE/ML | A072027 | 001 | Apr 13, 1989 |
| | 2.5MG/ML;EQ 0.05MG BASE/ML | A072028 | 001 | Apr 13, 1989 |
| HOSPIRA | 2.5MG/ML;EQ 0.05MG BASE/ML | A071982 | 001 | May 04, 1988 |
| INNOVAR | | | | |
| AKORN MFG | 2.5MG/ML;EQ 0.05MG BASE/ML | N016049 | 001 | |

DUTASTERIDE

CAPSULE; ORAL

DUTASTERIDE

| | | | | |
|------------------|-------|---------|-----|--------------|
| MYLAN PHARMS INC | 0.5MG | A203241 | 001 | Jun 14, 2016 |
|------------------|-------|---------|-----|--------------|

DYCLONINE HYDROCHLORIDE

SOLUTION; TOPICAL

DYCLONE

| | | | | |
|---|-------------|---------|---------|-----|
| + | ASTRAZENECA | 0.5% ** | N009925 | 002 |
| + | | 1% ** | N009925 | 001 |

DYDROGESTERONE

TABLET; ORAL

GYNOREST

| | | | | |
|--------|---------|---------|-----|--|
| SOLVAY | 5MG ** | N017388 | 001 | |
| | 10MG ** | N017388 | 002 | |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

DYPHYLLINE

| | | | |
|-----------------------|------------|--|-------------|
| ELIXIR; ORAL | | | |
| NEOTHYLLINE | | | |
| TEVA | 160MG/15ML | | N007794 003 |
| INJECTABLE; INJECTION | | | |
| NEOTHYLLINE | | | |
| TEVA | 250MG/ML | | N009088 001 |
| TABLET; ORAL | | | |
| DILOR | | | |
| SAVAGE LABS | 200MG | | A084514 001 |
| DILOR-400 | | | |
| SAVAGE LABS | 400MG | | A084751 001 |
| LUFYLLIN | | | |
| MYLAN SPECIALITY LP | 200MG | | A084566 001 |
| | 400MG | | A084566 002 |
| NEOTHYLLINE | | | |
| TEVA | 200MG | | N007794 001 |
| | 400MG | | N007794 002 |

ECHOTHIOPHATE IODIDE

| | | | |
|--------------------------|-------|--|-------------|
| FOR SOLUTION; OPHTHALMIC | | | |
| PHOSPHOLINE IODIDE | | | |
| WYETH PHARMS | 0.03% | | N011963 002 |
| | 0.06% | | N011963 004 |
| | 0.25% | | N011963 003 |

EDETATE CALCIUM DISODIUM

| | | | |
|----------------------------|-------|--|-------------|
| TABLET; ORAL | | | |
| CALCIUM DISODIUM VERSENATE | | | |
| MEDICIS | 500MG | | N008922 002 |

EDROPHONIUM CHLORIDE

| | | | |
|----------------------------------------|------------|--|--------------------------|
| INJECTABLE; INJECTION | | | |
| EDROPHONIUM CHLORIDE | | | |
| HOSPIRA | 10MG/ML | | A040131 001 Feb 24, 1998 |
| WATSON LABS | 10MG/ML | | A040044 001 Mar 20, 1996 |
| EDROPHONIUM CHLORIDE PRESERVATIVE FREE | | | |
| WATSON LABS | 10MG/ML | | A040043 001 Mar 20, 1996 |
| ENLON | | | |
| MYLAN INSTITUTIONAL | 10MG/ML | | A088873 001 Aug 06, 1985 |
| REVERSOL | | | |
| ORGANON USA INC | 10MG/ML | | A089624 001 May 13, 1988 |
| TENSILON | | | |
| + TELIGENT | 10MG/ML ** | | N007959 001 |
| TENSILON PRESERVATIVE FREE | | | |
| + TELIGENT | 10MG/ML ** | | N007959 002 |

EFAVIRENZ

| | | | |
|------------------------|----------|--|--------------------------|
| CAPSULE; ORAL | | | |
| SUSTIVA | | | |
| + BRISTOL MYERS SQUIBB | 100MG ** | | N020972 002 Sep 17, 1998 |
| TABLET; ORAL | | | |
| SUSTIVA | | | |
| + BRISTOL MYERS SQUIBB | 300MG ** | | N021360 001 Feb 01, 2002 |

EFAVIRENZ; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

| | | | |
|---------------------------------------------------------|---------------------|--|--------------------------|
| TABLET; ORAL | | | |
| EFAVIRENZ, LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE | | | |
| AUROBINDO PHARMA LTD | 600MG; 300MG; 300MG | | N022343 001 Aug 15, 2018 |

EFLORNITHINE HYDROCHLORIDE

| | | | |
|-----------------------|----------|--|--------------------------|
| INJECTABLE; INJECTION | | | |
| ORNIDYL | | | |
| SANOFI AVENTIS US | 200MG/ML | | N019879 002 Nov 28, 1990 |

ELVITEGRAVIR

| | | | |
|-----------------------|-------|--|--------------------------|
| TABLET; ORAL | | | |
| VITEKTA | | | |
| + GILEAD SCIENCES INC | 85MG | | N203093 001 Sep 24, 2014 |
| + GILEAD SCIENCES INC | 150MG | | N203093 002 Sep 24, 2014 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ENALAPRIL MALEATE

TABLET; ORAL

ENALAPRIL MALEATE

| | | | |
|----------------------|-------|-------------|--------------|
| APOTHECON | 2.5MG | A075583 001 | Aug 22, 2000 |
| | 5MG | A075583 002 | Aug 22, 2000 |
| | 10MG | A075583 003 | Aug 22, 2000 |
| | 20MG | A075583 004 | Aug 22, 2000 |
| IVAX SUB TEVA PHARMS | 2.5MG | A075482 001 | Aug 22, 2000 |
| | 5MG | A075482 002 | Aug 22, 2000 |
| | 10MG | A075482 003 | Aug 22, 2000 |
| | 20MG | A075482 004 | Aug 22, 2000 |
| KRKA DD NOVO MESTO | 2.5MG | A075370 001 | Aug 22, 2000 |
| | 5MG | A075370 002 | Aug 22, 2000 |
| | 10MG | A075369 001 | Aug 22, 2000 |
| | 20MG | A075369 002 | Aug 22, 2000 |
| MYLAN | 2.5MG | A075472 001 | Aug 22, 2000 |
| | 2.5MG | A075480 001 | Aug 22, 2000 |
| | 5MG | A075472 002 | Aug 22, 2000 |
| | 10MG | A075472 003 | Aug 22, 2000 |
| | 20MG | A075472 004 | Aug 22, 2000 |
| | 20MG | A075480 004 | Aug 22, 2000 |
| SANDOZ | 2.5MG | A075048 001 | Aug 22, 2000 |
| | 5MG | A075048 002 | Aug 22, 2000 |
| | 10MG | A075048 003 | Aug 22, 2000 |
| | 20MG | A075048 004 | Aug 22, 2000 |
| SANDOZ INC | 2.5MG | A075621 001 | Aug 22, 2000 |
| | 5MG | A075621 002 | Aug 22, 2000 |
| | 10MG | A075621 003 | Aug 22, 2000 |
| | 20MG | A075621 004 | Aug 22, 2000 |
| SUN PHARM INDS LTD | 2.5MG | A075556 001 | Aug 22, 2000 |
| | 5MG | A075556 002 | Aug 22, 2000 |
| | 10MG | A075556 003 | Aug 22, 2000 |
| | 20MG | A075556 004 | Aug 22, 2000 |
| WATSON LABS | 2.5MG | A075501 001 | Aug 22, 2000 |
| | 5MG | A075501 002 | Aug 22, 2000 |
| | 10MG | A075501 003 | Aug 22, 2000 |
| | 20MG | A075501 004 | Aug 22, 2000 |

ENALAPRIL MALEATE; FELODIPINE

TABLET, EXTENDED RELEASE; ORAL

LEXXEL

| | | | |
|-------------|------------|-------------|--------------|
| ASTRAZENECA | 5MG; 2.5MG | N020668 002 | Oct 28, 1998 |
| | 5MG; 5MG | N020668 001 | Dec 27, 1996 |

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE

| | | | |
|----------------------|-------------|-------------|--------------|
| IVAX SUB TEVA PHARMS | 5MG; 12.5MG | A075736 001 | Mar 25, 2003 |
| | 10MG; 25MG | A075736 002 | Mar 25, 2003 |
| UPSHER SMITH LABS | 5MG; 12.5MG | A076116 001 | Sep 19, 2001 |
| | 10MG; 25MG | A076116 002 | Sep 19, 2001 |

ENALAPRILAT

INJECTABLE; INJECTION

ENALAPRILAT

| | | | |
|--------------------------------|--------------|-------------|--------------|
| HOSPIRA | 1.25MG/ML | A075456 001 | Aug 22, 2000 |
| | 1.25MG/ML | A075571 001 | Aug 22, 2000 |
| VASOTEC + BIOVAIL LABS INTL | 1.25MG/ML ** | N019309 001 | Feb 09, 1988 |

ENFLURANE

LIQUID; INHALATION

ENFLURANE

| | | | |
|----------------------------|-------|-------------|--------------|
| ABBOTT | 99.9% | A070803 001 | Sep 08, 1987 |
| PIRAMAL CRITICAL | 99.9% | A074396 001 | Jul 29, 1994 |
| ETHRANE BAXTER HLTHCARE | 99.9% | N017087 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ENOXACIN

TABLET; ORAL

PENETREX

SANOFI AVENTIS US

200MG

N019616 004 Dec 31, 1991

400MG

N019616 005 Dec 31, 1991

ENOXAPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

LOVENOX (PRESERVATIVE FREE)

+ SANOFI AVENTIS US

90MG/0.6ML (150MG/ML) **

N020164 006 Jun 02, 2000

ENTACAPONE

TABLET; ORAL

ENTACAPONE

MYLAN PHARMS INC

200MG

A202394 001 May 13, 2013

EPINEPHRINE

AEROSOL, METERED; INHALATION

BRONKAID MIST

STERLING

0.25MG/INH

N016803 001

EPINEPHRINE

ARMSTRONG PHARMS

0.2MG/INH

A087907 001 May 23, 1984

PRIMATENE MIST

WYETH CONS

0.2MG/INH

N016126 001

INJECTABLE; INJECTION

SUS-PHRINE SULFITE FREE

FOREST LABS

1.5MG/AMP

N007942 003 Feb 05, 1999

5MG/ML

N007942 001

INJECTABLE; INTRAMUSCULAR

EPI E Z PEN JR

MYLAN SPECIALITY LP

0.15MG/DELIVERY

N019430 004 Aug 03, 1995

EPIPEN E Z PEN

MYLAN SPECIALITY LP

0.3MG/DELIVERY

N019430 003 Aug 03, 1995

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

TWINJECT 0.15

IMPAX

EQ 0.15MG/DELIVERY

N020800 002 May 28, 2004

TWINJECT 0.3

IMPAX

EQ 0.3MG/DELIVERY

N020800 001 May 30, 2003

EPINEPHRINE BITARTRATE

AEROSOL, METERED; INHALATION

BRONITIN MIST

WYETH CONS

0.3MG/INH

N016126 002

MEDIHALER-EPI

3M

0.3MG/INH

N010374 003

EPINEPHRINE BITARTRATE; ETIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DURANEST

+ ASTRAZENECA

0.005MG/ML; 1% **

N017751 006

+

0.005MG/ML; 1.5% **

N017751 007

+ DENTSPLY PHARM

0.005MG/ML; 1.5% **

N021384 001

EPINEPHRINE BITARTRATE; PRILLOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST FORTE

ASTRAZENECA

0.005MG/ML; 4%

N014763 008

EPINEPHRINE; ETIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DURANEST

+ ASTRAZENECA

0.005MG/ML; 0.5% **

N017751 004

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ALPHACAINE HYDROCHLORIDE W/ EPINEPHRINE

CARLISLE

0.01MG/ML; 2%

A084720 001

0.02MG/ML; 2%

A084732 001

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

BELMORA LLC

0.01MG/ML; 2%

A080504 004 Oct 19, 1983

0.02MG/ML; 2%

A080504 005 Oct 19, 1983

EASTMAN KODAK

0.01MG/ML; 2%

A040057 002 Feb 26, 1993

0.02MG/ML; 2%

A040057 001 Feb 26, 1993

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

| | | | |
|-----------------------------------------|--------------------------|---------|------------------|
| LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE | | | |
| HOSPIRA | 0.005MG/ML;1% | A089649 | 001 Jun 21, 1988 |
| | 0.005MG/ML;1.5% | A089650 | 001 Jun 21, 1988 |
| | 0.01MG/ML;2% | A078772 | 001 May 12, 2008 |
| | 0.02MG/ML;2% | A078772 | 002 May 12, 2008 |
| WEST-WARD PHARMS INT | 0.01MG/ML;1% | A080406 | 001 |
| | 0.01MG/ML;2% | A080406 | 002 |
| LIDOCAINE HYDROCHLORIDE W/ EPINEPHRINE | | | |
| ABBOTT | 0.01MG/ML;1% | A083154 | 001 |
| BEL MAR | 0.01MG/ML;1% | A080820 | 001 |
| | 0.01MG/ML;2% | A080757 | 001 |
| DELL LABS | 0.01MG/ML;1% | A083389 | 001 |
| | 0.01MG/ML;2% | A083390 | 001 |
| INTL MEDICATION | 0.01MG/ML;1% | A086402 | 001 |
| WATSON LABS | 0.01MG/ML;1% | A080377 | 003 |
| | 0.01MG/ML;1% | A085463 | 001 |
| | 0.01MG/ML;2% | A080377 | 004 |
| LIDOCATON | | | |
| PHARMATON | 0.01MG/ML;2% | A084729 | 001 Aug 17, 1983 |
| | 0.02MG/ML;2% | A084728 | 001 Aug 17, 1983 |
| OCTOCAINE | | | |
| SEPTODONT | 0.01MG/ML;2% | A084048 | 001 |
| | 0.02MG/ML;2% | A084048 | 002 |
| XYLOCAINE DENTAL WITH EPINEPHRINE | | | |
| DENTSPLY PHARM | 0.01MG/ML;2% | N021381 | 001 |
| | 0.02MG/ML;2% | N021381 | 002 |
| XYLOCAINE W/ EPINEPHRINE | | | |
| ASTRAZENECA | 0.005MG/ML;1% | N010418 | 006 |
| | 0.005MG/ML;1.5% | N010418 | 010 |
| | 0.005MG/ML;2% | N010418 | 008 |
| FRESENIUS KABI USA | 0.01MG/ML;2% | N006488 | 003 |
| PATCH; IONTOPHORESIS, TOPICAL | | | |
| LIDOSITE TOPICAL SYSTEM KIT | | | |
| VYTERIS | 1.05MG/PATCH;100MG/PATCH | N021504 | 001 May 06, 2004 |
| SOLUTION; IONTOPHORESIS | | | |
| IONTOCAINE | | | |
| IOMED | 0.01MG/ML;2% | N020530 | 001 Dec 21, 1995 |
| SOLUTION; IONTOPHORESIS, TOPICAL | | | |
| LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE | | | |
| EMPI | 0.01MG/ML;2% | N021486 | 001 Oct 26, 2004 |

EPINEPHRINE; PROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

| | | | |
|---------------------------------------|--------------|---------|-----|
| PROCAINE HYDROCHLORIDE W/ EPINEPHRINE | | | |
| BEL MAR | 0.02MG/ML;1% | A080758 | 001 |
| | 0.02MG/ML;2% | A080759 | 001 |

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

| | | | |
|--------------------------|----------------------|---------|------------------|
| EPIRUBICIN HYDROCHLORIDE | | | |
| EBEWE PHARMA | 50MG/25ML (2MG/ML) | A065339 | 001 Dec 22, 2009 |
| | 200MG/100ML (2MG/ML) | A065339 | 002 Dec 22, 2009 |
| HOSPIRA | 50MG/25ML (2MG/ML) | A065343 | 002 Apr 19, 2007 |
| MUSTAFA NEVSAT | 50MG/25ML (2MG/ML) | A090266 | 001 Apr 15, 2011 |
| | 200MG/100ML (2MG/ML) | A090266 | 002 Apr 15, 2011 |
| MYLAN INSTITUTIONAL | 50MG/25ML (2MG/ML) | A065371 | 001 Nov 28, 2007 |
| | 200MG/100ML (2MG/ML) | A065371 | 002 Nov 28, 2007 |
| POWDER; INTRAVENOUS | | | |
| EPIRUBICIN HYDROCHLORIDE | | | |
| HOSPIRA | 50MG/VIAL | N050807 | 001 Sep 15, 2006 |
| | 200MG/VIAL | N050807 | 002 Sep 15, 2006 |

EPLERENONE

TABLET; ORAL

| | | | |
|---------------|-------|---------|------------------|
| INSPRA | | | |
| GD SEARLE LLC | 100MG | N021437 | 003 Sep 27, 2002 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

EPROSARTAN MESYLATE

TABLET; ORAL

TEVETEN

| | | | |
|--------|---------------|-------------|--------------|
| ABBVIE | EQ 300MG BASE | N020738 004 | Dec 22, 1997 |
| + | EQ 400MG BASE | N020738 005 | Dec 22, 1997 |
| + | EQ 600MG BASE | N020738 006 | May 27, 1999 |

EPROSARTAN MESYLATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

TEVETEN HCT

| | | | |
|--------|--------------|-------------|--------------|
| ABBVIE | 600MG;12.5MG | N021268 001 | Nov 01, 2001 |
| | 600MG;25MG | N021268 002 | Nov 01, 2001 |

EPTIFIBATIDE

INJECTABLE; INJECTION

EPTIFIBATIDE

| | | | |
|-----------------|------------|-------------|--------------|
| TEVA PHARMS USA | 75MG/100ML | A091555 001 | Jun 05, 2015 |
|-----------------|------------|-------------|--------------|

ERGOCALCIFEROL

CAPSULE; ORAL

DELTALIN

| | | | |
|------------|-----------|-------------|--|
| LILLY | 50,000 IU | A080884 001 | |
| VITAMIN D | | | |
| CHASE CHEM | 50,000 IU | A080747 001 | |
| EVERYLIFE | 50,000 IU | A080956 001 | |
| IMPAX LABS | 50,000 IU | A080951 001 | |
| LANNETT | 50,000 IU | A080825 001 | |
| VITARINE | 50,000 IU | A084053 001 | |
| WEST WARD | 50,000 IU | A083102 001 | |

ERGOLOID MESYLATES

CAPSULE; ORAL

HYDERGINE LC

| | | | |
|----------|-----|-------------|--------------|
| NOVARTIS | 1MG | N018706 001 | Jan 18, 1983 |
|----------|-----|-------------|--------------|

SOLUTION; ORAL

HYDERGINE

| | | | |
|----------|--------|-------------|--|
| NOVARTIS | 1MG/ML | N018418 001 | |
|----------|--------|-------------|--|

TABLET; ORAL

ERGOLOID MESYLATES

| | | | |
|--------------|-----|-------------|--------------|
| MUTUAL PHARM | 1MG | A088891 001 | Nov 01, 1985 |
| WATSON LABS | 1MG | A086433 001 | May 27, 1982 |
| | 1MG | A087244 001 | Aug 16, 1982 |

GERIMAL

| | | | |
|-------------|-----|-------------|--------------|
| WATSON LABS | 1MG | A088207 001 | Mar 22, 1984 |
|-------------|-----|-------------|--------------|

HYDERGINE

| | | | |
|----------|-------|-------------|--|
| NOVARTIS | 0.5MG | N017993 003 | |
|----------|-------|-------------|--|

| | | | |
|---|-----|-------------|--|
| + | 1MG | N017993 001 | |
|---|-----|-------------|--|

TABLET; SUBLINGUAL

ALKERGOT

| | | | |
|--------|-------|-------------|--|
| SANDOZ | 0.5MG | A085153 001 | |
| | 1MG | A087417 001 | |

CIRCANOL

| | | | |
|----|-------|-------------|--|
| 3M | 0.5MG | A084868 001 | |
| | 1MG | A085809 001 | |

DEAPRIL-ST

| | | | |
|----------------------|-----|-------------|--|
| BRISTOL MYERS SQUIBB | 1MG | A085020 002 | |
|----------------------|-----|-------------|--|

ERGOLOID MESYLATES

| | | | |
|----------------------|-------|-------------|--------------|
| KV PHARM | 0.5MG | A085899 001 | |
| | 0.5MG | A086265 001 | |
| | 1MG | A085900 001 | |
| | 1MG | A086264 001 | |
| LEDERLE | 0.5MG | A086984 001 | |
| | 1MG | A086985 001 | |
| SUN PHARM INDUSTRIES | 0.5MG | A087407 001 | |
| | 1MG | A087552 001 | |
| SUPERPHARM | 0.5MG | A089233 001 | Sep 23, 1986 |
| | 1MG | A089234 001 | Sep 23, 1986 |
| VANGARD | 0.5MG | A088013 001 | Sep 20, 1982 |
| | 1MG | A088014 001 | Sep 20, 1982 |
| WATSON LABS | 0.5MG | A084930 001 | |
| | 0.5MG | A087233 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERGOLOID MESYLATES

TABLET;SUBLINGUAL

ERGOLOID MESYLATES

| | | |
|-----|---------|-----|
| 1MG | A085177 | 001 |
| 1MG | A087183 | 001 |

GERIMAL

WATSON LABS

| | | |
|-------|---------|-----|
| 0.5MG | A086189 | 001 |
| 1MG | A086188 | 001 |

HYDERGINE

NOVARTIS

| | | |
|-------|---------|-----|
| 0.5MG | N009087 | 002 |
| 1MG | N009087 | 001 |

HYDROGENATED ERGOT ALKALOIDS

IVAX PHARMS

| | | |
|-------|---------|-----|
| 0.5MG | A087186 | 001 |
| 1MG | A087185 | 001 |

ERGOTAMINE TARTRATE

AEROSOL, METERED; INHALATION

MEDIHALER ERGOTAMINE

3M

| | | |
|------------|---------|-----|
| 0.36MG/INH | N012102 | 001 |
|------------|---------|-----|

TABLET;SUBLINGUAL

ERGOSTAT

WATSON LABS INC

| | | | |
|-----|---------|-----|--------------|
| 2MG | A088337 | 001 | Jun 08, 1984 |
|-----|---------|-----|--------------|

WIGRETTES

ORGANON USA INC

| | | | |
|-----|---------|-----|--------------|
| 2MG | A086750 | 001 | Jul 29, 1982 |
|-----|---------|-----|--------------|

ERLOTINIB HYDROCHLORIDE

TABLET;ORAL

ERLOTINIB HYDROCHLORIDE

MYLAN PHARMS INC

| | | | |
|--------------|---------|-----|--------------|
| EQ 25MG BASE | A091002 | 001 | Jun 11, 2014 |
|--------------|---------|-----|--------------|

| | | | |
|---------------|---------|-----|--------------|
| EQ 100MG BASE | A091002 | 002 | Jun 11, 2014 |
|---------------|---------|-----|--------------|

| | | | |
|---------------|---------|-----|--------------|
| EQ 150MG BASE | A091002 | 003 | Jun 11, 2014 |
|---------------|---------|-----|--------------|

TEVA PHARMS USA

| | | | |
|---------------|---------|-----|--------------|
| EQ 100MG BASE | A091059 | 002 | Aug 28, 2015 |
|---------------|---------|-----|--------------|

| | | | |
|---------------|---------|-----|--------------|
| EQ 150MG BASE | A091059 | 003 | Aug 28, 2015 |
|---------------|---------|-----|--------------|

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS;ORAL

ERYC

PARKE DAVIS

| | | | |
|-------|---------|-----|--------------|
| 250MG | A062546 | 001 | Jul 25, 1985 |
|-------|---------|-----|--------------|

| | | | |
|-------|---------|-----|--------------|
| 250MG | A062618 | 001 | Sep 25, 1985 |
|-------|---------|-----|--------------|

WARNER CHILCOTT LLC

| | | | |
|-------|---------|-----|--|
| 250MG | A062338 | 001 | |
|-------|---------|-----|--|

ERYC 125

PARKE DAVIS

| | | | |
|-------|---------|-----|--------------|
| 125MG | A062648 | 001 | Oct 24, 1985 |
|-------|---------|-----|--------------|

ERYC SPRINKLES

HOSPIRA

| | | | |
|-------|---------|-----|--------------|
| 125MG | N050593 | 001 | Jul 22, 1985 |
|-------|---------|-----|--------------|

ERYTHROMYCIN

BARR

| | | | |
|-------|---------|-----|--------------|
| 250MG | A063098 | 001 | May 04, 1989 |
|-------|---------|-----|--------------|

GEL;TOPICAL

E-GLADES

MYLAN PHARMS INC

| | | | |
|----|---------|-----|--------------|
| 2% | A065009 | 001 | Mar 18, 2002 |
|----|---------|-----|--------------|

EMGEL

ALTANA

| | | | |
|----|---------|-----|--------------|
| 2% | A063107 | 001 | Aug 23, 1991 |
|----|---------|-----|--------------|

LOTION;TOPICAL

E-SOLVE 2

SYOSSET

| | | | |
|----|---------|-----|--------------|
| 2% | A062467 | 001 | Jul 03, 1985 |
|----|---------|-----|--------------|

OINTMENT;OPHTHALMIC

ERYTHROMYCIN

PHARMADERM

| | | | |
|--------|---------|-----|--------------|
| 5MG/GM | A062446 | 001 | Sep 26, 1983 |
|--------|---------|-----|--------------|

PHARMAFAIR

| | | | |
|--------|---------|-----|--------------|
| 5MG/GM | A062481 | 001 | Apr 05, 1984 |
|--------|---------|-----|--------------|

ILOTYCIN

DISTA

| | | | |
|------|---------|-----|--|
| 0.5% | N050368 | 001 | |
|------|---------|-----|--|

OINTMENT;TOPICAL

AKNE-MYCIN

+ DOW PHARM

| | | | |
|----|---------|-----|--------------|
| 2% | N050584 | 001 | Jan 10, 1985 |
|----|---------|-----|--------------|

POWDER;FOR RX COMPOUNDING

ERYTHROMYCIN

PADDOCK LLC

| | | | |
|------|---------|-----|--------------|
| 100% | N050610 | 001 | Nov 07, 1986 |
|------|---------|-----|--------------|

SOLUTION;TOPICAL

A/T/S

TARO

| | | | |
|----|---------|-----|--------------|
| 2% | A062405 | 001 | Nov 18, 1982 |
|----|---------|-----|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN

SOLUTION; TOPICAL

| | | | | |
|--------------------------------|---------|---------|-----|--------------|
| C-SOLVE-2 | | | | |
| FOUGERA PHARMS | 2% | A062468 | 001 | Jul 03, 1985 |
| ERYDERM | | | | |
| ARBOR PHARMS INC | 2% | A062290 | 001 | |
| ERYMAX | | | | |
| MERZ PHARMS | 2% | A062508 | 002 | Jul 11, 1985 |
| ERYTHRA-DERM | | | | |
| ANDA REPOSITORY | 2% | A062687 | 001 | Feb 05, 1988 |
| ERYTHRO-STATIN | | | | |
| HI TECH PHARMA | 2% | A064101 | 001 | Oct 22, 1996 |
| ERYTHROMYCIN | | | | |
| ALPHARMA US PHARMS | 1.5% | A062328 | 001 | Apr 19, 1982 |
| | 2% | A062326 | 001 | Apr 19, 1982 |
| | 2% | A062327 | 001 | Apr 19, 1982 |
| | 2% | A062342 | 001 | Feb 25, 1982 |
| | 2% | A062957 | 001 | Jul 21, 1988 |
| BAUSCH AND LOMB | 2% | A064039 | 001 | Jan 27, 1994 |
| FOUGERA PHARMS | 2% | A064187 | 001 | Sep 30, 1997 |
| LILLY | 2% | N050532 | 001 | |
| PHARMAFAIR | 1.5% | A062485 | 001 | Jul 11, 1984 |
| | 2% | A062616 | 001 | Jul 25, 1985 |
| RENAISSANCE PHARMA | 2% | A064127 | 001 | Feb 14, 1997 |
| SANSAC | | | | |
| DOW PHARM | 2% | A062522 | 001 | Jan 24, 1985 |
| STATICIN | | | | |
| + WESTWOOD SQUIBB | 1.5% ** | N050526 | 001 | |
| T-STAT | | | | |
| WESTWOOD SQUIBB | 2% ** | A062436 | 001 | Mar 09, 1983 |
| SWAB; TOPICAL | | | | |
| C-SOLVE-2 | | | | |
| IVAX SUB TEVA PHARMS | 2% | A062751 | 001 | Jul 30, 1993 |
| ERYCETTE | | | | |
| + JOHNSON AND JOHNSON | 2% ** | N050594 | 001 | Feb 15, 1985 |
| ERYTHROMYCIN | | | | |
| FOUGERA PHARMS | 2% | A065320 | 001 | Jul 25, 2006 |
| MYLAN PHARMS INC | 2% | A064128 | 001 | Jul 03, 1996 |
| T-STAT | | | | |
| WESTWOOD SQUIBB | 2% | A062748 | 001 | Jul 23, 1987 |
| TABLET, COATED PARTICLES; ORAL | | | | |
| PCE | | | | |
| + ARBOR PHARMS LLC | 333MG | N050611 | 001 | Sep 09, 1986 |
| + | 500MG | N050611 | 002 | Aug 22, 1990 |
| TABLET, DELAYED RELEASE; ORAL | | | | |
| E-BASE | | | | |
| BARR | 333MG | A063028 | 001 | May 15, 1990 |
| | 333MG | A063086 | 001 | May 15, 1990 |
| | 500MG | A062999 | 001 | Nov 25, 1988 |
| E-MYCIN | | | | |
| ARBOR PHARMS INC | 250MG | A060272 | 001 | |
| | 333MG | A060272 | 002 | |
| ILOTYCIN | | | | |
| DISTA | 250MG | A061910 | 001 | |
| R-P MYCIN | | | | |
| SOLVAY | 250MG | A061659 | 001 | |
| ROBIMYCIN | | | | |
| ROBINS AH | 250MG | A061633 | 001 | |

ERYTHROMYCIN ESTOLATE

CAPSULE; ORAL

ERYTHROMYCIN ESTOLATE

| | | | | |
|----------------------|---------------|---------|-----|--|
| BARR | EQ 125MG BASE | A062162 | 001 | |
| | EQ 250MG BASE | A062162 | 002 | |
| IVAX SUB TEVA PHARMS | EQ 250MG BASE | A062237 | 001 | |
| WATSON LABS | EQ 250MG BASE | A062087 | 001 | |
| ILOSONE | | | | |
| LILLY | EQ 125MG BASE | A061897 | 001 | |
| | EQ 250MG BASE | A061897 | 002 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN ESTOLATE

FOR SUSPENSION;ORAL

ILOSONE

DISTA EQ 125MG BASE/5ML A061893 001

SUSPENSION;ORAL

ERYTHROMYCIN ESTOLATE

ALPHARMA US PHARMS EQ 125MG BASE/5ML A062353 001 Nov 18, 1982

EQ 250MG BASE/5ML A062409 001 Dec 16, 1982

G AND W LABS INC EQ 125MG BASE/5ML A062169 001 Oct 17, 1990

EQ 250MG BASE/5ML A062169 002 Oct 17, 1990

LIFE LABS EQ 250MG BASE/5ML A062362 001 Dec 17, 1982

ILOSONE

LILLY EQ 125MG BASE/5ML A061894 001

EQ 125MG BASE/5ML N050010 001

EQ 250MG BASE/5ML A061894 002

EQ 250MG BASE/5ML N050010 002

SUSPENSION/DROPS;ORAL

ILOSONE

LILLY EQ 100MG BASE/ML A061894 003

TABLET;ORAL

ILOSONE

LILLY EQ 500MG BASE A061896 001

TABLET, CHEWABLE;ORAL

ILOSONE

DISTA EQ 125MG BASE A061895 001

EQ 250MG BASE A061895 002

ERYTHROMYCIN ESTOLATE; SULFISOXAZOLE ACETYL

SUSPENSION;ORAL

ILOSONE SULFA

LILLY EQ 125MG BASE/5ML;EQ 600MG BASE/5ML N050599 001 Sep 29, 1989

ERYTHROMYCIN ETHYLSUCCINATE

GRANULE;ORAL

PEDIAMYCIN

ROSS LABS EQ 200MG BASE/5ML A062305 001

SUSPENSION;ORAL

E-MYCIN E

PHARMACIA AND UPJOHN EQ 200MG BASE/5ML A062198 001

EQ 400MG BASE/5ML A062198 002

E.E.S. 200

ARBOR PHARMS LLC EQ 200MG BASE/5ML ** A061639 001

E.E.S. 400

ARBOR PHARMS LLC EQ 400MG BASE/5ML ** A061639 002

ERYTHROMYCIN ETHYLSUCCINATE

ALPHARMA US PHARMS EQ 200MG BASE/5ML A062200 001

EQ 400MG BASE/5ML A062200 002

DISTA EQ 200MG BASE/5ML A062177 001

EQ 400MG BASE/5ML A062177 002

NASKA EQ 400MG BASE/5ML A062674 001 Mar 10, 1987

PARKE DAVIS EQ 200MG BASE/5ML A062231 001

EQ 400MG BASE/5ML A062231 002

PHARMAFAIR EQ 200MG BASE/5ML A062559 001 Mar 15, 1985

EQ 400MG BASE/5ML A062558 001 Mar 15, 1985

PEDIAMYCIN

ARBOR PHARMS LLC EQ 200MG BASE/5ML A062304 001

PEDIAMYCIN 400

ARBOR PHARMS LLC EQ 400MG BASE/5ML A062304 002

WYAMYCIN E

WYETH AYERST EQ 200MG BASE/5ML A062123 002

EQ 400MG BASE/5ML A062123 001

SUSPENSION/DROPS;ORAL

PEDIAMYCIN

ROSS LABS EQ 100MG BASE/2.5ML A062305 002

TABLET;ORAL

E.E.S. 400

ARBOR PHARMS LLC EQ 400MG BASE A061905 001

ERYTHROMYCIN ETHYLSUCCINATE

BARR EQ 400MG BASE A062256 001

MYLAN EQ 400MG BASE A062847 001 Sep 14, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN ETHYLSUCCINATE

TABLET, CHEWABLE;ORAL

E.E.S.

ARBOR PHARMS INC EQ 200MG BASE N050297 002

ERYPED

ARBOR PHARMS INC EQ 200MG BASE N050297 003 Jul 05, 1988

PEDIAMYCIN

ROSS LABS EQ 200MG BASE A062306 001

ERYTHROMYCIN ETHYLSUCCINATE; SULFISOXAZOLE ACETYL

GRANULE;ORAL

ERYTHROMYCIN ETHYLSUCCINATE AND SULFISOXAZOLE ACETYL

BARR EQ 200MG BASE/5ML;EQ 600MG BASE/5ML A062759 001 May 20, 1988

ERYZOLE

ALRA EQ 200MG BASE/5ML;EQ 600MG BASE/5ML A062758 001 Jun 15, 1988

PEDIAZOLE

ROSS LABS EQ 200MG BASE/5ML;EQ 600MG BASE/5ML N050529 001

ERYTHROMYCIN GLUCEPTATE

INJECTABLE;INJECTION

ILOTYCIN GLUCEPTATE

DISTA EQ 250MG BASE/VIAL N050370 001

EQ 500MG BASE/VIAL N050370 002

EQ 1GM BASE/VIAL N050370 003

ERYTHROMYCIN LACTOBIONATE

INJECTABLE;INJECTION

ERYTHROCIN

ABBOTT EQ 500MG BASE/VIAL A062586 001 Jan 04, 1988

EQ 1GM BASE/VIAL A062586 002 Jan 04, 1988

HOSPIRA EQ 500MG BASE/VIAL N050182 002

EQ 1GM BASE/VIAL A062638 002 Oct 31, 1986

EQ 1GM BASE/VIAL N050182 003

+ EQ 1GM BASE/VIAL N050609 002 Sep 24, 1986

ERYTHROMYCIN

ELKINS SINN EQ 500MG BASE/VIAL A062563 001 Mar 28, 1985

EQ 1GM BASE/VIAL A062563 002 Mar 28, 1985

ERYTHROMYCIN LACTOBIONATE

ABRAXIS PHARM EQ 500MG BASE/VIAL A062604 001 Nov 24, 1986

EQ 1GM BASE/VIAL A062604 002 Nov 24, 1986

BAXTER HLTHCARE EQ 500MG BASE/VIAL A062993 001 May 09, 1989

EQ 1GM BASE/VIAL A062993 002 May 09, 1989

TEVA PARENTERAL EQ 500MG BASE/VIAL A063253 001 Jul 30, 1993

EQ 1GM BASE/VIAL A063253 002 Jul 30, 1993

ERYTHROMYCIN STEARATE

TABLET;ORAL

BRISTAMYCIN

BRISTOL EQ 250MG BASE A061304 001

EQ 250MG BASE A061887 001

ERYPAR

PARKE DAVIS EQ 250MG BASE A062032 001

EQ 500MG BASE A062032 002

WARNER CHILCOTT EQ 250MG BASE A062322 001

ERYTHROCIN STEARATE

ARBOR PHARMS LLC EQ 125MG BASE A060359 002

EQ 500MG BASE A060359 003

ERYTHROMYCIN STEARATE

ANI PHARMS INC EQ 250MG BASE A061461 001

EQ 250MG BASE A061591 001

EQ 500MG BASE A061461 002

EQ 500MG BASE A063179 001 May 15, 1990

LEDERLE EQ 250MG BASE A062089 001

EQ 500MG BASE A062089 002

MYLAN EQ 250MG BASE A061505 001

EQ 500MG BASE A061505 002

PUREPAC PHARM EQ 250MG BASE A061743 001

WATSON LABS EQ 250MG BASE A062121 002

EQ 500MG BASE A062121 001

ETHRIL 250

BRISTOL MYERS SQUIBB EQ 250MG BASE A061605 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ERYTHROMYCIN STEARATE

TABLET;ORAL

ETHRIL 500

BRISTOL MYERS SQUIBB EQ 500MG BASE

A061605 002

PFIZER-E

PFIZER EQ 250MG BASE

A061791 001

EQ 500MG BASE

A061791 002

WYAMYCIN S

WYETH AYERST EQ 250MG BASE

A061675 001

EQ 500MG BASE

A061675 002

ESCITALOPRAM OXALATE

CAPSULE;ORAL

ESCITALOPRAM OXALATE

MYLAN PHARMS INC EQ 5MG BASE

A077660 001 Jul 31, 2007

EQ 10MG BASE

A077660 002 Jul 31, 2007

EQ 20MG BASE

A077660 003 Jul 31, 2007

TABLET;ORAL

ESCITALOPRAM OXALATE

MYLAN PHARMS INC EQ 5MG BASE

A077550 001 May 14, 2015

EQ 10MG BASE

A077550 002 May 14, 2015

EQ 20MG BASE

A077550 003 May 14, 2015

ESMOLOL HYDROCHLORIDE

INJECTABLE;INJECTION

BREVIBLOC

BAXTER HLTHCARE 10MG/ML

N019386 003 Aug 15, 1988

20MG/ML

N019386 007 May 28, 2003

ESOMEPRAZOLE SODIUM

INJECTABLE;INTRAVENOUS

ESOMEPRAZOLE SODIUM

AUROBINDO PHARMA LTD EQ 20MG BASE/VIAL

A204657 001 Aug 10, 2016

MYLAN LABS LTD EQ 20MG BASE/VIAL

A202686 001 May 17, 2017

SUN PHARMA GLOBAL EQ 20MG BASE/VIAL

A200882 001 Mar 18, 2013

NEXIUM IV

+ ASTRAZENECA PHARMS EQ 20MG BASE/VIAL **

N021689 001 Mar 31, 2005

ESOMEPRAZOLE STRONTIUM

CAPSULE, DELAYED RELEASE;ORAL

ESOMEPRAZOLE STRONTIUM

+ R2 PHARMA LLC 24.65MG

N202342 001 Aug 06, 2013

ESTAZOLAM

TABLET;ORAL

PROSOM

+ ABBOTT 1MG **

N019080 001 Dec 26, 1990

+ 2MG **

N019080 002 Dec 26, 1990

ESTRADIOL

FILM, EXTENDED RELEASE;TRANSDERMAL

ESCLIM

WOMEN FIRST HLTHCARE 0.025MG/24HR

N020847 001 Aug 04, 1998

0.0375MG/24HR

N020847 002 Aug 04, 1998

0.05MG/24HR

N020847 003 Aug 04, 1998

0.075MG/24HR

N020847 004 Aug 04, 1998

0.1MG/24HR

N020847 005 Aug 04, 1998

ESTRADERM

+ NOVARTIS 0.05MG/24HR

N019081 002 Sep 10, 1986

+ 0.1MG/24HR

N019081 003 Sep 10, 1986

ESTRADIOL

ORTHO MCNEIL PHARM 0.05MG/24HR

N021048 001 Sep 20, 1999

0.075MG/24HR

N021048 002 Sep 20, 1999

0.1MG/24HR

N021048 003 Sep 20, 1999

FEMPATCH

PARKE DAVIS 0.025MG/24HR

N020417 001 Dec 03, 1996

VIVELLE

NOVARTIS 0.025MG/24HR

N020323 005 Aug 16, 2000

0.0375MG/24HR

N020323 001 Oct 28, 1994

0.05MG/24HR

N020323 002 Oct 28, 1994

0.075MG/24HR

N020323 003 Oct 28, 1994

0.1MG/24HR

N020323 004 Oct 28, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESTRADIOL

GEL;TOPICAL

ESTROGEL

ASCEND THERAPS US 0.06% N021166 001 Feb 09, 2004

TABLET;ORAL

ESTRACE

BRISTOL MYERS SQUIBB 0.5MG A081295 001 Jun 30, 1993
1MG A084499 001
2MG A084500 001

ESTRADIOL

LANNETT HOLDINGS INC 0.5MG A040138 001 Jan 30, 1998
1MG A040138 002 Jan 30, 1998
2MG A040138 003 Jan 30, 1998
USL PHARMA 0.5MG A040297 001 Apr 17, 2002
1MG A040297 002 Apr 17, 2002
2MG A040297 003 Apr 17, 2002

GYNODIOL

DURAMED PHARMS BARR 0.5MG A040212 001 Dec 29, 1997
1MG A040212 002 Dec 29, 1997
1.5MG A040212 003 Dec 29, 1997
2MG A040212 004 Dec 29, 1997

INNOFEM

NOVO NORDISK INC 0.5MG A040312 001 Nov 19, 1999
1MG A040312 002 Nov 19, 1999
2MG A040312 003 Nov 19, 1999

TABLET;VAGINAL

VAGIFEM

+ NOVO NORDISK INC 25MCG ** N020908 001 Mar 26, 1999

ESTRADIOL ACETATE

TABLET;ORAL

FEMTRACE

+ APIL 0.45MG N021633 001 Aug 20, 2004
+ 0.9MG N021633 002 Aug 20, 2004
+ 1.8MG N021633 003 Aug 20, 2004ESTRADIOL CYPIONATE

INJECTABLE;INJECTION

DEPO-ESTRADIOL

PHARMACIA AND UPJOHN 1MG/ML A085470 001
3MG/ML A085470 002

ESTRADIOL CYPIONATE

WATSON LABS 5MG/ML A085620 001

ESTRADIOL CYPIONATE; MEDROXYPROGESTERONE ACETATE

INJECTABLE;INTRAMUSCULAR

LUNELLE

PHARMACIA AND UPJOHN 5MG/0.5ML;25MG/0.5ML N020874 001 Oct 05, 2000

ESTRADIOL CYPIONATE; TESTOSTERONE CYPIONATE

INJECTABLE;INJECTION

DEPO-TESTADIOL

PHARMACIA AND UPJOHN 2MG/ML;50MG/ML N017968 001

TESTOSTERONE CYPIONATE-ESTRADIOL CYPIONATE

WATSON LABS 2MG/ML;50MG/ML A085603 001 Mar 13, 1986

ESTRADIOL HEMIHYDRATE

EMULSION;TOPICAL

ESTRASORB

+ EXELTIS USA INC 0.25% N021371 001 Oct 09, 2003

ESTRADIOL VALERATE

INJECTABLE;INJECTION

ESTRADIOL VALERATE

FOSUN PHARMA 10MG/ML A040628 001 Oct 04, 2007
20MG/ML A040628 002 Oct 04, 2007
40MG/ML A040628 003 Oct 04, 2007
WATSON LABS 10MG/ML A083546 001
40MG/ML A083714 001
WATSON LABS INC 20MG/ML A083547 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESTRADIOL VALERATE; TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

DITATE-DS

SAVAGE LABS 8MG/ML;180MG/ML A086423 001

TESTOSTERONE ENANTHATE AND ESTRADIOL VALERATE

WATSON LABS 4MG/ML;90MG/ML A085865 001

8MG/ML;180MG/ML A085860 001

ESTRADIOL; NORGESTIMATE

TABLET; ORAL

PREFEST

+ TEVA WOMENS 1MG,1MG;N/A,0.09MG ** N021040 001 Oct 22, 1999

ESTROGENS, CONJUGATED

TABLET; ORAL

PREMARIN

WYETH PHARMS 2.5MG N004782 002

ESTROGENS, CONJUGATED SYNTHETIC A

CREAM; VAGINAL

SYNTHETIC CONJUGATED ESTROGENS A

TEVA WOMENS 0.625MG/GM N021788 001 Nov 28, 2008

TABLET; ORAL

CENESTIN

+ TEVA BRANDED PHARM 0.3MG ** N020992 001 Jun 21, 2002

+ 0.45MG ** N020992 005 Feb 05, 2004

+ 0.625MG ** N020992 002 Mar 24, 1999

+ 0.9MG ** N020992 003 Mar 24, 1999

+ 1.25MG ** N020992 004 Mar 13, 2000

ESTROGENS, CONJUGATED SYNTHETIC B

TABLET; ORAL

ENJUVIA

TEVA BRANDED PHARM 0.3MG N021443 001 Dec 20, 2004

0.45MG N021443 002 Dec 20, 2004

0.625MG ** N021443 003 May 10, 2004

0.9MG N021443 005 Apr 27, 2007

1.25MG ** N021443 004 May 10, 2004

ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE

TABLET; ORAL-28

PREMPHASE (PREMARIN;CYCRIN 14/14)

WYETH PHARMS INC 0.625MG,0.625MG;N/A,5MG N020303 002 Dec 30, 1994

PREMPRO (PREMARIN;CYCRIN)

WYETH PHARMS INC 0.625MG,0.625MG;2.5MG,2.5MG N020303 001 Dec 30, 1994

ESTROGENS, CONJUGATED; MEPROBAMATE

TABLET; ORAL

MILPREM-200

MEDPOINTE PHARM HLC 0.45MG;200MG N011045 002

MILPREM-400

MEDPOINTE PHARM HLC 0.45MG;400MG N011045 001

PMB 200

WYETH AYERST 0.45MG;200MG N010971 005

PMB 400

WYETH AYERST 0.45MG;400MG N010971 003

ESTROGENS, ESTERIFIED

TABLET; ORAL

AMNESTROGEN

BRISTOL MYERS SQUIBB 0.3MG A083266 001

0.625MG A083266 002

1.25MG A083266 003

2.5MG A083266 004

ESTERIFIED ESTROGENS

PVT FORM 0.625MG A083414 001

1.25MG A083765 001

2.5MG A085907 001

SANDOZ 1.25MG A085302 001

ESTRATAB

SOLVAY 0.3MG A086715 001

0.625MG A083209 001

1.25MG A083856 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ESTROGENS, ESTERIFIED

TABLET; ORAL

ESTRATAB

2.5MG A083857 001

EVEX

ROCHE PALO 0.625MG A084215 001

1.25MG A083376 002

FEMOGEN

PVT FORM 0.625MG A085076 001

1.25MG A085008 001

2.5MG A085007 001

ESTRONE

INJECTABLE; INJECTION

ESTROGENIC SUBSTANCE

WYETH AYERST 2MG/ML A083488 001

ESTRONE

WATSON LABS 2MG/ML A083397 001

WATSON LABS TEVA 5MG/ML A085239 001

NATURAL ESTROGENIC SUBSTANCE-ESTRONE

WATSON LABS 2MG/ML A085237 001 Nov 23, 1982

THEELIN

PARKEDALE 1MG/ML N003977 001

2MG/ML N003977 002

5MG/ML N003977 003

ESTROPIPATE

CREAM; VAGINAL

OGEN

PHARMACIA AND UPJOHN 1.5MG/GM A084710 001

TABLET; ORAL

ESTROPIPATE

BARR 0.75MG A040135 001 Nov 27, 1996

1.5MG A040135 002 Nov 27, 1996

3MG A040135 003 Nov 27, 1996

DURAMED PHARMS BARR 0.75MG A040296 001 Nov 01, 1999

1.5MG A040296 002 Nov 01, 1999

3MG A040296 003 Nov 01, 1999

MYLAN 3MG A040359 003 Aug 26, 1999

WATSON LABS 0.75MG A081213 001 Sep 23, 1993

1.5MG A081214 001 Sep 23, 1993

6MG A081216 001 Sep 23, 1993

WATSON LABS TEVA 3MG A081215 001 Sep 23, 1993

OGEN .625

PHARMACIA AND UPJOHN 0.75MG A083220 001

OGEN 1.25

PHARMACIA AND UPJOHN 1.5MG A083220 002

OGEN 2.5

PHARMACIA AND UPJOHN 3MG A083220 003

ORTHO-EST

SUN PHARM INDS INC 0.75MG A089567 001 Feb 27, 1991

1.5MG A089582 001 Jul 17, 1991

ESZOPICLONE

TABLET; ORAL

ESZOPICLONE

WOCKHARDT LTD 1MG A091165 001 Jul 14, 2011

2MG A091165 002 Jul 14, 2011

3MG A091165 003 Jul 14, 2011

ETHACRYNIC ACID

TABLET; ORAL

EDECIN

ATON 50MG N016092 002

ETHAMBUTOL HYDROCHLORIDE

TABLET; ORAL

MYAMBUTOL

STI PHARMA LLC 200MG N016320 002

500MG N016320 004

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHCHLORVYNOL

CAPSULE; ORAL

ETHCHLORVYNOL

| | | |
|-------------------|-------|-------------|
| BANNER PHARMACAPS | 100MG | A084463 001 |
| | 200MG | A084463 002 |
| | 500MG | A084463 003 |
| | 750MG | A084463 004 |

PLACIDYL

| | | |
|--------|-------|-------------|
| ABBVIE | 100MG | N010021 004 |
| | 200MG | N010021 007 |
| | 500MG | N010021 002 |
| | 750MG | N010021 010 |

ETHINAMATE

CAPSULE; ORAL

VALMID

| | | |
|-------|-------|-------------|
| DISTA | 500MG | N009750 001 |
|-------|-------|-------------|

ETHINYL ESTRADIOL

TABLET; ORAL

ESTINYL

| | | |
|----------|--------|-------------|
| SCHERING | 0.02MG | N005292 001 |
| | 0.05MG | N005292 002 |
| | 0.5MG | N005292 003 |

FEMINONE

| | | |
|----------------------|--------|-------------|
| PHARMACIA AND UPJOHN | 0.05MG | N016649 001 |
|----------------------|--------|-------------|

LYNORAL

| | | |
|-----------------|--------|-------------|
| ORGANON USA INC | 0.01MG | N005490 003 |
| | 0.05MG | N005490 002 |

ETHINYL ESTRADIOL; ETHYNODIOL DIACETATE

TABLET; ORAL-21

DEMULEN 1/35-21

| | | |
|---------------|-----------------|-------------|
| GD SEARLE LLC | 0.035MG; 1MG ** | N018168 001 |
|---------------|-----------------|-------------|

DEMULEN 1/50-21

| | | |
|---------------|-------------|-------------|
| GD SEARLE LLC | 0.05MG; 1MG | N016927 001 |
|---------------|-------------|-------------|

ZOVIA 1/35E-21

| | | | |
|--------------------|--------------|-------------|--------------|
| WATSON PHARMS TEVA | 0.035MG; 1MG | A072720 001 | Dec 30, 1991 |
|--------------------|--------------|-------------|--------------|

ZOVIA 1/50E-21

| | | | |
|-------------|-------------|-------------|--------------|
| WATSON LABS | 0.05MG; 1MG | A072722 001 | Dec 30, 1991 |
|-------------|-------------|-------------|--------------|

TABLET; ORAL-28

DEMULEN 1/35-28

| | | |
|---------------|-----------------|-------------|
| GD SEARLE LLC | 0.035MG; 1MG ** | N018160 001 |
|---------------|-----------------|-------------|

DEMULEN 1/50-28

| | | |
|---------------|----------------|-------------|
| GD SEARLE LLC | 0.05MG; 1MG ** | N016936 001 |
|---------------|----------------|-------------|

ETHINYL ESTRADIOL; FERROUS FUMARATE; NORETHINDRONE

TABLET; ORAL-28

NORQUEST FE

| | | | |
|---------------|--------------------|-------------|--------------|
| GD SEARLE LLC | 0.035MG; 75MG; 1MG | N018926 001 | Jul 18, 1986 |
|---------------|--------------------|-------------|--------------|

ETHINYL ESTRADIOL; FERROUS FUMARATE; NORETHINDRONE ACETATE

TABLET; ORAL-28

NORLESTRIN FE 1/50

| | | |
|-------------|-------------------|-------------|
| PARKE DAVIS | 0.05MG; 75MG; 1MG | N016766 001 |
|-------------|-------------------|-------------|

NORLESTRIN FE 2.5/50

| | | |
|-------------|---------------------|-------------|
| PARKE DAVIS | 0.05MG; 75MG; 2.5MG | N016854 001 |
|-------------|---------------------|-------------|

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL

LYBREL

| | | | |
|--------------------|-------------------|-------------|--------------|
| + WYETH PHARMS INC | 0.02MG; 0.09MG ** | N021864 001 | May 22, 2007 |
|--------------------|-------------------|-------------|--------------|

PREVEN EMERGENCY CONTRACEPTIVE KIT

| | | | |
|--------------------|----------------|-------------|--------------|
| TEVA BRANDED PHARM | 0.05MG; 0.25MG | N020946 001 | Sep 01, 1998 |
|--------------------|----------------|-------------|--------------|

TABLET; ORAL-21

ALESSE

| | | | |
|------------------|------------------|-------------|--------------|
| + CADENCE HEALTH | 0.02MG; 0.1MG ** | N020683 001 | Mar 27, 1997 |
|------------------|------------------|-------------|--------------|

AVIANE-21

| | | | |
|---------------------|---------------|-------------|--------------|
| DURAMED PHARMS BARR | 0.02MG; 0.1MG | A075796 002 | Apr 30, 2001 |
|---------------------|---------------|-------------|--------------|

ENPRESSE-21

| | | | |
|---------------------|--------------------------------------------------|-------------|--------------|
| DURAMED PHARMS BARR | 0.03MG, 0.04MG, 0.03MG; 0.05MG, 0.075MG, 0.125MG | A075809 001 | Jul 16, 2001 |
|---------------------|--------------------------------------------------|-------------|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL-21

LESSINA-21

BARR 0.02MG;0.1MG A075803 001 Mar 20, 2002

LEVLITE

+ BAYER HLTHCARE 0.02MG;0.1MG ** N020860 001 Jul 13, 1998

LEVONORGESTREL AND ETHINYL ESTRADIOL

BARR 0.02MG;0.1MG A075862 001 Apr 29, 2003

LEVORA 0.15/30-21

WATSON LABS 0.03MG;0.15MG A073592 001 Dec 13, 1993

NORDETTE-21

TEVA BRANDED PHARM 0.03MG;0.15MG N018668 001 May 10, 1982

PORTIA-21

BARR 0.03MG;0.15MG A075866 001 May 23, 2002

TRIPHASIL-21

+ WYETH PHARMS 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG ** N019192 001 Nov 01, 1984

TRIVORA-21

MAYNE PHARMA 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG A074538 001 Dec 18, 1997

TABLET; ORAL-28

ALESSE

+ CADENCE HEALTH 0.02MG;0.1MG ** N020683 002 Mar 27, 1997

LEVLITE

+ BAYER HLTHCARE 0.02MG;0.1MG ** N020860 002 Jul 13, 1998

LEVONORGESTREL AND ETHINYL ESTRADIOL

BARR 0.02MG;0.1MG A075862 002 Apr 29, 2003

NORDETTE-28

+ TEVA BRANDED PHARM 0.03MG;0.15MG ** N018782 001 Jul 21, 1982

TRIPHASIL-28

+ WYETH PHARMS INC 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG ** N019190 001 Nov 01, 1984

ETHINYL ESTRADIOL; NORELGESTROMIN

FILM, EXTENDED RELEASE; TRANSDERMAL

ORTHO EVRA

+ JANSSEN PHARMS 0.035MG/24HR;0.15MG/24HR ** N021180 001 Nov 20, 2001

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21

BALZIVA-21

BARR 0.035MG;0.4MG A076198 001 Apr 22, 2004

BREVICON 21-DAY

ALLERGAN SALES LLC 0.035MG;0.5MG N017566 001

GENCEPT 10/11-21

BARR 0.035MG,0.035MG;0.5MG,1MG A072694 001 Feb 28, 1992

MODICON 21

ORTHO MCNEIL PHARM 0.035MG;0.5MG ** N017488 001

N.E.E. 1/35 21

LPI 0.035MG;1MG A071541 001 Dec 14, 1987

NORCEPT-E 1/35 21

ORTHO MCNEIL PHARM 0.035MG;1MG A071545 001 Feb 09, 1989

NORETHIN 1/35E-21

WATSON PHARMS TEVA 0.035MG;1MG A071480 001 Apr 12, 1988

NORETHINDRONE AND ETHINYL ESTRADIOL

WATSON LABS 0.035MG;0.4MG A078379 001 Feb 23, 2010

0.035MG;0.5MG A070684 001 Jan 29, 1987

WATSON PHARMS TEVA 0.035MG;1MG A070685 001 Jan 29, 1987

NORETHINDRONE AND ETHINYL ESTRADIOL (10/11)

WATSON LABS 0.035MG,0.035MG;0.5MG,1MG A071043 001 Apr 01, 1988

NORETHINDRONE AND ETHINYL ESTRADIOL (7/14)

WATSON LABS TEVA 0.035MG,0.035MG;0.5MG,1MG A071041 001 Sep 24, 1991

NORTREL 0.5/35-21

BARR 0.035MG;0.5MG A072692 001 Feb 28, 1992

ORTHO-NOVUM 1/35-21

ORTHO MCNEIL PHARM 0.035MG;1MG ** N017489 002

ORTHO-NOVUM 10/11-21

+ ORTHO MCNEIL JANSSEN 0.035MG,0.035MG;0.5MG,1MG ** N018354 001 Jan 11, 1982

ORTHO-NOVUM 7/14-21

ORTHO MCNEIL PHARM 0.035MG,0.035MG;0.5MG,1MG ** N019004 001 Apr 04, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; NORETHINDRONE

| | | | |
|--------------------------------------------|------------------------------------------|-------------|--------------|
| TABLET;ORAL-21 | | | |
| ORTHO-NOVUM 7/7/7-21 | | | |
| JANSSEN PHARMS | 0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG | N018985 001 | Apr 04, 1984 |
| | G | | |
| OVCON-35 | | | |
| + WARNER CHILCOTT | 0.035MG;0.4MG ** | N018127 001 | |
| OVCON-50 | | | |
| WARNER CHILCOTT | 0.05MG;1MG | N018128 001 | |
| TRI-NORINYL 21-DAY | | | |
| MAYNE PHARMA | 0.035MG,0.035MG,0.035MG;0.5MG,1MG,0.5MG | N018977 001 | Apr 13, 1984 |
| TABLET;ORAL-28 | | | |
| GENCEPT 10/11-28 | | | |
| BARR | 0.035MG,0.035MG;0.5MG,1MG | A072697 001 | Feb 28, 1992 |
| MODICON 28 | | | |
| + JANSSEN PHARMS | 0.035MG;0.5MG | N017735 001 | |
| N.E.E. 1/35 28 | | | |
| LPI | 0.035MG;1MG | A071542 001 | Dec 14, 1987 |
| NORCEPT-E 1/35 28 | | | |
| ORTHO MCNEIL PHARM | 0.035MG;1MG | A071546 001 | Feb 09, 1989 |
| NORETHIN 1/35E-28 | | | |
| WATSON LABS | 0.035MG;1MG | A071481 001 | Apr 12, 1988 |
| NORETHINDRONE AND ETHINYL ESTRADIOL | | | |
| MYLAN LABS LTD | 0.035MG;0.4MG | A200897 001 | May 11, 2015 |
| | 0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG | A200486 001 | Dec 28, 2015 |
| | G | | |
| | 0.035MG;0.5MG | A200488 001 | Oct 21, 2015 |
| | 0.035MG;1MG | A200489 001 | Oct 21, 2015 |
| WATSON LABS | 0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG | A076393 001 | Feb 04, 2010 |
| | G | | |
| NORETHINDRONE AND ETHINYL ESTRADIOL (7/14) | | | |
| WATSON LABS | 0.035MG,0.035MG;0.5MG,1MG | A071042 001 | Sep 24, 1991 |
| ORTHO-NOVUM 10/11-28 | | | |
| + ORTHO MCNEIL JANSSEN | 0.035MG,0.035MG;0.5MG,1MG | N018354 002 | Jan 11, 1982 |
| ORTHO-NOVUM 7/14-28 | | | |
| ORTHO MCNEIL PHARM | 0.035MG,0.035MG;0.5MG,1MG ** | N019004 002 | Apr 04, 1984 |
| OVCON-35 | | | |
| + WARNER CHILCOTT LLC | 0.035MG;0.4MG ** | N017716 001 | |
| OVCON-50 | | | |
| WARNER CHILCOTT LLC | 0.05MG;1MG ** | N017576 001 | |

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

| | | | |
|----------------------------------------------------------------------------------------|--------------------------------------|-------------|--------------|
| TABLET;ORAL | | | |
| FEMHRT | | | |
| + APIL | 0.005MG;1MG ** | N021065 002 | Oct 15, 1999 |
| NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE | | | |
| MYLAN LABS LTD | 0.01MG,0.01MG;1MG,N/A | A205049 001 | May 31, 2016 |
| NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE | | | |
| APOTEX INC | 0.02MG;1MG | A208639 001 | Mar 21, 2018 |
| TABLET;ORAL-21 | | | |
| ESTROSTEP 21 | | | |
| + APIL | 0.02MG,0.03MG,0.035MG;1MG,1MG,1MG ** | N020130 001 | Oct 09, 1996 |
| NORLESTRIN 21 1/50 | | | |
| PARKE DAVIS | 0.05MG;1MG | N016749 001 | |
| NORLESTRIN 21 2.5/50 | | | |
| PARKE DAVIS | 0.05MG;2.5MG | N016852 001 | |
| TABLET;ORAL-28 | | | |
| NORLESTRIN 28 1/50 | | | |
| PARKE DAVIS | 0.05MG;1MG | N016723 001 | |
| TABLET, CHEWABLE, TABLET;ORAL | | | |
| LO MINASTRIN FE | | | |
| + APIL | 0.01MG,0.01MG,N/A;1MG,N/A,N/A | N204654 001 | Jul 24, 2013 |

ETHINYL ESTRADIOL; NORGESTIMATE

| | | | |
|------------------|-----------------------------------------------|-------------|--------------|
| TABLET;ORAL-21 | | | |
| ORTHO CYCLEN-21 | | | |
| JANSSEN PHARMS | 0.035MG;0.25MG | N019653 001 | Dec 29, 1989 |
| ORTHO TRI-CYCLEN | | | |
| JANSSEN PHARMS | 0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG | N019697 002 | Jul 03, 1992 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

NORGESTIMATE AND ETHINYL ESTRADIOL

WATSON LABS

0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG,
0.25MG

A090479 001 Mar 09, 2011

0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG,
0.25MG

A076626 001 Aug 17, 2006

0.035MG; 0.25MG

A076627 001 Aug 17, 2006

ETHINYL ESTRADIOL; NORGESTREL

TABLET; ORAL-21

LO/OVRAL

CADENCE HEALTH

0.03MG; 0.3MG

N017612 001

LOW-OGESTREL-21

MAYNE PHARMA

0.03MG; 0.3MG

A075288 001 Jul 28, 1999

OGESTREL 0.5/50-21

WATSON LABS

0.05MG; 0.5MG

A075406 001 Dec 15, 1999

OVRAL

WYETH PHARMS

0.05MG; 0.5MG

N016672 001

TABLET; ORAL-28

LO/OVRAL-28

WYETH PHARMS

0.03MG; 0.3MG **

N017802 001

NORGESTREL AND ETHINYL ESTRADIOL

MYLAN LABS LTD

0.03MG; 0.3MG

A201828 001 Jun 21, 2016

0.05MG; 0.5MG

A202875 001 May 08, 2017

OVRAL-28

WYETH PHARMS

0.05MG; 0.5MG

N016806 001

ETHOPROPAZINE HYDROCHLORIDE

TABLET; ORAL

PARSIDOL

PARKE DAVIS

10MG

N009078 003

50MG

N009078 006

100MG

N009078 008

ETHOTOIN

TABLET; ORAL

PEGANONE

RECORDATI RARE

500MG

N010841 003

ETHOXZOLAMIDE

TABLET; ORAL

CARDRASE

PHARMACIA AND UPJOHN

62.5MG

N011047 002

125MG

N011047 001

ETHAMIDE

ALLERGAN

125MG

N016144 001

ETHYLESTRENOL

ELIXIR; ORAL

MAXIBOLIN

ORGANON USA INC

2MG/5ML

N014006 002

TABLET; ORAL

MAXIBOLIN

ORGANON USA INC

2MG

N014005 002

ETHYNODIOL DIACETATE; MESTRANOL

TABLET; ORAL-20

OVULEN

GD SEARLE LLC

1MG; 0.1MG

N016029 002

TABLET; ORAL-21

OVULEN-21

GD SEARLE LLC

1MG; 0.1MG

N016029 003

TABLET; ORAL-28

OVULEN-28

GD SEARLE LLC

1MG; 0.1MG

N016705 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DURANEST

| | | | | | |
|---|-------------|---------|---------|-----|--|
| + | ASTRAZENECA | 0.5% ** | N017751 | 003 | |
| + | | 1% ** | N017751 | 005 | |

ETIDRONATE DISODIUM

INJECTABLE; INJECTION

DIDRONEL

| | | | | | |
|--|----------------|---------|---------|-----|--------------|
| | MGI PHARMA INC | 50MG/ML | N019545 | 001 | Apr 20, 1987 |
|--|----------------|---------|---------|-----|--------------|

TABLET; ORAL

DIDRONEL

| | | | | | |
|---|------|----------|---------|-----|--|
| + | APIL | 200MG ** | N017831 | 001 | |
| + | | 400MG ** | N017831 | 002 | |

ETODOLAC

CAPSULE; ORAL

ETODOLAC

| | | | | | |
|--|---------------------|-------|---------|-----|--------------|
| | ANI PHARMS INC | 200MG | A074840 | 001 | Aug 29, 1997 |
| | | 200MG | A074899 | 001 | Jul 08, 1997 |
| | | 300MG | A074840 | 002 | Aug 29, 1997 |
| | | 300MG | A074899 | 002 | Jul 08, 1997 |
| | CHARTWELL MOLECULES | 200MG | A074842 | 001 | Jul 17, 1997 |
| | | 300MG | A074842 | 002 | Jul 17, 1997 |
| | ECI PHARMS LLC | 300MG | A074929 | 001 | Jan 30, 1998 |
| | MYLAN | 200MG | A074932 | 001 | May 16, 1997 |
| | | 200MG | A075071 | 001 | Sep 30, 1998 |
| | | 300MG | A074932 | 002 | May 16, 1997 |
| | | 300MG | A075071 | 002 | Sep 30, 1998 |
| | SANDOZ | 200MG | A074942 | 001 | Sep 30, 1997 |
| | | 300MG | A074942 | 002 | Sep 30, 1997 |
| | WATSON LABS | 200MG | A074844 | 001 | Dec 23, 1997 |
| | | 300MG | A074844 | 002 | Dec 23, 1997 |

LODINE

| | | | | | |
|---|------------------|----------|---------|-----|--------------|
| + | WYETH PHARMS INC | 200MG ** | N018922 | 002 | Jan 31, 1991 |
| + | | 300MG | N018922 | 003 | Jan 31, 1991 |

TABLET; ORAL

ETODOLAC

| | | | | | |
|--|----------------------|-------|---------|-----|--------------|
| | CHARTWELL MOLECULES | 400MG | A074841 | 001 | Jun 27, 1997 |
| | ECI PHARMS LLC | 400MG | A074927 | 001 | Oct 30, 1997 |
| | IVAX SUB TEVA PHARMS | 400MG | A074883 | 001 | Feb 28, 1997 |
| | | 500MG | A074883 | 002 | Nov 20, 1998 |
| | MYLAN | 400MG | A075012 | 001 | Sep 30, 1998 |
| | | 500MG | A075012 | 002 | Sep 30, 1998 |
| | MYLAN PHARMS INC | 400MG | A075104 | 001 | Feb 06, 1998 |
| | | 500MG | A075104 | 002 | Nov 20, 1998 |
| | OXFORD PHARMS | 400MG | A074819 | 001 | Feb 28, 1997 |
| | | 500MG | A074819 | 002 | Apr 28, 1998 |
| | RANBAXY LABS LTD | 400MG | A075226 | 001 | Nov 24, 1998 |
| | | 500MG | A075226 | 002 | Nov 24, 1998 |
| | SANDOZ | 400MG | A074839 | 001 | Jul 11, 1997 |
| | | 400MG | A074846 | 001 | Feb 28, 1997 |
| | TEVA | 400MG | A074847 | 001 | Apr 23, 1999 |
| | | 500MG | A074847 | 002 | Apr 23, 1999 |
| | WATSON LABS | 400MG | A074892 | 001 | Apr 16, 1997 |
| | | 400MG | A075069 | 001 | Apr 16, 1998 |
| | | 500MG | A074892 | 002 | Oct 29, 1998 |

LODINE

| | | | | | |
|---|------------------|----------|---------|-----|--------------|
| + | WYETH PHARMS INC | 400MG ** | N018922 | 004 | Jul 29, 1993 |
| + | | 500MG ** | N018922 | 005 | Jun 28, 1996 |

TABLET, EXTENDED RELEASE; ORAL

ETODOLAC

| | | | | | |
|--|---------------------|-------|---------|-----|--------------|
| | ACTAVIS ELIZABETH | 400MG | A075696 | 001 | Jul 31, 2000 |
| | ANI PHARMS INC | 400MG | A075943 | 001 | Jul 26, 2002 |
| | | 500MG | A075943 | 002 | Jul 26, 2002 |
| | | 600MG | A075943 | 003 | Jul 26, 2002 |
| | WATSON LABS FLORIDA | 400MG | A075829 | 001 | Nov 30, 2001 |
| | | 500MG | A075829 | 002 | Nov 30, 2001 |

LODINE XL

| | | | | | |
|--|------------------|----------|---------|-----|--------------|
| | WYETH PHARMS INC | 400MG ** | N020584 | 001 | Oct 25, 1996 |
|--|------------------|----------|---------|-----|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ETODOLACTABLET, EXTENDED RELEASE;ORAL
LODINE XL

| | | | |
|---|----------|-------------|--------------|
| | 500MG ** | N020584 003 | Jan 20, 1998 |
| + | 600MG ** | N020584 002 | Oct 25, 1996 |

ETONOGESTREL

IMPLANT; IMPLANTATION

IMPLANON

| | | | |
|-----------------|--------------|-------------|--------------|
| ORGANON USA INC | 68MG/IMPLANT | N021529 001 | Jul 17, 2006 |
|-----------------|--------------|-------------|--------------|

ETOPOSIDE

CAPSULE; ORAL

VEPESID

| | | | | |
|---|-----------------|-------|-------------|--------------|
| + | DAVA PHARMS INC | 50MG | N019557 001 | Dec 30, 1986 |
| + | | 100MG | N019557 002 | Dec 30, 1986 |

INJECTABLE; INJECTION

ETOPOSIDE

HOSPIRA

| | | | |
|--|---------|-------------|--------------|
| | 20MG/ML | A074320 001 | Aug 30, 1995 |
| | 20MG/ML | A074351 001 | Aug 30, 1995 |

| | | | |
|-----------------|---------|-------------|--------------|
| PHARMACHEMIE BV | 20MG/ML | A074227 001 | Feb 22, 1996 |
|-----------------|---------|-------------|--------------|

| | | | |
|--------------|---------|-------------|--------------|
| PIERRE FABRE | 20MG/ML | A074813 001 | Jul 09, 1997 |
|--------------|---------|-------------|--------------|

| | | | |
|-----------------|---------|-------------|--------------|
| TEVA PARENTERAL | 20MG/ML | A074510 001 | Jun 29, 1995 |
|-----------------|---------|-------------|--------------|

| | | | |
|-----------------|---------|-------------|--------------|
| TEVA PHARMS USA | 20MG/ML | A074284 001 | Feb 10, 1994 |
|-----------------|---------|-------------|--------------|

| | | | |
|-------------|---------|-------------|--------------|
| WATSON LABS | 20MG/ML | A074228 001 | Oct 15, 1996 |
|-------------|---------|-------------|--------------|

| | | | |
|-----------------|---------|-------------|--------------|
| WATSON LABS INC | 20MG/ML | A074968 001 | Jan 09, 1998 |
|-----------------|---------|-------------|--------------|

TOPOSAR

| | | | |
|-----------------|---------|-------------|--------------|
| TEVA PARENTERAL | 20MG/ML | A074166 001 | Feb 27, 1995 |
|-----------------|---------|-------------|--------------|

VEPESID

| | | | | |
|---|---------------|------------|-------------|--------------|
| + | CORDEN PHARMA | 20MG/ML ** | N018768 001 | Nov 10, 1983 |
|---|---------------|------------|-------------|--------------|

ETOPOSIDE PHOSPHATE

INJECTABLE; INJECTION

ETOPOPHOS PRESERVATIVE FREE

| | | | |
|----------------------|--------------------|-------------|--------------|
| BRISTOL MYERS SQUIBB | EQ 500MG BASE/VIAL | N020906 001 | Feb 27, 1998 |
|----------------------|--------------------|-------------|--------------|

| | | | |
|--|------------------|-------------|--------------|
| | EQ 1GM BASE/VIAL | N020906 002 | Feb 27, 1998 |
|--|------------------|-------------|--------------|

ETRETINATE

CAPSULE; ORAL

TEGISON

ROCHE

| | | | |
|--|------|-------------|--------------|
| | 10MG | N019369 001 | Sep 30, 1986 |
|--|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 25MG | N019369 002 | Sep 30, 1986 |
|--|------|-------------|--------------|

EVANS BLUE

INJECTABLE; INJECTION

EVANS BLUE

| | | | |
|-------------|---------|-------------|--|
| PARKE DAVIS | 0.5% ** | N008041 001 | |
|-------------|---------|-------------|--|

EZOGABINE

TABLET; ORAL

POTIGA

| | | | | |
|---|-----------------|------|-------------|--------------|
| + | GLAXOSMITHKLINE | 50MG | N022345 001 | Jun 10, 2011 |
|---|-----------------|------|-------------|--------------|

| | | | | |
|---|--|-------|-------------|--------------|
| + | | 200MG | N022345 002 | Jun 10, 2011 |
|---|--|-------|-------------|--------------|

| | | | | |
|---|--|-------|-------------|--------------|
| + | | 300MG | N022345 003 | Jun 10, 2011 |
|---|--|-------|-------------|--------------|

| | | | | |
|---|--|-------|-------------|--------------|
| + | | 400MG | N022345 004 | Jun 10, 2011 |
|---|--|-------|-------------|--------------|

FAMCICLOVIR

TABLET; ORAL

FAMVIR

| | | | | |
|---|----------|----------|-------------|--------------|
| + | NOVARTIS | 125MG ** | N020363 003 | Dec 11, 1995 |
|---|----------|----------|-------------|--------------|

| | | | | |
|---|--|----------|-------------|--------------|
| + | | 250MG ** | N020363 001 | Apr 26, 1996 |
|---|--|----------|-------------|--------------|

| | | | | |
|---|--|----------|-------------|--------------|
| + | | 500MG ** | N020363 002 | Jun 29, 1994 |
|---|--|----------|-------------|--------------|

FAMOTIDINE

INJECTABLE; INJECTION

FAMOTIDINE

APOTEX INC

| | | | |
|--|---------|-------------|--------------|
| | 10MG/ML | A075942 001 | Aug 02, 2002 |
|--|---------|-------------|--------------|

| | | | |
|-----------|---------|-------------|--------------|
| APOTHECON | 10MG/ML | A075707 001 | Apr 16, 2001 |
|-----------|---------|-------------|--------------|

| | | | |
|---------|---------|-------------|--------------|
| HOSPIRA | 10MG/ML | A075705 001 | Apr 16, 2001 |
|---------|---------|-------------|--------------|

| | | | |
|--|---------|-------------|--------------|
| | 10MG/ML | A075870 001 | Nov 23, 2001 |
|--|---------|-------------|--------------|

| | | | |
|--|---------|-------------|--------------|
| | 10MG/ML | A075905 001 | Nov 23, 2001 |
|--|---------|-------------|--------------|

| | | | |
|----------------------|---------|-------------|--------------|
| WEST-WARD PHARMS INT | 10MG/ML | A075799 001 | Apr 30, 2002 |
|----------------------|---------|-------------|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FAMOTIDINE

INJECTABLE; INJECTION

FAMOTIDINE PRESERVATIVE FREE

| | | | | |
|----------------------|---------|---------|-----|--------------|
| APOTEX INC | 10MG/ML | A076324 | 001 | Nov 27, 2002 |
| APOTHECON | 10MG/ML | A075708 | 001 | Apr 16, 2001 |
| HOSPIRA | 10MG/ML | A075669 | 001 | Apr 16, 2001 |
| WEST-WARD PHARMS INT | 10MG/ML | A075789 | 001 | Apr 30, 2002 |

FAMOTIDINE PRESERVATIVE FREE (PHARMACY BULK)

| | | | | |
|------------|---------|---------|-----|--------------|
| APOTEX INC | 10MG/ML | A076322 | 001 | Nov 27, 2002 |
|------------|---------|---------|-----|--------------|

FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER

| | | | | |
|--------|----------|---------|-----|--------------|
| ABBVIE | 0.4MG/ML | A075729 | 001 | Dec 17, 2001 |
|--------|----------|---------|-----|--------------|

PEPCID

| | | | | |
|---------|------------|---------|-----|--------------|
| + MERCK | 10MG/ML ** | N019510 | 001 | Nov 04, 1986 |
|---------|------------|---------|-----|--------------|

PEPCID PRESERVATIVE FREE

| | | | | |
|---------|------------|---------|-----|--------------|
| + MERCK | 10MG/ML ** | N019510 | 004 | Nov 04, 1986 |
|---------|------------|---------|-----|--------------|

PEPCID PRESERVATIVE FREE IN PLASTIC CONTAINER

| | | | | |
|---------------------|-------------|---------|-----|--------------|
| + MERCK SHARP DOHME | 0.4MG/ML ** | N020249 | 001 | Feb 18, 1994 |
|---------------------|-------------|---------|-----|--------------|

TABLET; ORAL

FAMOTIDINE

| | | | | |
|----------------------|------|---------|-----|--------------|
| ACTAVIS ELIZABETH | 20MG | A075650 | 001 | Sep 14, 2001 |
| | 40MG | A075650 | 002 | Sep 14, 2001 |
| APOTEX | 10MG | A075610 | 001 | Mar 12, 2002 |
| MYLAN PHARMS INC | 20MG | A075457 | 001 | Apr 18, 2001 |
| | 40MG | A075457 | 002 | Apr 18, 2001 |
| PLD ACQUISITIONS | 20MG | A075302 | 001 | Apr 16, 2001 |
| | 40MG | A075302 | 002 | Apr 16, 2001 |
| SANDOZ | 10MG | A076101 | 001 | Oct 21, 2002 |
| | 20MG | A075607 | 001 | May 10, 2001 |
| | 20MG | A075793 | 001 | Apr 16, 2001 |
| | 40MG | A075607 | 002 | May 10, 2001 |
| | 40MG | A075793 | 002 | Apr 16, 2001 |
| SUN PHARM INDUSTRIES | 20MG | A075639 | 002 | Dec 12, 2001 |
| | 40MG | A075639 | 001 | Dec 12, 2001 |
| WATSON LABS | 10MG | A075404 | 001 | Nov 28, 2001 |
| | 20MG | A075062 | 002 | Apr 16, 2001 |
| | 40MG | A075062 | 001 | Apr 16, 2001 |

TABLET, CHEWABLE; ORAL

PEPCID AC

| | | | | |
|------------------------|---------|---------|-----|--------------|
| + J AND J CONSUMER INC | 10MG ** | N020801 | 001 | Sep 24, 1998 |
|------------------------|---------|---------|-----|--------------|

TABLET, ORALLY DISINTEGRATING; ORAL

FLUXID

| | | | | |
|---------|------|---------|-----|--------------|
| UCB INC | 20MG | N021712 | 001 | Sep 24, 2004 |
| | 40MG | N021712 | 002 | Sep 24, 2004 |

PEPCID RPD

| | | | | |
|-------|------|---------|-----|--------------|
| MERCK | 20MG | N020752 | 001 | May 28, 1998 |
| | 40MG | N020752 | 002 | May 28, 1998 |

FELODIPINE

TABLET, EXTENDED RELEASE; ORAL

FELODIPINE

| | | | | |
|---------------|-------|---------|-----|--------------|
| WOCKHARDT LTD | 2.5MG | A091484 | 001 | Aug 15, 2012 |
| | 5MG | A091484 | 002 | Aug 15, 2012 |
| | 10MG | A091484 | 003 | Aug 15, 2012 |

PLENDIL

| | | | | |
|---------------|----------|---------|-----|--------------|
| + ASTRAZENECA | 2.5MG ** | N019834 | 004 | Sep 22, 1994 |
| | 5MG ** | N019834 | 001 | Jul 25, 1991 |
| | 10MG ** | N019834 | 002 | Jul 25, 1991 |

FENOFIBRATE

CAPSULE; ORAL

ANTARA (MICRONIZED)

| | | | | |
|----------------|------|---------|-----|--------------|
| LUPIN ATLANTIS | 87MG | N021695 | 002 | Nov 30, 2004 |
|----------------|------|---------|-----|--------------|

LIPIDIL

| | | | | |
|--------|-------|---------|-----|--------------|
| ABBVIE | 100MG | N019304 | 001 | Dec 31, 1993 |
|--------|-------|---------|-----|--------------|

LIPOFEN

| | | | | |
|-------------------|-------|---------|-----|--------------|
| CIPHER PHARMS INC | 100MG | N021612 | 002 | Jan 11, 2006 |
|-------------------|-------|---------|-----|--------------|

TRICOR (MICRONIZED)

| | | | | |
|----------|----------|---------|-----|--------------|
| + ABBVIE | 67MG ** | N019304 | 002 | Feb 09, 1998 |
| | 134MG ** | N019304 | 003 | Jun 30, 1999 |
| | 200MG ** | N019304 | 004 | Jun 30, 1999 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FENOFIBRATE

TABLET; ORAL

FENOFIBRATE

MYLAN

107MG

A076520 002 Dec 29, 2005

TRICOR

+ ABBVIE INC

54MG **

N021203 001 Sep 04, 2001

+

160MG **

N021203 003 Sep 04, 2001

TRIGLIDE

SKYE PHARMA AG

50MG

N021350 001 May 07, 2005

FENOLDOPAM MESYLATE

INJECTABLE; INJECTION

FENOLDOPAM MESYLATE

LUITPOLD

EQ 10MG BASE/ML

A076656 001 Dec 01, 2003

TEVA PARENTERAL

EQ 10MG BASE/ML

A077826 001 Mar 07, 2007

FENOPROFEN CALCIUM

CAPSULE; ORAL

FENOPROFEN CALCIUM

AM THERAP

EQ 200MG BASE

A072307 001 Aug 22, 1988

EQ 300MG BASE

A072308 001 Aug 22, 1988

AUROLIFE PHARMA LLC

EQ 200MG BASE

A072394 001 Oct 17, 1988

EQ 300MG BASE

A072395 001 Oct 17, 1988

HALSEY

EQ 200MG BASE

A072355 001 Aug 17, 1988

EQ 300MG BASE

A072356 001 Aug 17, 1988

PAR PHARM

EQ 200MG BASE

A072437 001 Aug 22, 1988

EQ 300MG BASE

A072438 001 Aug 22, 1988

QUANTUM PHARMICS

EQ 200MG BASE

A072214 001 Aug 17, 1988

EQ 300MG BASE

A071738 001 Aug 17, 1988

WARNER CHILCOTT

EQ 200MG BASE

A072946 001 Apr 30, 1991

EQ 300MG BASE

A072472 001 Apr 30, 1991

WATSON LABS

EQ 200MG BASE

A072294 001 Aug 17, 1988

EQ 200MG BASE

A072981 001 Aug 19, 1991

EQ 300MG BASE

A072293 001 Aug 17, 1988

EQ 300MG BASE

A072982 001 Aug 19, 1991

NALFON

XSPIRE PHARMA

EQ 300MG BASE

N017604 002

TABLET; ORAL

FENOPROFEN CALCIUM

ACTAVIS ELIZABETH

EQ 600MG BASE

A072274 001 May 02, 1988

AM THERAP

EQ 600MG BASE

A072309 001 Aug 17, 1988

AUROLIFE PHARMA LLC

EQ 600MG BASE

A072396 001 Oct 17, 1988

DAVA PHARMS INC

EQ 600MG BASE

A072326 001 Aug 17, 1988

HALSEY

EQ 600MG BASE

A072357 001 Aug 17, 1988

IVAX SUB TEVA PHARMS

EQ 600MG BASE

A072557 001 Aug 29, 1988

PAR PHARM

EQ 600MG BASE

A072429 001 Aug 17, 1988

QUANTUM PHARMICS

EQ 600MG BASE

A072194 001 Aug 17, 1988

SUN PHARM INDUSTRIES

EQ 600MG BASE

A072902 001 Dec 21, 1990

USL PHARMA

EQ 600MG BASE

A072362 001 Aug 17, 1988

WATSON LABS

EQ 600MG BASE

A072165 001 Aug 17, 1988

EQ 600MG BASE

A072602 001 Oct 11, 1988

WATSON LABS TEVA

EQ 600MG BASE

A072407 001 Aug 17, 1988

NALFON

DISTA

EQ 600MG BASE

N017710 001

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL

FENTANYL-100

ACTAVIS LABS UT INC

100MCG/HR

A076709 004 Aug 20, 2007

NOVEN

100MCG/HR

A077775 004 Oct 16, 2009

FENTANYL-25

ACTAVIS LABS UT INC

25MCG/HR

A076709 001 Aug 20, 2007

NOVEN

25MCG/HR

A077775 001 Oct 16, 2009

FENTANYL-50

ACTAVIS LABS UT INC

50MCG/HR

A076709 002 Aug 20, 2007

NOVEN

50MCG/HR

A077775 002 Oct 16, 2009

FENTANYL-75

ACTAVIS LABS UT INC

75MCG/HR

A076709 003 Aug 20, 2007

NOVEN

75MCG/HR

A077775 003 Oct 16, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FENTANYL CITRATE

FILM;BUCCAL

ONSOLIS

| | | | |
|------|---------------|-------------|--------------|
| BDSI | EQ 0.2MG BASE | N022266 001 | Jul 16, 2009 |
| | EQ 0.4MG BASE | N022266 002 | Jul 16, 2009 |
| | EQ 0.6MG BASE | N022266 003 | Jul 16, 2009 |
| | EQ 0.8MG BASE | N022266 004 | Jul 16, 2009 |
| | EQ 1.2MG BASE | N022266 005 | Jul 16, 2009 |

INJECTABLE;INJECTION

FENTANYL CITRATE

ABBOTT

| | | |
|-------------------|-------------|--------------|
| EQ 0.05MG BASE/ML | A070636 001 | Apr 30, 1990 |
| EQ 0.05MG BASE/ML | A070637 001 | Apr 30, 1990 |

WATSON LABS

| | | |
|-------------------|-------------|--------------|
| EQ 0.05MG BASE/ML | A073488 001 | Jun 30, 1992 |
|-------------------|-------------|--------------|

FENTANYL CITRATE PRESERVATIVE FREE

WATSON LABS INC

| | | |
|-------------------|-------------|--------------|
| EQ 0.05MG BASE/ML | A074917 001 | Feb 03, 1998 |
|-------------------|-------------|--------------|

TABLET;BUCCAL, SUBLINGUAL

FENTANYL CITRATE

WATSON LABS

| | | |
|---------------|-------------|--------------|
| EQ 0.1MG BASE | A079075 001 | Jan 07, 2011 |
| EQ 0.2MG BASE | A079075 002 | Jan 07, 2011 |
| EQ 0.4MG BASE | A079075 003 | Jan 07, 2011 |
| EQ 0.6MG BASE | A079075 004 | Jan 07, 2011 |
| EQ 0.8MG BASE | A079075 005 | Jan 07, 2011 |

FENTORA

+ CEPHALON

| | | |
|------------------|-------------|--------------|
| EQ 0.3MG BASE ** | N021947 006 | Mar 02, 2007 |
|------------------|-------------|--------------|

TROCHE/LOZENGE;ORAL

FENTANYL

CEPHALON

| | | |
|---------------|-------------|--------------|
| EQ 0.1MG BASE | N020195 007 | Oct 30, 1995 |
| EQ 0.2MG BASE | N020195 001 | Oct 04, 1993 |
| EQ 0.3MG BASE | N020195 002 | Oct 04, 1993 |
| EQ 0.4MG BASE | N020195 003 | Oct 04, 1993 |

TROCHE/LOZENGE;TRANSMUCOSAL

FENTANYL CITRATE

PAR PHARM

| | | |
|---------------|-------------|--------------|
| EQ 0.2MG BASE | A077312 001 | Oct 30, 2009 |
| EQ 0.4MG BASE | A077312 002 | Oct 30, 2009 |
| EQ 0.6MG BASE | A077312 003 | Oct 30, 2009 |
| EQ 0.8MG BASE | A077312 004 | Oct 30, 2009 |
| EQ 1.2MG BASE | A077312 005 | Oct 30, 2009 |
| EQ 1.6MG BASE | A077312 006 | Oct 30, 2009 |

FENTANYL HYDROCHLORIDE

SYSTEM;IONTOPHORESIS, TRANSDERMAL

IONSYS

+ THE MEDICINES CO

| | | |
|--------------------------|-------------|--------------|
| EQ 40MCG BASE/ACTIVATION | N021338 001 | May 22, 2006 |
|--------------------------|-------------|--------------|

FERRIC AMMONIUM CITRATE

FOR SOLUTION;ORAL

FERRISELTZ

OTSUKA

| | | |
|--------------|-------------|--------------|
| 600MG/PACKET | N020292 001 | Oct 14, 1997 |
|--------------|-------------|--------------|

FERRIC PYROPHOSPHATE CITRATE

SOLUTION;INTRAVENOUS

TRIFERIC

+ ROCKWELL MEDICAL INC

| | | |
|----------------------------------|-------------|--------------|
| 272MG IRON/50ML (5.44MG IRON/ML) | N206317 002 | Sep 04, 2015 |
|----------------------------------|-------------|--------------|

FERROUS CITRATE, FE-59

INJECTABLE;INJECTION

FERROUS CITRATE FE 59

MALLINCKRODT

| | | |
|----------|-------------|--|
| 25uCi/ML | N016729 001 | |
|----------|-------------|--|

FERROUS SULFATE; FOLIC ACID

CAPSULE;ORAL

FOLVRON

LEDERLE

| | | |
|--------------|-------------|--|
| 182MG;0.33MG | N006012 003 | |
|--------------|-------------|--|

FERUMOXIDES

INJECTABLE;INJECTION

FERIDEX I.V.

AMAG PHARMS INC

| | | |
|-------------------|-------------|--------------|
| EQ 11.2MG IRON/ML | N020416 001 | Aug 30, 1996 |
|-------------------|-------------|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FERUMOXSISUSPENSION;ORAL
GASTROMARK

AMAG PHARMS INC EQ 0.175MG IRON/ML N020410 001 Dec 06, 1996

FESOTERODINE FUMARATETABLET, EXTENDED RELEASE;ORAL
FESOTERODINE FUMARATEALKEM LABS LTD 4MG A204827 001 Dec 10, 2015
8MG A204827 002 Dec 10, 2015FEXOFENADINE HYDROCHLORIDE

CAPSULE;ORAL

ALLEGRA

SANOFI AVENTIS US 60MG ** N020625 001 Jul 25, 1996

FEXOFENADINE HYDROCHLORIDE

BARR 60MG A076169 001 Jul 13, 2005

SUSPENSION;ORAL

ALLEGRA

+ SANOFI AVENTIS US 30MG/5ML N021963 001 Oct 16, 2006

CHILDREN'S ALLEGRA HIVES

+ SANOFI AVENTIS US 30MG/5ML N201373 002 Jan 24, 2011

CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

TARO PHARM 30MG/5ML A208123 001 Nov 09, 2017

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

TARO PHARM 30MG/5ML A208123 002 Nov 09, 2017

FEXOFENADINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC 30MG/5ML A201311 001 Jul 25, 2012

TABLET;ORAL

ALLEGRA HIVES

+ SANOFI AVENTIS US 60MG N020872 008 Jan 24, 2011

+ 180MG N020872 009 Jan 24, 2011

CHILDREN'S ALLEGRA ALLERGY

+ SANOFI AVENTIS US 30MG N020872 005 Jan 24, 2011

CHILDREN'S ALLEGRA HIVES

+ SANOFI AVENTIS US 30MG N020872 006 Jan 24, 2011

TABLET, ORALLY DISINTEGRATING;ORAL

CHILDREN'S ALLEGRA ALLERGY

+ SANOFI AVENTIS US 30MG N021909 002 Jan 24, 2011

CHILDREN'S ALLEGRA HIVES

+ SANOFI AVENTIS US 30MG N021909 003 Jan 24, 2011

CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

DR REDDYS LABS LTD 30MG A202978 001 Jan 18, 2013

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

DR REDDYS LABS LTD 30MG A202978 002 Jan 18, 2013

FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

BARR 60MG;120MG A076236 001 Apr 14, 2005

IMPAX PHARMS 60MG;120MG A076298 001 Nov 12, 2010

FIBRINOGEN, I-125

INJECTABLE;INJECTION

IBRIN

GE HEALTHCARE 154uCi/VIAL N017879 001

RADIONUCLIDE-LABELED (125 I) FIBRINOGEN (HUMAN) SENSOR

ABBOTT 140uCi/ML N017787 001

FINASTERIDE

TABLET;ORAL

FINASTERIDE

GEDEON RICHTER USA 5MG A077251 001 Dec 22, 2006

IVAX SUB TEVA PHARMS 5MG A076340 001 Jun 19, 2006

MYLAN PHARMS INC 1MG A078161 001 Nov 05, 2013

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLAVOXATE HYDROCHLORIDE

TABLET; ORAL

FLAVOXATE HYDROCHLORIDE

IMPAX PHARMS 100MG A076234 001 Aug 28, 2003

URISPAS

ORTHO MCNEIL JANSSEN 100MG N016769 001

FLECAINIDE ACETATE

TABLET; ORAL

FLECAINIDE ACETATE

ANI PHARMS INC 50MG A076030 001 Oct 28, 2002

100MG A076030 002 Oct 28, 2002

150MG A076030 003 Oct 28, 2002

APOTEX INC

50MG A079164 001 Jul 09, 2009

100MG A079164 002 Jul 09, 2009

150MG A079164 003 Jul 09, 2009

TAMBOCOR

CNTY LINE PHARMS 200MG N018830 002 Oct 31, 1985

FLORBETAPIR F-18

SOLUTION; INTRAVENOUS

AMYVID

AVID RADIOPHARMS INC 10ML (13.5-51mCi/ML) N202008 001 Apr 06, 2012

FLOXURIDINE

INJECTABLE; INJECTION

FUDR

+ HOSPIRA 500MG/VIAL ** N016929 001

FLUCONAZOLE

FOR SUSPENSION; ORAL

FLUCONAZOLE

SUN PHARM INDS LTD 50MG/5ML A076332 001 Jul 29, 2004

200MG/5ML A076332 002 Jul 29, 2004

TARO PHARM INDS

50MG/5ML A076918 001 Dec 18, 2006

200MG/5ML A076918 002 Dec 18, 2006

INJECTABLE; INJECTION

DIFLUCAN IN DEXTROSE 5% IN PLASTIC CONTAINER

+ PFIZER 200MG/100ML (2MG/ML) N019950 003 Sep 29, 1992

+ 400MG/200ML (2MG/ML) N019950 005 Jul 08, 1994

DIFLUCAN IN SODIUM CHLORIDE 0.9%

+ PFIZER 200MG/100ML (2MG/ML) N019950 001 Jan 29, 1990

+ 400MG/200ML (2MG/ML) N019950 006 Jan 29, 1990

DIFLUCAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+ PFIZER 200MG/100ML (2MG/ML) N019950 002 Jan 29, 1990

+ 400MG/200ML (2MG/ML) N019950 004 Jan 29, 1990

FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER

MYLAN LABS LTD 200MG/100ML (2MG/ML) A076888 001 Mar 25, 2005

400MG/200ML (2MG/ML) A076888 002 Mar 25, 2005

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

TEVA PHARMS USA 200MG/100ML (2MG/ML) A076653 001 Jul 29, 2004

400MG/200ML (2MG/ML) A076653 002 Jul 29, 2004

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

HOSPIRA 200MG/100ML (2MG/ML) A076617 001 Jul 29, 2004

400MG/200ML (2MG/ML) A076617 002 Jul 29, 2004

MYLAN LABS LTD 200MG/100ML (2MG/ML) A076889 001 Mar 25, 2005

400MG/200ML (2MG/ML) A076889 002 Mar 25, 2005

TEVA PHARMS 200MG/100ML (2MG/ML) A076837 001 Jan 13, 2005

400MG/200ML (2MG/ML) A076837 002 Jan 13, 2005

TABLET; ORAL

FLUCONAZOLE

ANI PHARMS INC 50MG A076086 001 Jul 29, 2004

100MG A076086 002 Jul 29, 2004

150MG A076086 003 Jul 29, 2004

200MG A076086 004 Jul 29, 2004

GEDEON RICHTER USA

50MG A076432 001 Jul 29, 2004

100MG A076432 002 Jul 29, 2004

150MG A076432 003 Jul 29, 2004

200MG A076432 004 Jul 29, 2004

MYLAN PHARMS INC 50MG A076042 001 Jul 29, 2004

100MG A076042 002 Jul 29, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUCONAZOLETABLET; ORAL
FLUCONAZOLE

| | | | |
|------------------|-------|-------------|--------------|
| | 150MG | A076042 003 | Jul 29, 2004 |
| | 200MG | A076042 004 | Jul 29, 2004 |
| PLIVA | 50MG | A076424 001 | Jul 29, 2004 |
| | 100MG | A076424 002 | Jul 29, 2004 |
| | 150MG | A076424 003 | Jul 29, 2004 |
| | 200MG | A076424 004 | Jul 29, 2004 |
| RANBAXY LABS LTD | 50MG | A076386 001 | Jul 29, 2004 |
| | 100MG | A076386 002 | Jul 29, 2004 |
| | 150MG | A076386 003 | Jul 29, 2004 |
| | 200MG | A076386 004 | Jul 29, 2004 |
| ROXANE | 50MG | A076213 001 | Jul 29, 2004 |
| | 100MG | A076213 002 | Jul 29, 2004 |
| | 150MG | A076213 003 | Jul 29, 2004 |
| | 200MG | A076213 004 | Jul 29, 2004 |

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARA

+ GENZYME CORP

50MG/VIAL **

N020038 001 Apr 18, 1991

TABLET; ORAL

OFORTA

SANOFI AVENTIS US

10MG

N022273 001 Dec 18, 2008

FLUDEOXYGLUCOSE F-18

INJECTABLE; INJECTION

FLUDEOXYGLUCOSE F18

+ DOWNSTATE CLINCL

4-40mCi/ML **

N020306 001 Aug 19, 1994

+

4-90mCi/ML **

N020306 002 Sep 25, 2001

INJECTABLE; INTRAVENOUS

FLUDEOXYGLUCOSE F18

+ FEINSTEIN

20-200mCi/ML

N021870 001 Aug 19, 2005

MIDWEST MEDCL

20-200mCi/ML

A203736 001 Nov 19, 2015

WEILL MEDCL COLL

10-100mCi/ML **

N021768 001 Aug 05, 2004

FLUDROCORTISONE ACETATE

TABLET; ORAL

FLORINEF

+ CASPER PHARMA LLC

0.1MG **

N010060 001

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

BAXTER HLTHCARE CORP 0.5MG/5ML (0.1MG/ML)

A076755 002 Oct 12, 2004

1MG/10ML (0.1MG/ML)

A076755 001 Oct 12, 2004

TEVA PHARMS USA 0.5MG/5ML (0.1MG/ML)

A076589 002 Oct 12, 2004

1MG/10ML (0.1MG/ML)

A076589 001 Oct 12, 2004

ROMAZICON

+ HOFFMANN LA ROCHE

1MG/10ML (0.1MG/ML) **

N020073 001 Dec 20, 1991

+

0.5MG/5ML (0.1MG/ML) **

N020073 002 Dec 20, 1991

FLUMETHASONE PIVALATE

CREAM; TOPICAL

LOCORTEN

NOVARTIS

0.03%

N016379 001

FLUNISOLIDE

AEROSOL, METERED; INHALATION

AEROBID

ROCHE PALO

0.25MG/INH

N018340 001 Aug 17, 1984

SPRAY, METERED; NASAL

FLUNISOLIDE

APOTEX INC

0.029MG/SPRAY

A077436 001 Aug 09, 2007

NASALIDE

IVAX RES

0.025MG/SPRAY **

N018148 001

NASAREL

TEVA BRANDED PHARM

0.029MG/SPRAY

N020409 001 Mar 08, 1995

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUOCINOLONE ACETONIDE

CREAM;TOPICAL

FLUOCET

ALPHARMA US PHARMS 0.025% A088360 001 Jan 16, 1984

FLUOCINOLONE ACETONIDE

ALPHARMA US PHARMS 0.01% A088361 001 Jan 16, 1984

G AND W LABS 0.025% A089525 001 Jul 26, 1988

PERRIGO NEW YORK 0.01% A086810 001 Mar 04, 1982

0.025% A086811 001 Mar 04, 1982

PHARMADERM 0.01% A088047 001 Dec 16, 1982

0.025% A088045 001 Dec 16, 1982

PHARMAFAIR 0.01% A088499 001 Aug 02, 1984

0.025% A088506 001 Aug 02, 1984

TARO 0.01% A040035 001 Oct 31, 1994

0.01% A087102 001 Apr 27, 1982

0.025% A040042 001 Oct 31, 1994

USL PHARMA 0.01% A088757 001 Feb 11, 1985

0.025% A088756 001 Mar 28, 1985

FLUONID

ALLERGAN HERBERT 0.025% A087156 002 Sep 06, 1984

FLUOTREX

SAVAGE LABS 0.01% A088174 001 May 06, 1983

0.025% A088173 001 Mar 09, 1983

SYNALAR-HP

MEDIMETRIKS PHARMS 0.2% N016161 002

GEL;TOPICAL

FLUONID

ALLERGAN HERBERT 0.025% A087300 001 May 27, 1982

OINTMENT;TOPICAL

FLUOCINOLONE ACETONIDE

PHARMADERM 0.025% A088046 001 Dec 16, 1982

PHARMAFAIR 0.025% A088507 001 Feb 27, 1984

USL PHARMA 0.025% A088742 001 Feb 08, 1985

FLUONID

ALLERGAN HERBERT 0.025% A087157 001 Sep 06, 1984

FLUOTREX

SAVAGE LABS 0.025% A088172 001 Mar 09, 1983

SOLUTION;TOPICAL

FLUOCINOLONE ACETONIDE

ALPHARMA US PHARMS 0.01% A087159 001 Jun 16, 1982

BAUSCH AND LOMB 0.01% A040059 001 Dec 20, 1993

G AND W LABS INC 0.01% A207441 001 Sep 28, 2016

GLASSHOUSE PHARMS 0.01% A209596 001 Dec 26, 2017

MORTON GROVE 0.01% A088312 001 Jan 27, 1984

PHARMADERM 0.01% A088048 001 Dec 16, 1982

PHARMAFAIR 0.01% A088449 001 Feb 08, 1984

FLUONID

ALLERGAN HERBERT 0.01% A087158 001 Mar 17, 1983

FLUOTREX

SAVAGE LABS 0.01% A088171 001 Mar 09, 1983

FLUOCINONIDE

CREAM;TOPICAL

FLUOCINONIDE

PERRIGO NEW YORK 0.05% A071790 001 Jul 13, 1988

LIDEX

+ CNTY LINE PHARMS 0.05% N016908 002

SOLUTION;TOPICAL

FLUOCINONIDE

TARO 0.05% A072857 001 Aug 02, 1989

TEVA PHARMS 0.05% A072522 001 Sep 28, 1990

FLUORESCEIN SODIUM

INJECTABLE;INJECTION

FUNDUSCEIN-25

+ NOVARTIS 25% ** N017869 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUOROMETHOLONE

CREAM; TOPICAL

OXYLONE

PHARMACIA AND UPJOHN 0.025%

N011748 001

SUSPENSION/DROPS; OPHTHALMIC

FLUOR-OP

NOVARTIS 0.1%

A070185 001 Feb 27, 1986

FLUOROMETHOLONE ACETATE; TOBRAMYCIN

SUSPENSION/DROPS; OPHTHALMIC

TOBRASONE

ALCON 0.1%; 0.3%

N050628 001 Jul 21, 1989

FLUOROMETHOLONE; SULFACETAMIDE SODIUM

SUSPENSION/DROPS; OPHTHALMIC

FML-S

ALLERGAN 0.1%; 10%

N019525 001 Sep 29, 1989

FLUOROURACIL

INJECTABLE; INJECTION

ADRUCIL

PHARMACIA AND UPJOHN 50MG/ML

A081222 001 Jun 28, 1991

50MG/ML

N017959 001

TEVA PARENTERAL 50MG/ML

A040023 001 Oct 18, 1991

50MG/ML

A081225 001 Aug 28, 1991

FLUOROURACIL

ABIC 50MG/ML

A088929 001 Mar 04, 1986

ABRAXIS PHARM 50MG/ML

A089152 001 Mar 21, 1986

50MG/ML

A089428 001 Jan 12, 1987

50MG/ML

A089519 001 Mar 12, 1987

50MG/ML

A089508 001 Jan 26, 1988

BEDFORD 50MG/ML

A040772 001 Aug 11, 2008

EBEWE PHARMA 500MG/10ML (50MG/ML)

A040291 001 Mar 24, 1999

FRESENIUS KABI USA 50MG/ML

A040379 001 Nov 15, 2000

50MG/ML

A087791 001 Jan 18, 1983

MARCHAR 50MG/ML

A091299 001 May 02, 2011

SANDOZ 2.5GM/50ML (50MG/ML)

A091299 002 May 02, 2011

5GM/100ML (50MG/ML)

A088766 001 Dec 28, 1984

SMITH AND NEPHEW 50MG/ML

A088767 001 Dec 28, 1984

50MG/ML

A089434 001 Mar 26, 1987

50MG/ML

A087792 001 Oct 13, 1982

SPECTRUM PHARMS 50MG/ML

N012209 001

+ 500MG/10ML (50MG/ML) **

N012209 002 Jul 29, 2016

+ 2.5GM/50ML (50MG/ML)

SOLUTION; TOPICAL

FLUOROPLEX

ELORAC 1%

N016765 001

FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL

FLUOXETINE

SUN PHARM INDUSTRIES EQ 10MG BASE

A075787 001 Jan 29, 2002

EQ 20MG BASE

A075787 002 Jan 29, 2002

WATSON LABS EQ 10MG BASE

A075662 001 Jan 29, 2002

EQ 20MG BASE

A075662 002 Jan 29, 2002

FLUOXETINE HYDROCHLORIDE

ANI PHARMS INC EQ 10MG BASE

A076287 001 May 20, 2008

EQ 20MG BASE

A076287 002 May 20, 2008

BARR EQ 40MG BASE

A076251 001 May 18, 2005

CARLSBAD EQ 10MG BASE

A076022 001 Jan 30, 2002

EQ 20MG BASE

A076022 002 Jan 30, 2002

CR DOUBLE CRANE EQ 10MG BASE

A076165 001 Feb 01, 2002

EQ 20MG BASE

A076165 002 Feb 01, 2002

MYLAN EQ 10MG BASE

A075207 001 Jan 30, 2002

EQ 20MG BASE

A075207 002 Jan 30, 2002

EQ 40MG BASE

A075207 003 May 25, 2007

MYLAN PHARMS INC EQ 10MG BASE

A075577 001 Jan 29, 2002

EQ 20MG BASE

A075577 002 Jan 29, 2002

PAR PHARM EQ 10MG BASE

A076922 001 Dec 16, 2004

EQ 20MG BASE

A076922 002 Dec 16, 2004

SANDOZ EQ 10MG BASE

A075807 001 Jan 29, 2002

EQ 10MG BASE

A077469 001 Nov 17, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL

FLUOXETINE HYDROCHLORIDE

| | | |
|---------------|--------------|--------------|
| EQ 20MG BASE | A075807 002 | Jan 29, 2002 |
| EQ 20MG BASE | A077469 002 | Nov 17, 2008 |
| WOCKHARDT LTD | EQ 10MG BASE | A078143 001 |
| EQ 20MG BASE | A078143 002 | Jan 16, 2008 |
| EQ 40MG BASE | A078143 003 | Jan 16, 2008 |

PROZAC

ELI LILLY AND CO

EQ 60MG BASE N018936 004 Jun 15, 1999

SARAFEM

+ ELI LILLY AND CO

EQ 10MG BASE ** N018936 007 Jul 06, 2000

+

EQ 20MG BASE ** N018936 008 Jul 06, 2000

SOLUTION; ORAL

FLUOXETINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC

EQ 20MG BASE/5ML A075690 001 Jan 31, 2002

APOTEX INC

EQ 20MG BASE/5ML A075292 001 Feb 07, 2002

AUROBINDO PHARMA LTD

EQ 20MG BASE/5ML A079209 001 Mar 20, 2009

HI TECH PHARMA

EQ 20MG BASE/5ML A075525 001 Jun 27, 2002

LANNETT CO INC

EQ 20MG BASE/5ML A076458 001 May 14, 2004

PROZAC

+ LILLY

EQ 20MG BASE/5ML ** N020101 001 Apr 24, 1991

TABLET; ORAL

FLUOXETINE HYDROCHLORIDE

BARR

EQ 10MG BASE A075810 001 Feb 01, 2002

FOSUN PHARMA

EQ 10MG BASE A076024 001 Jan 29, 2002

IVAX SUB TEVA PHARMS

EQ 10MG BASE A075865 001 Feb 28, 2002

EQ 40MG BASE A075865 003 Aug 30, 2004

PROZAC

+ LILLY

EQ 10MG BASE ** N020974 001 Mar 09, 1999

+

EQ 20MG BASE ** N020974 002 Mar 09, 1999

FLUOXYMESTERONE

TABLET; ORAL

ANDROID-F

VALEANT PHARM INTL

10MG A087196 001

FLUOXYMESTERONE

VALEANT PHARM INTL

10MG A088221 001 May 05, 1983

WATSON LABS

2MG A088260 001 Dec 06, 1983

5MG A088265 001 Dec 06, 1983

10MG A088309 001 Dec 06, 1983

HALOTESTIN

PHARMACIA AND UPJOHN

2MG N010611 002

5MG N010611 006

10MG N010611 010

ORA-TESTRYL

BRISTOL MYERS SQUIBB

2MG N011359 001

5MG N011359 002

FLUPHENAZINE DECANOATE

INJECTABLE; INJECTION

FLUPHENAZINE DECANOATE

HOSPIRA 25MG/ML A074966 001 Apr 16, 1998

TEVA PARENTERAL 25MG/ML A074795 001 Sep 10, 1996

PROLIXIN DECANOATE

+ BRISTOL MYERS SQUIBB 25MG/ML ** N016727 001

FLUPHENAZINE ENANTHATE

INJECTABLE; INJECTION

PROLIXIN ENANTHATE

APOTHECON 25MG/ML ** N016110 001

FLUPHENAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

FLUPHENAZINE HYDROCHLORIDE

ANI PHARMS INC

5MG/ML A073058 001 Aug 30, 1991

PERMITIL

SCHERING 5MG/ML ** N016008 001

PROLIXIN

APOTHECON

5MG/ML A070533 001 Nov 07, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUPHENAZINE HYDROCHLORIDE

ELIXIR; ORAL

FLUPHENAZINE HYDROCHLORIDE

ANI PHARMS INC

2.5MG/5ML

A081310 001 Apr 29, 1993

PROLIXIN

+ APOTHECON

2.5MG/5ML **

N012145 003

INJECTABLE; INJECTION

PROLIXIN

APOTHECON

2.5MG/ML **

N011751 005

TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

WATSON LABS

1MG

A088555 001 Dec 18, 1987

2.5MG

A088544 001 Dec 18, 1987

5MG

A088527 001 Dec 18, 1987

10MG

A088550 001 Dec 18, 1987

PERMITIL

SCHERING

0.25MG

N012034 001

2.5MG

N012034 004

5MG

N012034 005

10MG

N012034 006

PROLIXIN

+ APOTHECON

1MG **

N011751 004

+

2.5MG **

N011751 001

+

5MG **

N011751 003

+

10MG **

N011751 002

TABLET, EXTENDED RELEASE; ORAL

PERMITIL

SCHERING

1MG

N012419 004

FLUPREDNISOLONE

TABLET; ORAL

ALPHADROL

PHARMACIA AND UPJOHN

1.5MG

N012259 002

FLURANDRENOLIDE

LOTION; TOPICAL

FLURANDRENOLIDE

ALPHARMA US PHARMS

0.05%

A087203 001 Apr 29, 1982

OINTMENT; TOPICAL

CORDRAN

+ AQUA PHARMS

0.025% **

N012806 004

FLURANDRENOLIDE; NEOMYCIN SULFATE

CREAM; TOPICAL

CORDRAN N

LILLY

0.05%;EQ 3.5MG BASE/GM

N050346 001

OINTMENT; TOPICAL

CORDRAN N

LILLY

0.05%;EQ 3.5MG BASE/GM

N050345 001

FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL

DALMANE

VALEANT PHARM INTL

15MG **

N016721 001

+

30MG **

N016721 002

FLURAZEPAM HYDROCHLORIDE

AUROLIFE PHARMA LLC

15MG

A071717 002 Jul 31, 1991

30MG

A071717 001 Jul 31, 1991

HALSEY

15MG

A071808 001 Jan 07, 1988

30MG

A071809 001 Jan 07, 1988

HIKMA INTL PHARMS

15MG

A071107 001 Dec 08, 1986

HIKMA PHARMS

30MG

A071108 001 Dec 08, 1986

PAR PHARM

15MG

A070444 001 Mar 20, 1986

30MG

A070445 001 Mar 20, 1986

PUREPAC PHARM

15MG

A071927 001 Sep 09, 1987

30MG

A071551 001 Sep 09, 1987

SUN PHARM INDUSTRIES

15MG

A070454 001 Aug 04, 1986

30MG

A070455 001 Aug 04, 1986

SUPERPHARM

15MG

A071659 001 Aug 04, 1988

30MG

A071660 001 Aug 04, 1988

USL PHARMA

15MG

A070562 001 Jul 09, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL

FLURAZEPAM HYDROCHLORIDE

| | | | | |
|-----------------|------|---------|-----|--------------|
| | 30MG | A070563 | 001 | Jul 09, 1987 |
| WARNER CHILCOTT | 15MG | A071767 | 001 | Dec 04, 1987 |
| | 30MG | A071768 | 001 | Dec 04, 1987 |
| WATSON LABS | 15MG | A071205 | 001 | Nov 25, 1986 |
| | 15MG | A072368 | 001 | Mar 30, 1989 |
| | 30MG | A071068 | 001 | Nov 25, 1986 |
| | 30MG | A072369 | 001 | Mar 30, 1989 |

FLURBIPROFEN

TABLET; ORAL

ANSAID

| | | | | |
|----------------------|-------|---------|-----|--------------|
| PHARMACIA AND UPJOHN | 50MG | N018766 | 002 | Oct 31, 1988 |
| | 100MG | N018766 | 003 | Oct 31, 1988 |

FLURBIPROFEN

| | | | | |
|----------------------|-------|---------|-----|--------------|
| AUROLIFE PHARMA LLC | 50MG | A074448 | 001 | Jul 28, 1995 |
| | 100MG | A074448 | 002 | Jul 28, 1995 |
| IVAX SUB TEVA PHARMS | 50MG | A074411 | 001 | May 31, 1995 |
| | 100MG | A074411 | 002 | May 31, 1995 |
| PLIVA | 50MG | A074647 | 001 | Apr 01, 1997 |
| | 100MG | A074647 | 002 | Apr 01, 1997 |
| TEVA | 50MG | A074405 | 002 | May 24, 1995 |
| | 100MG | A074405 | 001 | May 24, 1995 |
| THERAGEN | 100MG | A074560 | 002 | May 16, 1997 |

FLUTAMIDE

CAPSULE; ORAL

EULEXIN

+ SCHERING

125MG

N018554 001 Jan 27, 1989

FLUTAMIDE

MYLAN

125MG

A076224 001 May 09, 2003

YAOPHARMA CO LTD

125MG

A075818 001 Sep 18, 2001

FLUTEMETAMOL F-18

INJECTABLE; INTRAVENOUS

VIZAMYL

+ GE HEALTHCARE

40.5mCi/10ML (4.05mCi/ML)

N203137 001 Oct 25, 2013

FLUTICASON PROPIONATE

AEROSOL, METERED; INHALATION

FLOVENT

GLAXOSMITHKLINE 0.044MG/INH

N020548 001 Mar 27, 1996

0.11MG/INH

N020548 002 Mar 27, 1996

0.22MG/INH

N020548 003 Mar 27, 1996

CREAM; TOPICAL

CUTIVATE

+ FOUGERA PHARMS

0.05% **

N019958 001 Dec 18, 1990

FLUTICASON PROPIONATE

NESHER PHARMS

0.05%

A076865 001 Sep 10, 2004

OINTMENT; TOPICAL

CUTIVATE

+ FOUGERA PHARMS

0.005%

N019957 001 Dec 14, 1990

FLUTICASON PROPIONATE

FOUGERA PHARMS

0.005%

A076300 001 May 14, 2004

TARO PHARM INDS

0.005%

A077145 001 Jun 14, 2005

POWDER; INHALATION

FLOVENT

GLAXOSMITHKLINE 0.044MG/INH

N020549 001 Nov 07, 1997

0.088MG/INH

N020549 002 Nov 07, 1997

0.22MG/INH

N020549 003 Nov 07, 1997

SPRAY, METERED; NASAL

FLONASE

+ GLAXOSMITHKLINE

0.05MG/SPRAY **

N020121 001 Oct 19, 1994

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FLUVASTATIN SODIUM

CAPSULE;ORAL

LESCOL

| | | | | | |
|---|----------|-----------------|---------|-----|--------------|
| + | NOVARTIS | EQ 20MG BASE ** | N020261 | 001 | Dec 31, 1993 |
| + | | EQ 40MG BASE ** | N020261 | 002 | Dec 31, 1993 |

FLUVOXAMINE MALEATE

CAPSULE, EXTENDED RELEASE;ORAL

LUVOX CR

| | | | | | |
|---|-------------|----------|---------|-----|--------------|
| + | JAZZ PHARMS | 100MG ** | N022033 | 001 | Feb 28, 2008 |
| + | | 150MG ** | N022033 | 002 | Feb 28, 2008 |

TABLET;ORAL

FLUVOXAMINE MALEATE

| | | | | | |
|----------------------|--------|----------|---------|--------------|--------------|
| ACTAVIS ELIZABETH | 25MG | A075901 | 001 | Dec 28, 2000 | |
| | 50MG | A075901 | 002 | Dec 28, 2000 | |
| | 100MG | A075901 | 003 | Dec 28, 2000 | |
| ANI PHARMS INC | 25MG | A075898 | 001 | Mar 12, 2001 | |
| | 50MG | A075898 | 002 | Mar 12, 2001 | |
| | 100MG | A075898 | 003 | Mar 12, 2001 | |
| ECI PHARMS LLC | 25MG | A075900 | 001 | Feb 23, 2006 | |
| | 50MG | A075900 | 002 | Feb 23, 2006 | |
| | 100MG | A075900 | 003 | Feb 23, 2006 | |
| MYLAN | 50MG | A075950 | 001 | Oct 15, 2001 | |
| | 100MG | A075950 | 002 | Oct 15, 2001 | |
| SUN PHARM INDUSTRIES | 25MG | A076125 | 001 | Apr 29, 2002 | |
| | 50MG | A076125 | 002 | Apr 29, 2002 | |
| | 100MG | A076125 | 003 | Apr 29, 2002 | |
| SYNTHON PHARMS | 25MG | A075899 | 001 | Jan 17, 2001 | |
| | 50MG | A075899 | 002 | Jan 17, 2001 | |
| | 100MG | A075899 | 003 | Jan 17, 2001 | |
| UPSHER SMITH LABS | 25MG | A075887 | 001 | Jan 05, 2001 | |
| | 50MG | A075887 | 002 | Jan 05, 2001 | |
| | 100MG | A075887 | 003 | Jan 05, 2001 | |
| WATSON LABS | 25MG | A075894 | 001 | Apr 18, 2001 | |
| | 50MG | A075894 | 002 | Apr 18, 2001 | |
| | 100MG | A075894 | 003 | Apr 18, 2001 | |
| LUVOX | | | | | |
| + | SOLVAY | 25MG ** | N020243 | 001 | Dec 05, 1994 |
| + | | 50MG ** | N020243 | 002 | Dec 05, 1994 |
| + | | 100MG ** | N020243 | 003 | Dec 05, 1994 |
| + | | 150MG ** | N020243 | 004 | Dec 05, 1994 |

FOLIC ACID

INJECTABLE;INJECTION

FOLIC ACID

| | | | | |
|-----------|--------|---------|-----|--------------|
| BEN VENUE | 5MG/ML | A081066 | 001 | Dec 29, 1993 |
|-----------|--------|---------|-----|--------------|

FOLVITE

| | | | | |
|------------------|--------|---------|-----|--|
| WYETH PHARMS INC | 5MG/ML | N005897 | 008 | |
|------------------|--------|---------|-----|--|

TABLET;ORAL

FOLIC ACID

| | | | | |
|----------------------|-----|---------|-----|--------------|
| BARR | 1MG | A089177 | 001 | Jan 08, 1986 |
| CONTRACT PHARMACAL | 1MG | A085061 | 001 | |
| EVERYLIFE | 1MG | A080755 | 001 | |
| HALSEY | 1MG | A083598 | 001 | |
| IMPAX LABS | 1MG | A080686 | 001 | |
| IVAX SUB TEVA PHARMS | 1MG | A083000 | 001 | |
| JUBILANT CADISTA | 1MG | A040514 | 001 | Jun 14, 2005 |
| LANNETT | 1MG | A080816 | 001 | |
| LILLY | 1MG | N006135 | 003 | |
| MK LABS | 1MG | A083526 | 001 | |
| NEXGEN PHARMA INC | 1MG | A084915 | 001 | |
| PHARMERAL | 1MG | A084158 | 001 | |
| PIONEER PHARMS | 1MG | A088949 | 001 | Sep 13, 1985 |
| PUREPAC PHARM | 1MG | A080784 | 001 | |
| SANDOZ | 1MG | A084472 | 001 | |
| SUN PHARM INDUSTRIES | 1MG | A040582 | 001 | Jul 18, 2005 |
| TABLICAPS | 1MG | A083133 | 002 | |
| UDL | 1MG | A088199 | 001 | Mar 29, 1983 |
| USL PHARMA | 1MG | A087828 | 001 | May 13, 1982 |
| VALEANT PHARM INTL | 1MG | A080903 | 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FOLIC ACID

TABLET; ORAL

FOLIC ACID

| | | | |
|----------------------|-----|-------------|--------------|
| VANGARD | 1MG | A088730 001 | Mar 23, 1984 |
| VINTAGE PHARMS | 1MG | A086296 001 | |
| WATSON LABS | 1MG | A083141 001 | |
| | 1MG | A085141 002 | |
| WHITEWORTH TOWN PLSN | 1MG | A080691 002 | |
| FOLICET | | | |
| MISSION PHARMA | 1MG | A087438 001 | |
| FOLVITE | | | |
| WYETH PHARMS INC | 1MG | N005897 004 | |

FOLLITROPIN ALFA/BETA

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

FOLLISTIM

| | | | |
|-----------------|-------------|-------------|--------------|
| ORGANON USA INC | 75 IU/VIAL | N020582 001 | Sep 29, 1997 |
| | 150 IU/VIAL | N020582 002 | Sep 29, 1997 |

INJECTABLE; SUBCUTANEOUS

FOLLISTIM AQ

| | | | |
|-----------------|---------------|-------------|--------------|
| ORGANON USA INC | 75 IU/0.5ML | N021273 001 | Aug 26, 2005 |
| | 150 IU/0.18ML | N021211 003 | Feb 11, 2004 |
| | 150 IU/0.5ML | N021273 002 | Aug 26, 2005 |

GONAL-F

| | | | |
|------------|--------------|-------------|--------------|
| EMD SERONO | 37.5 IU/VIAL | N020378 003 | May 25, 2000 |
| | 37.5 IU/VIAL | N021765 001 | Mar 25, 2004 |
| | 75 IU/VIAL | N020378 001 | Sep 29, 1997 |
| | 150 IU/VIAL | N020378 002 | Sep 29, 1997 |
| | 150 IU/VIAL | N021765 003 | Mar 25, 2004 |

FOMEPIZOLE

INJECTABLE; INJECTION

FOMEPIZOLE

| | | | |
|---------------------|----------------------|-------------|--------------|
| MYLAN INSTITUTIONAL | 1.5GM/1.5ML (1GM/ML) | A079033 001 | Apr 07, 2009 |
|---------------------|----------------------|-------------|--------------|

FOMIVIRSEN SODIUM

INJECTABLE; INJECTION

VITRAVENE PRESERVATIVE FREE

| | | | |
|----------|----------|-------------|--------------|
| NOVARTIS | 6.6MG/ML | N020961 001 | Aug 26, 1998 |
|----------|----------|-------------|--------------|

FORMOTEROL FUMARATE

POWDER; INHALATION

FORADIL

+ NOVARTIS

0.012MG/INH

N020831 001 Feb 16, 2001

FORADIL CERTIHALER

NOVARTIS

0.0085MG/INH

N021592 001 Dec 15, 2006

FOSAPREPITANT DIMEGLUMINE

POWDER; INTRAVENOUS

EMEND

+ MERCK AND CO INC

EQ 115MG BASE/VIAL **

N022023 001 Jan 25, 2008

FOSCARNET SODIUM

INJECTABLE; INJECTION

FOSCARNET SODIUM

HOSPIRA

2.4GM/100ML

A077174 001 May 31, 2005

FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

ACTAVIS LABS FL INC

10MG

A076620 001 Oct 15, 2004

20MG

A076620 002 Oct 15, 2004

40MG

A076620 003 Oct 15, 2004

RANBAXY LABS LTD

10MG

A076580 001 Apr 23, 2004

20MG

A076580 002 Apr 23, 2004

40MG

A076580 003 Apr 23, 2004

UPSHER SMITH LABS

10MG

A076188 001 Oct 08, 2004

20MG

A076188 002 Oct 08, 2004

40MG

A076188 003 Oct 08, 2004

WATSON LABS

10MG

A076987 001 Dec 23, 2004

10MG

A077531 001 Aug 31, 2006

20MG

A076987 002 Dec 23, 2004

20MG

A077531 002 Aug 31, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

40MG

A076987 003 Dec 23, 2004

40MG

A077531 003 Aug 31, 2006

MONOPRIL

+ BRISTOL MYERS SQUIBB 10MG **

N019915 002 May 16, 1991

+ 20MG **

N019915 003 May 16, 1991

+ 40MG **

N019915 004 Mar 28, 1995

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE

ACTAVIS LABS FL INC 10MG;12.5MG

A076608 001 Dec 03, 2004

20MG;12.5MG

A076608 002 Dec 03, 2004

MYLAN 10MG;12.5MG

A077705 001 Aug 14, 2006

20MG;12.5MG

A077705 002 Aug 14, 2006

SUN PHARM INDS LTD 10MG;12.5MG

A076739 001 Dec 17, 2004

20MG;12.5MG

A076739 002 Dec 17, 2004

TEVA 10MG;12.5MG

A076945 001 Jul 05, 2006

20MG;12.5MG

A076945 002 Jul 05, 2006

WATSON LABS 10MG;12.5MG

A077144 001 Aug 16, 2005

20MG;12.5MG

A077144 002 Aug 16, 2005

MONOPRIL-HCT

+ BRISTOL MYERS SQUIBB 10MG;12.5MG **

N020286 002 Nov 30, 1994

+ 20MG;12.5MG **

N020286 001 Nov 30, 1994

FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

FOSPHENYTOIN SODIUM

APOTEX INC EQ 50MG PHENYTOIN NA/ML

A078126 001 Aug 06, 2007

HOSPIRA EQ 50MG PHENYTOIN NA/ML

A078158 001 Aug 06, 2007

TEVA PHARMS USA EQ 50MG PHENYTOIN NA/ML

A076886 001 Aug 06, 2007

FOSPROPOFOL DISODIUM

SOLUTION; INTRAVENOUS

LUSEDRA

EISAI INC 1050MG/30ML (35MG/ML)

N022244 001 Dec 12, 2008

FURAZOLIDONE

SUSPENSION; ORAL

FUROXONE

SHIRE 50MG/15ML

N011323 002

TABLET; ORAL

FUROXONE

SHIRE 100MG

N011270 002

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

ABRAXIS PHARM 10MG/ML

N018507 001 Jul 30, 1982

10MG/ML

N019036 001 Aug 13, 1984

ACCORD HLTHCARE 10MG/ML

A070017 001 Dec 15, 1986

ASTRAZENECA 10MG/ML

A070014 001 Sep 09, 1985

HOSPIRA 10MG/ML

A070578 001 Jul 08, 1987

10MG/ML

A072080 001 Aug 13, 1991

10MG/ML

A074337 001 Oct 31, 1994

IGI LABS INC 10MG/ML

A070095 001 Sep 09, 1985

10MG/ML

A070096 001 Sep 09, 1985

INTL MEDICATION 10MG/ML

N018025 001

+ LUITPOLD 10MG/ML **

N018579 001 Nov 30, 1983

MARSAM PHARMS LLC 10MG/ML

A074017 001 Jun 30, 1994

SMITH AND NEPHEW 10MG/ML

A070023 001 Feb 05, 1986

10MG/ML

A070078 001 Feb 05, 1986

WARNER CHILCOTT 10MG/ML

N018420 001 Feb 26, 1982

WATSON LABS 10MG/ML

A070019 001 Sep 22, 1986

10MG/ML

A070604 001 Jan 02, 1987

WEST-WARD PHARMS INT 10MG/ML

A071439 001 Sep 14, 1990

10MG/ML

N018267 001

WYETH AYERST 10MG/ML

N018670 001 Jul 20, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

FUROSEMIDE

INJECTABLE; INJECTION

LASIX

+ SANOFI AVENTIS US 10MG/ML ** N016363 001

SOLUTION; ORAL

LASIX

SANOFI AVENTIS US 10MG/ML N017688 001

TABLET; ORAL

FUROSEMIDE

| | | | |
|----------------------|------|-------------|--------------|
| DAVA PHARMS INC | 20MG | N018415 001 | Jul 27, 1982 |
| | 40MG | N018415 002 | Jul 27, 1982 |
| | 80MG | N018415 003 | Nov 26, 1984 |
| INTL MEDICATION | 20MG | N018753 001 | Feb 28, 1984 |
| | 40MG | N018753 002 | Feb 28, 1984 |
| KALAPHARM | 20MG | N018868 001 | Jun 28, 1983 |
| | 40MG | N018868 002 | Jun 28, 1983 |
| SANDOZ | 40MG | N018750 002 | Jul 30, 1984 |
| SUN PHARM INDS INC | 20MG | A091258 001 | Apr 01, 2014 |
| | 40MG | A091258 002 | Apr 01, 2014 |
| | 40MG | N018790 001 | Nov 29, 1983 |
| | 80MG | A091258 003 | Apr 01, 2014 |
| SUN PHARM INDUSTRIES | 20MG | A070043 001 | Sep 26, 1985 |
| | 80MG | A070100 001 | Jan 26, 1988 |
| SUPERPHARM | 20MG | N018370 002 | Jun 26, 1984 |
| | 40MG | N018370 001 | Feb 10, 1983 |
| WARNER CHILCOTT | 20MG | N018419 001 | Jan 31, 1983 |
| | 40MG | N018419 002 | Jan 31, 1983 |
| | 80MG | N018419 003 | Nov 13, 1984 |
| WATSON LABS | 20MG | A070412 001 | Feb 26, 1986 |
| | 20MG | A071379 001 | Jan 02, 1987 |
| | 20MG | N018369 001 | May 14, 1982 |
| | 40MG | A070413 001 | Feb 26, 1986 |
| | 40MG | A070450 001 | Nov 22, 1985 |
| | 40MG | N018369 002 | May 14, 1982 |
| | 80MG | A071594 001 | Feb 09, 1988 |
| WATSON LABS TEVA | 20MG | A070449 001 | Nov 22, 1985 |
| | 80MG | A070528 001 | Jan 07, 1986 |

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

| | | | |
|----------------------|-------|-------------|--------------|
| CSPC OUYI PHARM CO | 100MG | A075477 001 | Mar 23, 2005 |
| | 300MG | A075477 002 | Mar 23, 2005 |
| | 400MG | A075477 003 | Mar 23, 2005 |
| HIKMA | 100MG | A078150 001 | Sep 25, 2007 |
| | 300MG | A078150 002 | Sep 25, 2007 |
| | 400MG | A078150 003 | Sep 25, 2007 |
| SANDOZ | 100MG | A075428 001 | Jan 24, 2006 |
| | 100MG | A075539 001 | Apr 06, 2005 |
| | 300MG | A075428 002 | Jan 24, 2006 |
| | 300MG | A075539 002 | Apr 06, 2005 |
| | 400MG | A075428 003 | Jan 24, 2006 |
| | 400MG | A075539 003 | Apr 06, 2005 |
| SUN PHARM INDS LTD | 100MG | A076606 001 | Oct 07, 2005 |
| | 300MG | A076606 002 | Oct 07, 2005 |
| | 400MG | A076606 003 | Oct 07, 2005 |
| SUN PHARM INDUSTRIES | 100MG | A076537 001 | Jun 30, 2005 |
| | 300MG | A076537 002 | Jun 30, 2005 |
| | 400MG | A076537 003 | Jun 30, 2005 |
| WATSON LABS | 100MG | A075485 003 | May 11, 2007 |
| | 300MG | A075485 002 | May 11, 2007 |
| | 400MG | A075485 001 | May 11, 2007 |

TABLET; ORAL

GABAPENTIN

| | | | |
|--------------|-------|-------------|--------------|
| HIKMA PHARMS | 600MG | A078782 001 | Jul 21, 2011 |
| | 800MG | A078782 002 | Jul 21, 2011 |
| RANBAXY | 600MG | A076605 001 | Sep 14, 2005 |
| | 800MG | A076605 002 | Sep 14, 2005 |
| SANDOZ | 600MG | A076120 001 | Jan 27, 2006 |
| | 600MG | A076877 001 | Jul 06, 2006 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GABAPENTINTABLET; ORAL
GABAPENTIN

| | | | |
|------|-------|-------------|--------------|
| | 800MG | A076120 002 | Jan 27, 2006 |
| | 800MG | A076877 002 | Jul 06, 2006 |
| TEVA | 600MG | A075827 001 | Dec 15, 2004 |
| | 800MG | A075827 002 | Dec 15, 2004 |

GADODIAMIDEINJECTABLE; INJECTION
OMNISCAN

| | | | |
|---------------|-------------------------|-------------|--------------|
| GE HEALTHCARE | 14.35GM/50ML (287MG/ML) | N022066 001 | Sep 05, 2007 |
|---------------|-------------------------|-------------|--------------|

GADOFOSVESET TRISODIUMSOLUTION; INTRAVENOUS
ABLAVAR

| | | | |
|----------------|------------------------|-------------|--------------|
| LANTHEUS MEDCL | 2440MG/10ML (244MG/ML) | N021711 001 | Dec 22, 2008 |
| | 3660MG/15ML (244MG/ML) | N021711 002 | Dec 22, 2008 |

GADOVERSETAMIDE

INJECTABLE; INJECTION

OPTIMARK

| | | | | |
|---|------------------|----------------------------|-------------|--------------|
| + | LIEBEL-FLARSHEIM | 1654.5MG/5ML (330.9MG/ML) | N020937 001 | Dec 08, 1999 |
| + | | 3309MG/10ML (330.9MG/ML) | N020937 002 | Dec 08, 1999 |
| + | | 4963.5MG/15ML (330.9MG/ML) | N020937 003 | Dec 08, 1999 |
| + | | 6618MG/20ML (330.9MG/ML) | N020937 004 | Dec 08, 1999 |
| + | | 16.545GM/50ML (330.9MG/ML) | N020975 001 | Dec 08, 1999 |

OPTIMARK IN PLASTIC CONTAINER

| | | | | |
|---|------------------|----------------------------|-------------|--------------|
| + | LIEBEL-FLARSHEIM | 3309MG/10ML (330.9MG/ML) | N020976 002 | Dec 08, 1999 |
| + | | 4963.5MG/15ML (330.9MG/ML) | N020976 003 | Dec 08, 1999 |
| + | | 6618MG/20ML (330.9MG/ML) | N020976 004 | Dec 08, 1999 |
| + | | 9927MG/30ML (330.9MG/ML) | N020976 001 | Dec 08, 1999 |

GALANTAMINE HYDROBROMIDE

CAPSULE, EXTENDED RELEASE; ORAL

GALANTAMINE HYDROBROMIDE

| | | | |
|------------|--------------|-------------|--------------|
| IMPAX LABS | EQ 8MG BASE | A078484 001 | May 27, 2009 |
| | EQ 16MG BASE | A078484 002 | May 27, 2009 |
| | EQ 24MG BASE | A078484 003 | May 27, 2009 |
| MYLAN | EQ 8MG BASE | A090900 001 | Jan 24, 2011 |
| | EQ 16MG BASE | A090900 002 | Jan 24, 2011 |
| | EQ 24MG BASE | A090900 003 | Jan 24, 2011 |

SOLUTION; ORAL

RAZADYNE

| | | | |
|----------------|-----------|-------------|--------------|
| JANSSEN PHARMS | 4MG/ML ** | N021224 001 | Jun 22, 2001 |
|----------------|-----------|-------------|--------------|

TABLET; ORAL

GALANTAMINE HYDROBROMIDE

| | | | |
|-------------------|--------------|-------------|--------------|
| ACTAVIS ELIZABETH | EQ 4MG BASE | A077585 001 | Sep 15, 2009 |
| | EQ 8MG BASE | A077585 002 | Sep 15, 2009 |
| | EQ 12MG BASE | A077585 003 | Sep 15, 2009 |
| MYLAN | EQ 4MG BASE | A077603 001 | Aug 28, 2008 |
| | EQ 8MG BASE | A077603 002 | Aug 28, 2008 |
| | EQ 12MG BASE | A077603 003 | Aug 28, 2008 |

GALLAMINE TRIETHIODIDE

INJECTABLE; INJECTION

FLAXEDIL

| | | |
|----------------|----------|-------------|
| DAVIS AND GECK | 20MG/ML | N007842 001 |
| | 100MG/ML | N007842 002 |

GALLIUM CITRATE GA-67

INJECTABLE; INJECTION

GALLIUM CITRATE GA 67

| | | |
|---------------|---------|-------------|
| GE HEALTHCARE | 1mCi/ML | N017700 001 |
|---------------|---------|-------------|

NEOSCAN

| | | |
|---------------|---------|-------------|
| GE HEALTHCARE | 2mCi/ML | N017655 001 |
|---------------|---------|-------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GALLIUM NITRATE

INJECTABLE; INJECTION

GANITE

CHAPTER 7 TRUSTEE

25MG/ML **

N019961 002 Jan 17, 1991

GANCICLOVIR

CAPSULE; ORAL

CYTOVENE

+ ROCHE PALO

250MG **

N020460 001 Dec 22, 1994

+

500MG **

N020460 002 Dec 12, 1997

GANCICLOVIR

RANBAXY LABS LTD

250MG

A076457 001 Jun 27, 2003

500MG

A076457 002 Jun 27, 2003

IMPLANT; IMPLANTATION

VITRASERT

BAUSCH AND LOMB

4.5MG

N020569 001 Mar 04, 1996

GANCICLOVIR SODIUM

INJECTABLE; INJECTION

GANCICLOVIR SODIUM

WEST-WARD PHARMS INT EQ 500MG BASE/VIAL

A076222 001 Jul 16, 2003

GATIFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

GATIFLOXACIN

APOTEX INC

0.3%

A079084 001 Aug 19, 2011

GEFITINIB

TABLET; ORAL

IRESSA

ASTRAZENECA

250MG

N021399 001 May 05, 2003

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

HAMELN RDS GMBH

EQ 200MG BASE/VIAL

A090663 001 Sep 10, 2012

EQ 1GM BASE/VIAL

A090663 002 Sep 10, 2012

SAGENT PHARMS

EQ 200MG BASE/VIAL

A091597 001 May 07, 2013

EQ 1GM BASE/VIAL

A091597 002 May 07, 2013

GEMFIBROZIL

CAPSULE; ORAL

GEMFIBROZIL

MYLAN

300MG

A073466 001 Jan 25, 1993

PUREPAC PHARM

300MG

A072929 001 Jan 29, 1993

LOPID

PFIZER PHARMS

200MG

N018422 001

300MG

N018422 002

TABLET; ORAL

GEMFIBROZIL

MYLAN

600MG

A074452 001 Feb 16, 1995

PUREPAC PHARM

600MG

A074360 001 Aug 31, 1994

WATSON LABS

600MG

A074156 001 Oct 24, 1994

600MG

A074442 001 Apr 28, 1995

YAOPHARMA CO LTD

600MG

A074615 001 Sep 29, 1995

GEMTUZUMAB OZOGAMICIN

INJECTABLE; INJECTION

MYLOTARG

WYETH PHARMS INC

5MG/VIAL

N021174 001 May 17, 2000

GENTAMICIN SULFATE

CREAM; TOPICAL

GARAMYCIN

SCHERING

EQ 0.1% BASE **

A060462 001

GENTAFAIR

PHARMAFAIR

EQ 0.1% BASE

A062458 001 Sep 01, 1983

GENTAMICIN SULFATE

ALPHARMA US PHARMS

EQ 0.1% BASE

A062471 001 Sep 27, 1983

FOUGERA PHARMS INC

EQ 0.1% BASE

A062531 001 Jul 05, 1984

PHARMADERM

EQ 1MG BASE/GM

A062530 001 Jul 05, 1984

TARO

EQ 0.1% BASE

A062427 001 May 26, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GENTAMICIN SULFATE

INJECTABLE; INJECTION

APOGEN

| | | | | |
|-------------|-----------------|---------|-----|--|
| KING PHARMS | EQ 10MG BASE/ML | A062289 | 001 | |
| | EQ 40MG BASE/ML | A062289 | 002 | |

BRISTAGEN

| | | | | |
|---------|-----------------|---------|-----|--|
| BRISTOL | EQ 40MG BASE/ML | A062288 | 001 | |
|---------|-----------------|---------|-----|--|

GARAMYCIN

| | | | | |
|----------|--------------------|---------|-----|--|
| SCHERING | EQ 1MG BASE/ML ** | A061716 | 002 | |
| | EQ 10MG BASE/ML ** | A061739 | 001 | |
| | EQ 40MG BASE/ML ** | A061716 | 001 | |

GENTAFAIR

| | | | | |
|------------|-----------------|---------|-----|--------------|
| PHARMAFAIR | EQ 40MG BASE/ML | A062493 | 001 | Aug 28, 1985 |
|------------|-----------------|---------|-----|--------------|

GENTAMICIN

| | | | | |
|-----------------|---------------------|---------|-----|--------------|
| INTL MEDICATION | EQ 1MG BASE/ML | A062325 | 003 | Jun 23, 1982 |
| | EQ 40MG BASE/ML | A062325 | 001 | |
| | EQ 100MG BASE/100ML | A062325 | 004 | Jun 23, 1982 |

GENTAMICIN SULFATE

ABBOTT

| | | | | |
|--|---------------------|---------|-----|--------------|
| | EQ 1.2MG BASE/ML | A062413 | 001 | Aug 11, 1983 |
| | EQ 1.4MG BASE/ML | A062413 | 002 | Aug 11, 1983 |
| | EQ 1.6MG BASE/ML | A062413 | 003 | Aug 11, 1983 |
| | EQ 1.8MG BASE/ML | A062413 | 004 | Aug 11, 1983 |
| | EQ 2MG BASE/ML | A062413 | 005 | Aug 11, 1983 |
| | EQ 60MG BASE/100ML | A062413 | 006 | Aug 11, 1983 |
| | EQ 70MG BASE/100ML | A062413 | 007 | Aug 11, 1983 |
| | EQ 80MG BASE/100ML | A062413 | 008 | Aug 11, 1983 |
| | EQ 90MG BASE/100ML | A062413 | 009 | Aug 11, 1983 |
| | EQ 100MG BASE/100ML | A062413 | 010 | Aug 11, 1983 |

FRESENIUS KABI USA

| | | | | |
|--|-----------------|---------|-----|--------------|
| | EQ 10MG BASE/ML | A062356 | 001 | Mar 04, 1982 |
| | EQ 40MG BASE/ML | A062356 | 002 | Mar 04, 1982 |

KALAPHARM

| | | | | |
|--|-----------------|---------|-----|--------------|
| | EQ 40MG BASE/ML | A062354 | 001 | Apr 05, 1982 |
|--|-----------------|---------|-----|--------------|

PHARM SPEC

| | | | | |
|--|-----------------|---------|-----|--------------|
| | EQ 40MG BASE/ML | A062340 | 001 | Mar 28, 1983 |
|--|-----------------|---------|-----|--------------|

SOLOPAK

| | | | | |
|--|-----------------|---------|-----|--------------|
| | EQ 10MG BASE/ML | A062507 | 001 | Jun 06, 1985 |
|--|-----------------|---------|-----|--------------|

| | | | | |
|--|-----------------|---------|-----|--------------|
| | EQ 40MG BASE/ML | A062507 | 002 | Jun 06, 1985 |
|--|-----------------|---------|-----|--------------|

TEVA PARENTERAL

| | | | | |
|--|-----------------|---------|-----|--------------|
| | EQ 10MG BASE/ML | A063149 | 001 | Nov 21, 1991 |
|--|-----------------|---------|-----|--------------|

| | | | | |
|--|-----------------|---------|-----|--------------|
| | EQ 40MG BASE/ML | A063106 | 002 | Nov 21, 1991 |
|--|-----------------|---------|-----|--------------|

WATSON LABS

| | | | | |
|--|-----------------|---------|-----|--|
| | EQ 10MG BASE/ML | A062318 | 002 | |
|--|-----------------|---------|-----|--|

| | | | | |
|--|-----------------|---------|-----|--|
| | EQ 40MG BASE/ML | A062318 | 001 | |
|--|-----------------|---------|-----|--|

| | | | | |
|----------------------|-----------------|---------|-----|--|
| WEST-WARD PHARMS INT | EQ 10MG BASE/ML | A062251 | 002 | |
|----------------------|-----------------|---------|-----|--|

| | | | | |
|--|-----------------|---------|-----|--|
| | EQ 40MG BASE/ML | A062251 | 001 | |
|--|-----------------|---------|-----|--|

WYETH AYERST

| | | | | |
|--|-----------------|---------|-----|--|
| | EQ 10MG BASE/ML | A062264 | 001 | |
|--|-----------------|---------|-----|--|

| | | | | |
|--|-----------------|---------|-----|--|
| | EQ 40MG BASE/ML | A062264 | 002 | |
|--|-----------------|---------|-----|--|

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN

| | | | | |
|--|------------------|---------|-----|--------------|
| | EQ 0.8MG BASE/ML | A062814 | 001 | Aug 28, 1987 |
|--|------------------|---------|-----|--------------|

| | | | | |
|--|------------------|---------|-----|--------------|
| | EQ 1.2MG BASE/ML | A062814 | 002 | Aug 28, 1987 |
|--|------------------|---------|-----|--------------|

| | | | | |
|--|------------------|---------|-----|--------------|
| | EQ 1.4MG BASE/ML | A062814 | 003 | Aug 28, 1987 |
|--|------------------|---------|-----|--------------|

| | | | | |
|--|------------------|---------|-----|--------------|
| | EQ 1.6MG BASE/ML | A062814 | 004 | Aug 28, 1987 |
|--|------------------|---------|-----|--------------|

| | | | | |
|--|------------------|---------|-----|--------------|
| | EQ 1.8MG BASE/ML | A062814 | 005 | Aug 28, 1987 |
|--|------------------|---------|-----|--------------|

| | | | | |
|--|----------------|---------|-----|--------------|
| | EQ 2MG BASE/ML | A062814 | 006 | Aug 28, 1987 |
|--|----------------|---------|-----|--------------|

| | | | | |
|--|------------------|---------|-----|--------------|
| | EQ 2.4MG BASE/ML | A062814 | 007 | Aug 28, 1987 |
|--|------------------|---------|-----|--------------|

| | | | | |
|--|--------------------|---------|-----|--------------|
| | EQ 40MG BASE/100ML | A062814 | 008 | Aug 28, 1987 |
|--|--------------------|---------|-----|--------------|

| | | | | |
|--|--------------------|---------|-----|--------------|
| | EQ 60MG BASE/100ML | A062814 | 009 | Aug 28, 1987 |
|--|--------------------|---------|-----|--------------|

| | | | | |
|--|--------------------|---------|-----|--------------|
| | EQ 70MG BASE/100ML | A062814 | 010 | Aug 28, 1987 |
|--|--------------------|---------|-----|--------------|

| | | | | |
|--|--------------------|---------|-----|--------------|
| | EQ 80MG BASE/100ML | A062814 | 011 | Aug 28, 1987 |
|--|--------------------|---------|-----|--------------|

| | | | | |
|--|--------------------|---------|-----|--------------|
| | EQ 90MG BASE/100ML | A062814 | 012 | Aug 28, 1987 |
|--|--------------------|---------|-----|--------------|

| | | | | |
|--|---------------------|---------|-----|--------------|
| | EQ 100MG BASE/100ML | A062814 | 013 | Aug 28, 1987 |
|--|---------------------|---------|-----|--------------|

| | | | | |
|--|---------------------|---------|-----|--------------|
| | EQ 120MG BASE/100ML | A062814 | 014 | Aug 28, 1987 |
|--|---------------------|---------|-----|--------------|

| | | | | |
|-----------------|------------------|---------|-----|--------------|
| BAXTER HLTHCARE | EQ 0.8MG BASE/ML | A062373 | 001 | Sep 07, 1982 |
|-----------------|------------------|---------|-----|--------------|

| | | | | |
|--|------------------|---------|-----|--------------|
| | EQ 2.4MG BASE/ML | A062373 | 010 | Sep 07, 1982 |
|--|------------------|---------|-----|--------------|

| | | | | |
|--|--------------------|---------|-----|--------------|
| | EQ 40MG BASE/100ML | A062373 | 003 | Sep 07, 1982 |
|--|--------------------|---------|-----|--------------|

| | | | | |
|--|--------------------|---------|-----|--------------|
| | EQ 60MG BASE/100ML | A062373 | 004 | Sep 07, 1982 |
|--|--------------------|---------|-----|--------------|

HOSPIRA

| | | | | |
|--|------------------|---------|-----|--------------|
| | EQ 1.2MG BASE/ML | A062588 | 001 | Jan 06, 1986 |
|--|------------------|---------|-----|--------------|

| | | | | |
|--|------------------|---------|-----|--------------|
| | EQ 1.4MG BASE/ML | A062414 | 002 | Aug 15, 1983 |
|--|------------------|---------|-----|--------------|

| | | | | |
|--|------------------|---------|-----|--------------|
| | EQ 1.4MG BASE/ML | A062588 | 002 | Jan 06, 1986 |
|--|------------------|---------|-----|--------------|

| | | | | |
|--|------------------|---------|-----|--------------|
| | EQ 1.6MG BASE/ML | A062588 | 003 | Jan 06, 1986 |
|--|------------------|---------|-----|--------------|

| | | | | |
|--|------------------|---------|-----|--------------|
| | EQ 1.8MG BASE/ML | A062414 | 004 | Aug 15, 1983 |
|--|------------------|---------|-----|--------------|

| | | | | |
|--|------------------|---------|-----|--------------|
| | EQ 1.8MG BASE/ML | A062588 | 004 | Jan 06, 1986 |
|--|------------------|---------|-----|--------------|

| | | | | |
|--|----------------|---------|-----|--------------|
| | EQ 2MG BASE/ML | A062414 | 005 | Aug 15, 1983 |
|--|----------------|---------|-----|--------------|

| | | | | |
|--|----------------|---------|-----|--------------|
| | EQ 2MG BASE/ML | A062588 | 005 | Jan 06, 1986 |
|--|----------------|---------|-----|--------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GENTAMICIN SULFATE

INJECTABLE; INJECTION

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

| | | | |
|---------------------|---------|-----|--------------|
| EQ 60MG BASE/100ML | A062414 | 006 | Aug 15, 1983 |
| EQ 60MG BASE/100ML | A062588 | 006 | Jan 06, 1986 |
| EQ 70MG BASE/100ML | A062414 | 007 | Aug 15, 1983 |
| EQ 70MG BASE/100ML | A062588 | 007 | Jan 06, 1986 |
| EQ 80MG BASE/100ML | A062588 | 008 | Jan 06, 1986 |
| EQ 90MG BASE/100ML | A062414 | 009 | Aug 15, 1983 |
| EQ 90MG BASE/100ML | A062588 | 009 | Jan 06, 1986 |
| EQ 100MG BASE/100ML | A062588 | 010 | Jan 06, 1986 |

U-GENCIN

| | | | |
|--------------------------------------|---------|-----|--|
| PHARMACIA AND UPJOHN EQ 10MG BASE/ML | A062248 | 001 | |
| EQ 40MG BASE/ML | A062248 | 002 | |

INJECTABLE; INTRATHECAL

GARAMYCIN

| | | | |
|------------------------------|---------|-----|--|
| + SCHERING EQ 2MG BASE/ML ** | N050505 | 001 | |
|------------------------------|---------|-----|--|

OINTMENT; OPHTHALMIC

GARAMYCIN

| | | | |
|-----------------------|---------|-----|--|
| SCHERING EQ 0.3% BASE | N050425 | 001 | |
|-----------------------|---------|-----|--|

GENTACIDIN

| | | | |
|-----------------------|---------|-----|--------------|
| NOVARTIS EQ 0.3% BASE | A062501 | 001 | Jul 26, 1984 |
|-----------------------|---------|-----|--------------|

GENTAFAIR

| | | | |
|---------------------------|---------|-----|--------------|
| PHARMAFAIR EQ 3MG BASE/GM | A062443 | 001 | May 26, 1983 |
|---------------------------|---------|-----|--------------|

OINTMENT; TOPICAL

GARAMYCIN

| | | | |
|--------------------------|---------|-----|--|
| SCHERING EQ 0.1% BASE ** | A060463 | 001 | |
|--------------------------|---------|-----|--|

GENTAFAIR

| | | | |
|-------------------------|---------|-----|--------------|
| PHARMAFAIR EQ 0.1% BASE | A062444 | 001 | May 26, 1983 |
|-------------------------|---------|-----|--------------|

GENTAMICIN SULFATE

| | | | |
|---------------------------------|---------|-----|--------------|
| ALPHARMA US PHARMS EQ 0.1% BASE | A062496 | 001 | Mar 14, 1984 |
|---------------------------------|---------|-----|--------------|

| | | | |
|-------------------------|---------|-----|--------------|
| PHARMADERM EQ 0.1% BASE | A062534 | 001 | Oct 10, 1984 |
|-------------------------|---------|-----|--------------|

SOLUTION/DROPS; OPHTHALMIC

GARAMYCIN

| | | | |
|----------------------------|---------|-----|--|
| + SCHERING EQ 0.3% BASE ** | N050039 | 002 | |
|----------------------------|---------|-----|--|

GENTACIDIN

| | | | |
|-----------------------|---------|-----|--------------|
| NOVARTIS EQ 0.3% BASE | A062480 | 001 | Mar 30, 1984 |
|-----------------------|---------|-----|--------------|

GENTAFAIR

| | | | |
|-------------------------|---------|-----|--------------|
| PHARMAFAIR EQ 0.3% BASE | A062440 | 001 | May 03, 1983 |
|-------------------------|---------|-----|--------------|

GENTAMICIN SULFATE

| | | | |
|-------------------------------|---------|-----|--------------|
| ALCON PHARMS LTD EQ 0.3% BASE | A062523 | 001 | Nov 25, 1985 |
|-------------------------------|---------|-----|--------------|

| | | | |
|---------------------|---------|-----|--------------|
| PACO EQ 3MG BASE/ML | A062932 | 001 | Nov 07, 1988 |
|---------------------|---------|-----|--------------|

GENTIAN VIOLET

SUPPOSITORY; VAGINAL

GVS

| | | | |
|------------------|---------|-----|--|
| SAVAGE LABS 0.4% | A083513 | 001 | |
|------------------|---------|-----|--|

TAMPON; VAGINAL

GENAPAX

| | | | |
|----------------|---------|-----|--|
| KEY PHARMS 5MG | A085017 | 001 | |
|----------------|---------|-----|--|

GLATIRAMER ACETATE

FOR SOLUTION; SUBCUTANEOUS

COPAXONE

| | | | |
|---------------------------|---------|-----|--------------|
| TEVA PHARMS USA 20MG/VIAL | N020622 | 001 | Dec 20, 1996 |
|---------------------------|---------|-----|--------------|

GLIMEPIRIDE

TABLET; ORAL

GLIMEPIRIDE

| | | | |
|-------------------------|---------|-----|--------------|
| ACTAVIS LABS FL INC 1MG | A076995 | 001 | Apr 27, 2010 |
|-------------------------|---------|-----|--------------|

| | | | |
|-----|---------|-----|--------------|
| 2MG | A076995 | 002 | Apr 27, 2010 |
|-----|---------|-----|--------------|

| | | | |
|-----|---------|-----|--------------|
| 4MG | A076995 | 003 | Apr 27, 2010 |
|-----|---------|-----|--------------|

| | | | |
|---------------------|---------|-----|--------------|
| EPIC PHARMA LLC 1MG | A077274 | 001 | Oct 06, 2005 |
|---------------------|---------|-----|--------------|

| | | | |
|-----|---------|-----|--------------|
| 2MG | A077274 | 002 | Oct 06, 2005 |
|-----|---------|-----|--------------|

| | | | |
|-----|---------|-----|--------------|
| 4MG | A077274 | 003 | Oct 06, 2005 |
|-----|---------|-----|--------------|

| | | | |
|------------------|---------|-----|--------------|
| HIKMA PHARMS 1MG | A078952 | 001 | Aug 01, 2013 |
|------------------|---------|-----|--------------|

| | | | |
|-----|---------|-----|--------------|
| 2MG | A078952 | 002 | Aug 01, 2013 |
|-----|---------|-----|--------------|

| | | | |
|-----|---------|-----|--------------|
| 4MG | A078952 | 003 | Aug 01, 2013 |
|-----|---------|-----|--------------|

| | | | |
|-----------|---------|-----|--------------|
| MYLAN 1MG | A077486 | 001 | Feb 10, 2006 |
|-----------|---------|-----|--------------|

| | | | |
|-----|---------|-----|--------------|
| 2MG | A077486 | 002 | Feb 10, 2006 |
|-----|---------|-----|--------------|

| | | | |
|-----|---------|-----|--------------|
| 4MG | A077486 | 003 | Feb 10, 2006 |
|-----|---------|-----|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GLIMEPIRIDE

TABLET; ORAL

GLIMEPIRIDE

| | | | |
|------------------|-----|-------------|--------------|
| RANBAXY | 3MG | A077366 001 | Oct 06, 2005 |
| | 6MG | A077366 002 | Oct 06, 2005 |
| RANBAXY LABS LTD | 1MG | A076875 001 | Oct 06, 2005 |
| | 2MG | A076875 002 | Oct 06, 2005 |
| | 4MG | A076875 003 | Oct 06, 2005 |
| | 8MG | A076875 004 | Oct 06, 2005 |
| WATSON LABS | 1MG | A077280 001 | Feb 03, 2006 |
| | 2MG | A077280 002 | Feb 03, 2006 |
| | 4MG | A077280 003 | Feb 03, 2006 |

GLIMEPIRIDE; ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDARYL

| | | | |
|--------------|-------------|-------------|--------------|
| + SB PHARMCO | 1MG; 4MG ** | N021700 001 | Nov 23, 2005 |
| + | 2MG; 4MG ** | N021700 002 | Nov 23, 2005 |
| + | 2MG; 8MG ** | N021700 004 | Mar 30, 2007 |
| + | 4MG; 4MG ** | N021700 003 | Nov 23, 2005 |
| + | 4MG; 8MG ** | N021700 005 | Mar 30, 2007 |

ROSIGLITAZONE MALEATE AND GLIMEPIRIDE

| | | | |
|-----------------|----------|-------------|--------------|
| TEVA PHARMS USA | 1MG; 4MG | A078709 001 | Apr 01, 2016 |
| | 2MG; 4MG | A078709 002 | Apr 01, 2016 |
| | 2MG; 8MG | A078709 004 | Apr 01, 2016 |
| | 4MG; 4MG | A078709 003 | Apr 01, 2016 |
| | 4MG; 8MG | A078709 005 | Apr 01, 2016 |

GLIPIZIDE

TABLET; ORAL

GLIPIZIDE

| | | | |
|----------------|-------|-------------|--------------|
| ANI PHARMS INC | 5MG | A074387 001 | Mar 04, 1996 |
| | 10MG | A074387 002 | Mar 04, 1996 |
| BARR LABS INC | 5MG | A074619 001 | Apr 04, 1997 |
| | 10MG | A074619 002 | Apr 04, 1997 |
| MYLAN | 5MG | A074438 001 | Jun 20, 1995 |
| | 10MG | A074438 002 | Jun 20, 1995 |
| OXFORD PHARMS | 5MG | A074378 001 | Nov 28, 1994 |
| | 10MG | A074378 002 | Nov 28, 1994 |
| SANDOZ | 5MG | A074542 001 | Jun 20, 1995 |
| | 10MG | A074542 002 | Jun 20, 1995 |
| WATSON LABS | 5MG | A074370 001 | Nov 22, 1994 |
| | 10MG | A074370 002 | Nov 22, 1994 |
| GLUCOTROL | | | |
| PFIZER | 2.5MG | N017783 003 | May 11, 1993 |

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

METAGLIP

| | | | |
|------------------------|-----------------|-------------|--------------|
| + BRISTOL MYERS SQUIBB | 2.5MG; 250MG ** | N021460 001 | Oct 21, 2002 |
| + | 2.5MG; 500MG ** | N021460 002 | Oct 21, 2002 |
| + | 5MG; 500MG ** | N021460 003 | Oct 21, 2002 |

GLUCAGON HYDROCHLORIDE

INJECTABLE; INJECTION

GLUCAGON

| | | | |
|---------|----------------------|-------------|--|
| + LILLY | EQ 1MG BASE/VIAL ** | N012122 001 | |
| + | EQ 10MG BASE/VIAL ** | N012122 002 | |

GLUTETHIMIDE

CAPSULE; ORAL

DORIDEN

| | | | |
|-------------------|-------|-------------|--|
| SANOFI AVENTIS US | 500MG | N009519 008 | |
|-------------------|-------|-------------|--|

TABLET; ORAL

DORIDEN

| | | | |
|-------------------|-------|-------------|--|
| SANOFI AVENTIS US | 250MG | N009519 002 | |
| | 500MG | N009519 005 | |

GLUTETHIMIDE

| | | | |
|---------|-------|-------------|--------------|
| HALSEY | 250MG | A089458 001 | Oct 10, 1986 |
| | 500MG | A089459 001 | Oct 10, 1986 |
| LANNETT | 250MG | A083475 001 | |
| | 500MG | A085571 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GLUTETHIMIDE

TABLET; ORAL

GLUTETHIMIDE

| | | | |
|-------------------|-------|---------|-----|
| UCB INC | 500MG | A085171 | 001 |
| UPSHER SMITH LABS | 500MG | A083234 | 002 |
| VITARINE | 500MG | A087297 | 001 |
| WATSON LABS | 500MG | A084362 | 001 |
| | 500MG | A085763 | 001 |

GLYBURIDE

TABLET; ORAL

GLYBURIDE

| | | | | |
|-------------------|-------|---------|-----|--------------|
| ACTAVIS ELIZABETH | 1.5MG | A075947 | 001 | Nov 14, 2002 |
| | 3MG | A075947 | 002 | Nov 14, 2002 |
| | 6MG | A075947 | 003 | Nov 14, 2002 |

GLYBURIDE (MICRONIZED)

| | | | | |
|-------------------|-------|---------|-----|--------------|
| SANOFI AVENTIS US | 1.5MG | N020055 | 001 | Apr 17, 1992 |
| | 3MG | N020055 | 002 | Apr 17, 1992 |
| | 6MG | N020055 | 003 | Mar 08, 2000 |
| YAOPHARMA CO LTD | 1.5MG | A075174 | 001 | Jun 22, 1998 |
| | 3MG | A075174 | 002 | Jun 22, 1998 |

GLYNASE

| | | | | |
|------------------------|----------|---------|-----|--------------|
| + PHARMACIA AND UPJOHN | 4.5MG ** | N020051 | 003 | Sep 24, 1993 |
|------------------------|----------|---------|-----|--------------|

MICRONASE

| | | | | |
|------------------------|-----------|---------|-----|--------------|
| + PHARMACIA AND UPJOHN | 1.25MG ** | N017498 | 001 | May 01, 1984 |
| | 2.5MG | N017498 | 002 | May 01, 1984 |
| + PHARMACIA AND UPJOHN | 5MG ** | N017498 | 003 | May 01, 1984 |

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLUCOVANCE

| | | | | |
|------------------------|------------------|---------|-----|--------------|
| + BRISTOL MYERS SQUIBB | 1.25MG; 250MG ** | N021178 | 001 | Jul 31, 2000 |
| + BRISTOL MYERS SQUIBB | 2.5MG; 500MG ** | N021178 | 002 | Jul 31, 2000 |
| + BRISTOL MYERS SQUIBB | 5MG; 500MG ** | N021178 | 003 | Jul 31, 2000 |

GLYBURIDE AND METFORMIN HYDROCHLORIDE

| | | | | |
|----------------|---------------|---------|-----|--------------|
| IMPAX LABS INC | 1.25MG; 250MG | A076731 | 001 | Nov 19, 2004 |
| | 2.5MG; 500MG | A076731 | 002 | Nov 19, 2004 |
| | 5MG; 500MG | A076731 | 003 | Nov 19, 2004 |
| TEVA | 1.25MG; 250MG | A076821 | 001 | Jan 27, 2005 |
| | 2.5MG; 500MG | A076821 | 002 | Jan 27, 2005 |
| | 5MG; 500MG | A076821 | 003 | Jan 27, 2005 |

GLYCINE

SOLUTION; IRRIGATION

GLYCINE 1.5% IN PLASTIC CONTAINER

| | | | | |
|-----------------|-------------|---------|-----|--------------|
| BAXTER HLTHCARE | 1.5GM/100ML | N018522 | 001 | Feb 19, 1982 |
| HOSPIRA | 1.5GM/100ML | N017633 | 001 | |

GLYCOPYRROLATE

INJECTABLE; INJECTION

GLYCOPYRROLATE

| | | | | |
|-----------------|----------|---------|-----|--------------|
| ABRAXIS PHARM | 0.2MG/ML | A088475 | 001 | Jun 12, 1984 |
| HOSPIRA | 0.2MG/ML | A089393 | 001 | Jun 15, 1988 |
| TEVA PARENTERAL | 0.2MG/ML | A081169 | 001 | Sep 10, 1991 |
| WATSON LABS | 0.2MG/ML | A086947 | 001 | Jun 24, 1983 |

ROBINUL

| | | | |
|------------------------|-------------|---------|-----|
| ROBINS AH | 0.2MG/ML | N014764 | 001 |
| + WEST-WARD PHARMS INT | 0.2MG/ML ** | N017558 | 001 |

TABLET; ORAL

GLYCOPYRROLATE

| | | | | |
|-------------------|-----|---------|-----|--------------|
| HIKMA INTL PHARMS | 1MG | A040836 | 001 | Mar 05, 2009 |
| | 2MG | A040836 | 002 | Mar 05, 2009 |
| RENATA | 1MG | A040568 | 001 | Dec 22, 2004 |
| | 2MG | A040568 | 002 | Dec 22, 2004 |
| WATSON LABS | 1MG | A085562 | 001 | |
| | 1MG | A086902 | 001 | |
| | 2MG | A085563 | 001 | |
| | 2MG | A086178 | 001 | |
| | 2MG | A086900 | 001 | |

ROBINUL

| | | | |
|---------------------|-----|---------|-----|
| + CASPER PHARMA LLC | 1MG | N012827 | 001 |
|---------------------|-----|---------|-----|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GLYCOPYRROLATE

TABLET; ORAL

ROBINUL FORTE

+ CASPER PHARMA LLC 2MG

N012827 002

GONADORELIN ACETATE

INJECTABLE; INJECTION

LUTREPULSE KIT

FERRING 0.8MG/VIAL

N019687 001 Oct 10, 1989

3.2MG/VIAL

N019687 002 Oct 10, 1989

GONADORELIN HYDROCHLORIDE

INJECTABLE; INJECTION

FACTREL

WEST-WARD PHARMS INT EQ 0.1MG BASE/VIAL

N018123 001 Sep 30, 1982

EQ 0.2MG BASE/VIAL

N018123 002 Sep 30, 1982

EQ 0.5MG BASE/VIAL

N018123 003 Sep 30, 1982

GONADOTROPIN, CHORIONIC

INJECTABLE; INJECTION

A.P.L.

FERRING 5,000 UNITS/VIAL

N017055 001

10,000 UNITS/VIAL

N017055 002

20,000 UNITS/VIAL

N017055 003

CHORIONIC GONADOTROPIN

BEL MAR 5,000 UNITS/VIAL

N017054 001

10,000 UNITS/VIAL

N017054 002

FERRING 2,000 UNITS/VIAL

N017016 009 Dec 27, 1984

2,000 UNITS/VIAL

N017016 011 Feb 16, 1990

15,000 UNITS/VIAL

N017016 010 Feb 15, 1985

20,000 UNITS/VIAL

N017016 004

FRESENIUS KABI USA 5,000 UNITS/VIAL

N017067 001

15,000 UNITS/VIAL

N017067 004

20,000 UNITS/VIAL

N017067 003

FOLLUTEIN

BRISTOL MYERS SQUIBB 10,000 UNITS/VIAL

N017056 001

GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OPHTHALMIC

NEO-POLYCIN

DOW PHARM 0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML

A060427 001

NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN

IPHARM 0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML

A062818 001 Oct 11, 1988

WATSON LABS 0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML

A062788 001 Jun 11, 1987

NEOMYCIN SULFATE AND POLYMYXIN B SULFATE GRAMICIDIN

PHARMAFAIR 0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML

A062383 001 Aug 31, 1982

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

GRANISETRON HYDROCHLORIDE

BAXTER HLTHCARE CORP EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)

A078197 001 Dec 31, 2007

EQ 1MG BASE/ML (EQ 1MG BASE/ML)

A078198 001 Jun 30, 2008

EQ 4MG BASE/4ML (EQ 1MG BASE/ML)

A078198 002 Jun 30, 2008

SANDOZ INC EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)

A078808 001 Apr 29, 2008

TEVA PHARMS USA EQ 1MG BASE/ML (EQ 1MG BASE/ML)

A077963 001 Jan 03, 2008

GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE

TEVA PHARMS USA EQ 1MG BASE/ML (EQ 1MG BASE/ML)

A077165 001 Dec 31, 2007

KYTRIL

+ ROCHE EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML) **

N020239 003 Sep 17, 2004

+ EQ 1MG BASE/ML (EQ 1MG BASE/ML) **

N020239 004 Mar 11, 1994

+ EQ 3MG BASE/ML **

N020239 001 Dec 29, 1993

+ EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **

N020239 002 Mar 11, 1994

SOLUTION; ORAL

GRANISOL

PEDIATRAX EQ 2MG BASE/10ML

A078334 001 Feb 28, 2008

KYTRIL

+ ROCHE EQ 2MG BASE/10ML **

N021238 001 Jun 27, 2001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GRANISETRON HYDROCHLORIDE

TABLET; ORAL

GRANISETRON HYDROCHLORIDE

| | | | |
|-----------------|-------------|-------------|--------------|
| BARR | EQ 1MG BASE | A078221 001 | Dec 31, 2007 |
| EPIC PHARMA LLC | EQ 1MG BASE | A078260 001 | Dec 31, 2007 |

KYTRIL

| | | | |
|---------|----------------|-------------|--------------|
| + ROCHE | EQ 1MG BASE ** | N020305 001 | Mar 16, 1995 |
| + | EQ 2MG BASE ** | N020305 002 | Jun 15, 1998 |

GREPAFLOXACIN HYDROCHLORIDE

TABLET; ORAL

RAXAR

| | | | |
|--------|---------------|-------------|--------------|
| OTSUKA | EQ 200MG BASE | N020695 001 | Nov 06, 1997 |
| | EQ 400MG BASE | N020695 002 | May 14, 1998 |
| | EQ 600MG BASE | N020695 003 | May 14, 1998 |

GRISEOFULVIN, MICROCRYSTALLINE

CAPSULE; ORAL

GRISACTIN

| | | | |
|--------------|-------|-------------|--|
| WYETH AYERST | 125MG | N050051 002 | |
| | 250MG | N050051 001 | |

SUSPENSION; ORAL

GRIFULVIN V

| | | | |
|-----------------------|--------------|-------------|--|
| + JOHNSON AND JOHNSON | 125MG/5ML ** | N050448 001 | |
|-----------------------|--------------|-------------|--|

TABLET; ORAL

FULVICIN-U/F

| | | | |
|--------------|-------|-------------|--|
| CHARTWELL RX | 250MG | A060569 002 | |
| | 500MG | A060569 001 | |

GRIFULVIN V

| | | | |
|--------------------|----------|-------------|--|
| J AND J | 125MG | A060618 001 | |
| | 250MG | A060618 002 | |
| | 500MG | A060618 003 | |
| VALEANT LUXEMBOURG | 125MG | A062279 001 | |
| | 250MG ** | A062279 002 | |

GRISACTIN

| | | | |
|--------------|-------|-------------|--|
| WYETH AYERST | 500MG | A060212 001 | |
|--------------|-------|-------------|--|

GRISEOFULVIN, MICROSIZE

SUSPENSION; ORAL

GRIFULVIN V

| | | | |
|--------------------|--------------|-------------|--------------|
| VALEANT LUXEMBOURG | 125MG/5ML ** | A062483 001 | Jan 26, 1984 |
|--------------------|--------------|-------------|--------------|

TABLET; ORAL

GRIFULVIN V

| | | | |
|--------------------|-------|-------------|--|
| VALEANT LUXEMBOURG | 500MG | A062279 003 | |
|--------------------|-------|-------------|--|

GRISEOFULVIN, ULTRAMICROCRYSTALLINE

TABLET; ORAL

FULVICIN P/G

| | | | |
|--------------|-------|-------------|--|
| CHARTWELL RX | 125MG | A061996 001 | |
| | 250MG | A061996 002 | |

FULVICIN P/G 165

| | | | |
|--------------|-------|-------------|--------------|
| CHARTWELL RX | 165MG | A061996 003 | Apr 06, 1982 |
|--------------|-------|-------------|--------------|

FULVICIN P/G 330

| | | | |
|--------------|-------|-------------|--------------|
| CHARTWELL RX | 330MG | A061996 004 | Apr 06, 1982 |
|--------------|-------|-------------|--------------|

GRISACTIN ULTRA

| | | | |
|--------------|-------|-------------|--------------|
| WYETH AYERST | 125MG | A062178 001 | |
| | 165MG | A062438 001 | Nov 17, 1983 |
| | 250MG | A062178 002 | |
| | 330MG | A062438 002 | Nov 17, 1983 |

ULTRAGRIS-165

| | | | |
|-------|-------|-------------|--------------|
| PLIVA | 165MG | A062645 001 | Jun 30, 1992 |
|-------|-------|-------------|--------------|

ULTRAGRIS-330

| | | | |
|-------|-------|-------------|--------------|
| PLIVA | 330MG | A062646 001 | Jun 30, 1992 |
|-------|-------|-------------|--------------|

GUAIFENESIN; HYDROCODONE BITARTRATE

SOLUTION; ORAL

FLOWTUSS

| | | | |
|------------|----------------------|-------------|--------------|
| BKK PHARMS | 200MG/5ML; 2.5MG/5ML | N022424 001 | May 14, 2015 |
|------------|----------------------|-------------|--------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

GUAIFENESIN; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

HYCOFENIX

+ BKK PHARMS

200MG/5ML; 2.5MG/5ML; 30MG/5ML

N022279 001 May 14, 2015

GUANABENZ ACETATE

TABLET; ORAL

GUANABENZ ACETATE

ANI PHARMS INC

EQ 4MG BASE

A074267 001 Jun 01, 1994

EQ 8MG BASE

A074267 002 Jun 01, 1994

WATSON LABS

EQ 4MG BASE

A074025 001 Feb 28, 1994

EQ 8MG BASE

A074025 002 Feb 28, 1994

YAOPHARMA CO LTD

EQ 4MG BASE

A074517 001 Sep 30, 1998

EQ 8MG BASE

A074517 002 Sep 30, 1998

WYTENSIN

WYETH AYERST

EQ 4MG BASE

N018587 001 Sep 07, 1982

EQ 8MG BASE

N018587 002 Sep 07, 1982

EQ 16MG BASE

N018587 003 Sep 07, 1982

GUANADREL SULFATE

TABLET; ORAL

HYLOREL

PHARMACIA AND UPJOHN 10MG

N018104 001 Dec 29, 1982

25MG

N018104 002 Dec 29, 1982

GUANETHIDINE MONOSULFATE

TABLET; ORAL

GUANETHIDINE MONOSULFATE

WATSON LABS

EQ 10MG SULFATE

A086113 001 Mar 26, 1985

EQ 25MG SULFATE

A086114 001 Mar 26, 1985

ISMELIN

NOVARTIS

EQ 10MG SULFATE

N012329 001

EQ 25MG SULFATE

N012329 002

GUANETHIDINE MONOSULFATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ESIMIL

NOVARTIS

10MG; 25MG

N013553 001

GUANFACINE HYDROCHLORIDE

TABLET; ORAL

GUANFACINE HYDROCHLORIDE

EPIC PHARMA LLC

EQ 1MG BASE

A074673 001 Feb 28, 1997

EQ 2MG BASE

A074673 002 Feb 28, 1997

WATSON LABS

EQ 1MG BASE

A074762 001 Jun 25, 1997

EQ 2MG BASE

A074762 002 Jun 25, 1997

TENEX

+ PROMIUS PHARMA

EQ 1MG BASE

N019032 001 Oct 27, 1986

+ EQ 2MG BASE

N019032 002 Nov 07, 1988

EQ 3MG BASE

N019032 003 Nov 07, 1988

TABLET, EXTENDED RELEASE; ORAL

GUANFACINE HYDROCHLORIDE

IMPAX LABS INC

EQ 1MG BASE

A202238 001 Oct 20, 2015

EQ 2MG BASE

A202238 002 Oct 20, 2015

EQ 3MG BASE

A202238 003 Oct 20, 2015

EQ 4MG BASE

A202238 004 Oct 20, 2015

HALAZEPAM

TABLET; ORAL

PAXIPAM

SCHERING

20MG

N017736 003

40MG

N017736 004

HALCINONIDE

CREAM; TOPICAL

HALOG

WESTWOOD SQUIBB

0.025%

N017818 001

HALOG-E

SUN PHARM INDS INC

0.1%

N018234 001

OINTMENT; TOPICAL

HALOG

BRISTOL MYERS SQUIBB 0.025%

N018125 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HALCINONIDE

SOLUTION;TOPICAL

HALOG

SUN PHARM INDS INC 0.1% N017823 001

HALOBETASOL PROPIONATE

OINTMENT;TOPICAL

HALOBETASOL PROPIONATE

FOUGERA PHARMS 0.05% A076903 001 Dec 16, 2004

HALOFANTRINE HYDROCHLORIDE

TABLET;ORAL

HALFAN

GLAXOSMITHKLINE 250MG N020250 001 Jul 24, 1992

HALOPERIDOL

TABLET;ORAL

HALDOL

| | | | | |
|---|--------------|----------|-------------|--------------|
| + | ORTHO MCNEIL | 0.5MG ** | N015921 001 | |
| + | | 1MG ** | N015921 002 | |
| + | | 2MG ** | N015921 003 | |
| + | | 5MG ** | N015921 004 | |
| + | | 10MG ** | N015921 005 | |
| + | | 20MG ** | N015921 006 | Feb 02, 1982 |

HALDOL SOLUTAB

ORTHO MCNEIL PHARM 1MG N017079 001

HALOPERIDOL

| | | | | |
|--|---------------------|-------|-------------|--------------|
| | ANDA REPOSITORY | 0.5MG | A071156 001 | Jan 02, 1987 |
| | | 1MG | A071157 001 | Jan 02, 1987 |
| | | 2MG | A071172 001 | Jan 02, 1987 |
| | | 5MG | A071212 001 | Jan 07, 1988 |
| | | 10MG | A071173 001 | Jan 07, 1988 |
| | | 20MG | A071177 001 | Jan 07, 1988 |
| | CYCLE PHARMS LTD | 0.5MG | A071128 001 | Feb 17, 1987 |
| | | 1MG | A071129 001 | Feb 17, 1987 |
| | | 2MG | A071130 001 | Feb 17, 1987 |
| | | 5MG | A071131 001 | Feb 17, 1987 |
| | | 10MG | A071132 001 | May 12, 1987 |
| | | 20MG | A071133 001 | May 12, 1987 |
| | DURAMED PHARMS BARR | 0.5MG | A071216 001 | Dec 04, 1986 |
| | | 1MG | A071217 001 | Dec 04, 1986 |
| | | 2MG | A071218 001 | Dec 04, 1986 |
| | | 5MG | A071219 001 | Dec 04, 1986 |
| | | 10MG | A071220 001 | Jul 07, 1987 |
| | | 20MG | A071221 001 | Jul 07, 1987 |
| | LEDERLE | 0.5MG | A072727 001 | Sep 19, 1989 |
| | | 1MG | A072728 001 | Sep 19, 1989 |
| | | 2MG | A072729 001 | Sep 19, 1989 |
| | | 5MG | A072730 001 | Sep 19, 1989 |
| | | 10MG | A072731 001 | Sep 19, 1989 |
| | | 20MG | A072732 001 | Sep 19, 1989 |
| | PAR PHARM | 20MG | A071328 001 | Jul 20, 1987 |
| | PUREPAC PHARM | 0.5MG | A071071 001 | Nov 03, 1986 |
| | | 1MG | A071072 001 | Nov 03, 1986 |
| | | 2MG | A071073 001 | Nov 03, 1986 |
| | | 5MG | A071074 001 | Nov 03, 1986 |
| | | 10MG | A071075 001 | Aug 04, 1987 |
| | | 20MG | A071076 001 | Aug 04, 1987 |
| | QUANTUM PHARMICS | 0.5MG | A071255 001 | Feb 17, 1987 |
| | | 1MG | A071269 001 | Feb 17, 1987 |
| | | 2MG | A071256 001 | Feb 17, 1987 |
| | | 5MG | A071257 001 | Feb 17, 1987 |
| | ROYCE LABS | 0.5MG | A071722 001 | Dec 24, 1987 |
| | | 1MG | A071723 001 | Dec 24, 1987 |
| | | 2MG | A071724 001 | Dec 24, 1987 |
| | | 5MG | A071725 001 | Dec 24, 1987 |
| | | 10MG | A072121 001 | Dec 24, 1987 |
| | | 20MG | A072122 001 | Dec 24, 1987 |
| | SCS | 0.5MG | A070720 001 | Jun 10, 1986 |
| | | 1MG | A070721 001 | Jun 10, 1986 |
| | | 2MG | A070722 001 | Jun 10, 1986 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HALOPERIDOL

TABLET; ORAL

HALOPERIDOL

| | | | | |
|-------------|-------|---------|-----|--------------|
| | 5MG | A070723 | 001 | Jun 10, 1986 |
| | 10MG | A070724 | 001 | Jun 10, 1986 |
| | 20MG | A070725 | 001 | Sep 24, 1986 |
| VINTAGE | 0.5MG | A071235 | 002 | Nov 03, 1986 |
| | 1MG | A071235 | 003 | Nov 03, 1986 |
| | 2MG | A071235 | 001 | Nov 03, 1986 |
| | 5MG | A071235 | 004 | Nov 03, 1986 |
| | 10MG | A071235 | 005 | Jul 20, 1987 |
| WATSON LABS | 0.5MG | A070981 | 001 | Mar 06, 1987 |
| | 0.5MG | A071571 | 001 | Jun 03, 1988 |
| | 1MG | A070982 | 001 | Mar 06, 1987 |
| | 1MG | A071572 | 001 | Jun 03, 1988 |
| | 2MG | A070983 | 001 | Mar 06, 1987 |
| | 2MG | A071573 | 001 | Jun 03, 1988 |
| | 5MG | A070984 | 001 | Mar 06, 1987 |
| | 5MG | A071374 | 001 | Jun 03, 1988 |
| | 10MG | A071375 | 001 | Jun 03, 1988 |
| | 10MG | A072113 | 001 | Aug 27, 1991 |
| | 20MG | A071376 | 001 | Jun 03, 1988 |
| | 20MG | A072353 | 001 | Aug 27, 1991 |

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

HALOPERIDOL DECANOATE

| | | | | |
|------------|------------------|---------|-----|--------------|
| HOSPIRA | EQ 50MG BASE/ML | A075176 | 001 | Feb 09, 2000 |
| | EQ 100MG BASE/ML | A075176 | 002 | Feb 09, 2000 |
| SANDOZ INC | EQ 50MG BASE/ML | A076463 | 001 | Jun 24, 2005 |
| | EQ 100MG BASE/ML | A076463 | 002 | Jun 24, 2005 |

HALOPERIDOL LACTATE

CONCENTRATE; ORAL

HALDOL

| | | | | |
|-----------------------|-------------------|---------|-----|--------------|
| ORTHO MCNEIL | EQ 2MG BASE/ML ** | N015922 | 001 | |
| HALOPERIDOL | | | | |
| ALPHARMA | EQ 2MG BASE/ML | A070318 | 001 | Apr 11, 1986 |
| MORTON GROVE | EQ 2MG BASE/ML | A070710 | 001 | Mar 07, 1986 |
| SCS | EQ 2MG BASE/ML | A070726 | 001 | Jun 10, 1986 |
| TEVA | EQ 2MG BASE/ML | A071015 | 001 | Aug 25, 1987 |
| HALOPERIDOL INTENSOL | | | | |
| CYCLE PHARMS LTD | EQ 2MG BASE/ML | A072045 | 001 | Apr 12, 1988 |
| INJECTABLE; INJECTION | | | | |
| HALOPERIDOL | | | | |
| ABRAXIS PHARM | EQ 5MG BASE/ML | A071187 | 001 | Jan 20, 1987 |
| BAXTER HLTHCARE CORP | EQ 5MG BASE/ML | A076791 | 001 | Aug 25, 2004 |
| | EQ 5MG BASE/ML | A076828 | 001 | Aug 25, 2004 |
| FOSUN PHARMA | EQ 5MG BASE/ML | A076464 | 001 | Sep 29, 2004 |
| MARSAM PHARMS LLC | EQ 5MG BASE/ML | A072516 | 001 | Feb 25, 1993 |
| | EQ 5MG BASE/ML | A072517 | 001 | Feb 25, 1993 |
| SMITH AND NEPHEW | EQ 5MG BASE/ML | A070802 | 001 | Dec 14, 1987 |
| SOLOPAK | EQ 5MG BASE/ML | A070800 | 001 | Dec 14, 1987 |
| | EQ 5MG BASE/ML | A070801 | 001 | Dec 14, 1987 |
| | EQ 5MG BASE/ML | A070864 | 001 | Dec 14, 1987 |
| WATSON LABS | EQ 5MG BASE/ML | A070713 | 001 | May 17, 1988 |
| | EQ 5MG BASE/ML | A070714 | 001 | May 17, 1988 |
| | EQ 5MG BASE/ML | A070744 | 001 | May 17, 1988 |
| SOLUTION; ORAL | | | | |
| HALOPERIDOL LACTATE | | | | |
| ACTAVIS MID ATLANTIC | EQ 1MG BASE/ML | A074536 | 001 | Nov 02, 1995 |

HALOPROGIN

CREAM; TOPICAL

HALOTEX

| | | | | |
|-------------------|----|---------|-----|--|
| WESTWOOD SQUIBB | 1% | N016942 | 001 | |
| SOLUTION; TOPICAL | | | | |
| HALOTEX | | | | |
| WESTWOOD SQUIBB | 1% | N016943 | 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HALOTHANE

LIQUID; INHALATION

FLUOTHANE

WYETH AYERST 99.99% N011338 001

HALOTHANE

BH 99.99% A084977 001

HALOCARBON 99.99% A080810 001

HOSPIRA 99.99% A083254 001

HEPARIN CALCIUM

INJECTABLE; INJECTION

CALCIPARINE

SANOFI AVENTIS US 25,000 UNITS/ML N018237 001

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN LOCK FLUSH

HOSPIRA 100 UNITS/ML N005264 010

INTL MEDICATION 10 UNITS/ML A086357 001

500 UNITS/ML A086357 002

LUITPOLD 10 UNITS/ML A089063 001

100 UNITS/ML A089064 001 Oct 09, 1985

PARKE DAVIS 10 UNITS/ML N017346 006

SMITH AND NEPHEW 10 UNITS/ML A087904 001 Apr 20, 1983

10 UNITS/ML A087958 001 Apr 20, 1983

10 UNITS/ML A088458 001 Jul 26, 1984

10 UNITS/ML A088580 001 Oct 25, 1984

100 UNITS/ML A087906 001 Apr 20, 1983

100 UNITS/ML A087959 001 Apr 20, 1983

100 UNITS/ML A088460 001 Jul 26, 1984

100 UNITS/ML A088581 001 Oct 25, 1984

SOLOPAK 10 UNITS/ML A087903 001 Apr 20, 1983

10 UNITS/ML A088457 001 Oct 25, 1984

100 UNITS/ML A087905 001 Apr 20, 1983

100 UNITS/ML A088459 001 Jul 26, 1984

HEPARIN SODIUM

ABRAXIS PHARM 1,000 UNITS/ML N017033 001

1,000 UNITS/ML N017979 001

5,000 UNITS/ML N017979 003

10,000 UNITS/ML N017979 002

AKORN 1,000 UNITS/ML N017486 001

5,000 UNITS/ML N017486 002

10,000 UNITS/ML N017486 003

20,000 UNITS/ML N017486 004

40,000 UNITS/ML N017486 005

CHAMBERLIN PARENTERL 1,000 UNITS/ML N017130 001

5,000 UNITS/ML N017130 002

10,000 UNITS/ML N017130 003

20,000 UNITS/ML N017130 004

DELL LABS 1,000 UNITS/ML N017540 001

5,000 UNITS/ML N017540 002

10,000 UNITS/ML N017540 003

20,000 UNITS/ML N017540 004

40,000 UNITS/ML N017540 005

FRESENIUS KABI USA 1,000 UNITS/ML N017651 005

5,000 UNITS/ML N017029 002

10,000 UNITS/ML N017651 003

20,000 UNITS/ML N017651 008

HOSPIRA 2,500 UNITS/ML A088099 001 Apr 28, 1983

10,000 UNITS/ML A040095 001 Jul 26, 1996

LILLY 1,000 UNITS/ML N005521 001

10,000 UNITS/ML N005521 002

20,000 UNITS/ML N005521 004

LUITPOLD 1,000 UNITS/ML A087452 001 Oct 31, 1983

ORGANON USA INC 1,000 UNITS/ML N000552 008

5,000 UNITS/ML N000552 009

10,000 UNITS/ML N000552 010

PARKE DAVIS 1,000 UNITS/ML N017346 001

5,000 UNITS/ML N017346 002

7,500 UNITS/ML N017346 003

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

| | | | |
|-----------------------------|-----------------------------------------------|-------------|--------------|
| | 10,000 UNITS/ML | N017346 004 | |
| | 20,000 UNITS/ML | N017346 005 | |
| PHARM SPEC | 1,000 UNITS/ML | N017780 001 | |
| | 5,000 UNITS/ML | N017780 002 | |
| | 10,000 UNITS/ML | N017780 003 | |
| | 20,000 UNITS/ML | N017780 004 | |
| | 40,000 UNITS/ML | N017780 005 | |
| PHARMACIA AND UPJOHN | 1,000 UNITS/ML | N004570 001 | |
| | 5,000 UNITS/ML | N004570 002 | |
| | 10,000 UNITS/ML | N004570 003 | |
| SMITH AND NEPHEW | 1,000 UNITS/ML | A088239 001 | Jul 26, 1984 |
| SOLOPAK | 1,000 UNITS/ML | A087043 001 | |
| | 5,000 UNITS/ML | A087077 001 | |
| | 5,000 UNITS/0.5ML | A087395 001 | |
| | 10,000 UNITS/ML | A087107 001 | |
| | 10,000 UNITS/0.5ML | A087363 001 | |
| WATSON LABS | 1,000 UNITS/ML | N017064 002 | |
| | 2,500 UNITS/ML | N017064 015 | |
| | 3,000 UNITS/ML | N017064 016 | |
| | 4,000 UNITS/ML | N017064 017 | |
| | 5,000 UNITS/ML | N017064 003 | |
| | 6,000 UNITS/ML | N017064 018 | |
| | 7,500 UNITS/ML | N017064 019 | |
| | 10,000 UNITS/ML | N017064 004 | |
| | 20,000 UNITS/ML | N017064 005 | |
| | 40,000 UNITS/ML | N017064 006 | |
| WATSON LABS INC | 1,000 UNITS/ML | A040007 001 | Jun 07, 1996 |
| | 1,000 UNITS/ML | A040008 001 | Oct 10, 1995 |
| + WEST-WARD PHARMS INT | 1,000 UNITS/ML ** | N017007 001 | |
| + | 2,500 UNITS/ML ** | N017007 007 | |
| + | 5,000 UNITS/ML ** | N017007 002 | |
| + | 5,000 UNITS/0.5ML ** | N017007 010 | |
| | 5,000 UNITS/0.5ML | N017037 013 | Apr 07, 1986 |
| + | 7,500 UNITS/ML ** | N017007 003 | |
| + | 10,000 UNITS/ML ** | N017007 004 | |
| + | 15,000 UNITS/ML ** | N017007 005 | |
| + | 20,000 UNITS/ML ** | N017007 006 | |
| HEPARIN SODIUM 1,000 UNITS | IN DEXTROSE 5% IN PLASTIC CONTAINER | | |
| MCGAW | 200 UNITS/100ML | N019130 001 | Dec 31, 1984 |
| HEPARIN SODIUM 1,000 UNITS | IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | |
| B BRAUN | 200 UNITS/100ML | N019042 001 | Mar 29, 1985 |
| HEPARIN SODIUM 10,000 UNITS | AND DEXTROSE 5% IN PLASTIC CONTAINER | | |
| BAXTER HLTHCARE | 2,000 UNITS/100ML | N018814 002 | Jul 09, 1985 |
| HEPARIN SODIUM 10,000 UNITS | IN DEXTROSE 5% | | |
| HOSPIRA | 10,000 UNITS/100ML | N018911 006 | Jan 30, 1985 |
| HEPARIN SODIUM 10,000 UNITS | IN SODIUM CHLORIDE 0.45% | | |
| HOSPIRA | 10,000 UNITS/100ML | N018911 001 | Jan 30, 1985 |
| | 10,000 UNITS/100ML | N018916 005 | Jan 31, 1984 |
| HEPARIN SODIUM 10,000 UNITS | IN SODIUM CHLORIDE 0.9% | | |
| HOSPIRA | 10,000 UNITS/100ML | N018911 003 | Jan 30, 1985 |
| | 10,000 UNITS/100ML | N018916 002 | Jan 31, 1984 |
| HEPARIN SODIUM 12,500 UNITS | IN DEXTROSE 5% | | |
| HOSPIRA | 5,000 UNITS/100ML | N018911 007 | Jan 30, 1985 |
| HEPARIN SODIUM 12,500 UNITS | IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER | | |
| B BRAUN | 5,000 UNITS/100ML | N019802 001 | Jul 20, 1992 |
| HEPARIN SODIUM 12,500 UNITS | IN SODIUM CHLORIDE 0.9% | | |
| HOSPIRA | 5,000 UNITS/100ML | N018911 005 | Jan 30, 1985 |
| | 5,000 UNITS/100ML | N018916 003 | Jan 31, 1984 |
| HEPARIN SODIUM 2,000 UNITS | IN DEXTROSE 5% IN PLASTIC CONTAINER | | |
| MCGAW | 200 UNITS/100ML | N019130 003 | Dec 31, 1984 |
| HEPARIN SODIUM 2,000 UNITS | IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | |
| B BRAUN | 200 UNITS/100ML | N019042 002 | Mar 29, 1985 |
| HEPARIN SODIUM 20,000 UNITS | AND DEXTROSE 5% IN PLASTIC CONTAINER | | |
| BAXTER HLTHCARE | 4,000 UNITS/100ML | N018814 001 | Oct 31, 1983 |
| HEPARIN SODIUM 25,000 UNITS | AND DEXTROSE 5% IN PLASTIC CONTAINER | | |
| BAXTER HLTHCARE | 5,000 UNITS/100ML | N018814 003 | Jul 09, 1985 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HEPARIN SODIUM

INJECTABLE; INJECTION

| | | | | |
|---------------------------------------------------------------------------|--------------------|---------|-----|--------------|
| HEPARIN SODIUM 25,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER | | | | |
| | 10,000 UNITS/100ML | N018814 | 004 | Jul 02, 1987 |
| HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% | | | | |
| HOSPIRA | 5,000 UNITS/100ML | N018911 | 009 | Jan 30, 1985 |
| | 10,000 UNITS/100ML | N018911 | 008 | Jan 30, 1985 |
| HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 5,000 UNITS/100ML | N019134 | 001 | Mar 29, 1985 |
| HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 5,000 UNITS/100ML | N019802 | 005 | Jul 20, 1992 |
| | 10,000 UNITS/100ML | N019802 | 002 | Jul 20, 1992 |
| HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.9% | | | | |
| HOSPIRA | 5,000 UNITS/100ML | N018911 | 004 | Jan 30, 1985 |
| HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 5,000 UNITS/100ML | N019135 | 001 | Mar 29, 1985 |
| | 5,000 UNITS/100ML | N019802 | 003 | Jul 20, 1992 |
| HOSPIRA | 5,000 UNITS/100ML | N018916 | 009 | Jan 31, 1984 |
| HEPARIN SODIUM 5,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | | |
| BAXTER HLTHCARE | 500 UNITS/100ML | N018609 | 003 | Apr 28, 1982 |
| HEPARIN SODIUM 5,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER | | | | |
| MCGAW | 1,000 UNITS/100ML | N019130 | 002 | Dec 31, 1984 |
| HEPARIN SODIUM 5,000 UNITS IN SODIUM CHLORIDE 0.45% | | | | |
| HOSPIRA | 100 UNITS/ML | N018911 | 002 | Jan 30, 1985 |
| | 100 UNITS/ML | N018916 | 004 | Jan 31, 1984 |
| HEPARIN SODIUM 5,000 UNITS IN SODIUM CHLORIDE 0.9% | | | | |
| HOSPIRA | 1,000 UNITS/100ML | N018916 | 001 | Jan 31, 1984 |
| HEPARIN SODIUM 5,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 1,000 UNITS/100ML | N019042 | 004 | Mar 29, 1985 |
| HEPARIN SODIUM PRESERVATIVE FREE | | | | |
| HOSPIRA | 2,000 UNITS/ML | N005264 | 013 | Apr 07, 1986 |
| | 2,500 UNITS/ML | N005264 | 014 | Apr 07, 1986 |
| PHARMA SERVE NY | 1,000 UNITS/ML | A086129 | 001 | |
| WATSON LABS INC | 1,000 UNITS/ML | A089464 | 001 | Jun 03, 1986 |
| LIPO-HEPIN | | | | |
| 3M | 1,000 UNITS/0.5ML | N017027 | 001 | |
| | 1,000 UNITS/ML | N017027 | 006 | |
| | 5,000 UNITS/0.5ML | N017027 | 002 | |
| | 5,000 UNITS/ML | N017027 | 008 | |
| | 7,500 UNITS/0.5ML | N017027 | 010 | |
| | 10,000 UNITS/0.5ML | N017027 | 003 | |
| | 10,000 UNITS/ML | N017027 | 009 | |
| | 15,000 UNITS/0.5ML | N017027 | 011 | |
| | 20,000 UNITS/0.5ML | N017027 | 004 | |
| | 20,000 UNITS/ML | N017027 | 007 | |
| | 40,000 UNITS/ML | N017027 | 005 | |
| LIQUAEMIN LOCK FLUSH | | | | |
| ORGANON USA INC | 100 UNITS/ML | N000552 | 007 | |
| LIQUAEMIN SODIUM | | | | |
| ORGANON USA INC | 1,000 UNITS/ML | N000552 | 004 | |
| | 5,000 UNITS/ML | N000552 | 003 | |
| | 10,000 UNITS/ML | N000552 | 005 | |
| | 20,000 UNITS/ML | N000552 | 001 | |
| | 40,000 UNITS/ML | N000552 | 002 | |
| LIQUAEMIN SODIUM PRESERVATIVE FREE | | | | |
| ORGANON USA INC | 1,000 UNITS/ML | N000552 | 011 | Apr 11, 1986 |
| | 5,000 UNITS/ML | N000552 | 012 | Apr 11, 1986 |
| | 10,000 UNITS/ML | N000552 | 013 | Apr 11, 1986 |
| PANHEPRIN | | | | |
| HOSPIRA | 1,000 UNITS/ML | N005264 | 004 | |
| | 5,000 UNITS/ML | N005264 | 006 | |
| | 10,000 UNITS/ML | N005264 | 007 | |
| | 20,000 UNITS/ML | N005264 | 008 | |
| | 40,000 UNITS/ML | N005264 | 009 | |
| SODIUM HEPARIN | | | | |
| ABRAXIS PHARM | 5,000 UNITS/ML | N017033 | 002 | |
| | 10,000 UNITS/ML | N017033 | 003 | |
| | 20,000 UNITS/ML | N017033 | 004 | |
| BAXTER HLTHCARE | 1,000 UNITS/ML | N017036 | 001 | Mar 04, 1988 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HETACILLIN

FOR SUSPENSION;ORAL

VERSAPEN

| | | |
|---------|------------------------|-------------|
| BRISTOL | EQ 112.5MG AMPICIL/ML | A061398 001 |
| | EQ 112.5MG AMPICIL/5ML | N050060 001 |
| | EQ 112.5MG AMPICIL/ML | N050060 003 |
| | EQ 225MG AMPICIL/5ML | A061398 002 |

HETACILLIN POTASSIUM

CAPSULE;ORAL

VERSAPEN-K

| | | |
|---------|------------------|-------------|
| BRISTOL | EQ 225MG AMPICIL | A061396 001 |
| | EQ 450MG AMPICIL | A061396 002 |

HEXACHLOROPHENE

AEROSOL;TOPICAL

SEPTISOL

| | | |
|-------------|-------|-------------|
| VESTAL LABS | 0.23% | N017424 001 |
|-------------|-------|-------------|

TURGEX

| | | |
|----------|----|-------------|
| XTRTRIUM | 3% | N018375 001 |
|----------|----|-------------|

EMULSION;TOPICAL

HEXA-GERM

| | | |
|-----------------|----|-------------|
| HUNTINGTON LABS | 3% | N017411 001 |
|-----------------|----|-------------|

PHISOHEX

| | | |
|-------------------|----|-------------|
| SANOFI AVENTIS US | 3% | N006882 001 |
|-------------------|----|-------------|

| | | |
|--|----|-------------|
| | 3% | N008402 001 |
|--|----|-------------|

SOY-DOME

| | | |
|--------------|----|-------------|
| BAYER PHARMS | 3% | N017405 001 |
|--------------|----|-------------|

TURGEX

| | | |
|----------|----|-------------|
| XTRTRIUM | 3% | N019055 001 |
|----------|----|-------------|

SOAP;TOPICAL

GAMOPHEN

| | | |
|---------|----|-------------|
| ARBROOK | 2% | N006270 003 |
|---------|----|-------------|

SOLUTION;TOPICAL

DIAL

| | | |
|------|-------|-------------|
| DIAL | 0.25% | N017421 002 |
|------|-------|-------------|

GERMA-MEDICA

| | | |
|-----------------|----|-------------|
| HUNTINGTON LABS | 1% | N017412 001 |
|-----------------|----|-------------|

GERMA-MEDICA "MG"

| | | |
|-----------------|-------|-------------|
| HUNTINGTON LABS | 0.25% | N017412 002 |
|-----------------|-------|-------------|

SEPTI-SOFT

| | | |
|--------|-------|-------------|
| CALGON | 0.25% | N017460 001 |
|--------|-------|-------------|

SEPTISOL

| | | |
|-------------|-------|-------------|
| VESTAL LABS | 0.25% | N017423 001 |
|-------------|-------|-------------|

SPONGE;TOPICAL

E-Z SCRUB

| | | |
|------------------|-------|-------------|
| BECTON DICKINSON | 450MG | N017452 001 |
|------------------|-------|-------------|

HEXASCRUB

| | | |
|------------|----|-------------|
| PROF DSPLS | 3% | N018363 001 |
|------------|----|-------------|

PHISO-SCRUB

| | | |
|-------------------|----|-------------|
| SANOFI AVENTIS US | 3% | N017446 001 |
|-------------------|----|-------------|

SCRUBTEAM SURGICAL SPONGEBRUSH

| | | |
|----|-------|-------------|
| 3M | 330MG | N017413 001 |
|----|-------|-------------|

HEXAFLUORENIUM BROMIDE

INJECTABLE;INJECTION

MYLAXEN

| | | |
|---------------------|---------|-------------|
| MEDPOINTE PHARM HLC | 20MG/ML | N009789 003 |
|---------------------|---------|-------------|

HEXOCYLIUM METHYLSULFATE

TABLET;ORAL

TRAL

| | | |
|--------|------|-------------|
| ABBVIE | 25MG | N010599 001 |
|--------|------|-------------|

HEXYLCAINE HYDROCHLORIDE

SOLUTION;TOPICAL

CYCLAINE

| | | |
|-------|----|-------------|
| MERCK | 5% | N008472 001 |
|-------|----|-------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HISTAMINE PHOSPHATE

INJECTABLE; INJECTION

HISTAMINE PHOSPHATE

LILLY

EQ 0.1MG BASE/ML

N000734 003

EQ 0.2MG BASE/ML

N000734 002

EQ 1MG BASE/ML

N000734 001

HISTRELIN ACETATE

INJECTABLE; INJECTION

SUPPRELIN

SHIRE

EQ 0.2MG BASE/ML

N019836 001 Dec 24, 1991

EQ 0.5MG BASE/ML

N019836 002 Dec 24, 1991

EQ 1MG BASE/ML

N019836 003 Dec 24, 1991

HOMATROPINE METHYLBROMIDE

TABLET; ORAL

HOMAPIN-10

MISSION PHARMA

10MG

A086308 001

HOMAPIN-5

MISSION PHARMA

5MG

A086309 001

TABLET, CHEWABLE; ORAL

EQUIPIN

MISSION PHARMA

3MG

A086310 001

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

HYCODAN

+ GENUS LIFESCIENCES 1.5MG/5ML; 5MG/5ML **

N005213 002 Jul 26, 1988

HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE

BIO-PHARM INC

1.5MG/5ML; 5MG/5ML

A204765 001 Mar 06, 2017

IVAX SUB TEVA PHARMS

1.5MG/5ML; 5MG/5ML

A040285 001 Jul 19, 1999

HYDROPANE

HALSEY

1.5MG/5ML; 5MG/5ML

A088066 001 Jun 28, 1985

TABLET; ORAL

HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE

ACTAVIS ELIZABETH

1.5MG; 5MG

A040295 001 Dec 01, 2000

HYCODAN

+ GENUS LIFESCIENCES 1.5MG; 5MG **

N005213 001 Jul 26, 1988

HYALURONIDASE

INJECTABLE; INJECTION

HYDASE

AKORN INC

150 UNITS/ML

N021716 001 Oct 25, 2005

VITRASE

BAUSCH AND LOMB

6,200 UNITS/VIAL

N021640 001 May 05, 2004

WYDASE

BAXTER HLTHCARE

150 UNITS/ML **

N006343 002

150 UNITS/VIAL **

N006343 006

1,500 UNITS/VIAL **

N006343 005

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

APRESOLINE

+ NOVARTIS

20MG/ML **

N008303 003

HYDRALAZINE HYDROCHLORIDE

ABRAXIS PHARM

20MG/ML

A089532 001 Aug 11, 1987

SMITH AND NEPHEW

20MG/ML

A088518 001 Apr 20, 1984

SOLOPAK

20MG/ML

A088517 001 Aug 22, 1985

TEVA PARENTERAL

20MG/ML

A040373 001 Feb 23, 2000

TABLET; ORAL

APRESOLINE

+ NOVARTIS

10MG **

N008303 004

+

25MG **

N008303 001

+

50MG **

N008303 002

+

100MG **

N008303 005

DRALZINE

TEVA

25MG

A084301 001

HYDRALAZINE HYDROCHLORIDE

ACTAVIS ELIZABETH

25MG

A088560 001 Oct 04, 1984

50MG

A088649 001 Oct 18, 1984

ACTAVIS GRP PTC

10MG

A091679 001 Mar 04, 2013

25MG

A091679 002 Mar 04, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDRALAZINE HYDROCHLORIDE

TABLET; ORAL

HYDRALAZINE HYDROCHLORIDE

| | | | |
|----------------------|-------|-------------|--------------|
| | 50MG | A091679 003 | Mar 04, 2013 |
| | 100MG | A091679 004 | Mar 04, 2013 |
| ANDA REPOSITORY | 10MG | A089359 001 | Jul 25, 1986 |
| | 25MG | A089258 001 | May 05, 1986 |
| | 50MG | A089259 001 | May 05, 1986 |
| | 100MG | A088729 001 | Apr 11, 1985 |
| ASCOT | 25MG | A088310 001 | Dec 19, 1984 |
| | 50MG | A088311 001 | Dec 19, 1984 |
| CHARTWELL RX | 10MG | A088846 001 | Feb 26, 1985 |
| | 25MG | A088847 001 | Feb 26, 1985 |
| | 50MG | A088848 001 | Feb 26, 1985 |
| | 100MG | A088849 001 | Feb 26, 1985 |
| HALSEY | 10MG | A089218 001 | Jan 22, 1986 |
| | 25MG | A089130 001 | Jan 15, 1986 |
| | 50MG | A089222 001 | Jan 22, 1986 |
| | 100MG | A089178 001 | Jan 15, 1986 |
| HERITAGE PHARMS INC | 10MG | A040858 001 | Feb 26, 2010 |
| | 25MG | A040858 002 | Feb 26, 2010 |
| | 50MG | A040858 003 | Feb 26, 2010 |
| | 100MG | A040858 004 | Feb 26, 2010 |
| IMPAX LABS | 25MG | A084922 001 | |
| | 50MG | A084923 001 | |
| IVAX SUB TEVA PHARMS | 10MG | A084443 001 | |
| | 25MG | A084437 001 | |
| | 50MG | A084469 002 | |
| | 100MG | A084581 001 | |
| MUTUAL PHARM | 10MG | A088728 001 | Apr 11, 1985 |
| | 25MG | A084106 002 | |
| | 50MG | A084107 002 | |
| MYLAN | 10MG | A090413 001 | Dec 08, 2010 |
| | 25MG | A090413 002 | Dec 08, 2010 |
| | 50MG | A090413 003 | Dec 08, 2010 |
| | 100MG | A090413 004 | Dec 08, 2010 |
| PUREPAC PHARM | 25MG | A088177 001 | Jul 29, 1983 |
| | 50MG | A088178 001 | Aug 15, 1983 |
| QUANTUM PHARMICS | 10MG | A088671 001 | May 01, 1984 |
| | 25MG | A088657 001 | Jun 15, 1984 |
| | 50MG | A088652 001 | May 08, 1984 |
| | 100MG | A088686 001 | May 01, 1984 |
| SUPERPHARM | 10MG | A088787 001 | Aug 28, 1984 |
| | 25MG | A088788 001 | Aug 28, 1984 |
| | 50MG | A088789 001 | Aug 28, 1984 |
| UPSHER SMITH LABS | 10MG | A083241 001 | |
| | 25MG | A083560 001 | |
| | 50MG | A083561 001 | |
| UPSHER-SMITH LABS | 50MG | A085088 001 | |
| USL PHARMA | 25MG | A087780 001 | Mar 29, 1982 |
| | 50MG | A087751 001 | Mar 29, 1982 |
| VANGARD | 25MG | A087712 001 | |
| | 50MG | A087908 001 | May 07, 1982 |
| VITARINE | 25MG | A086088 001 | |
| WATSON LABS | 25MG | A084504 001 | |
| | 25MG | A085532 002 | May 24, 1982 |
| | 50MG | A084503 001 | |
| | 50MG | A085533 002 | May 25, 1982 |
| WEST WARD | 25MG | A088240 001 | May 27, 1983 |
| | 50MG | A088241 001 | May 27, 1983 |

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

APRESAZIDE

| | | | |
|------------|-------------|-------------|--------------|
| NOVARTIS | 25MG; 25MG | A084735 001 | |
| | 50MG; 50MG | A084810 001 | |
| | 100MG; 50MG | A084811 001 | |
| HYDRA-ZIDE | | | |
| PAR PHARM | 100MG; 50MG | A088961 001 | Oct 21, 1985 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDRALAZINE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

| | | | | |
|-------------|------------|---------|-----|--------------|
| SOLVAY | 25MG;25MG | A087608 | 001 | Feb 08, 1982 |
| | 50MG;50MG | A087213 | 001 | Feb 08, 1982 |
| | 100MG;50MG | A087609 | 001 | Feb 08, 1982 |
| SUPERPHARM | 25MG;25MG | A089200 | 001 | Feb 09, 1987 |
| | 50MG;50MG | A089201 | 001 | Feb 09, 1987 |
| WATSON LABS | 25MG;25MG | A085457 | 001 | Mar 04, 1982 |
| | 50MG;50MG | A085446 | 001 | Mar 04, 1982 |
| | 100MG;50MG | A085440 | 001 | Mar 04, 1982 |

HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 100/50

| | | | | |
|-------------|------------|---------|-----|--------------|
| IVAX PHARMS | 100MG;50MG | A088358 | 001 | Apr 10, 1984 |
|-------------|------------|---------|-----|--------------|

HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 25/25

| | | | | |
|-------------|-----------|---------|-----|--------------|
| IVAX PHARMS | 25MG;25MG | A088356 | 001 | Apr 10, 1984 |
|-------------|-----------|---------|-----|--------------|

HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 50/50

| | | | | |
|-------------|-----------|---------|-----|--------------|
| IVAX PHARMS | 50MG;50MG | A088357 | 001 | Apr 10, 1984 |
|-------------|-----------|---------|-----|--------------|

TABLET; ORAL

APRESOLINE-ESIDRIX

| | | | | |
|----------|-----------|---------|-----|--|
| NOVARTIS | 25MG;15MG | N012026 | 002 | |
|----------|-----------|---------|-----|--|

HYDRALAZINE AND HYDROCHLOROTHIAZIDE

| | | | | |
|-------------|-----------|---------|-----|--|
| WATSON LABS | 25MG;15MG | A085827 | 001 | |
|-------------|-----------|---------|-----|--|

HYDROCHLOROTHIAZIDE W/ HYDRALAZINE

| | | | | |
|-------------|-----------|---------|-----|--|
| WATSON LABS | 25MG;15MG | A085373 | 001 | |
|-------------|-----------|---------|-----|--|

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CAM-AP-ES

| | | | | |
|--------------|-----------------|---------|-----|--|
| CHARTWELL RX | 25MG;15MG;0.1MG | A084897 | 001 | |
|--------------|-----------------|---------|-----|--|

HYDRALAZINE HYDROCHLORIDE, HYDROCHLOROTHIAZIDE AND RESERPINE

| | | | | |
|----------------------|-----------------|---------|-----|--|
| IVAX SUB TEVA PHARMS | 25MG;15MG;0.1MG | A084291 | 001 | |
|----------------------|-----------------|---------|-----|--|

HYDRALAZINE HYDROCHLORIDE-HYDROCHLOROTHIAZIDE-RESERPINE

| | | | | |
|-------|-----------------|---------|-----|--|
| MYLAN | 25MG;15MG;0.1MG | A087085 | 001 | |
|-------|-----------------|---------|-----|--|

HYDRALAZINE, HYDROCHLOROTHIAZIDE W/ RESERPINE

| | | | | |
|-------------|-----------------|---------|-----|--|
| WATSON LABS | 25MG;15MG;0.1MG | A085771 | 001 | |
|-------------|-----------------|---------|-----|--|

HYDRAP-ES

| | | | | |
|--------|-----------------|---------|-----|--|
| SANDOZ | 25MG;15MG;0.1MG | A084876 | 001 | |
|--------|-----------------|---------|-----|--|

HYDROCHLOROTHIAZIDE W/ RESERPINE AND HYDRALAZINE

| | | | | |
|-------------|-----------------|---------|-----|--|
| WATSON LABS | 25MG;15MG;0.1MG | A083770 | 001 | |
|-------------|-----------------|---------|-----|--|

HYDROSERPINE PLUS (R-H-H)

| | | | | |
|----------------------|-----------------|---------|-----|--|
| IVAX SUB TEVA PHARMS | 25MG;15MG;0.1MG | A083877 | 001 | |
|----------------------|-----------------|---------|-----|--|

RESERPINE, HYDRALAZINE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

| | | | | |
|--------|-----------------|---------|-----|--------------|
| SOLVAY | 25MG;15MG;0.1MG | A088376 | 001 | Oct 28, 1983 |
|--------|-----------------|---------|-----|--------------|

| | | | | |
|----------------------|-----------------|---------|-----|--------------|
| SUN PHARM INDUSTRIES | 25MG;15MG;0.1MG | A088570 | 001 | Apr 10, 1984 |
|----------------------|-----------------|---------|-----|--------------|

| | | | | |
|-------------|-----------------|---------|-----|--|
| WATSON LABS | 25MG;15MG;0.1MG | A085549 | 001 | |
|-------------|-----------------|---------|-----|--|

| | | | | |
|--|-----------------|---------|-----|--|
| | 25MG;15MG;0.1MG | A087556 | 001 | |
|--|-----------------|---------|-----|--|

RESERPINE, HYDROCHLOROTHIAZIDE, AND HYDRALAZINE HYDROCHLORIDE

| | | | | |
|---------|-----------------|---------|-----|--------------|
| LEDERLE | 25MG;15MG;0.1MG | A087709 | 001 | May 13, 1982 |
|---------|-----------------|---------|-----|--------------|

SER-A-GEN

| | | | | |
|--------|-----------------|---------|-----|--|
| SOLVAY | 25MG;15MG;0.1MG | A087210 | 001 | |
|--------|-----------------|---------|-----|--|

SER-AP-ES

| | | | | |
|----------|-----------------|---------|-----|--|
| NOVARTIS | 25MG;15MG;0.1MG | N012193 | 005 | |
|----------|-----------------|---------|-----|--|

UNIPRES

| | | | | |
|--------|-----------------|---------|-----|--|
| SOLVAY | 25MG;15MG;0.1MG | A085893 | 001 | |
|--------|-----------------|---------|-----|--|

| | | | | |
|--|-----------------|---------|-----|--|
| | 25MG;15MG;0.1MG | A086298 | 001 | |
|--|-----------------|---------|-----|--|

HYDRALAZINE HYDROCHLORIDE; RESERPINE

TABLET; ORAL

DRALSERP

| | | | | |
|--------|------------|---------|-----|--|
| SANDOZ | 25MG;0.1MG | A084617 | 001 | |
|--------|------------|---------|-----|--|

SERPASIL-APRESOLINE

| | | | | |
|----------|------------|---------|-----|--|
| NOVARTIS | 25MG;0.1MG | N009296 | 004 | |
|----------|------------|---------|-----|--|

| | | | | |
|--|------------|---------|-----|--|
| | 50MG;0.2MG | N009296 | 002 | |
|--|------------|---------|-----|--|

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

| | | | | |
|--------|--------|---------|-----|--------------|
| APOTEX | 12.5MG | A078389 | 001 | May 16, 2008 |
|--------|--------|---------|-----|--------------|

| | | | | |
|-------------------|--------|---------|-----|--------------|
| HIKMA INTL PHARMS | 12.5MG | A077885 | 001 | Nov 26, 2007 |
|-------------------|--------|---------|-----|--------------|

| | | | | |
|----------------|--------|---------|-----|--------------|
| LANNETT CO INC | 12.5MG | A091662 | 001 | Jan 27, 2012 |
|----------------|--------|---------|-----|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE

SOLUTION; ORAL

HYDROCHLOROTHIAZIDE

| | | | | |
|--------------|----------|---------|-----|--------------|
| MORTON GROVE | 50MG/5ML | A089661 | 001 | Jun 20, 1988 |
| ROXANE | 50MG/5ML | A088587 | 001 | Jul 02, 1984 |

HYDROCHLOROTHIAZIDE INTENSOL

| | | | | |
|--------|----------|---------|-----|--------------|
| ROXANE | 100MG/ML | A088588 | 001 | Jul 02, 1984 |
|--------|----------|---------|-----|--------------|

TABLET; ORAL

ESIDRIX

| | | | | |
|----------|-------|---------|-----|--|
| NOVARTIS | 25MG | N011793 | 005 | |
| | 50MG | N011793 | 008 | |
| | 100MG | N011793 | 009 | |

HYDRO-D

| | | | | |
|--------|------|---------|-----|--|
| HALSEY | 25MG | A086504 | 001 | |
| | 50MG | A083891 | 002 | |

HYDROCHLOROTHIAZIDE

| | | | | |
|----------------------|--------|---------|-----|--------------|
| ABC HOLDING | 50MG | A085672 | 001 | |
| ACTAVIS ELIZABETH | 25MG | A085054 | 002 | |
| | 50MG | A085208 | 001 | |
| ALRA | 25MG | A086369 | 001 | |
| | 50MG | A083554 | 001 | |
| APOTEX | 25MG | A040774 | 001 | Oct 03, 2007 |
| | 50MG | A040774 | 002 | Oct 03, 2007 |
| ASCOT | 25MG | A087539 | 001 | Feb 03, 1982 |
| | 50MG | A087540 | 001 | Feb 03, 1982 |
| AUROLIFE PHARMA LLC | 25MG | A083899 | 001 | |
| | 50MG | A085219 | 001 | |
| BARR | 50MG | A084771 | 001 | |
| CHARTWELL RX | 25MG | A085683 | 001 | |
| | 50MG | A083965 | 001 | |
| DAVA PHARMS INC | 100MG | A087060 | 001 | |
| ELKINS SINN | 50MG | A085152 | 002 | |
| HEATHER | 50MG | A084135 | 001 | |
| HIKMA INTL PHARMS | 25MG | A084878 | 002 | Jul 12, 2006 |
| IMPAX LABS | 25MG | A084029 | 001 | |
| | 50MG | A083607 | 002 | |
| | 100MG | A085098 | 001 | |
| INWOOD LABS | 25MG | A084776 | 001 | |
| | 25MG | A085067 | 001 | |
| | 50MG | A084776 | 002 | |
| IVAX SUB TEVA PHARMS | 50MG | A084658 | 001 | |
| | 100MG | A085022 | 001 | |
| JUBILANT CADISTA | 25MG | A040809 | 001 | Sep 04, 2007 |
| | 50MG | A040809 | 002 | Sep 04, 2007 |
| LANNETT CO INC | 25MG | A084325 | 001 | |
| | 50MG | A084324 | 001 | |
| MAST MM | 25MG | A086192 | 001 | |
| | 50MG | A086192 | 002 | |
| MYLAN | 25MG | A084880 | 001 | |
| | 50MG | A085112 | 001 | |
| MYLAN PHARMS INC | 12.5MG | A040770 | 001 | Jan 23, 2007 |
| PVT FORM | 50MG | A086597 | 001 | |
| ROXANE | 25MG | A085004 | 001 | |
| | 50MG | A084536 | 002 | |
| | 50MG | A085005 | 001 | |
| SOLVAY | 25MG | A085323 | 001 | |
| SUN PHARM INDUSTRIES | 25MG | A083972 | 001 | |
| | 50MG | A083972 | 002 | |
| | 100MG | A083972 | 003 | |
| SUPERPHARM | 25MG | A088827 | 001 | Dec 28, 1984 |
| | 50MG | A088828 | 001 | Dec 28, 1984 |
| | 100MG | A088829 | 001 | Dec 28, 1984 |
| TEVA | 25MG | A088924 | 001 | Feb 07, 1985 |
| | 50MG | A088923 | 001 | Feb 07, 1985 |
| USL PHARMA | 25MG | A087827 | 001 | Apr 19, 1982 |
| | 50MG | A087752 | 001 | Apr 19, 1982 |
| VANGARD | 25MG | A087638 | 001 | |
| | 50MG | A087610 | 001 | |
| WARNER CHILCOTT | 25MG | A087586 | 001 | May 03, 1982 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE

TABLET; ORAL

HYDROCHLOROTHIAZIDE

| | | | |
|----------------------|----------|-------------|--------------|
| | 50MG | A087587 001 | May 03, 1982 |
| WATSON LABS | 25MG | A081189 001 | Jan 24, 1992 |
| | 25MG | A083458 001 | |
| | 25MG | A085232 002 | |
| | 50MG | A083456 001 | |
| | 50MG | A085233 001 | |
| | 50MG | A086087 001 | |
| | 50MG | A086594 001 | |
| | 100MG | A081190 001 | Jan 24, 1992 |
| | 100MG | A085099 001 | |
| | 100MG | A087002 001 | |
| WATSON LABS TEVA | 50MG | A083232 001 | |
| WEST WARD | 25MG | A084899 001 | |
| WHITEWORTH TOWN PLSN | 25MG | A083809 002 | |
| | 50MG | A083809 001 | |
| | 100MG | A085347 001 | |
| YAOPHARMA CO LTD | 25MG | A087565 001 | Mar 09, 1982 |
| | 50MG | A084912 001 | |
| HYDRODIURIL | | | |
| + MERCK | 25MG ** | N011835 003 | |
| + | 50MG ** | N011835 006 | |
| + | 100MG ** | N011835 007 | |
| ORETIC | | | |
| ABBVIE | 25MG | N011971 001 | |
| | 50MG | N011971 002 | |
| ZIDE | | | |
| SOLVAY | 50MG | A083925 001 | |

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

AVALIDE

| | | | |
|------------------------------------|-----------------|-------------|--------------|
| + SANOFI AVENTIS US | 12.5MG; 75MG ** | N020758 001 | Sep 30, 1997 |
| + | 25MG; 300MG ** | N020758 004 | Mar 15, 2005 |
| IRBESARTAN AND HYDROCHLOROTHIAZIDE | | | |
| ATLAS PHARMS LLC | 12.5MG; 150MG | A203036 001 | Jan 15, 2016 |
| | 12.5MG; 300MG | A203036 002 | Jan 15, 2016 |
| | 25MG; 300MG | A203036 003 | Jan 15, 2016 |
| MYLAN PHARMS INC | 12.5MG; 150MG | A077969 001 | Sep 27, 2012 |
| | 12.5MG; 300MG | A077969 002 | Sep 27, 2012 |
| | 25MG; 300MG | A077969 003 | Jul 20, 2016 |
| TEVA | 25MG; 300MG | A077369 003 | Mar 30, 2012 |
| WATSON LABS INC | 12.5MG; 150MG | A091539 001 | Oct 22, 2012 |
| | 12.5MG; 300MG | A091539 002 | Oct 22, 2012 |

HYDROCHLOROTHIAZIDE; LABETALOL HYDROCHLORIDE

TABLET; ORAL

NORMOZIDE

| | | | |
|-----------------|-------------|-------------|--------------|
| SCHERING | 25MG; 100MG | N019046 001 | Apr 06, 1987 |
| | 25MG; 200MG | N019046 002 | Apr 06, 1987 |
| | 25MG; 300MG | N019046 003 | Apr 06, 1987 |
| | 25MG; 400MG | N019046 004 | Apr 06, 1987 |
| TRANDATE HCT | | | |
| GLAXOSMITHKLINE | 25MG; 100MG | N019174 001 | Apr 10, 1987 |
| | 25MG; 200MG | N019174 002 | Apr 10, 1987 |
| | 25MG; 300MG | N019174 003 | Apr 10, 1987 |
| | 25MG; 400MG | N019174 004 | Apr 10, 1987 |

HYDROCHLOROTHIAZIDE; LISINAPRIL

TABLET; ORAL

LISINAPRIL AND HYDROCHLOROTHIAZIDE

| | | | |
|------------|--------------|-------------|--------------|
| APOTEX INC | 12.5MG; 10MG | A076674 001 | Oct 05, 2004 |
| | 12.5MG; 20MG | A076674 002 | Oct 05, 2004 |
| | 25MG; 20MG | A076674 003 | Oct 05, 2004 |
| SANDOZ | 12.5MG; 10MG | A075926 001 | Jul 01, 2002 |
| | 12.5MG; 20MG | A075926 002 | Jul 01, 2002 |
| | 25MG; 20MG | A075926 003 | Jul 01, 2002 |
| TEVA | 12.5MG; 10MG | A075869 001 | Jul 01, 2002 |
| | 12.5MG; 20MG | A075869 002 | Jul 01, 2002 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; LISINOPRIL

TABLET; ORAL

LISINOPRIL AND HYDROCHLOROTHIAZIDE

| | | | | |
|----------|----------------|---------|-----|--------------|
| | 25MG;20MG | A075869 | 003 | Jul 01, 2002 |
| PRINZIDE | | | | |
| + MERCK | 12.5MG;10MG ** | N019778 | 003 | Nov 18, 1993 |
| + | 12.5MG;20MG ** | N019778 | 001 | Feb 16, 1989 |
| + | 25MG;20MG ** | N019778 | 002 | Feb 16, 1989 |

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET; ORAL

HYZAAR

| | | | | |
|---------------------|-------------|---------|-----|--------------|
| + MERCK SHARP DOHME | 12.5MG;50MG | N020387 | 001 | Apr 28, 1995 |
| + | 25MG;100MG | N020387 | 002 | Nov 10, 1998 |

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE

| | | | | |
|-------------|--------------|---------|-----|--------------|
| APOTEX | 12.5MG;50MG | A090150 | 001 | Oct 06, 2010 |
| | 12.5MG;100MG | A090150 | 002 | Aug 11, 2010 |
| | 25MG;100MG | A090150 | 003 | Oct 06, 2010 |
| WATSON LABS | 12.5MG;50MG | A200180 | 001 | Jan 12, 2011 |
| | 12.5MG;100MG | A200180 | 002 | Jan 12, 2011 |
| | 25MG;100MG | A200180 | 003 | Jan 12, 2011 |

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

ALDORIL 15

| | | | | |
|-------|------------|---------|-----|--|
| MERCK | 15MG;250MG | N013402 | 001 | |
|-------|------------|---------|-----|--|

ALDORIL 25

| | | | | |
|-------|------------|---------|-----|--|
| MERCK | 25MG;250MG | N013402 | 002 | |
|-------|------------|---------|-----|--|

ALDORIL D30

| | | | | |
|-------|------------|---------|-----|--|
| MERCK | 30MG;500MG | N013402 | 003 | |
|-------|------------|---------|-----|--|

ALDORIL D50

| | | | | |
|-------|------------|---------|-----|--|
| MERCK | 50MG;500MG | N013402 | 004 | |
|-------|------------|---------|-----|--|

METHYLDOPA AND HYDROCHLOROTHIAZIDE

| | | | | |
|----------------------|------------|---------|-----|--------------|
| DAVA PHARMS INC | 15MG;250MG | A072507 | 001 | Jun 02, 1989 |
| | 25MG;250MG | A072508 | 001 | Jun 02, 1989 |
| | 30MG;500MG | A072509 | 001 | Jun 02, 1989 |
| | 50MG;500MG | A072510 | 001 | Jun 02, 1989 |
| IVAX SUB TEVA PHARMS | 15MG;250MG | A071458 | 001 | Mar 08, 1988 |
| | 25MG;250MG | A071459 | 001 | Mar 08, 1988 |
| | 30MG;500MG | A071460 | 001 | Mar 08, 1988 |
| | 50MG;500MG | A071461 | 001 | Mar 08, 1988 |
| PAR PHARM | 15MG;250MG | A070616 | 001 | Feb 02, 1987 |
| | 25MG;250MG | A070612 | 001 | Feb 02, 1987 |
| | 30MG;500MG | A070613 | 001 | Feb 02, 1987 |
| | 50MG;500MG | A070614 | 001 | Feb 02, 1987 |
| PARKE DAVIS | 15MG;250MG | A071897 | 001 | Nov 23, 1987 |
| | 25MG;250MG | A071898 | 001 | Nov 23, 1987 |
| | 30MG;500MG | A071899 | 001 | Nov 23, 1987 |
| | 50MG;500MG | A071900 | 001 | Nov 23, 1987 |
| PUREPAC PHARM | 15MG;250MG | A070853 | 001 | Oct 08, 1986 |
| | 25MG;250MG | A070688 | 001 | Apr 24, 1986 |
| | 30MG;500MG | A070854 | 001 | Oct 08, 1986 |
| | 50MG;500MG | A070689 | 001 | Apr 24, 1986 |
| SANDOZ | 15MG;250MG | A070829 | 001 | Mar 09, 1987 |
| | 25MG;250MG | A070830 | 001 | Mar 09, 1987 |
| TEVA | 15MG;250MG | A071819 | 001 | Apr 08, 1988 |
| | 25MG;250MG | A071820 | 001 | Apr 08, 1988 |
| | 30MG;500MG | A071821 | 001 | Apr 08, 1988 |
| | 50MG;500MG | A071822 | 001 | Apr 08, 1988 |
| WATSON LABS | 15MG;250MG | A070365 | 001 | Mar 19, 1986 |
| | 15MG;250MG | A070958 | 001 | Feb 06, 1989 |
| | 15MG;250MG | A071920 | 001 | Aug 29, 1988 |
| | 25MG;250MG | A070366 | 001 | Apr 16, 1986 |
| | 25MG;250MG | A070959 | 001 | Jan 19, 1989 |
| | 25MG;250MG | A071921 | 001 | Aug 29, 1988 |
| | 30MG;500MG | A070367 | 001 | Mar 19, 1986 |
| | 30MG;500MG | A071069 | 001 | Jan 19, 1989 |
| | 30MG;500MG | A071922 | 001 | Aug 29, 1988 |
| | 50MG;500MG | A070368 | 001 | Apr 16, 1986 |
| | 50MG;500MG | A070960 | 001 | Feb 06, 1989 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

METHYLDOPA AND HYDROCHLOROTHIAZIDE

| | | | | |
|------------------|-------------|---------|-----|--------------|
| | 50MG; 500MG | A071923 | 001 | Aug 29, 1988 |
| YAOPHARMA CO LTD | 15MG; 250MG | A070182 | 001 | Jan 15, 1986 |
| | 25MG; 250MG | A070183 | 001 | Jan 15, 1986 |
| | 30MG; 500MG | A070543 | 001 | Jan 15, 1986 |
| | 50MG; 500MG | A070544 | 001 | Jan 15, 1986 |

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET; ORAL

LOPRESSOR HCT

| | | | | |
|------------------------|----------------|---------|-----|--------------|
| + US PHARMS HOLDINGS I | 50MG; 100MG ** | N018303 | 003 | Dec 31, 1984 |
|------------------------|----------------|---------|-----|--------------|

HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

| | | | | |
|--------------|------------------|---------|-----|--------------|
| CHARTWELL RX | 12.5MG; 7.5MG | A090096 | 001 | Sep 25, 2008 |
| | 12.5MG; 15MG | A090096 | 002 | Sep 25, 2008 |
| | 25MG; 15MG | A090096 | 003 | Sep 25, 2008 |
| UNIRETIC | | | | |
| UCB INC | 12.5MG; 7.5MG ** | N020729 | 001 | Jun 27, 1997 |
| | 12.5MG; 15MG ** | N020729 | 003 | Feb 14, 2002 |
| | 25MG; 15MG ** | N020729 | 002 | Jun 27, 1997 |

HYDROCHLOROTHIAZIDE; PINDOLOL

TABLET; ORAL

VISKAZIDE

| | | | | |
|----------|------------|---------|-----|--------------|
| NOVARTIS | 25MG; 5MG | N018872 | 001 | Jul 22, 1987 |
| | 25MG; 10MG | N018872 | 002 | Jul 22, 1987 |

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

INDERIDE LA 120/50

| | | | | |
|--------------------|-------------|---------|-----|--------------|
| WYETH AYERST | 50MG; 120MG | N019059 | 002 | Jul 03, 1985 |
| INDERIDE LA 160/50 | | | | |
| WYETH AYERST | 50MG; 160MG | N019059 | 003 | Jul 03, 1985 |
| INDERIDE LA 80/50 | | | | |
| WYETH AYERST | 50MG; 80MG | N019059 | 001 | Jul 03, 1985 |

TABLET; ORAL

INDERIDE-40/25

| | | | | |
|--------------------|---------------|---------|-----|--|
| + WYETH PHARMS INC | 25MG; 40MG ** | N018031 | 001 | |
|--------------------|---------------|---------|-----|--|

INDERIDE-80/25

| | | | | |
|--------------------|---------------|---------|-----|--|
| + WYETH PHARMS INC | 25MG; 80MG ** | N018031 | 002 | |
|--------------------|---------------|---------|-----|--|

PROPRANOLOL HYDROCHLORIDE & HYDROCHLOROTHIAZIDE

| | | | | |
|---------------------|------------|---------|-----|--------------|
| DURAMED PHARMS BARR | 25MG; 40MG | A071126 | 001 | Mar 02, 1987 |
| | 25MG; 80MG | A071127 | 001 | Mar 02, 1987 |

PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

| | | | | |
|----------------------|------------|---------|-----|--------------|
| ACTAVIS ELIZABETH | 25MG; 40MG | A070851 | 001 | May 15, 1986 |
| | 25MG; 80MG | A070852 | 001 | May 15, 1986 |
| ANI PHARMS INC | 25MG; 40MG | A070704 | 001 | Oct 01, 1986 |
| | 25MG; 40MG | A072042 | 001 | Mar 14, 1988 |
| | 25MG; 80MG | A070705 | 001 | Oct 01, 1986 |
| | 25MG; 80MG | A072043 | 001 | Mar 14, 1988 |
| IVAX SUB TEVA PHARMS | 25MG; 40MG | A071552 | 001 | Dec 01, 1988 |
| | 25MG; 80MG | A071553 | 001 | Dec 01, 1988 |
| WARNER CHILCOTT | 25MG; 40MG | A071771 | 001 | Jan 26, 1988 |
| | 25MG; 80MG | A071772 | 001 | Jan 26, 1988 |
| WATSON LABS | 25MG; 40MG | A070301 | 001 | Apr 18, 1986 |
| | 25MG; 40MG | A071498 | 001 | Dec 18, 1991 |
| | 25MG; 80MG | A070305 | 001 | Apr 18, 1986 |
| | 25MG; 80MG | A071501 | 001 | Dec 18, 1991 |
| YAOPHARMA CO LTD | 25MG; 40MG | A071060 | 001 | Aug 26, 1987 |
| | 25MG; 80MG | A071061 | 001 | Aug 26, 1987 |

HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

| | | | | |
|--------------------|----------------------|---------|-----|--------------|
| SUN PHARM INDS LTD | 12.5MG; EQ 10MG BASE | A078211 | 001 | Mar 04, 2009 |
| | 12.5MG; EQ 20MG BASE | A078211 | 002 | Mar 04, 2009 |
| | 25MG; EQ 20MG BASE | A078211 | 003 | Mar 04, 2009 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

H.R.-50

| | | | | |
|--------------------------------------|--------------|---------|-----|--------------|
| WHITEWORTH TOWN PLSN | 50MG;0.125MG | A085338 | 001 | |
| HYDRO-RESERP | | | | |
| ABC HOLDING | 50MG;0.125MG | A084714 | 002 | Jun 29, 1982 |
| HYDRO-SERP "25" | | | | |
| SANDOZ | 25MG;0.125MG | A084827 | 001 | |
| HYDRO-SERP "50" | | | | |
| SANDOZ | 50MG;0.125MG | A085213 | 001 | |
| HYDROCHLOROTHIAZIDE W/ RESERPINE | | | | |
| IVAX SUB TEVA PHARMS | 25MG;0.1MG | A083572 | 001 | |
| | 25MG;0.125MG | A083571 | 001 | |
| | 50MG;0.1MG | A083568 | 001 | |
| | 50MG;0.125MG | A083573 | 001 | |
| PHARMERAL | 25MG;0.125MG | A085421 | 001 | |
| | 50MG;0.125MG | A085420 | 001 | |
| ROXANE | 50MG;0.125MG | A084603 | 001 | |
| WATSON LABS | 25MG;0.125MG | A084466 | 001 | |
| | 25MG;0.125MG | A085317 | 001 | |
| | 25MG;0.125MG | A086330 | 002 | |
| | 50MG;0.125MG | A083666 | 001 | |
| | 50MG;0.125MG | A084467 | 001 | |
| | 50MG;0.125MG | A086331 | 001 | |
| HYDROPRES 25 | | | | |
| MERCK | 25MG;0.125MG | N011958 | 002 | |
| HYDROPRES 50 | | | | |
| MERCK | 50MG;0.125MG | N011958 | 003 | |
| RESERPINE AND HYDROCHLOROTHIAZIDE | | | | |
| BARR | 25MG;0.125MG | A084580 | 001 | |
| | 50MG;0.125MG | A084579 | 001 | |
| SANDOZ | 50MG;0.125MG | A088200 | 001 | Jan 31, 1984 |
| RESERPINE AND HYDROCHLOROTHIAZIDE-50 | | | | |
| WEST WARD | 50MG;0.125MG | A088189 | 001 | May 10, 1984 |
| SERPASIL-ESIDRIX #1 | | | | |
| NOVARTIS | 25MG;0.1MG | N011878 | 003 | |
| SERPASIL-ESIDRIX #2 | | | | |
| NOVARTIS | 50MG;0.1MG | N011878 | 005 | |

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

TABLET; ORAL

SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE

| | | | | |
|---------------------------------------|-----------|---------|-----|--------------|
| ASCOT | 25MG;25MG | A088025 | 001 | Nov 23, 1984 |
| MUTUAL PHARM | 25MG;25MG | A087267 | 001 | |
| PUREPAC PHARM | 25MG;25MG | A087999 | 001 | Nov 06, 1985 |
| SUPERPHARM | 25MG;25MG | A089137 | 001 | Aug 26, 1985 |
| WATSON LABS | 25MG;25MG | A087398 | 001 | |
| YAOPHARMA CO LTD | 25MG;25MG | A086881 | 001 | |
| SPIRONOLACTONE W/ HYDROCHLOROTHIAZIDE | | | | |
| IVAX PHARMS | 25MG;25MG | A087004 | 002 | May 24, 1982 |
| LEDERLE | 25MG;25MG | A087511 | 001 | |
| PARKE DAVIS | 25MG;25MG | A087948 | 001 | Feb 22, 1983 |
| PUREPAC PHARM | 25MG;25MG | A088054 | 001 | Aug 18, 1983 |
| UPSHER SMITH | 25MG;25MG | A087553 | 001 | |
| USL PHARMA | 25MG;25MG | A087651 | 001 | |
| VANGARD | 25MG;25MG | A087655 | 001 | |
| WATSON LABS | 25MG;25MG | A085974 | 001 | |
| | 25MG;25MG | A086026 | 001 | |

HYDROCHLOROTHIAZIDE; TIMOLOL MALEATE

TABLET; ORAL

TIMOLIDE 10-25

| | | | | |
|-------|-----------|---------|-----|--|
| MERCK | 25MG;10MG | N018061 | 001 | |
|-------|-----------|---------|-----|--|

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

DYAZIDE

| | | | | |
|-------------------------------------|-------------|---------|-----|--------------|
| GLAXOSMITHKLINE LLC | 25MG;50MG | N016042 | 002 | |
| TRIAMTERENE AND HYDROCHLOROTHIAZIDE | | | | |
| ANI PHARMS INC | 25MG;37.5MG | A074970 | 001 | Jan 06, 1998 |
| NOVARTIS | 25MG;37.5MG | A074857 | 001 | Sep 09, 1997 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

VITARINE 25MG; 50MG

A071737 001 Feb 12, 1988

TABLET; ORAL

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

AM THERAP 50MG; 75MG

A072022 001 Apr 17, 1988

QUANTUM PHARMICS 50MG; 75MG

A071980 001 Apr 17, 1988

WATSON LABS 50MG; 75MG

A071969 001 Apr 17, 1988

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

VALSARTAN AND HYDROCHLOROTHIAZIDE

APOTEX INC 12.5MG; 80MG

A203026 001 Mar 21, 2013

12.5MG; 160MG

A203026 002 Mar 21, 2013

12.5MG; 320MG

A203026 003 Mar 21, 2013

25MG; 160MG

A203026 004 Mar 21, 2013

25MG; 320MG

A203026 005 Mar 21, 2013

WATSON LABS TEVA 12.5MG; 80MG

A091519 001 Mar 21, 2013

12.5MG; 160MG

A091519 002 Mar 21, 2013

12.5MG; 320MG

A091519 003 Mar 21, 2013

25MG; 160MG

A091519 004 Mar 21, 2013

25MG; 320MG

A091519 005 Mar 21, 2013

HYDROCODONE BITARTRATE

TABLET, EXTENDED RELEASE; ORAL

VANTRELA ER

+ TEVA BRANDED PHARM 15MG

N207975 001 Jan 17, 2017

+ 30MG

N207975 002 Jan 17, 2017

+ 45MG

N207975 003 Jan 17, 2017

+ 60MG

N207975 004 Jan 17, 2017

+ 90MG

N207975 005 Jan 17, 2017

HYDROCODONE BITARTRATE; IBUPROFEN

TABLET; ORAL

HYDROCODONE BITARTRATE AND IBUPROFEN

ACTAVIS LABS FL INC 5MG; 200MG

A077454 001 Jun 23, 2010

VICOPROFEN

+ ABBVIE 7.5MG; 200MG

N020716 001 Sep 23, 1997

HYDROCODONE BITARTRATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

SYRUP; ORAL

CODAMINE

ALPHARMA US PHARMS 5MG/5ML; 25MG/5ML

A075103 001 Sep 29, 2000

HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

TRIS PHARMA INC 5MG/5ML; 60MG/5ML

A203839 001 Oct 28, 2014

HYDROCORTAMATE HYDROCHLORIDE

OINTMENT; TOPICAL

MAGNACORT

PFIZER 0.5%

N010554 001

HYDROCORTISONE

AEROSOL; TOPICAL

AEROSEB-HC

ALLERGAN HERBERT 0.5%

A085805 001

CREAM; TOPICAL

CORT-DOME

BAYER PHARMS 0.5%

N009585 003

1%

N009585 001

DERMACORT

MONARCH PHARMS 1%

A083011 002

ELDECORT

VALEANT PHARM INTL 1%

A080459 001

2.5%

A084055 001

FLEXICORT

WESTWOOD SQUIBB 0.5%

A087136 003 Apr 08, 1982

1%

A087136 002 Apr 08, 1982

2.5%

A087136 001 Apr 08, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE

CREAM;TOPICAL

H-CORT

| | | | |
|---------------------|------|---------|-----|
| PHARM ASSOC | 0.5% | A086823 | 001 |
| HC #1 | | | |
| BAYER PHARMS | 0.5% | A080438 | 001 |
| HC #4 | | | |
| BAYER PHARMS | 1% | A080438 | 002 |
| HC (HYDROCORTISONE) | | | |
| C AND M PHARMA | 0.5% | A080482 | 003 |
| | 1% | A080482 | 004 |

HI-COR

| | | | |
|----------------|------|---------|-----|
| C AND M PHARMA | 2.5% | A080483 | 001 |
|----------------|------|---------|-----|

HYDROCORTISONE

| | | | | |
|----------------------|------|---------|-----|--------------|
| ALPHARMA US PHARMS | 2.5% | A089754 | 001 | Feb 01, 1989 |
| ALTANA | 0.5% | A080848 | 002 | |
| | 1% | A080848 | 003 | |
| AMBIX | 1% | A086080 | 001 | |
| | 2.5% | A086271 | 001 | |
| EVERYLIFE | 0.5% | A080452 | 001 | |
| | 1% | A080452 | 002 | |
| G AND W LABS | 1% | A084059 | 001 | |
| INGRAM PHARM | 0.5% | A080456 | 002 | |
| | 1% | A080456 | 003 | |
| IVAX PHARMS | 1% | A085733 | 001 | |
| NASKA | 1% | A089706 | 001 | Mar 10, 1988 |
| PERRIGO NEW YORK | 0.5% | A084970 | 002 | |
| | 1% | A085026 | 001 | |
| PHARMADERM | 1% | A088845 | 001 | Feb 27, 1986 |
| | 2.5% | A089413 | 001 | Dec 16, 1986 |
| PHARMAFAIR | 1% | A087838 | 001 | Jul 28, 1982 |
| STIEFEL | 1% | A086170 | 001 | |
| SYOSSET | 0.5% | A085527 | 001 | |
| TARO | 0.5% | A086154 | 001 | |
| TARO PHARM INDS LTD | 1% | A086155 | 001 | |
| TEVA | 0.5% | A080400 | 002 | |
| | 1% | A080400 | 003 | |
| | 1% | A085191 | 001 | |
| | 2.5% | A080400 | 004 | |
| TOPIDERM | 1% | A089273 | 001 | Feb 17, 1989 |
| USL PHARMA | 1% | A088027 | 001 | Sep 27, 1983 |
| | 2.5% | A088029 | 001 | Sep 27, 1983 |
| WHITEWORTH TOWN PLSN | 1% | A080496 | 002 | |

HYTONE

| | | | |
|--------------|---------|---------|-----|
| VALEANT INTL | 1% ** | A080472 | 003 |
| | 2.5% ** | A080472 | 004 |

NOGENIC HC

| | | | | |
|-------------|----|---------|-----|--------------|
| IVAX PHARMS | 1% | A087427 | 001 | Apr 04, 1988 |
|-------------|----|---------|-----|--------------|

NUTRACORT

| | | | |
|-----------|------|---------|-----|
| DOW PHARM | 0.5% | A080442 | 002 |
| | 1% | A080442 | 003 |

PENECORT

| | | | | |
|------------------|----|---------|-----|--------------|
| ALLERGAN HERBERT | 1% | A088216 | 001 | Jun 06, 1984 |
|------------------|----|---------|-----|--------------|

PROCTOCORT

| | | | |
|----------------|----|---------|-----|
| MONARCH PHARMS | 1% | A083011 | 001 |
|----------------|----|---------|-----|

SYNACORT

| | | | |
|---------|------|---------|-----|
| MEDICIS | 0.5% | A087459 | 001 |
| | 1% | A087458 | 001 |
| | 2.5% | A087457 | 001 |

GEL;TOPICAL

NUTRACORT

| | | | |
|-------------|----|---------|-----|
| HEALTHPOINT | 1% | A084698 | 001 |
|-------------|----|---------|-----|

PENECORT

| | | | | |
|------------------|----|---------|-----|--------------|
| ALLERGAN HERBERT | 1% | A088215 | 001 | Jun 06, 1984 |
|------------------|----|---------|-----|--------------|

INJECTABLE;INJECTION

CORTEF

| | | | |
|----------------------|---------|---------|-----|
| PHARMACIA AND UPJOHN | 50MG/ML | N009864 | 001 |
|----------------------|---------|---------|-----|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE

LOTION; TOPICAL

| | | | | |
|---------------------|---------|---------|-----|--------------|
| ACTICORT | | | | |
| BAKER NORTON | 1% | A086535 | 001 | |
| ALA-CORT | | | | |
| CROWN LABS | 1% | A083201 | 001 | |
| BALNEOL-HC | | | | |
| SOLVAY | 1% | A088041 | 001 | Dec 03, 1982 |
| BETA-HC | | | | |
| BETA DERMAC | 1% | A089495 | 001 | Jan 25, 1988 |
| CETACORT | | | | |
| DOW PHARM | 0.5% | A080426 | 002 | |
| | 1% | A080426 | 001 | |
| CORT-DOME | | | | |
| BAYER PHARMS | 0.5% | N009895 | 003 | |
| | 1% | N009895 | 001 | |
| DERMACORT | | | | |
| SOLVAY | 0.5% | A084573 | 002 | |
| | 1% | A086462 | 001 | |
| EPICORT | | | | |
| BLULINE | 0.5% | A083219 | 002 | |
| GLYCORT | | | | |
| HERAN | 1% | A087489 | 001 | Oct 03, 1983 |
| H-CORT | | | | |
| PHARM ASSOC | 0.5% | A086824 | 001 | |
| HYDROCORTISONE | | | | |
| ALPHARMA US PHARMS | 0.5% | A087317 | 001 | Jun 07, 1982 |
| | 1% | A087315 | 001 | Jun 07, 1982 |
| MERICON | 0.5% | A085282 | 001 | |
| | 1% | A085282 | 002 | Feb 26, 1987 |
| NASKA | 1% | A089705 | 001 | Apr 25, 1988 |
| PERRIGO NEW YORK | 0.5% | A085662 | 001 | |
| | 1% | A085663 | 001 | |
| TARO | 1% | A089024 | 001 | Feb 12, 1986 |
| HYTONE | | | | |
| VALEANT INTL | 1% ** | A080473 | 003 | |
| | 2.5% ** | A080473 | 004 | Nov 30, 1982 |
| NUTRACORT | | | | |
| DOW PHARM | 0.5% | A080443 | 002 | |
| | 1% | A080443 | 003 | |
| | 2.5% | A087644 | 001 | Aug 24, 1982 |
| STIE-CORT | | | | |
| PERRIGO CO | 1% | A089066 | 001 | Nov 25, 1985 |
| OINTMENT; TOPICAL | | | | |
| CORTRIL | | | | |
| PFIZER GLOBAL | 1% | N009176 | 001 | |
| | 2.5% | N009176 | 002 | |
| HC (HYDROCORTISONE) | | | | |
| C AND M PHARMA | 0.5% | A080481 | 001 | |
| | 1% | A080481 | 002 | |
| HYDROCORTISONE | | | | |
| ALTANA | 0.5% | A080489 | 002 | |
| | 1% | A080489 | 003 | |
| AMBIX | 1% | A086079 | 001 | |
| | 2.5% | A086272 | 001 | |
| NASKA | 1% | A089704 | 001 | Mar 10, 1988 |
| PERRIGO NEW YORK | 0.5% | A084969 | 003 | |
| | 1% | A085028 | 001 | |
| PHARMADERM | 1% | A088842 | 001 | Feb 09, 1987 |
| TARO | 0.5% | A086256 | 001 | |
| | 2.5% | A040310 | 001 | Dec 29, 2000 |
| USL PHARMA | 1% | A088061 | 001 | Sep 27, 1983 |
| | 2.5% | A088039 | 001 | Sep 27, 1983 |
| HYTONE | | | | |
| DERMIK LABS | 1% ** | A080474 | 003 | |
| | 2.5% ** | A080474 | 004 | |
| PENECORT | | | | |
| ALLERGAN HERBERT | 2.5% | A088217 | 001 | Jun 06, 1984 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE

POWDER; FOR RX COMPOUNDING

H-CORT

TORCH 100% A087834 001 Mar 29, 1982

HYDRO-RX

X GEN PHARMS 100% A085982 001

HYDROCORTISONE

PADDOCK LLC 100% A088082 001 Apr 08, 1983

SOLUTION; TOPICAL

PENECORT

ALLERGAN HERBERT 1% A088214 001 Jun 06, 1984

TEXACORT

MISSION PHARMA 1% A080425 001

TABLET; ORAL

CORTRIL

PFIZER 10MG N009127 005

20MG N009127 003

HYDROCORTISONE

BARR 20MG A083999 001

ELKINS SINN 20MG A080624 001

FERRANTE 10MG A080568 001

20MG A080568 002

IMPAX LABS 20MG A080781 001

INWOOD LABS 20MG A080732 001

LANNETT 20MG A085070 001

NEXGEN PHARMA INC 20MG A083140 001

PANRAY 10MG N009659 001

20MG N009659 002

PARKE DAVIS 20MG A084243 001

PUREPAC PHARM 10MG A084247 003 Aug 31, 1982

20MG A080395 001

20MG A084247 002

ROXANE 10MG A088539 001 Mar 21, 1984

SANDOZ 20MG A080642 002

WATSON LABS 20MG A080355 001

WHITEWORTH TOWN PLSN 10MG A080344 001

20MG A080344 002

HYDROCORTONE

MERCK 10MG N008506 007

20MG N008506 011

TABLET; VAGINAL

CORTRIL

PFIPHARMECS 10MG N009796 001

HYDROCORTISONE ACETATE

CREAM; TOPICAL

HEMSOL-HC

ABLE 1% A081274 001 Jun 19, 1992

HYDROCORTISONE ACETATE

CENCI 1% A080419 001 Jan 25, 1982

FERNDAL LABS 2.5% A040259 001 Jul 29, 1999

PARKE DAVIS 1% A089914 001 Jan 03, 1989

PUREPAC PHARM 0.5% A086050 001

1% A086052 001

MICORT-HC

SEBELA IRELAND LTD 2% A040398 001 Mar 29, 2002

INJECTABLE; INJECTION

CORTEF ACETATE

PHARMACIA AND UPJOHN 50MG/ML N009378 002

CORTRIL

PFIZER 25MG/ML N009164 001

HYDROCORTISONE ACETATE

AKORN 25MG/ML N009637 001

50MG/ML N009637 002

BEL MAR 25MG/ML A083739 001

50MG/ML A083739 002

WATSON LABS 25MG/ML A083128 001

25MG/ML A083759 001

50MG/ML A083759 002

50MG/ML A085214 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE ACETATE

INJECTABLE; INJECTION

HYDROCORTONE

MERCCK

25MG/ML

N008228 001

50MG/ML

N008228 004

LOTION; TOPICAL

DRICORT

INGRAM PHARM

0.5%

A086207 001

OINTMENT; OPHTHALMIC

HYDROCORTISONE ACETATE

FERA PHARMS

0.5%

A080828 001

OINTMENT; OPHTHALMIC, OTIC

HYDROCORTONE

MERCCK

1.5%

N009018 003

OINTMENT; TOPICAL

CORTEF ACETATE

PHARMACIA AND UPJOHN

1%

N008917 002

+

2.5% **

N008917 001

PASTE; TOPICAL

ORABASE HCA

COLGATE

0.5%

A083205 001

POWDER; FOR RX COMPOUNDING

HYDROCORTISONE ACETATE

X GEN PHARMS

100%

A085981 001

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-CORTEF

PHARMACIA AND UPJOHN

1%;EQ 3.5MG BASE/GM

A061049 001

2.5%;EQ 3.5MG BASE/GM

A061049 002

OINTMENT; OPHTHALMIC

NEO-CORTEF

PHARMACIA AND UPJOHN

0.5%;EQ 3.5MG BASE/GM

A060610 001

1.5%;EQ 3.5MG BASE/GM

A060610 002

OINTMENT; TOPICAL

NEO-CORTEF

PHARMACIA AND UPJOHN

0.5%;EQ 3.5MG BASE/GM

A060751 001

1%;EQ 3.5MG BASE/GM

A060751 002

2.5%;EQ 3.5MG BASE/GM

A060751 003

SUSPENSION/DROPS; OPHTHALMIC

COR-OTICIN

AKORN

1.5%;EQ 3.5MG BASE/ML

A060188 001

NEO-CORTEF

PHARMACIA AND UPJOHN

0.5%;EQ 3.5MG BASE/ML

A060612 002

1.5%;EQ 3.5MG BASE/ML

A060612 001

HYDROCORTISONE ACETATE; OXYTETRACYCLINE HYDROCHLORIDE

SUSPENSION; OPHTHALMIC

TERRA-CORTRIL

PFIZER

1.5%;EQ 5MG BASE/ML

A061016 001

HYDROCORTISONE ACETATE; PRAMOXINE HYDROCHLORIDE

AEROSOL, METERED; TOPICAL

HYDROCORTISONE ACETATE 1% AND PRAMOXINE HYDROCHLORIDE 1%

GENUS LIFESCIENCES

1%;1%

A089440 001 May 17, 1988

LOTION; TOPICAL

PRAMOSONE

FERNDALE LABS

0.5%;1%

A083213 002

HYDROCORTISONE ACETATE; UREA

CREAM; TOPICAL

CARMOL HC

FOUGERA PHARMS

1%;10%

A080505 001

HYDROCORTISONE BUTYRATE

CREAM; TOPICAL

LOCOID

YAMANOUCHI

0.1%

N018795 001 Jan 07, 1983

OINTMENT; TOPICAL

LOCOID

YAMANOUCHI

0.1%

N019106 001 Jul 03, 1984

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE BUTYRATESOLUTION; TOPICAL
LOCOID

YAMAOUCHI 0.1% N019819 001 Sep 15, 1988

HYDROCORTISONE CYPIONATESUSPENSION; ORAL
CORTEF

PHARMACIA AND UPJOHN EQ 10MG BASE/5ML N009900 001

HYDROCORTISONE SODIUM PHOSPHATEINJECTABLE; INJECTION
HYDROCORTONE

MERCK EQ 50MG BASE/ML N012052 001

HYDROCORTISONE SODIUM SUCCINATEINJECTABLE; INJECTION
A-HYDROCORT

| | | | | |
|---------|--------------------|---------|-----|--------------|
| ABBOTT | EQ 100MG BASE/VIAL | A085928 | 001 | |
| | EQ 100MG BASE/VIAL | A089577 | 001 | Apr 11, 1989 |
| | EQ 250MG BASE/VIAL | A089578 | 001 | Apr 11, 1989 |
| | EQ 500MG BASE/VIAL | A089579 | 001 | Apr 11, 1989 |
| | EQ 1GM BASE/VIAL | A089580 | 001 | Apr 11, 1989 |
| HOSPIRA | EQ 100MG BASE/VIAL | A040666 | 001 | Apr 06, 2006 |
| | EQ 100MG BASE/VIAL | A085929 | 001 | |
| | EQ 250MG BASE/VIAL | A085930 | 001 | |
| | EQ 500MG BASE/VIAL | A085931 | 001 | |
| | EQ 1GM BASE/VIAL | A085932 | 001 | |

HYDROCORTISONE SODIUM SUCCINATE

| | | | | |
|-----------------|--------------------|---------|-----|--------------|
| ABRAXIS PHARM | EQ 100MG BASE/VIAL | A088667 | 001 | Jun 08, 1984 |
| | EQ 100MG BASE/VIAL | A088712 | 001 | Jun 08, 1984 |
| | EQ 250MG BASE/VIAL | A088668 | 001 | Jun 08, 1984 |
| | EQ 500MG BASE/VIAL | A088669 | 001 | Jun 08, 1984 |
| | EQ 1GM BASE/VIAL | A088670 | 001 | Jun 08, 1984 |
| BAXTER HLTHCARE | EQ 100MG BASE/VIAL | A086619 | 001 | |
| | EQ 250MG BASE/VIAL | A087567 | 001 | |
| | EQ 500MG BASE/VIAL | A087568 | 001 | |
| | EQ 1GM BASE/VIAL | A087569 | 001 | |
| INTL MEDICATION | EQ 100MG BASE/VIAL | A087532 | 001 | Mar 19, 1982 |
| WATSON LABS | EQ 100MG BASE/VIAL | A084737 | 002 | |
| | EQ 100MG BASE/VIAL | A084738 | 001 | |
| | EQ 250MG BASE/VIAL | A084737 | 001 | |
| | EQ 500MG BASE/VIAL | A084747 | 001 | |
| | EQ 1GM BASE/VIAL | A084748 | 001 | |

HYDROCORTISONE VALERATE

CREAM; TOPICAL

HYDROCORTISONE VALERATE

G AND W LABS INC 0.2% A074489 001 Aug 12, 1998

WESTCORT

+ SUN PHARM INDS INC 0.2% ** N017950 001

OINTMENT; TOPICAL

HYDROCORTISONE VALERATE

FOUGERA PHARMS 0.2% A075085 001 Jul 31, 2001

WESTCORT

+ SUN PHARM INDS INC 0.2% ** N018726 001 Aug 08, 1983

HYDROCORTISONE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-CORT-DOME

BAYER PHARMS 0.5%;EQ 3.5MG BASE/GM N050237 006 Jun 05, 1984

1%;EQ 3.5MG BASE/GM N050237 005 Jun 05, 1984

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OTIC

CORTISPORIN

+ MONARCH PHARMS 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML N050479 001

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

AMRING PHARMS 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML A065216 001 Oct 31, 2005

NEOMYCIN SULFATE-POLYMYXIN B SULFATE-HYDROCORTISONE

PHARMAFAIR 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML A062394 001 Sep 29, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS;OTIC

OTOCORT

WATSON LABS 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML A060730 002

SUSPENSION/DROPS;OPHTHALMIC

CORTISPORIN

MONARCH PHARMS 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML N050169 001

NEOMYCIN SULFATE-POLYMYXIN B SULFATE-HYDROCORTISONE

PHARMAFAIR 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML A062623 001 Sep 24, 1985

SUSPENSION/DROPS;OTIC

NEOMYCIN SULFATE, POLYMYXIN B SULFATE & HYDROCORTISONE

PHARMAFAIR 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML A062617 001 Sep 18, 1985

OTICAIR

PHARMAFAIR 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML A062399 001 Nov 18, 1982

OTOBIONE

SCHERING 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML A061816 001

OTOCORT

ACTAVIS LABS FL INC 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML A062521 001 Jul 11, 1985

PEDIOTIC

MONARCH PHARMS 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML A062822 001 Sep 29, 1987

HYDROCORTISONE; POLYMYXIN B SULFATE

SOLUTION/DROPS;OTIC

OTOBiotic

SCHERING 5MG/ML;EQ 10,000 UNITS BASE/ML A062302 001

PYOCIDIN

FOREST LABS 5MG/ML;EQ 10,000 UNITS BASE/ML A061606 001

HYDROCORTISONE; TETRACYCLINE HYDROCHLORIDE

OINTMENT;OPHTHALMIC

ACHROMYCIN

LEDERLE 1.5%;1% N050272 001

HYDROCORTISONE; UREA

CREAM;TOPICAL

ALPHADERM

BIOGLAN 1%;10% A086008 001

CALMURID HC

PHARMACIA AND UPJOHN 1%;10% A083947 001

HYDROFLUMETHIAZIDE

TABLET;ORAL

DIUCARDIN

WYETH AYERST 50MG A083383 001

HYDROFLUMETHIAZIDE

PAR PHARM 50MG A088850 001 May 31, 1985

WATSON LABS 50MG A088031 001 Apr 06, 1983

50MG A088528 001 Aug 15, 1984

SALURON

+ SHIRE LLC 50MG N011949 001

HYDROFLUMETHIAZIDE; RESERPINE

TABLET;ORAL

HYDROFLUMETHIAZIDE AND RESERPINE

USL PHARMA 50MG;0.125MG A088195 001 Oct 26, 1983

WATSON LABS 25MG;0.125MG A088127 001 Mar 22, 1983

50MG;0.125MG A088110 001 Mar 22, 1983

RESERPINE AND HYDROFLUMETHIAZIDE

IVAX PHARMS 50MG;0.125MG A088932 001 Jan 11, 1985

PAR PHARM 50MG;0.125MG A088907 001 Sep 20, 1985

SALUTENSIN

SHIRE 50MG;0.125MG N012359 003

SALUTENSIN-DEMI

SHIRE 25MG;0.125MG N012359 004

HYDROMORPHONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

PALLADONE

PURDUE PHARMA LP 12MG N021044 001 Sep 24, 2004

16MG N021044 002 Sep 24, 2004

24MG N021044 003 Sep 24, 2004

32MG N021044 004 Sep 24, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION

DILAUDID

+ FRESENIUS KABI USA 4MG/ML N019034 005 Apr 30, 2009

DILAUDID-HP

+ FRESENIUS KABI USA 10MG/ML N019034 001 Jan 11, 1984
250MG/VIAL N019034 002 Aug 04, 1994

HYDROMORPHONE HYDROCHLORIDE

HOSPIRA 10MG/ML A074598 001 Jun 19, 1997

WATSON LABS 10MG/ML A074317 001 Aug 23, 1995

TABLET; ORAL

HYDROMORPHONE HYDROCHLORIDE

NESHER PHARMS 2MG A077311 001 Nov 09, 2005

4MG A077311 002 Nov 09, 2005

8MG A077311 003 Nov 09, 2005

TABLET, EXTENDED RELEASE; ORAL

HYDROMORPHONE HYDROCHLORIDE

ACTAVIS LABS FL INC 8MG A202144 001 May 12, 2014

12MG A202144 002 May 12, 2014

16MG A202144 003 May 12, 2014

32MG A202144 004 Jun 30, 2016

HYDROXOCOBALAMIN

INJECTABLE; INJECTION

ALPHAREDISOL

MERCK 1MG/ML A080778 001

CYANOKIT

SERB SA 2.5GM/VIAL (5GM/KIT) N022041 002 Dec 15, 2006

HYDROXOCOBALAMIN

ABRAXIS PHARM 1MG/ML A084921 001

WATSON LABS 1MG/ML A085528 001

HYDROXOMIN

BEL MAR 1MG/ML A084629 001

HYDROXYAMPHETAMINE HYDROBROMIDE

SOLUTION/DROPS; OPHTHALMIC

PAREDRIE

PHARMICS 1% N000004 004

HYDROXYCHLOROQUINE SULFATE

TABLET; ORAL

HYDROXYCHLOROQUINE SULFATE

SANDOZ 200MG A040150 001 Jan 27, 1996

HYDROXYPROGESTERONE CAPROATE

INJECTABLE; INJECTION

HYDROXYPROGESTERONE CAPROATE

AKORN 125MG/ML N018004 001

ALLERGAN SALES LLC 125MG/ML N017439 001

250MG/ML N017439 002

SOLUTION; INTRAMUSCULAR

DELALUTIN

+ BRISTOL MYERS SQUIBB 125MG/ML (125MG/ML) ** N010347 004

+ 125MG/ML (125MG/ML) ** N016911 001

+ 250MG/ML (250MG/ML) ** N010347 002

+ 250MG/ML (250MG/ML) ** N016911 002

HYDROXYSTILBAMIDINE ISETHIONATE

INJECTABLE; INJECTION

HYDROXYSTILBAMIDINE ISETHIONATE

SANOFI AVENTIS US 225MG/AMP N009166 001

HYDROXYUREA

CAPSULE; ORAL

HYDROXYUREA

BARR 250MG A075143 002 Sep 21, 2000

BARR LABS INC 250MG A075020 002 Jun 26, 2000

500MG A075020 001 Jul 30, 1998

ROXANE 500MG A074476 001 Aug 18, 1995

TABLET; ORAL

HYDROXYUREA

BARR 1GM A075734 001 Aug 29, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE

BAXTER HLTHCARE 50MG/ML

A085551 002

HYDROXYZINE HYDROCHLORIDE

ALTANA 25MG/ML

A087273 001 Apr 20, 1982

50MG/ML

A087273 002 Apr 20, 1982

BAXTER HLTHCARE 25MG/ML

A085551 001

FRESENIUS KABI USA 25MG/ML

A087329 001

25MG/ML

A088184 001 Mar 31, 1983

50MG/ML

A087329 002

50MG/ML

A088185 001 Mar 31, 1983

HOSPIRA 25MG/ML

A087416 001

50MG/ML

A086821 001

50MG/ML

A087546 001

PHARMAFAIR 25MG/ML

A088862 001 Feb 14, 1986

25MG/ML

A089106 001 Feb 14, 1986

50MG/ML

A088881 001 Feb 14, 1986

50MG/ML

A089107 001 Feb 14, 1986

SMITH AND NEPHEW 25MG/ML

A087592 001

SOLOPAK 25MG/ML

A086822 001

25MG/ML

A087591 001

50MG/ML

A087310 001

50MG/ML

A087593 001

50MG/ML

A087595 001

50MG/ML

A087596 001

WATSON LABS 25MG/ML

A085778 001

25MG/ML

A087274 001

50MG/ML

A085779 001

50MG/ML

A087274 002

WYETH AYERST 25MG/ML

A086258 001

50MG/ML

A086258 002

ORGATRAK

ORGANON USA INC 25MG/ML

A087014 001

50MG/ML

A087014 002

VISTARIL

+ PFIZER 25MG/ML **

N011111 001

+ 50MG/ML **

N011111 002

SYRUP; ORAL

ATARAX

ROERIG 10MG/5ML **

N010485 001

HYDROXYZINE HYDROCHLORIDE

ALPHARMA US PHARMS 10MG/5ML

A088785 001 Feb 03, 1988

KV PHARM 10MG/5ML

A087730 001 Jul 01, 1982

STI PHARMA LLC 10MG/5ML

A086880 001

TABLET; ORAL

ATARAX

+ PFIZER 10MG **

N010392 001

+ 25MG **

N010392 004

+ 50MG **

N010392 006

+ 100MG **

N010392 005

HYDROXYZINE HYDROCHLORIDE

ABLE 10MG

A040559 001 Jul 22, 2004

25MG

A040562 001 Jul 22, 2004

50MG

A040563 001 Jul 22, 2004

ACTAVIS ELIZABETH 10MG

A089071 001 Jul 22, 1986

25MG

A089072 001 Jul 22, 1986

50MG

A089073 001 Jul 22, 1986

AUROLIFE PHARMA LLC 10MG

A087871 002 Dec 20, 1982

25MG

A087871 003 Dec 20, 1982

50MG

A087871 001 Dec 20, 1982

HALSEY 10MG

A089366 001 May 02, 1988

25MG

A089117 001 May 02, 1988

50MG

A089396 001 May 02, 1988

IVAX PHARMS 10MG

A087216 001

25MG

A087410 001

50MG

A087411 001

KV PHARM 10MG

A087819 001 Jun 23, 1982

25MG

A087820 001 Jun 23, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROXYZINE HYDROCHLORIDE

TABLET;ORAL

HYDROXYZINE HYDROCHLORIDE

| | | | | |
|----------------------|-------|---------|-----|--------------|
| | 50MG | A087821 | 001 | Jun 23, 1982 |
| | 100MG | A087822 | 001 | Jun 23, 1982 |
| MUTUAL PHARM | 10MG | A088409 | 001 | Nov 15, 1983 |
| | 25MG | A087857 | 001 | Apr 18, 1983 |
| | 50MG | A087860 | 001 | Apr 18, 1983 |
| PLIVA | 100MG | A081054 | 001 | Sep 25, 1995 |
| PUREPAC PHARM | 10MG | A088120 | 001 | Sep 25, 1984 |
| | 25MG | A088121 | 001 | Sep 25, 1984 |
| | 50MG | A088122 | 001 | Sep 25, 1984 |
| QUANTUM PHARMICS | 10MG | A088540 | 001 | Oct 22, 1985 |
| | 25MG | A088551 | 001 | Oct 22, 1985 |
| | 50MG | A088529 | 001 | Oct 22, 1985 |
| SANDOZ | 10MG | A087246 | 002 | |
| | 25MG | A085247 | 001 | |
| | 50MG | A087245 | 001 | |
| SUN PHARM INDS INC | 10MG | A040899 | 001 | Jun 10, 2008 |
| | 25MG | A040899 | 002 | Jun 10, 2008 |
| | 50MG | A040899 | 003 | Jun 10, 2008 |
| SUN PHARM INDUSTRIES | 10MG | A089381 | 001 | May 19, 1986 |
| | 25MG | A089382 | 001 | May 19, 1986 |
| | 50MG | A089383 | 001 | May 19, 1986 |
| | 100MG | A087862 | 001 | Apr 18, 1983 |
| SUPERPHARM | 10MG | A088794 | 001 | Dec 05, 1984 |
| | 25MG | A088795 | 001 | Dec 05, 1984 |
| | 50MG | A088796 | 001 | Dec 05, 1984 |
| USL PHARMA | 10MG | A089121 | 001 | Mar 20, 1986 |
| | 25MG | A089122 | 001 | Mar 20, 1986 |
| | 50MG | A089123 | 001 | Mar 20, 1986 |
| VINTAGE | 10MG | A087602 | 001 | Jan 22, 1982 |
| | 25MG | A087603 | 001 | Jan 22, 1982 |
| | 50MG | A087604 | 001 | Jan 22, 1982 |
| WATSON LABS | 10MG | A081149 | 001 | Mar 18, 1994 |
| | 10MG | A086827 | 001 | |
| | 10MG | A088348 | 001 | Sep 15, 1983 |
| | 25MG | A081150 | 001 | Mar 18, 1994 |
| | 25MG | A086829 | 001 | |
| | 25MG | A088349 | 001 | Sep 15, 1983 |
| | 50MG | A081151 | 001 | Mar 18, 1994 |
| | 50MG | A086836 | 001 | |
| | 50MG | A088350 | 001 | Sep 15, 1983 |

HYDROXYZINE PAMOATE

CAPSULE;ORAL

HY-PAM "25"

TEVA

EQ 25MG HYDROCHLORIDE

A088713 001 Mar 04, 1985

HYDROXYZINE PAMOATE

DURAMED PHARMS BARR

EQ 25MG HYDROCHLORIDE

A088593 001 Feb 29, 1984

EQ 50MG HYDROCHLORIDE

A088594 001 Feb 29, 1984

EQ 100MG HYDROCHLORIDE

A088595 001 Feb 29, 1984

IVAX SUB TEVA PHARMS

EQ 25MG HYDROCHLORIDE

A087761 001 Mar 05, 1982

EQ 50MG HYDROCHLORIDE

A087760 001 Mar 05, 1982

PAR PHARM

EQ 25MG HYDROCHLORIDE

A087656 001 Jun 11, 1982

EQ 25MG HYDROCHLORIDE

A089145 001 Mar 17, 1986

EQ 50MG HYDROCHLORIDE

A087657 001 Jun 11, 1982

EQ 50MG HYDROCHLORIDE

A089146 001 Mar 17, 1986

EQ 100MG HYDROCHLORIDE

A087658 001 Jun 11, 1982

SANDOZ

EQ 25MG HYDROCHLORIDE

A081127 001 Jun 28, 1991

EQ 50MG HYDROCHLORIDE

A081128 001 Jun 28, 1991

EQ 100MG HYDROCHLORIDE

A081129 001 Jun 28, 1991

SUPERPHARM

EQ 25MG HYDROCHLORIDE

A089031 001 Jan 02, 1987

EQ 50MG HYDROCHLORIDE

A089032 001 Jan 02, 1987

EQ 100MG HYDROCHLORIDE

A089033 001 Jan 02, 1987

VANGARD

EQ 25MG HYDROCHLORIDE

A088392 001 Sep 19, 1983

EQ 50MG HYDROCHLORIDE

A088393 001 Sep 19, 1983

WATSON LABS

EQ 25MG HYDROCHLORIDE

A081165 001 Jul 31, 1991

EQ 25MG HYDROCHLORIDE

A086698 001

EQ 25MG HYDROCHLORIDE

A086840 001 Jul 01, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

HYDROXYZINE PAMOATE

CAPSULE;ORAL

HYDROXYZINE PAMOATE

| | | | |
|------------------------|---------|-----|--------------|
| EQ 50MG HYDROCHLORIDE | A086695 | 001 | |
| EQ 50MG HYDROCHLORIDE | A086705 | 001 | Jul 01, 1982 |
| EQ 50MG HYDROCHLORIDE | A087767 | 001 | Aug 16, 1982 |
| EQ 100MG HYDROCHLORIDE | A086697 | 001 | |
| EQ 100MG HYDROCHLORIDE | A086728 | 001 | Oct 05, 1982 |
| EQ 100MG HYDROCHLORIDE | A087790 | 001 | Aug 16, 1982 |

VISTARIL

PFIZER

EQ 100MG HYDROCHLORIDE ** N011459 006

SUSPENSION;ORAL

VISTARIL

PFIZER

EQ 25MG HYDROCHLORIDE/5ML N011795 001

IBANDRONATE SODIUM

TABLET;ORAL

BONIVA

+ HOFFMANN LA ROCHE

EQ 2.5MG BASE ** N021455 001 May 16, 2003

IBANDRONATE SODIUM

MYLAN PHARMS INC

EQ 150MG BASE A078995 001 Mar 19, 2012

IBUPROFEN

CAPSULE;ORAL

IBUPROFEN

CONTRACT PHARMACAL

200MG A074782 001 Jul 06, 1998

MIDOL

BAYER

200MG ** A070626 001 Sep 02, 1987

200MG ** A071002 001 Sep 02, 1987

SOLUTION;INTRAVENOUS

CALDOLOR

CUMBERLAND PHARMS

400MG/4ML (100MG/ML) N022348 001 Jun 11, 2009

SUSPENSION;ORAL

CHILDREN'S ADVIL

WYETH CONS

100MG/5ML N019833 002 Sep 19, 1989

IBU

ABBOTT

100MG/5ML N019784 001 Dec 18, 1989

MOTRIN

+ MCNEIL CONSUMER

100MG/5ML ** N019842 001 Sep 19, 1989

SUSPENSION/DROPS;ORAL

MOTRIN

MCNEIL

40MG/ML N020476 001 May 25, 1995

TABLET;ORAL

ACHES-N-PAIN

LEDERLE

200MG A071065 001 May 28, 1987

CAP-PROFEN

PERRIGO

200MG A072097 001 Dec 08, 1987

IBU

BASF

400MG A070083 001 Feb 22, 1985

400MG N018197 001

600MG A070088 001 Feb 08, 1985

600MG A070099 001 Mar 29, 1985

800MG A070745 001 Jul 23, 1986

IBU-TAB

ALRA

800MG A071965 001 Aug 11, 1988

IBUPRIN

PLIVA

200MG A071773 001 Jul 16, 1987

IBUPROFEN

ABBOTT

600MG A070556 001 Jun 14, 1985

800MG A071264 001 Jul 25, 1986

ANI PHARMS INC

200MG A071144 001 Jan 20, 1987

200MG A072901 001 Dec 19, 1991

200MG A072903 001 Dec 19, 1991

AUROLIFE PHARMA LLC

300MG A070736 002 Jun 12, 1986

400MG A070736 003 Jun 12, 1986

600MG A070736 001 Jun 12, 1986

800MG A071938 001 Jan 14, 1988

CONTRACT PHARMACAL

200MG A071265 001 Oct 15, 1986

200MG A071265 002 Sep 10, 1987

200MG A071735 001 Sep 10, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IBUPROFEN

TABLET; ORAL

IBUPROFEN

| | | | | |
|----------------------|-------|---------|-----|--------------|
| | 200MG | A073691 | 001 | Feb 25, 1994 |
| | 200MG | A074931 | 001 | Jul 20, 1998 |
| HALSEY | 200MG | A071027 | 001 | Sep 29, 1987 |
| | 300MG | A071028 | 001 | Mar 23, 1987 |
| | 400MG | A071029 | 001 | Mar 23, 1987 |
| | 600MG | A071030 | 001 | Mar 23, 1987 |
| | 800MG | A072137 | 001 | Feb 05, 1988 |
| IVAX SUB TEVA PHARMS | 200MG | A071154 | 001 | Oct 27, 1987 |
| | 200MG | A072040 | 001 | Apr 29, 1988 |
| | 400MG | A071145 | 001 | Sep 23, 1986 |
| | 600MG | A071146 | 001 | Sep 23, 1986 |
| | 800MG | A071769 | 001 | May 08, 1987 |
| J AND J CONSUMER INC | 400MG | A070081 | 001 | Jun 16, 1986 |
| LEDERLE | 400MG | A070629 | 001 | Sep 19, 1986 |
| | 600MG | A070630 | 001 | Sep 19, 1986 |
| LEINER | 300MG | A071266 | 001 | Oct 15, 1986 |
| LNK | 100MG | A076741 | 001 | Jun 17, 2004 |
| MCNEIL | 600MG | A070476 | 001 | Jun 16, 1986 |
| MYLAN | 200MG | A071870 | 001 | May 05, 1988 |
| | 600MG | A070057 | 001 | Sep 24, 1985 |
| | 800MG | A071999 | 001 | Dec 03, 1987 |
| MYLAN PHARMS INC | 400MG | A070045 | 001 | Sep 24, 1985 |
| NORTHSTAR HLTHCARE | 400MG | A078132 | 001 | Sep 10, 2007 |
| | 600MG | A078132 | 002 | Sep 10, 2007 |
| | 800MG | A078132 | 003 | Sep 10, 2007 |
| OHM LABS | 400MG | A070818 | 001 | Dec 26, 1985 |
| P AND L DEV LLC | 200MG | A070733 | 001 | Sep 19, 1986 |
| PAR PHARM | 200MG | A071575 | 001 | May 08, 1987 |
| | 300MG | A070328 | 001 | Aug 06, 1985 |
| | 400MG | A070329 | 001 | Aug 06, 1985 |
| | 600MG | A070330 | 001 | Aug 06, 1985 |
| | 800MG | A070986 | 001 | Jul 25, 1986 |
| PERRIGO | 200MG | A072098 | 001 | Dec 08, 1987 |
| PLIVA | 400MG | A071666 | 001 | Jun 18, 1987 |
| | 600MG | A071667 | 001 | Jun 18, 1987 |
| | 800MG | A071668 | 001 | Jun 18, 1987 |
| PUREPAC PHARM | 200MG | A071122 | 001 | Oct 03, 1986 |
| | 200MG | A071664 | 001 | Feb 03, 1987 |
| | 300MG | A071123 | 001 | Sep 19, 1986 |
| | 400MG | A071124 | 001 | Sep 19, 1986 |
| | 600MG | A071125 | 001 | Sep 19, 1986 |
| | 800MG | A071964 | 001 | Feb 01, 1988 |
| SANDOZ | 200MG | A071807 | 001 | Feb 25, 1988 |
| | 200MG | A074525 | 001 | Dec 15, 1995 |
| | 200MG | A074533 | 001 | Dec 15, 1995 |
| | 400MG | A072064 | 001 | Jan 14, 1988 |
| | 600MG | A072065 | 001 | Jan 14, 1988 |
| | 800MG | A072169 | 001 | Dec 11, 1987 |
| SUN PHARM INDUSTRIES | 200MG | A070493 | 001 | Dec 24, 1985 |
| | 200MG | A070908 | 001 | Sep 26, 1986 |
| | 200MG | A071462 | 001 | Oct 02, 1986 |
| | 400MG | A070079 | 001 | Jul 24, 1985 |
| | 600MG | A070080 | 001 | Jul 24, 1985 |
| | 800MG | A071448 | 001 | Feb 18, 1987 |
| SUPERPHARM | 600MG | A070709 | 001 | Apr 25, 1986 |
| TEVA | 200MG | A073141 | 001 | May 29, 1992 |
| | 400MG | A073343 | 001 | Jun 30, 1992 |
| | 600MG | A073344 | 001 | Jun 30, 1992 |
| | 800MG | A073345 | 001 | Jun 30, 1992 |
| VINTAGE PHARMS | 200MG | A072249 | 001 | Jan 10, 1989 |
| | 300MG | A071230 | 001 | Oct 22, 1986 |
| | 400MG | A071231 | 001 | Oct 22, 1986 |
| | 600MG | A071232 | 001 | Oct 22, 1986 |
| | 800MG | A072004 | 001 | Nov 18, 1987 |
| WATSON LABS | 200MG | A070435 | 001 | Mar 05, 1986 |
| | 200MG | A071765 | 001 | Sep 04, 1987 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IBUPROFEN

TABLET; ORAL

| | | | |
|------------------------|----------|-------------|--------------|
| IBUPROFEN | 200MG | A071905 001 | Mar 08, 1988 |
| | 300MG | A071338 001 | Dec 01, 1986 |
| | 400MG | A070038 001 | Sep 06, 1985 |
| | 400MG | A070436 001 | Aug 21, 1985 |
| | 600MG | A070041 001 | Sep 06, 1985 |
| | 600MG | A070437 001 | Aug 21, 1985 |
| | 800MG | A071547 001 | Jul 02, 1987 |
| | 800MG | A071911 001 | Oct 13, 1987 |
| IBUPROHM | | | |
| OHM LABS | 400MG | A070469 001 | Aug 29, 1985 |
| MEDIPREN | | | |
| MCNEIL | 200MG | A070475 001 | Feb 06, 1986 |
| | 200MG | A071215 001 | Jun 26, 1986 |
| MIDOL | | | |
| BAYER | 200MG | A070591 001 | Sep 02, 1987 |
| | 200MG | A071001 001 | Sep 02, 1987 |
| MOTRIN | | | |
| + MCNEIL CONSUMER | 300MG ** | N017463 003 | |
| + | 400MG ** | N017463 002 | |
| + | 600MG ** | N017463 004 | |
| + | 800MG ** | N017463 005 | May 22, 1985 |
| MCNEIL PED | 100MG | N020418 001 | Nov 16, 1994 |
| MOTRIN MIGRAINE PAIN | | | |
| J AND J CONSUMER INC | 200MG | N019012 004 | Feb 25, 2000 |
| NUPRIN | | | |
| BRISTOL MYERS | 200MG | A072035 001 | Feb 16, 1988 |
| | 200MG | A072036 001 | Feb 16, 1988 |
| J AND J CONSUMER INC | 200MG | N019012 001 | May 18, 1984 |
| | 200MG | N019012 002 | Jul 29, 1987 |
| RUFEN | | | |
| BASF | 600MG | N018197 002 | Mar 05, 1984 |
| TABLET, CHEWABLE; ORAL | | | |
| CHILDREN'S MOTRIN | | | |
| + J AND J CONSUMER INC | 50MG | N020601 001 | Nov 15, 1996 |
| IBUPROFEN | | | |
| PERRIGO | 50MG | A076359 001 | Jan 16, 2004 |
| JUNIOR STRENGTH MOTRIN | | | |
| + J AND J CONSUMER INC | 100MG | N020601 003 | Nov 15, 1996 |
| MOTRIN | | | |
| MCNEIL PED | 50MG | N020135 001 | Nov 16, 1994 |
| | 100MG | N020135 002 | Nov 16, 1994 |

IBUPROFEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

| | | | |
|---------------------------------------|---------------|-------------|--------------|
| COMBUNOX | | | |
| FOREST LABS | 400MG; 5MG ** | N021378 001 | Nov 26, 2004 |
| OXYCODONE HYDROCHLORIDE AND IBUPROFEN | | | |
| WATSON LABS | 400MG; 5MG | A078394 001 | Nov 26, 2007 |

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET; ORAL

| | | | |
|---------------------------------------------|-------------|-------------|--------------|
| IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE | | | |
| CONTRACT PHARMACAL | 200MG; 30MG | A075588 001 | Apr 08, 2002 |

IBUTILIDE FUMARATE

INJECTABLE; INJECTION

| | | | |
|---------------------|----------|-------------|--------------|
| IBUTILIDE FUMARATE | | | |
| MYLAN INSTITUTIONAL | 0.1MG/ML | A090924 001 | Jan 11, 2010 |

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

| | | | |
|--------------------------|-----------|-------------|--------------|
| IDAMYCIN | | | |
| PHARMACIA AND UPJOHN | 5MG/VIAL | N050661 002 | Sep 27, 1990 |
| | 10MG/VIAL | N050661 001 | Sep 27, 1990 |
| | 20MG/VIAL | N050661 003 | Apr 25, 1995 |
| IDARUBICIN HYDROCHLORIDE | | | |
| SANDOZ | 1MG/ML | A091293 001 | Mar 29, 2011 |
| TEVA PARENTERAL | 5MG/VIAL | A065037 003 | May 01, 2002 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDARUBICIN HYDROCHLORIDE

10MG/VIAL

A065037 002 May 01, 2002

20MG/VIAL

A065037 001 May 01, 2002

IDOXURIDINE

OINTMENT; OPHTHALMIC

STOXIL

GLAXOSMITHKLINE 0.5%

N015868 001

SOLUTION/DROPS; OPHTHALMIC

DENDRID

+ ALCON 0.1%

N014169 001

HERPLEX

ALLERGAN 0.1%

N013935 002

STOXIL

GLAXOSMITHKLINE 0.1%

N013934 001

IFOSFAMIDE

INJECTABLE; INJECTION

IFOSFAMIDE

FRESENIUS KABI USA 1GM/20ML (50MG/ML)

A090181 001 Sep 22, 2009

3GM/60ML (50MG/ML)

A090181 002 Sep 22, 2009

IFOSFAMIDE; MESNA

INJECTABLE; INJECTION

IFEX/MESNEX KIT

BAXTER HLTHCARE 1GM/VIAL; 100MG/ML

N019763 003 Oct 10, 1992

3GM/VIAL; 100MG/ML

N019763 004 Oct 10, 1992

INJECTABLE; INTRAVENOUS

IFOSFAMIDE/MESNA KIT

TEVA PHARMS USA 1GM/20ML; 1GM/10ML (50MG/ML; 100MG/ML)

A075874 001 Feb 26, 2002

3GM/60ML; 1GM/10ML (50MG/ML; 100MG/ML)

A075874 002 Feb 26, 2002

ILOPROST

SOLUTION; INHALATION

VENTAVIS

ACTELION PHARMS LTD 20MCG/2ML (10MCG/ML)

N021779 001 Dec 29, 2004

IMATINIB MESYLATE

CAPSULE; ORAL

GLEEVEC

+ NOVARTIS EQ 50MG BASE **

N021335 001 May 10, 2001

+ EQ 100MG BASE **

N021335 002 May 10, 2001

IMIPRAMINE HYDROCHLORIDE

CONCENTRATE; ORAL

IMIPRAMINE HYDROCHLORIDE

NOVARTIS 25MG/ML

A086765 001

INJECTABLE; INJECTION

TOFRANIL

NOVARTIS 12.5MG/ML

N011838 002

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

LEDERLE 10MG

A086269 001

25MG

A086267 001

50MG

A086268 001

OXFORD PHARMS 50MG

A040751 001 Feb 28, 2008

PAR PHARM 10MG

A089422 001 Jul 14, 1987

25MG

A089497 001 Jul 14, 1987

ROXANE 10MG

A083799 001

25MG

A083799 002

50MG

A083799 003

SANDOZ 10MG

A085200 001

25MG

A084869 002

50MG

A085133 001

TEVA 10MG

A083729 001

25MG

A083729 004

50MG

A083729 003

USL PHARMA 25MG

A087776 001 Feb 10, 1982

VANGARD 10MG

A088036 001 Nov 03, 1982

25MG

A087619 001 Feb 09, 1982

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

| | | | |
|-------------------|------|-------------|--------------|
| | 50MG | A087631 001 | Jan 04, 1982 |
| WATSON LABS | 10MG | A085220 001 | |
| | 10MG | A085875 001 | |
| | 25MG | A084252 002 | |
| | 25MG | A085878 001 | |
| | 50MG | A085221 001 | |
| | 50MG | A085877 001 | |
| WEST WARD | 25MG | A088222 001 | May 26, 1983 |
| | 50MG | A088223 001 | May 26, 1983 |
| JANIMINE | | | |
| ABBOTT | 10MG | N017895 001 | |
| | 25MG | N017895 002 | |
| | 50MG | N017895 003 | |
| PRAMINE | | | |
| ALRA | 10MG | A083827 001 | |
| | 25MG | A083827 002 | |
| | 50MG | A083827 003 | |
| PRESAMINE | | | |
| SANOFI AVENTIS US | 10MG | N011836 006 | |
| | 25MG | N011836 003 | |
| | 50MG | N011836 007 | |

IMIPRAMINE PAMOATE

CAPSULE; ORAL

IMIPRAMINE PAMOATE

| | | | |
|------------------|------------------------|---------------------------|--------------|
| MYLAN PHARMS INC | EQ 75MG HYDROCHLORIDE | A202338 001 | Jun 28, 2013 |
| | EQ 100MG HYDROCHLORIDE | A202338 002 | Jun 28, 2013 |
| | EQ 125MG HYDROCHLORIDE | A202338 003 | Jun 28, 2013 |
| | EQ 150MG HYDROCHLORIDE | A202338 004 | Jun 28, 2013 |
| TOFRANIL-PM | | | |
| + | SPECGX LLC | EQ 75MG HYDROCHLORIDE ** | N017090 001 |
| + | | EQ 100MG HYDROCHLORIDE ** | N017090 004 |
| + | | EQ 125MG HYDROCHLORIDE ** | N017090 003 |
| + | | EQ 150MG HYDROCHLORIDE ** | N017090 002 |

IMIQUIMOD

CREAM; TOPICAL

IMIQUIMOD

| | | | |
|------------------|----|-------------|--------------|
| G AND W LABS INC | 5% | A200481 001 | Apr 18, 2011 |
| STRIDES PHARMA | 5% | A202002 001 | Jun 24, 2014 |

INAMRINONE LACTATE

INJECTABLE; INJECTION

AMRINONE LACTATE

| | | | |
|----------------------|----------------|-------------|--------------|
| BAXTER HLTHCARE CORP | EQ 5MG BASE/ML | A075542 001 | May 10, 2000 |
| HOSPIRA | EQ 5MG BASE/ML | A074616 001 | Aug 03, 1998 |
| INOCOR | | | |
| SANOFI AVENTIS US | EQ 5MG BASE/ML | N018700 001 | Jul 31, 1984 |

INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

| | | | |
|------------------|-------------------|-------------|--------------|
| ANI PHARMS INC | 1.25MG | A074498 002 | Feb 12, 1998 |
| | 2.5MG | A074498 001 | Oct 31, 1996 |
| MYLAN PHARMS INC | 1.25MG | A075105 001 | Jul 23, 1998 |
| | 2.5MG | A075105 002 | Jul 23, 1998 |
| TEVA | 1.25MG | A074665 001 | Apr 04, 1997 |
| | 2.5MG | A074665 002 | Apr 04, 1997 |
| WATSON LABS | 1.25MG | A074585 001 | Sep 26, 1996 |
| | 2.5MG | A074585 002 | Sep 26, 1996 |
| YAOPHARMA CO LTD | 1.25MG | A074594 001 | May 23, 1996 |
| | 2.5MG | A074594 002 | May 23, 1996 |
| LOZOL | | | |
| + | SANOFI AVENTIS US | 1.25MG ** | N018538 002 |
| + | | 2.5MG ** | N018538 001 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INDECAINIDE HYDROCHLORIDETABLET, EXTENDED RELEASE;ORAL
DECABID

| | | | |
|-------|---------------|-------------|--------------|
| LILLY | EQ 50MG BASE | N019693 001 | Dec 29, 1989 |
| | EQ 75MG BASE | N019693 002 | Dec 29, 1989 |
| | EQ 100MG BASE | N019693 003 | Dec 29, 1989 |

INDINAVIR SULFATE

CAPSULE;ORAL

CRIXIVAN

| | | | |
|-------------------|---------------|-------------|--------------|
| MERCK SHARP DOHME | EQ 100MG BASE | N020685 006 | Apr 19, 2000 |
| | EQ 333MG BASE | N020685 005 | Dec 17, 1998 |

INDIUM IN-111 OXYQUINOLINE

INJECTABLE;INJECTION

INDIUM IN 111 OXYQUINOLINE

| | | | |
|----------|---------|-------------|--------------|
| BWXT ITG | 1mCi/ML | A202586 001 | Jul 25, 2018 |
|----------|---------|-------------|--------------|

INDOCYANINE GREEN

INJECTABLE;INJECTION

IC-GREEN

| | | | |
|-------|--------------|-------------|--|
| AKORN | 10MG/VIAL ** | N011525 003 | |
| | 40MG/VIAL ** | N011525 004 | |
| | 50MG/VIAL ** | N011525 002 | |

INDOMETHACIN

CAPSULE;ORAL

INDO-LEMMON

| | | | |
|------|------|-------------|--------------|
| TEVA | 25MG | A070266 001 | Nov 07, 1985 |
| | 50MG | A070267 001 | Nov 07, 1985 |

INDOCIN

| | | | |
|--------------------|---------|-------------|--|
| + IROKO PHARMS LLC | 25MG ** | N016059 001 | |
| + | 50MG ** | N016059 002 | |

INDOMETHACIN

| | | | |
|----------------------|------|-------------|--------------|
| ABLE | 25MG | A076666 001 | Dec 17, 2003 |
| | 50MG | A076666 002 | Dec 17, 2003 |
| CHARTWELL MOLECULES | 25MG | N018829 002 | Aug 06, 1984 |
| | 50MG | A070651 001 | Mar 05, 1986 |
| | 50MG | N018829 001 | Aug 06, 1984 |
| CYCLE PHARMS LTD | 25MG | A070353 001 | Jun 18, 1985 |
| | 50MG | A070354 001 | Jun 18, 1985 |
| DURAMED PHARMS BARR | 25MG | A070326 001 | Oct 18, 1985 |
| | 50MG | A070327 001 | Oct 18, 1985 |
| HALSEY | 25MG | A070782 001 | Jun 03, 1987 |
| | 50MG | A070635 001 | Jun 03, 1987 |
| IVAX SUB TEVA PHARMS | 25MG | N018730 001 | May 04, 1984 |
| | 50MG | N018730 002 | May 04, 1984 |
| MUTUAL PHARM | 25MG | A070067 001 | Oct 03, 1986 |
| | 50MG | A070068 001 | Oct 03, 1986 |
| MYLAN | 50MG | N018858 002 | Apr 20, 1984 |
| PARKE DAVIS | 25MG | N018806 001 | Nov 23, 1984 |
| | 50MG | N018806 002 | Nov 23, 1984 |
| PIONEER PHARMS | 25MG | A070813 001 | Aug 11, 1986 |
| | 50MG | A070592 001 | Aug 11, 1986 |
| PLIVA | 25MG | A071148 001 | Mar 18, 1987 |
| | 50MG | A071149 001 | Mar 18, 1987 |
| SUN PHARM INDUSTRIES | 25MG | A070900 002 | Feb 09, 1987 |
| | 50MG | A070900 001 | Feb 09, 1987 |
| SUPERPHARM | 25MG | A070487 001 | Oct 10, 1986 |
| | 50MG | A070488 001 | Oct 10, 1986 |
| TEVA | 25MG | A071342 001 | Apr 18, 1988 |
| | 50MG | A071343 001 | Apr 18, 1988 |
| WATSON LABS | 25MG | A070529 001 | Oct 18, 1985 |
| | 25MG | A070784 001 | Aug 20, 1986 |
| | 25MG | A072996 001 | Jul 31, 1991 |
| | 25MG | N018690 001 | Jul 31, 1984 |
| | 50MG | A070530 001 | Oct 18, 1985 |
| | 50MG | A070785 001 | Aug 20, 1986 |
| | 50MG | A071635 001 | May 18, 1987 |
| | 50MG | A072997 001 | Jul 31, 1991 |
| | 50MG | N018690 002 | Jul 31, 1984 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INDOMETHACIN

CAPSULE, EXTENDED RELEASE;ORAL

INDOCIN SR

+ IROKO PHARMS 75MG ** N018185 001 Feb 23, 1982

INDOMETHACIN

ABLE 75MG A076114 001 Feb 06, 2002

INWOOD LABS 75MG A072410 001 Mar 15, 1989

WATSON LABS INC 75MG A202572 001 Dec 09, 2013

SUPPOSITORY;RECTAL

INDOCIN

+ IROKO PHARMS 50MG ** N017814 001 Aug 13, 1984

SUSPENSION;ORAL

INDOMETHACIN

CYCLE PHARMS LTD 25MG/5ML A071412 001 Mar 18, 1987

INSULIN ASPART PROTAMINE RECOMBINANT; INSULIN ASPART RECOMBINANT

INJECTABLE;SUBCUTANEOUS

NOVOLOG MIX 50/50

NOVO NORDISK INC 50 UNITS/ML;50 UNITS/ML N021810 001 Aug 26, 2008

NOVOLOG MIX 70/30 PENFILL

NOVO NORDISK INC 210 UNITS/3ML;90 UNITS/3ML (70 UNITS/ML; 30 UNITS/ML) N021172 002 Nov 01, 2001

210 UNITS/3ML;90 UNITS/3ML (70 UNITS/ML; 30 UNITS/ML) N021172 003 Nov 01, 2001

INSULIN ASPART RECOMBINANT

INJECTABLE;SUBCUTANEOUS

NOVOLOG FLEXTOUCH

+ NOVO NORDISK INC 300 UNITS/3ML (100 UNITS/ML) N020986 005 Oct 31, 2013

NOVOLOG INNOLET

NOVO NORDISK INC 300 UNITS/3ML (100 UNITS/ML) N020986 004 Apr 23, 2004

INSULIN DETEMIR RECOMBINANT

INJECTABLE;SUBCUTANEOUS

LEVEMIR FLEXPEN

NOVO NORDISK INC 300 UNITS/3ML (100 UNITS/ML) N021536 002 Jun 16, 2005

LEVEMIR INNOLET

NOVO NORDISK INC 300 UNITS/3ML (100 UNITS/ML) N021536 003 Jun 16, 2005

LEVEMIR PENFILL

NOVO NORDISK INC 300 UNITS/3ML (100 UNITS/ML) N021536 004 Jun 16, 2005

INSULIN GLULISINE RECOMBINANT

INJECTABLE;INTRAVENOUS, SUBCUTANEOUS

APIDRA

+ SANOFI AVENTIS US 300 UNITS/3ML (100 UNITS/ML) N021629 002 Dec 20, 2005

INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT

INJECTABLE;INJECTION

HUMALOG MIX 50/50 PEN

LILLY 50 UNITS/ML;50 UNITS/ML N021018 003 Dec 22, 1999

HUMALOG MIX 75/25 PEN

LILLY 75 UNITS/ML;25 UNITS/ML N021017 003 Dec 22, 1999

INSULIN LISPRO RECOMBINANT

INJECTABLE;INJECTION

HUMALOG PEN

LILLY 100 UNITS/ML N020563 002 Aug 06, 1998

INSULIN PORK

INJECTABLE;INJECTION

ILETIN I

LILLY 500 UNITS/ML N017931 001

INSULIN

NOVO NORDISK INC 40 UNITS/ML N017926 001

REGULAR INSULIN

NOVO NORDISK INC 100 UNITS/ML N017926 003

INSULIN PURIFIED BEEF

INJECTABLE;INJECTION

REGULAR ILETIN II

LILLY 100 UNITS/ML N018478 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INSULIN PURIFIED PORK

INJECTABLE; INJECTION

Iletin II

LILLY 500 UNITS/ML N018344 002

REGULAR Iletin II (PORK)

LILLY 100 UNITS/ML N018344 001

REGULAR PURIFIED PORK INSULIN

NOVO NORDISK INC 100 UNITS/ML N018381 001

VELOSULIN

NOVO NORDISK INC 100 UNITS/ML N018193 001

INSULIN PURIFIED PORK; INSULIN SUSP ISOPHANE PURIFIED PORK

INJECTABLE; INJECTION

INSULIN NORDISK MIXTARD (PORK)

NOVO NORDISK INC 30 UNITS/ML; 70 UNITS/ML N018195 001

INSULIN RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN BR

LILLY 100 UNITS/ML N019529 001 Apr 28, 1986

VELOSULIN BR

NOVO NORDISK INC 100 UNITS/ML N021028 001 Jul 19, 1999

POWDER; INHALATION

EXUBERA

PFIZER 1MG/INH N021868 001 Jan 27, 2006

3MG/INH N021868 002 Jan 27, 2006

INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN 50/50

LILLY 50 UNITS/ML; 50 UNITS/ML N020100 001 Apr 29, 1992

INSULIN RECOMBINANT PURIFIED HUMAN

INJECTABLE; INJECTION

NOVOLIN R

NOVO NORDISK INC 100 UNITS/ML N018778 001 Aug 30, 1983

VELOSULIN BR HUMAN

NOVO NORDISK INC 100 UNITS/ML N019450 001 May 30, 1986

INSULIN RECOMBINANT PURIFIED HUMAN; INSULIN SUSP ISOPHANE SEMISYNTHETIC PURIFIED HUMAN

INJECTABLE; INJECTION

MIXTARD HUMAN 70/30

BAYER PHARMS 30 UNITS/ML; 70 UNITS/ML N019585 001 Mar 11, 1988

NOVOLIN 70/30

NOVO NORDISK INC 30 UNITS/ML; 70 UNITS/ML N019441 001 Jul 11, 1986

INSULIN SUSP ISOPHANE BEEF

INJECTABLE; INJECTION

NPH INSULIN

NOVO NORDISK INC 40 UNITS/ML N017929 001

100 UNITS/ML N017929 003

INSULIN SUSP ISOPHANE BEEF/PORK

INJECTABLE; INJECTION

NPH Iletin I (BEEF-PORK)

LILLY 40 UNITS/ML N017936 001

100 UNITS/ML N017936 002

INSULIN SUSP ISOPHANE PURIFIED BEEF

INJECTABLE; INJECTION

NPH Iletin II

LILLY 100 UNITS/ML N018479 001

INSULIN SUSP ISOPHANE PURIFIED PORK

INJECTABLE; INJECTION

INSULIN INSULATARD NPH NORDISK

NOVO NORDISK INC 100 UNITS/ML N018194 001

NPH Iletin II (PORK)

LILLY 100 UNITS/ML N018345 001

NPH PURIFIED PORK ISOPHANE INSULIN

NOVO NORDISK INC 100 UNITS/ML N018623 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INSULIN SUSP ISOPHANE SEMISYNTHETIC PURIFIED HUMAN

INJECTABLE; INJECTION

INSULATARD NPH HUMAN

NOVO NORDISK INC 100 UNITS/ML

N019449 001 May 30, 1986

NOVOLIN N

NOVO NORDISK INC 100 UNITS/ML

N019065 001 Jan 23, 1985

INSULIN SUSP PROTAMINE ZINC BEEF/PORK

INJECTABLE; INJECTION

PROTAMINE ZINC & ILETIN I (BEEF-PORK)

LILLY 40 UNITS/ML

N017932 001

100 UNITS/ML

N017932 002

INSULIN SUSP PROTAMINE ZINC PURIFIED BEEF

INJECTABLE; INJECTION

PROTAMINE ZINC AND ILETIN II

LILLY 100 UNITS/ML

N018476 001

PROTAMINE ZINC INSULIN

BRISTOL MYERS SQUIBB 40 UNITS/ML

N017928 001

100 UNITS/ML

N017928 003

INSULIN SUSP PROTAMINE ZINC PURIFIED PORK

INJECTABLE; INJECTION

PROTAMINE ZINC AND ILETIN II (PORK)

LILLY 100 UNITS/ML

N018346 001

INSULIN ZINC SUSP BEEF

INJECTABLE; INJECTION

LENTE INSULIN

NOVO NORDISK INC 40 UNITS/ML

N017998 001

100 UNITS/ML

N017998 003

INSULIN ZINC SUSP EXTENDED BEEF

INJECTABLE; INJECTION

ULTRALENTE INSULIN

NOVO NORDISK INC 100 UNITS/ML

N017997 003

INSULIN ZINC SUSP EXTENDED PURIFIED BEEF

INJECTABLE; INJECTION

ULTRALENTE

NOVO NORDISK INC 100 UNITS/ML

N018385 001

INSULIN ZINC SUSP EXTENDED RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN U

LILLY 40 UNITS/ML

N019571 001 Jun 10, 1987

100 UNITS/ML

N019571 002 Jun 10, 1987

INSULIN ZINC SUSP PROMPT BEEF

INJECTABLE; INJECTION

SEMILENTE INSULIN

NOVO NORDISK INC 100 UNITS/ML

N017996 003

INSULIN ZINC SUSP PROMPT PURIFIED PORK

INJECTABLE; INJECTION

SEMILENTE

NOVO NORDISK INC 100 UNITS/ML

N018382 001

INSULIN ZINC SUSP PURIFIED BEEF

INJECTABLE; INJECTION

LENTE ILETIN II

LILLY 100 UNITS/ML

N018477 001

INSULIN ZINC SUSP PURIFIED BEEF/PORK

INJECTABLE; INJECTION

LENTARD

NOVO NORDISK INC 100 UNITS/ML

N018384 001

INSULIN ZINC SUSP PURIFIED PORK

INJECTABLE; INJECTION

LENTE

NOVO NORDISK INC 100 UNITS/ML

N018383 001

LENTE ILETIN II (PORK)

LILLY 100 UNITS/ML

N018347 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

INSULIN ZINC SUSP RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN L

LILLY

100 UNITS/ML

N019377 002 Sep 30, 1985

NOVOLIN L

NOVO NORDISK INC

100 UNITS/ML

N019965 001 Jun 25, 1991

INSULIN ZINC SUSP SEMISYNTHETIC PURIFIED HUMAN

INJECTABLE; INJECTION

NOVOLIN L

NOVO NORDISK INC

100 UNITS/ML

N018777 001 Aug 30, 1983

INULIN

INJECTABLE; INJECTION

INULIN AND SODIUM CHLORIDE

ISO TEX

100MG/ML

N002282 001

INVERT SUGAR

INJECTABLE; INJECTION

TRAVERT 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE

10GM/100ML

N016717 001

IOBENGUANE SULFATE I-131

INJECTABLE; INJECTION

IOBENGUANE SULFATE I 131

PHARMALUCENCE

2.3mCi/ML

N020084 001 Mar 25, 1994

IO CETAMIC ACID

TABLET; ORAL

CHOLEBRINE

MALLINCKRODT

750MG

N017129 001

IODAMIDE MEGLUMINE

INJECTABLE; INJECTION

RENOVUE-65

BRACCO

65%

N017902 001

RENOVUE-DIP

BRACCO

24%

N017903 001

IODIPAMIDE MEGLUMINE

INJECTABLE; INJECTION

CHOLOGRAFIN MEGLUMINE

BRACCO

10.3%

N009321 007

+

52%

N009321 003

IODIPAMIDE SODIUM

INJECTABLE; INJECTION

CHOLOGRAFIN SODIUM

BRACCO

20%

N009321 001

IODIXANOL

INJECTABLE; INJECTION

VISIPAQUE 270

GE HEALTHCARE

55%

N020808 001 Aug 29, 1997

IODOHIPPURATE SODIUM I-123

INJECTABLE; INJECTION

NEPHROFLOW

GE HEALTHCARE

1mCi/ML

N018289 001 Dec 28, 1984

IODOHIPPURATE SODIUM I-131

INJECTABLE; INJECTION

HIPURAN I 131

MALLINCKRODT

0.25mCi/ML

N016666 001

HIPPUTOPE

BRACCO

1-2mCi/VIAL

N015419 002

IODOHIPPURATE SODIUM I 131

PHARMALUCENCE

0.2mCi/ML

N017313 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IODOXAMATE MEGLUMINE

INJECTABLE; INJECTION

CHOLOVUE

| | | |
|--------|-------|-------------|
| BRACCO | 9.9% | N018077 001 |
| | 40.3% | N018076 001 |

IOFETAMINE HYDROCHLORIDE I-123

INJECTABLE; INJECTION

SPECTAMINE

| | | | |
|-----|---------|-------------|--------------|
| IMP | 1mCi/ML | N019432 001 | Dec 24, 1987 |
|-----|---------|-------------|--------------|

IOHEXOL

INJECTABLE; INJECTION

OMNIPAQUE 210

| | | | |
|---------------|-------|-------------|--------------|
| GE HEALTHCARE | 45.3% | N018956 006 | Jun 30, 1989 |
|---------------|-------|-------------|--------------|

SOLUTION; INJECTION, ORAL, RECTAL

OMNIPAQUE 240

| | | | |
|---------------|-------|-------------|--------------|
| GE HEALTHCARE | 51.8% | N020608 001 | Oct 24, 1995 |
|---------------|-------|-------------|--------------|

SOLUTION; URETHRAL

OMNIPAQUE 70

| | | | |
|---------------|-------|-------------|--------------|
| GE HEALTHCARE | 15.1% | N018956 007 | Jun 01, 1994 |
|---------------|-------|-------------|--------------|

IOPAMIDOL

INJECTABLE; INJECTION

IOPAMIDOL

| | | | |
|-----------------|-----|-------------|--------------|
| BAXTER HLTHCARE | 41% | A074629 001 | Nov 06, 1996 |
|-----------------|-----|-------------|--------------|

51%

A074629 004

Mar 31, 1998

61%

A074629 002

Nov 06, 1996

76%

A074629 003

Nov 06, 1996

HOSPIRA

61%

A074734 001

Dec 10, 1996

76%

A074734 002

Dec 10, 1996

IOPAMIDOL-200

| | | | |
|--------------|-----|-------------|--------------|
| COOK IMAGING | 41% | A074881 001 | Jul 28, 2000 |
|--------------|-----|-------------|--------------|

| | | | |
|---------|-----|-------------|--------------|
| HOSPIRA | 41% | A074898 001 | Dec 30, 1997 |
|---------|-----|-------------|--------------|

IOPAMIDOL-200 IN PLASTIC CONTAINER

HOSPIRA

41%

A074636 001

Dec 30, 1997

IOPAMIDOL-250

| | | | |
|--------------|-----|-------------|--------------|
| COOK IMAGING | 51% | A074881 002 | Jul 28, 2000 |
|--------------|-----|-------------|--------------|

| | | | |
|--------------------|-----|-------------|--------------|
| FRESENIUS KABI USA | 51% | A074679 001 | Apr 02, 1997 |
|--------------------|-----|-------------|--------------|

| | | | |
|---------|-----|-------------|--------------|
| HOSPIRA | 51% | A074898 002 | Dec 30, 1997 |
|---------|-----|-------------|--------------|

| | | | |
|--|-----|-------------|--------------|
| | 51% | A075005 001 | Feb 24, 1998 |
|--|-----|-------------|--------------|

IOPAMIDOL-250 IN PLASTIC CONTAINER

HOSPIRA

51%

A074636 002

Dec 30, 1997

IOPAMIDOL-300

| | | | |
|--------|-----|-------------|--------------|
| ABBVIE | 61% | A074638 001 | Apr 30, 1997 |
|--------|-----|-------------|--------------|

| | | | |
|--------------|-----|-------------|--------------|
| COOK IMAGING | 61% | A074881 003 | Jul 28, 2000 |
|--------------|-----|-------------|--------------|

| | | | |
|--------------------|-----|-------------|--------------|
| FRESENIUS KABI USA | 61% | A074679 002 | Apr 02, 1997 |
|--------------------|-----|-------------|--------------|

| | | | |
|---------|-----|-------------|--------------|
| HOSPIRA | 61% | A074898 003 | Dec 30, 1997 |
|---------|-----|-------------|--------------|

| | | | |
|--|-----|-------------|--------------|
| | 61% | A075005 002 | Feb 24, 1998 |
|--|-----|-------------|--------------|

IOPAMIDOL-300 IN PLASTIC CONTAINER

HOSPIRA

61%

A074636 003

Dec 30, 1997

61%

A074637 001

Apr 03, 1997

IOPAMIDOL-370

| | | | |
|--------------|-----|-------------|--------------|
| COOK IMAGING | 76% | A074881 004 | Jul 28, 2000 |
|--------------|-----|-------------|--------------|

| | | | |
|--------------------|-----|-------------|--------------|
| FRESENIUS KABI USA | 76% | A074679 003 | Apr 02, 1997 |
|--------------------|-----|-------------|--------------|

| | | | |
|---------|-----|-------------|--------------|
| HOSPIRA | 76% | A074898 004 | Dec 30, 1997 |
|---------|-----|-------------|--------------|

| | | | |
|--|-----|-------------|--------------|
| | 76% | A075005 003 | Feb 24, 1998 |
|--|-----|-------------|--------------|

IOPAMIDOL-370 IN PLASTIC CONTAINER

HOSPIRA

76%

A074636 004

Dec 30, 1997

ISOVUE-128

| | | | |
|--------|-----|-------------|--------------|
| BRACCO | 26% | N018735 005 | Oct 21, 1986 |
|--------|-----|-------------|--------------|

ISOVUE-200

| | | | |
|--------|-----|-------------|--------------|
| BRACCO | 41% | N020327 001 | Oct 12, 1994 |
|--------|-----|-------------|--------------|

IOPANOIC ACID

TABLET; ORAL

TELEPAQUE

| | | |
|---------------|-------|-------------|
| GE HEALTHCARE | 500MG | N008032 001 |
|---------------|-------|-------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IOPHENDYLATE

INJECTABLE; INJECTION

PANTOPAQUE

ALCON

100%

N005319 001

IOPROMIDE

INJECTABLE; INJECTION

ULTRAVIST 150

+ BAYER HLTHCARE 31.2%

N020220 004 May 10, 1995

ULTRAVIST 300 IN PLASTIC CONTAINER

+ BAYER HLTHCARE 62.3%

N020220 005 Nov 18, 2008

IOTHALAMATE MEGLUMINE

INJECTABLE; INJECTION

CONRAY 30

+ LIEBEL-FLARSHEIM 30%

N016983 001

IOTHALAMATE MEGLUMINE; IOTHALAMATE SODIUM

INJECTABLE; INJECTION

VASCORAY

MALLINCKRODT

52%;26%

N016783 001

IOTHALAMATE SODIUM

INJECTABLE; INJECTION

ANGIO-CONRAY

MALLINCKRODT

80%

N013319 001

CONRAY 325

MALLINCKRODT

54.3%

N017685 001

CONRAY 400

MALLINCKRODT

66.8%

N014295 001

IOTROLAN

INJECTABLE; INTRATHECAL

OSMOVIST 190

BAYER HLTHCARE 40.6%

N019580 001 Dec 07, 1989

OSMOVIST 240

BAYER HLTHCARE 51.3%

N019580 002 Dec 07, 1989

IOVERSOL

INJECTABLE; INJECTION

OPTIRAY 160

LIEBEL-FLARSHEIM 34%

N019710 003 Dec 30, 1988

OPTIRAY 240

LIEBEL-FLARSHEIM 51%

N020923 001 May 28, 1998

IOXAGLATE MEGLUMINE; IOXAGLATE SODIUM

INJECTABLE; INJECTION

HEXABRIX

GUERBET

39.3%;19.6%

N018905 002 Jul 26, 1985

IOXILAN

INJECTABLE; INJECTION

OXILAN-300

GUERBET

62%

N020316 001 Dec 21, 1995

OXILAN-350

GUERBET

73%

N020316 002 Dec 21, 1995

IPODATE CALCIUM

GRANULE; ORAL

ORAGRAFIN CALCIUM

BRACCO

3GM/PACKET

N012968 001

IPODATE SODIUM

CAPSULE; ORAL

BILIVIST

BAYER HLTHCARE 500MG

A087768 001 Aug 11, 1982

ORAGRAFIN SODIUM

BRACCO

500MG

N012967 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

IPRATROPIUM BROMIDE

AEROSOL, METERED; INHALATION

ATROVENT

BOEHRINGER INGELHEIM 0.018MG/INH

N019085 001 Dec 29, 1986

SOLUTION; INHALATION

ATROVENT

+ BOEHRINGER INGELHEIM 0.02% **

N020228 001 Sep 29, 1993

IPRATROPIUM BROMIDE

ACTAVIS MID ATLANTIC 0.02%

A075111 001 Apr 22, 1999

APOTEX INC 0.02%

A075441 001 Mar 28, 2001

BAUSCH AND LOMB INC 0.02%

A075835 001 Oct 15, 2001

MYLAN SPECIALITY LP 0.02%

A074755 001 Jan 10, 1997

PHARMASCIENCE INC 0.02%

A075507 001 Jan 19, 2001

ROXANE 0.02%

A075867 001 Jul 22, 2002

TEVA PHARMS USA 0.02%

A075313 001 Feb 07, 2000

SPRAY, METERED; NASAL

ATROVENT

+ BOEHRINGER INGELHEIM 0.021MG/SPRAY

N020393 001 Oct 20, 1995

+ 0.042MG/SPRAY

N020394 001 Oct 20, 1995

IPRATROPIUM BROMIDE

APOTEX INC 0.021MG/SPRAY

A076156 001 Apr 18, 2003

IRBESARTAN

TABLET; ORAL

IRBESARTAN

AJANTA PHARMA LTD 75MG

A203685 001 Dec 10, 2015

150MG

A203685 002 Dec 10, 2015

300MG

A203685 003 Dec 10, 2015

APOTEX INC 75MG

A200832 001 Oct 15, 2012

150MG

A200832 002 Oct 15, 2012

300MG

A200832 003 Oct 15, 2012

MYLAN PHARMS INC 75MG

A200461 001 Sep 27, 2012

150MG

A200461 002 Sep 27, 2012

300MG

A200461 003 Sep 27, 2012

WATSON LABS INC 75MG

A090720 001 Oct 12, 2012

150MG

A090720 002 Oct 12, 2012

300MG

A090720 003 Oct 12, 2012

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

IRINOTECAN HYDROCHLORIDE

SANDOZ 40MG/2ML (20MG/ML)

A077994 001 Feb 27, 2008

100MG/5ML (20MG/ML)

A077994 002 Feb 27, 2008

SANDOZ INC 40MG/2ML (20MG/ML)

A090137 001 Nov 12, 2009

100MG/5ML (20MG/ML)

A090137 002 Nov 12, 2009

SUN PHARMA GLOBAL 40MG/2ML (20MG/ML)

A078805 001 Apr 21, 2008

100MG/5ML (20MG/ML)

A078805 002 Apr 21, 2008

IRON DEXTRAN

INJECTABLE; INJECTION

IRON DEXTRAN

SANOFI AVENTIS US EQ 50MG IRON/ML

N010787 002

IRON SUCROSE

INJECTABLE; INTRAVENOUS

VENOFER

LUITPOLD EQ 65MG BASE/3.25ML (EQ 20MG BASE/ML)

N021135 005 Mar 29, 2013

EQ 75MG BASE/3.75ML (EQ 20MG BASE/ML)

N021135 003 Mar 29, 2005

ISOETHARINE HYDROCHLORIDE

SOLUTION; INHALATION

BETA-2

NEPHRON 1%

A086711 001

BRONKOSOL

SANOFI AVENTIS US 0.25%

N012339 009

1%

N012339 008

ISOETHARINE HYDROCHLORIDE

ALPHARMA US PHARMS 1%

A087101 001

ASTRAZENECA 0.062%

A087937 001 Nov 15, 1982

0.062%

A089614 001 Jun 13, 1991

0.125%

A087938 001 Nov 15, 1982

0.125%

A089615 001 Jun 13, 1991

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOETHARINE HYDROCHLORIDE

SOLUTION; INHALATION

ISOETHARINE HYDROCHLORIDE

| | | | | |
|-------------------------------|--------|---------|-----|--------------|
| | 0.167% | A088470 | 001 | Mar 14, 1984 |
| | 0.167% | A089616 | 001 | Jun 13, 1991 |
| | 0.2% | A088471 | 001 | Mar 14, 1984 |
| | 0.2% | A089617 | 001 | Jun 13, 1991 |
| | 0.25% | A088472 | 001 | Mar 14, 1984 |
| | 0.25% | A089618 | 001 | Jun 13, 1991 |
| BAXTER HLTHCARE | 0.08% | A088144 | 001 | Jul 29, 1983 |
| | 0.14% | A088145 | 001 | Mar 26, 1984 |
| | 0.25% | A088146 | 001 | Aug 01, 1983 |
| DEY | 0.08% | A088187 | 001 | Dec 03, 1982 |
| | 0.1% | A087389 | 001 | |
| | 0.17% | A087390 | 001 | |
| | 0.25% | A088188 | 001 | Dec 03, 1982 |
| | 1% | A086763 | 001 | |
| INTL MEDICATION | 0.077% | A086651 | 001 | |
| | 0.08% | A086651 | 002 | |
| | 0.1% | A086651 | 003 | |
| | 0.143% | A086651 | 004 | |
| | 0.167% | A086651 | 005 | |
| | 0.2% | A086651 | 006 | |
| | 0.25% | A086651 | 007 | |
| | 1% | A086651 | 008 | |
| PARKE DAVIS | 0.5% | A085997 | 001 | |
| | 1% | A085889 | 001 | |
| ROXANE | 0.1% | A087396 | 001 | |
| | 0.125% | A087025 | 001 | |
| | 0.167% | A088226 | 001 | Sep 16, 1983 |
| | 0.2% | A087324 | 001 | |
| | 0.25% | A088275 | 001 | Jun 03, 1983 |
| | 1% | A086899 | 001 | |
| ISOETHARINE HYDROCHLORIDE S/F | | | | |
| DEY | 0.08% | A089817 | 001 | Nov 22, 1988 |
| | 0.1% | A089818 | 001 | Nov 22, 1988 |
| | 0.17% | A089819 | 001 | Nov 22, 1988 |
| | 0.25% | A089820 | 001 | Nov 22, 1988 |
| | 1% | A089252 | 001 | Sep 15, 1986 |

ISOETHARINE MESYLATE

AEROSOL, METERED; INHALATION

BRONKOMETER

| | | | | |
|----------------------|------------|---------|-----|--------------|
| SANOFI AVENTIS US | 0.34MG/INH | N012339 | 007 | |
| ISOETHARINE MESYLATE | | | | |
| ALPHARMA US PHARMS | 0.34MG/INH | A087858 | 001 | Aug 21, 1984 |

ISOFLURANE

LIQUID; INHALATION

ISOFLURANE

| | | | | |
|-----------------|-------|---------|-----|--------------|
| HOSPIRA | 99.9% | A074097 | 001 | Jan 25, 1993 |
| WATSON LABS INC | 99.9% | A074393 | 001 | May 12, 1995 |

ISOFLUROPHATE

OINTMENT; OPHTHALMIC

FLOROPRYL

| | | | | |
|-------|--------|---------|-----|--|
| MERCK | 0.025% | N010656 | 001 | |
|-------|--------|---------|-----|--|

ISONIAZID

INJECTABLE; INJECTION

NYDRAZID

| | | | | |
|--------|-------------|---------|-----|--|
| SANDOZ | 100MG/ML ** | N008662 | 001 | |
|--------|-------------|---------|-----|--|

RIMIFON

| | | | | |
|-------|----------|---------|-----|--|
| ROCHE | 25MG/ML | N008420 | 002 | |
| | 100MG/ML | N008420 | 003 | |

SYRUP; ORAL

ISONIAZID

| | | | | |
|--------|----------|---------|-----|--------------|
| MIKART | 50MG/5ML | A081118 | 001 | Jul 21, 1997 |
|--------|----------|---------|-----|--------------|

LANIAZID

| | | | | |
|---------|----------|---------|-----|--------------|
| LANNETT | 50MG/5ML | A089243 | 001 | Feb 03, 1986 |
|---------|----------|---------|-----|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISONIAZID

SYRUP;ORAL

RIMIFON

ROCHE 50MG/5ML N008420 001

TABLET;ORAL

DOW-ISONIAZID

DOW PHARM 300MG A080330 002

HYZYD

MEDPOINTE PHARM HLC 100MG A080134 003

300MG A080134 004

INH

NOVARTIS 300MG A080935 001

ISONIAZID

DURAMED PHARMS BARR 100MG A088231 001 Mar 17, 1983

300MG A088119 001 Mar 17, 1983

HALSEY 50MG A083632 001

HIKMA INTL PHARMS 100MG A080212 001

300MG A087425 001

IMPAX LABS 100MG A080153 001

IVAX SUB TEVA PHARMS 100MG A080270 001

300MG A083610 001

LILLY 100MG N008499 002

300MG N008499 003

MK LABS 100MG A080941 001

NEXGEN PHARMA INC 100MG A084050 001

PANRAY 50MG N008428 001

100MG N008428 002

300MG N008428 003

PERRIGO 100MG A083060 001

PHARMAVITE 100MG A085091 001

PHOENIX LABS NY 50MG A080368 001

100MG A080368 002

PUREPAC PHARM 50MG A080132 003 Jul 14, 1982

100MG A080132 004 Jul 14, 1982

SUN PHARM INDUSTRIES 100MG A080136 001

300MG A083633 001

WATSON LABS 50MG A080522 001

100MG A080401 001

100MG A080523 001

100MG A085790 001

300MG A080521 001

300MG A083178 001

300MG A085784 001

WHITEWORTH TOWN PLSN 100MG A080120 002

LANIAZID

LANNETT 50MG A080140 001

100MG A080140 002

NYDRAZID

BRISTOL MYERS SQUIBB 100MG N008392 003

STANOZIDE

EVERYLIFE 100MG A080126 001

300MG A080126 002

ISONIAZID; RIFAMPIN

CAPSULE;ORAL

RIFAMPIN AND ISONIAZID

HIKMA INTL PHARMS 150MG;300MG A065221 001 Jul 29, 2005

ISOPROPAMIDE IODIDE

TABLET;ORAL

DARBID

GLAXOSMITHKLINE EQ 5MG BASE N010744 001

ISOPROTERENOL HYDROCHLORIDE

AEROSOL, METERED; INHALATION

ISOPROTERENOL HYDROCHLORIDE

3M 0.12MG/INH N010375 004

ALPHARMA US PHARMS 0.12MG/INH A085904 001

ISUPREL

SANOFI AVENTIS US 0.103MG/INH N011178 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOPROTERENOL HYDROCHLORIDE

DISC; INHALATION

NORISODRINE AEROTROL

ABBOTT 0.25% N016814 001

INJECTABLE; INJECTION

ISOPROTERENOL HYDROCHLORIDE

ABRAXIS PHARM 0.2MG/ML A083431 001

BAXTER HLTHCARE 0.2MG/ML A083486 001

HOSPIRA 0.02MG/ML A083283 001

0.2MG/ML A083346 001

INTL MEDICATION 0.2MG/ML A083724 001

SOLUTION; INHALATION

AEROLONE

LILLY 0.25% N007245 001

ISOPROTERENOL HYDROCHLORIDE

ARMOUR PHARM 0.031% A087935 001 Nov 18, 1982

0.062% A087936 001 Nov 18, 1982

DEY 0.5% A086764 001 Jan 04, 1982

PARKE DAVIS 0.25% A085994 001

0.5% A085540 001

ISUPREL

SANOFI AVENTIS US 0.5% N006327 002

1% N006327 003

VAPO-ISO

FISONS 0.5% N016813 001

TABLET; RECTAL, SUBLINGUAL

ISUPREL

SANOFI AVENTIS US 10MG N006328 001

15MG N006328 002

ISOPROTERENOL HYDROCHLORIDE; PHENYLEPHRINE BITARTRATE

AEROSOL, METERED; INHALATION

DUO-MEDIAHALER

3M 0.16MG/INH; 0.24MG/INH N013296 001

ISOPROTERENOL SULFATE

AEROSOL, METERED; INHALATION

MEDIHALER-ISO

3M 0.08MG/INH N010375 003

POWDER; INHALATION

NORISODRINE

ABBVIE 10% N006905 003

25% N006905 002

ISOSORBIDE

SOLUTION; ORAL

ISMOTIC

ALCON 100GM/220ML N017063 001

ISOSORBIDE DINITRATE

CAPSULE, EXTENDED RELEASE; ORAL

ISORDIL

WYETH AYERST 40MG N012882 002 Jul 29, 1988

TABLET; ORAL

ISORDIL

+ VALEANT PHARMS NORTH 10MG ** N012093 002 Jul 29, 1988

+ 20MG ** N012093 006 Jul 29, 1988

+ 30MG ** N012093 005 Jul 29, 1988

ISOSORBIDE DINITRATE

SUN PHARM INDUSTRIES 5MG A086166 002 Sep 19, 1986

10MG A086169 001 Sep 19, 1986

20MG A086167 001 Sep 19, 1986

30MG A087564 001 Sep 18, 1986

SUPERPHARM 5MG A089190 001 Feb 17, 1987

10MG A089191 001 Feb 17, 1987

20MG A089192 001 Feb 17, 1987

WATSON LABS 5MG A086034 001 Jan 06, 1988

10MG A086032 001 Jan 07, 1988

SORBITRATE

ASTRAZENECA 5MG N016192 001 Apr 01, 1996

10MG N016192 002 Apr 01, 1996

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISOSORBIDE DINITRATETABLET; ORAL
SORBITRATE

| | | | |
|------|---------|-----|--------------|
| 20MG | A086405 | 002 | Aug 21, 1990 |
| 30MG | A088124 | 001 | Aug 21, 1990 |
| 40MG | A088125 | 001 | Aug 21, 1990 |

TABLET; SUBLINGUAL

ISORDIL

| | | | | |
|-----------|----------|---------|-----|--------------|
| + BIOVAIL | 2.5MG ** | N012940 | 004 | Jul 29, 1988 |
| + | 5MG ** | N012940 | 003 | Jul 29, 1988 |
| + | 10MG ** | N012940 | 005 | Jul 29, 1988 |

ISOSORBIDE DINITRATE

| | | | | |
|----------------------|----------|---------|-----|--------------|
| HIKMA INTL PHARMS | 2.5MG | A086054 | 001 | Oct 29, 1987 |
| | 5MG | A086055 | 001 | Nov 02, 1987 |
| SANDOZ | 2.5MG | A086225 | 001 | Feb 19, 1988 |
| | 5MG | A086222 | 001 | Feb 19, 1988 |
| SUN PHARM INDUSTRIES | 2.5MG | A084204 | 001 | Sep 18, 1986 |
| | 5MG | A086168 | 001 | Sep 18, 1986 |
| | 10MG | A087545 | 001 | Sep 18, 1986 |
| WATSON LABS | 2.5MG ** | A086033 | 001 | Feb 26, 1988 |
| WATSON LABS TEVA | 5MG ** | A086031 | 001 | Sep 29, 1987 |

SORBITRATE

| | | | | |
|-------------|-------|---------|-----|--------------|
| ASTRAZENECA | 2.5MG | N016191 | 002 | Apr 01, 1996 |
| | 5MG | N016191 | 001 | Apr 01, 1996 |

TABLET, CHEWABLE; ORAL

SORBITRATE

| | | | | |
|-------------|------|---------|-----|--------------|
| ASTRAZENECA | 5MG | N016776 | 002 | Apr 01, 1996 |
| | 10MG | N016776 | 003 | Apr 01, 1996 |

TABLET, EXTENDED RELEASE; ORAL

ISORDIL

| | | | | |
|--------------|------|---------|-----|--------------|
| WYETH AYERST | 40MG | N012882 | 001 | Jul 29, 1988 |
|--------------|------|---------|-----|--------------|

ISOSORBIDE DINITRATE

| | | | | |
|----------------|------|---------|-----|--------------|
| IMPAX LABS INC | 40MG | A040723 | 001 | Mar 17, 2008 |
|----------------|------|---------|-----|--------------|

ISOSORBIDE MONONITRATE

TABLET; ORAL

ISMO

| | | | | |
|----------------|------|---------|-----|--------------|
| PROMIUS PHARMA | 20MG | N019091 | 001 | Dec 30, 1991 |
|----------------|------|---------|-----|--------------|

TABLET, EXTENDED RELEASE; ORAL

IMDUR

| | | | | |
|-------------------|----------|---------|-----|--------------|
| + SCHERING PLOUGH | 30MG ** | N020225 | 001 | Aug 12, 1993 |
| + | 60MG ** | N020225 | 002 | Aug 12, 1993 |
| + | 120MG ** | N020225 | 003 | Mar 30, 1995 |

ISOSORBIDE MONONITRATE

| | | | | |
|----------------------|-------|---------|-----|--------------|
| ACCORD HLTHCARE | 30MG | A209684 | 001 | Oct 24, 2017 |
| | 60MG | A209684 | 002 | Oct 24, 2017 |
| | 120MG | A209684 | 003 | Oct 24, 2017 |
| ACTAVIS ELIZABETH | 30MG | A075306 | 001 | Dec 31, 1998 |
| | 60MG | A075306 | 002 | Dec 31, 1998 |
| ALKERMES GAINESVILLE | 60MG | A075041 | 001 | Sep 22, 1998 |
| IVAX SUB TEVA PHARMS | 30MG | A075448 | 002 | Aug 07, 2001 |
| | 60MG | A075448 | 001 | Jun 19, 2000 |
| | 120MG | A075448 | 003 | Aug 07, 2001 |
| SKYEPHARMA AG | 60MG | A075166 | 001 | Oct 07, 1999 |

ISOSULFAN BLUE

INJECTABLE; INJECTION

LYMPHAZURIN

| | | | | |
|------------|-------|---------|-----|--|
| + COVIDIEN | 1% ** | N018310 | 001 | |
|------------|-------|---------|-----|--|

ISOTRETINOIN

CAPSULE; ORAL

ACCUTANE

| | | | | |
|---------------------|---------|---------|-----|--------------|
| + HOFFMANN LA ROCHE | 10MG ** | N018662 | 002 | May 07, 1982 |
| + | 20MG ** | N018662 | 004 | Mar 28, 1983 |
| + | 40MG ** | N018662 | 003 | May 07, 1982 |

SOTRET

| | | | | |
|--------------------|------|---------|-----|--------------|
| SUN PHARM INDS LTD | 10MG | A076041 | 001 | Dec 24, 2002 |
| | 20MG | A076041 | 002 | Dec 24, 2002 |
| | 30MG | A076503 | 001 | Jun 20, 2003 |
| | 40MG | A076041 | 003 | Dec 24, 2002 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ISRADIPINE

CAPSULE; ORAL

DYNACIRC

+ SMITHKLINE BEECHAM 2.5MG

N019546 001 Dec 20, 1990

+ 5MG

N019546 002 Dec 20, 1990

TABLET, EXTENDED RELEASE; ORAL

DYNACIRC CR

+ GLAXOSMITHKLINE LLC 5MG **

N020336 001 Jun 01, 1994

+ 10MG **

N020336 002 Jun 01, 1994

ISRADIPINE

MYLAN PHARMS INC 5MG

A201067 001 Nov 27, 2015

10MG

A201067 002 Nov 27, 2015

ITRACONAZOLE

INJECTABLE; INJECTION

SPORANOX

JANSSEN PHARMS 10MG/ML

N020966 001 Mar 30, 1999

IVERMECTIN

TABLET; ORAL

STROMEKTOL

MERCK SHARP DOHME 6MG

N050742 001 Nov 22, 1996

KANAMYCIN SULFATE

CAPSULE; ORAL

KANTREX

APOTHECON EQ 500MG BASE

A060516 001

EQ 500MG BASE

A061911 001

EQ 500MG BASE

A062726 001 Mar 06, 1987

INJECTABLE; INJECTION

KANAMYCIN

WEST-WARD PHARMS INT EQ 75MG BASE/2ML

A062324 001

EQ 500MG BASE/2ML

A062324 002

EQ 1GM BASE/3ML

A062324 003

KANAMYCIN SULFATE

ABRAXIS PHARM EQ 75MG BASE/2ML

A062504 001 Apr 05, 1984

EQ 500MG BASE/2ML

A062504 002 Apr 05, 1984

EQ 1GM BASE/3ML

A062504 003 Apr 05, 1984

FRESENIUS KABI USA EQ 500MG BASE/2ML

A065111 001 Dec 17, 2002

EQ 1GM BASE/3ML

A065111 002 Dec 17, 2002

INTL MEDICATION EQ 500MG BASE/2ML

A062466 001 Sep 30, 1983

EQ 1GM BASE/3ML

A062466 002 Sep 30, 1983

LOCH EQ 75MG BASE/2ML

A063021 001 Jul 31, 1992

EQ 500MG BASE/2ML

A063022 001 Jul 31, 1992

EQ 1GM BASE/3ML

A063025 001 Jul 31, 1992

PHARMAFAIR EQ 75MG BASE/2ML

A062668 001 May 07, 1987

EQ 500MG BASE/2ML

A062672 001 May 07, 1987

EQ 1GM BASE/3ML

A062669 001 May 07, 1987

SOLOPAK EQ 75MG BASE/2ML

A062605 003 Feb 26, 1986

EQ 500MG BASE/2ML

A062605 001 Feb 26, 1986

EQ 1GM BASE/3ML

A062605 002 Feb 26, 1986

WARNER CHILCOTT EQ 1GM BASE/3ML

A063092 001 Oct 11, 1989

WATSON LABS EQ 1GM BASE/3ML

A062520 003 May 09, 1985

KANTREX

APOTHECON EQ 75MG BASE/2ML

A061655 003

EQ 75MG BASE/2ML

A061901 003

EQ 75MG BASE/2ML

A062564 001 Sep 21, 1984

EQ 500MG BASE/2ML

A061655 001

EQ 500MG BASE/2ML

A061901 001

EQ 500MG BASE/2ML

A062564 002 Sep 21, 1984

EQ 1GM BASE/3ML

A061655 002

EQ 1GM BASE/3ML

A061901 002

EQ 1GM BASE/3ML

A062564 003 Sep 21, 1984

KLEBCIL

KING PHARMS EQ 75MG BASE/2ML

A062170 001

EQ 500MG BASE/2ML

A062170 002

EQ 1GM BASE/3ML

A062170 003

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

KETOCONAZOLE

CREAM;TOPICAL

NIZORAL

+ JANSSEN PHARMA 2%

N019084 001 Dec 31, 1985

SUSPENSION;ORAL

NIZORAL

JANSSEN PHARMA 100MG/5ML

A070767 001 Nov 07, 1986

TABLET;ORAL

KETOCONAZOLE

AAIPHARMA LLC 200MG

A075341 001 Jul 27, 1999

APOTEX 200MG

A075912 001 Jan 10, 2002

PLIVA 200MG

A075362 001 Jun 15, 1999

SUN PHARM INDUSTRIES 200MG

A075314 001 Jun 15, 1999

TEVA 200MG

A074971 001 Jun 15, 1999

NIZORAL

+ JANSSEN PHARMS 200MG **

N018533 001

KETOPROFEN

CAPSULE;ORAL

KETOPROFEN

AUROLIFE PHARMA LLC 50MG

A074024 001 Dec 29, 1995

75MG

A074024 002 Dec 29, 1995

MYLAN 50MG

A074035 002 Dec 31, 1996

75MG

A074035 003 Dec 31, 1996

TEVA 25MG

A073515 001 Dec 22, 1992

ORUDIS

+ WYETH AYERST 25MG **

N018754 001 Jul 31, 1987

+ 50MG **

N018754 002 Jan 09, 1986

+ 75MG **

N018754 003 Jan 09, 1986

CAPSULE, EXTENDED RELEASE;ORAL

KETOPROFEN

ACTAVIS LABS FL INC 100MG

A075270 002 Mar 24, 1999

150MG

A075270 003 Mar 24, 1999

200MG

A075270 001 Mar 24, 1999

ALKERMES GAINESVILLE 200MG

A074879 001 Dec 10, 1997

MYLAN 100MG

A075679 003 Feb 20, 2002

150MG

A075679 002 Feb 20, 2002

ORUVAIL

+ WYETH PHARMS INC 100MG **

N019816 003 Feb 08, 1995

+ 150MG **

N019816 002 Feb 08, 1995

+ 200MG **

N019816 001 Sep 24, 1993

FILM;ORAL

NEXCEDE

NOVARTIS 12.5MG

N022470 001 Nov 25, 2009

TABLET;ORAL

ACTRON

BAYER 12.5MG

N020499 001 Oct 06, 1995

KETOPROFEN

PERRIGO 12.5MG

A075364 001 Feb 07, 2002

ORUDIS KT

+ WYETH CONS 12.5MG **

N020429 001 Oct 06, 1995

KETOROLAC TROMETHAMINE

INJECTABLE;INJECTION

KETOROLAC TROMETHAMINE

APOTEX INC 30MG/ML

A075626 001 Jul 24, 2001

30MG/ML

A077201 001 Oct 14, 2005

APOTHECON 15MG/ML

A075348 001 Nov 28, 2000

30MG/ML

A075348 002 Nov 28, 2000

BAXTER HLTHCARE CORP 15MG/ML

A075631 002 Jun 29, 2001

30MG/ML

A075631 001 Jun 29, 2001

BEDFORD 15MG/ML

A075230 002 Oct 25, 1999

30MG/ML

A075230 001 Oct 25, 1999

GLAND PHARMA LTD 15MG/ML

A076722 001 Jul 27, 2004

30MG/ML

A076722 002 Jul 27, 2004

HOSPIRA 15MG/ML

A074801 001 Jun 05, 1997

30MG/ML

A074801 002 Jun 05, 1997

LUITPOLD 15MG/ML

A078145 001 Jan 14, 2008

30MG/ML

A078145 002 Jan 14, 2008

MYLAN LABS LTD 15MG/ML

A078299 001 Jul 16, 2007

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

| | | | |
|----------------------------|------------|-------------|--------------|
| | 15MG/ML | A201155 001 | Aug 04, 2014 |
| | 30MG/ML | A078299 002 | Jul 16, 2007 |
| | 30MG/ML | A201155 002 | Aug 04, 2014 |
| SANDOZ INC | 15MG/ML | A076271 001 | Oct 06, 2004 |
| SUN PHARMA GLOBAL | 15MG/ML | A078737 001 | Oct 06, 2008 |
| | 30MG/ML | A078737 002 | Oct 06, 2008 |
| WEST-WARD PHARMS INT | 15MG/ML ** | A075222 001 | Apr 26, 1999 |
| | 15MG/ML | A075299 001 | Nov 03, 1999 |
| | 15MG/ML | A075772 001 | Jul 21, 2004 |
| | 30MG/ML ** | A075222 002 | Apr 26, 1999 |
| | 30MG/ML ** | A075228 001 | Apr 26, 1999 |
| | 30MG/ML | A075299 002 | Nov 03, 1999 |
| | 30MG/ML | A075772 002 | Jul 21, 2004 |
| WOCKHARDT | 30MG/ML | A077943 001 | Mar 27, 2007 |
| TORADOL | | | |
| + ROCHE PALO | 15MG/ML ** | N019698 001 | Nov 30, 1989 |
| + | 30MG/ML ** | N019698 002 | Nov 30, 1989 |
| SOLUTION/DROPS; OPHTHALMIC | | | |
| ACULAR PRESERVATIVE FREE | | | |
| ALLERGAN | 0.5% | N020811 001 | Nov 03, 1997 |
| KETOROLAC TROMETHAMINE | | | |
| AKORN | 0.45% | A203376 001 | Feb 10, 2014 |
| TABLET; ORAL | | | |
| KETOROLAC TROMETHAMINE | | | |
| CYCLE PHARMS LTD | 10MG | A074790 001 | Jun 26, 1997 |
| WATSON LABS | 10MG | A074955 001 | Sep 19, 1997 |
| TORADOL | | | |
| + ROCHE PALO | 10MG ** | N019645 001 | Dec 20, 1991 |

KETOTIFEN FUMARATE

SOLUTION/DROPS; OPHTHALMIC

KETOTIFEN FUMARATE

| | | | |
|----------------|-------------------|-------------|--------------|
| APOTEX INC | EQ 0.025% BASE | A077354 001 | May 09, 2006 |
| ZADITOR | | | |
| + ALCON PHARMA | EQ 0.025% BASE ** | N021066 002 | Oct 19, 2006 |

KRYPTON, KR-81M

GAS; INHALATION

MPI KRYPTON 81M GENERATOR

| | | | |
|---------------|-----|-------------|--|
| GE HEALTHCARE | N/A | N018088 001 | |
|---------------|-----|-------------|--|

LABETALOL HYDROCHLORIDE

INJECTABLE; INJECTION

LABETALOL HYDROCHLORIDE

| | | | |
|-------------------------|-----------|-------------|--------------|
| AKORN INC | 5MG/ML | A075524 001 | Nov 29, 1999 |
| APOTHECON | 5MG/ML | A075355 001 | Nov 29, 1999 |
| BAXTER HLTHCARE CORP | 5MG/ML | A076051 001 | Jul 05, 2002 |
| HOSPIRA | 5MG/ML | A075242 001 | Sep 30, 1999 |
| NORMODYNE | | | |
| SCHERING | 5MG/ML | N018686 001 | Aug 01, 1984 |
| TRANDATE | | | |
| + SEBELA IRELAND LTD | 5MG/ML ** | N019425 001 | Dec 31, 1985 |
| TABLET; ORAL | | | |
| LABETALOL HYDROCHLORIDE | | | |
| APOTHECON | 100MG | A075223 001 | Nov 20, 1998 |
| | 200MG | A075223 002 | Nov 20, 1998 |
| | 300MG | A075223 003 | Nov 20, 1998 |
| TEVA | 100MG | A074989 001 | Sep 30, 1998 |
| | 200MG | A074989 002 | Sep 30, 1998 |
| | 300MG | A074989 003 | Sep 30, 1998 |
| NORMODYNE | | | |
| + SCHERING | 100MG ** | N018687 001 | Aug 31, 1987 |
| + | 200MG ** | N018687 002 | Aug 01, 1984 |
| + | 300MG ** | N018687 003 | Aug 01, 1984 |
| + | 400MG ** | N018687 004 | Aug 01, 1984 |
| TRANDATE | | | |
| + CNTY LINE PHARMS | 400MG ** | N018716 004 | Aug 01, 1984 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LACTULOSE

SOLUTION; ORAL

CHRONULAC

+ SANOFI AVENTIS US 10GM/15ML **

N017884 001

CONSTULOSE

ACTAVIS MID ATLANTIC 10GM/15ML

A070288 001 Aug 15, 1988

DUPHALAC

SOLVAY 10GM/15ML

A072372 001 Mar 22, 1989

EVALOSE

TEVA PHARMS 10GM/15ML

A073497 001 May 28, 1993

LACTULOSE

APOTEX INC 10GM/15ML

A075911 001 Feb 21, 2002

MORTON GROVE 10GM/15ML

A071841 001 Sep 22, 1988

PACO 10GM/15ML

A073160 001 Aug 25, 1992

LAXILOSE

NOSTRUM LABS 10GM/15ML

A073686 001 May 28, 1993

SOLUTION; ORAL, RECTAL

ACILAC

NOSTRUM LABS 10GM/15ML

A073685 001 May 28, 1993

CEPHULAC

+ SANOFI AVENTIS US 10GM/15ML **

N017657 001

GENERLAC

MORTON GROVE 10GM/15ML

A071842 001 Sep 27, 1988

HEPTALAC

TEVA PHARMS 10GM/15ML

A073504 001 May 28, 1993

LACTULOSE

APOTEX INC 10GM/15ML

A076645 001 Jul 28, 2003

PACO 10GM/15ML

A072029 001 Aug 25, 1992

ROXANE 10GM/15ML

A073590 001 May 29, 1992

SOLVAY 10GM/15ML

N017906 001

PORTALAC

SOLVAY 10GM/15ML

A072374 001 Mar 22, 1989

LAMIVUDINE; NEVIRAPINE; ZIDOVUDINE

TABLET; ORAL

LAMIVUDINE, NEVIRAPINE AND ZIDOVUDINE

+ MICRO LABS 150MG; 200MG; 300MG

N205626 001 Aug 13, 2018

LAMIVUDINE; RALTEGRAVIR POTASSIUM

TABLET; ORAL

DUTREBIS

MERCK SHARP DOHME 150MG; EQ 300MG BASE

N206510 001 Feb 06, 2015

LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

LAMIVUDINE AND TENOFOVIR DISOPROXIL FUMARATE

AUROBINDO PHARMA LTD 300MG; 300MG

N022344 001 May 15, 2018

LAMIVUDINE; ZIDOVUDINE

TABLET; ORAL

LAMIVUDINE AND ZIDOVUDINE

PHARMACARE 150MG; 300MG

N022018 001 Mar 17, 2017

TEVA PHARMS 150MG; 300MG

A079081 001 May 25, 2011

LAMOTRIGINE

TABLET; ORAL

LAMICTAL

+ GLAXOSMITHKLINE LLC 50MG **

N020241 006 Dec 27, 1994

+ 250MG **

N020241 004 Dec 27, 1994

LAMOTRIGINE

ACTAVIS TOTOWA 25MG

A078669 001 Apr 08, 2011

100MG

A078669 002 Apr 08, 2011

150MG

A078669 003 Apr 08, 2011

200MG

A078669 004 Apr 08, 2011

HIKMA PHARMS 25MG

A078134 001 Apr 19, 2011

100MG

A078134 002 Apr 19, 2011

150MG

A078134 003 Apr 19, 2011

200MG

A078134 004 Apr 19, 2011

MYLAN 25MG

A077428 001 Jan 27, 2009

100MG

A077428 002 Jan 27, 2009

150MG

A077428 003 Jan 27, 2009

200MG

A077428 004 Jan 27, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LAMOTRIGINE

TABLET;ORAL

LAMOTRIGINE

| | | | |
|-------------------|-------|-------------|--------------|
| MYLAN LABS LTD | 25MG | A078443 001 | Feb 11, 2009 |
| | 100MG | A078443 002 | Feb 11, 2009 |
| | 150MG | A078443 003 | Feb 11, 2009 |
| | 200MG | A078443 004 | Feb 11, 2009 |
| PHARMASCIENCE INC | 25MG | A078310 001 | Feb 04, 2009 |
| | 100MG | A078310 002 | Feb 04, 2009 |
| | 150MG | A078310 003 | Feb 04, 2009 |
| | 200MG | A078310 004 | Feb 04, 2009 |
| ROXANE | 25MG | A077392 001 | Jan 27, 2009 |
| | 100MG | A077392 002 | Jan 27, 2009 |
| | 150MG | A077392 003 | Jan 27, 2009 |
| | 200MG | A077392 004 | Jan 27, 2009 |
| SANDOZ | 25MG | A078645 001 | Jan 27, 2009 |
| | 100MG | A078645 002 | Jan 27, 2009 |
| | 150MG | A078645 003 | Jan 27, 2009 |
| | 200MG | A078645 004 | Jan 27, 2009 |
| WOCKHARDT | 25MG | A078982 001 | Jan 27, 2009 |
| | 100MG | A078982 002 | Jan 27, 2009 |
| | 150MG | A078982 003 | Jan 27, 2009 |
| | 200MG | A078982 004 | Jan 27, 2009 |

TABLET, CHEWABLE;ORAL

LAMICTAL CD

| | | | |
|---------------------|-------|-------------|--------------|
| GLAXOSMITHKLINE LLC | 100MG | N020764 003 | Aug 24, 1998 |
| LAMOTRIGINE | | | |
| MYLAN | 5MG | A076630 001 | Jan 22, 2009 |
| | 25MG | A076630 002 | Jan 22, 2009 |
| SANDOZ | 5MG | A078409 002 | Jan 22, 2009 |
| | 25MG | A078409 003 | Jan 22, 2009 |

TABLET, EXTENDED RELEASE;ORAL

LAMOTRIGINE

| | | | |
|------------------|------|-------------|--------------|
| HANDA PHARMS LLC | 25MG | A202887 001 | Jun 17, 2013 |
| | 50MG | A202887 002 | Jun 17, 2013 |

LANSOPRAZOLE

FOR SUSPENSION, DELAYED RELEASE;ORAL

PREVACID

| | | | |
|------------------|-------------|-------------|--------------|
| TAKEDA PHARMS NA | 15MG/PACKET | N021281 001 | May 03, 2001 |
| | 30MG/PACKET | N021281 002 | May 03, 2001 |

INJECTABLE;INTRAVENOUS

PREVACID IV

| | | | |
|--------------------|--------------|-------------|--------------|
| + TAKEDA PHARMS NA | 30MG/VIAL ** | N021566 001 | May 27, 2004 |
|--------------------|--------------|-------------|--------------|

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

LANSOPRAZOLE

| | | | |
|----------------|------|-------------|--------------|
| ANI PHARMS INC | 15MG | A078730 001 | Oct 15, 2010 |
| | 30MG | A078730 002 | Oct 15, 2010 |

LANSOPRAZOLE; NAPROXEN

CAPSULE, DELAYED REL PELLETS, TABLET;ORAL

PREVACID NAPRAPAC 250 (COPACKAGED)

| | | | |
|--------------------|-----------------------|-------------|--------------|
| + TAKEDA PHARMS NA | 15MG,N/A;N/A,250MG ** | N021507 002 | Nov 14, 2003 |
|--------------------|-----------------------|-------------|--------------|

PREVACID NAPRAPAC 375 (COPACKAGED)

| | | | |
|------------------|--------------------|-------------|--------------|
| TAKEDA PHARMS NA | 15MG,N/A;N/A,375MG | N021507 003 | Nov 14, 2003 |
|------------------|--------------------|-------------|--------------|

PREVACID NAPRAPAC 500 (COPACKAGED)

| | | | |
|------------------|--------------------|-------------|--------------|
| TAKEDA PHARMS NA | 15MG,N/A;N/A,500MG | N021507 004 | Nov 14, 2003 |
|------------------|--------------------|-------------|--------------|

LANTHANUM CARBONATE

TABLET, CHEWABLE;ORAL

FOSRENOL

| | | | |
|-----------|---------------|-------------|--------------|
| SHIRE LLC | EQ 250MG BASE | N021468 001 | Oct 26, 2004 |
|-----------|---------------|-------------|--------------|

LAPYRIUM CHLORIDE; UNDECOYLIIUM CHLORIDE IODINE COMPLEX

SOLUTION;TOPICAL

VIRAC REX

| | | | |
|-------------------|-----------|-------------|--|
| CHESEBROUGH PONDS | 0.5%;1.8% | N011914 001 | |
|-------------------|-----------|-------------|--|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LATANOPROST

SOLUTION/DROPS;OPHTHALMIC

LATANOPROST

APOTEX INC

0.005%

A077697 001 Mar 22, 2011

LEFLUNOMIDE

TABLET;ORAL

LEFLUNOMIDE

FOSUN PHARMA

10MG

A077087 001 Sep 13, 2005

SANDOZ

20MG

A077087 002 Sep 13, 2005

SANDOZ

10MG

A077085 001 Sep 13, 2005

SANDOZ

20MG

A077085 002 Sep 13, 2005

LEPIRUDIN RECOMBINANT

INJECTABLE;INJECTION

REFLUDAN

BAYER HLTHCARE

50MG/VIAL

N020807 001 Mar 06, 1998

LETROZOLE

TABLET;ORAL

LETROZOLE

ACTAVIS TOTOWA

2.5MG

A090292 001 Jul 13, 2011

IMPAX LABS

2.5MG

A091638 001 Jun 03, 2011

LANNETT CO INC

2.5MG

A091098 001 Jun 03, 2011

LANNETT CO INC

2.5MG

A202048 001 Oct 29, 2014

MYLAN

2.5MG

A078190 001 Dec 24, 2008

SUN PHARM INDS LTD

2.5MG

A091466 001 Jun 03, 2011

SYNTHON PHARMS

2.5MG

A090196 001 Jun 03, 2011

LEUCOVORIN CALCIUM

FOR SOLUTION;ORAL

LEUCOVORIN CALCIUM

HOSPIRA

EQ 60MG BASE/VIAL

N008107 003 Jan 30, 1987

INJECTABLE;INJECTION

LEUCOVORIN CALCIUM

ABIC

EQ 3MG BASE/ML

A089352 001 Jun 01, 1988

ABIC

EQ 50MG BASE/VIAL

A089353 001 Jun 01, 1988

ABRAXIS PHARM

EQ 50MG BASE/VIAL

A088939 001 Dec 01, 1986

ELKINS SINN

EQ 50MG BASE/VIAL

A070480 001 Jan 02, 1987

ELKINS SINN

EQ 100MG BASE/VIAL

A081224 001 Jun 03, 1994

+ HOSPIRA

EQ 3MG BASE/ML **

N008107 001

+

EQ 50MG BASE/VIAL **

N008107 002

+

EQ 100MG BASE/VIAL **

N008107 004 May 23, 1988

+

EQ 350MG BASE/VIAL **

N008107 005 Apr 05, 1989

PHARMACHEMIE

EQ 350MG BASE/VIAL

A040262 001 Dec 15, 1999

PHARMACHEMIE USA

EQ 50MG BASE/VIAL

A089628 001 Apr 17, 1997

PHARMACHEMIE USA

EQ 100MG BASE/VIAL

A089915 001 Apr 17, 1997

TEVA PARENTERAL

EQ 50MG BASE/VIAL

A081278 001 Sep 28, 1993

LEUCOVORIN CALCIUM PRESERVATIVE FREE

HOSPIRA

EQ 10MG BASE/ML **

A040147 001 Jun 25, 1997

LUITPOLD

EQ 50MG BASE/VIAL

A040338 001 Jan 31, 2001

TEVA PARENTERAL

EQ 10MG BASE/ML

A040332 001 Jun 28, 1999

WELLCOVORIN

GLAXOSMITHKLINE

EQ 5MG BASE/ML

A087439 001 Oct 19, 1982

GLAXOSMITHKLINE

EQ 25MG BASE/VIAL

A089833 001 Jan 23, 1989

GLAXOSMITHKLINE

EQ 50MG BASE/VIAL

A089465 001 Jan 23, 1989

GLAXOSMITHKLINE

EQ 100MG BASE/VIAL

A089834 001 Jan 23, 1989

TABLET;ORAL

LEUCOVORIN CALCIUM

ANI PHARMS INC

EQ 15MG BASE

A075327 001 Mar 24, 1999

EPIC PHARMA LLC

EQ 5MG BASE

A074544 001 Aug 28, 1997

EPIC PHARMA LLC

EQ 25MG BASE

A074544 002 Aug 28, 1997

PAR PHARM

EQ 5MG BASE

A071600 001 Oct 14, 1987

PAR PHARM

EQ 25MG BASE

A071598 001 Oct 14, 1987

PHARMACHEMIE

EQ 5MG BASE

A073099 001 Mar 28, 1997

PHARMACHEMIE

EQ 25MG BASE

A073101 001 Mar 28, 1997

XANODYNE PHARM

EQ 5MG BASE

N018459 001 Jan 30, 1986

XANODYNE PHARM

EQ 10MG BASE

A071962 001 Nov 19, 1987

XANODYNE PHARM

EQ 15MG BASE

A071104 001 Mar 04, 1987

WELLCOVORIN

+

GLAXOSMITHKLINE

EQ 5MG BASE **

N018342 001 Jul 08, 1983

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEUCOVORIN CALCIUM

TABLET; ORAL
WELLCOVORIN
+

EQ 25MG BASE **

N018342 002 Jul 08, 1983

LEUPROLIDE ACETATE

IMPLANT; IMPLANTATION
VIADUR

ORTHO MCNEIL JANSSEN EQ 65MG BASE

N021088 001 Mar 03, 2000

INJECTABLE; INJECTION

LEUPROLIDE ACETATE

GENZYME 1MG/0.2ML

A075721 001 Nov 29, 2001

LUPRON

+ ABBVIE ENDOCRINE INC 1MG/0.2ML

N019010 001 Apr 09, 1985

LUPRON DEPOT

+ ABBVIE ENDOCRINE INC 3.75MG/VIAL **

N020011 001 Oct 22, 1990

LUPRON DEPOT-PED

+ ABBVIE ENDOCRINE INC 3.75MG/VIAL, 7.5MG/VIAL **

N020263 003 Apr 16, 1993

+ 7.5MG/VIAL, 7.5MG/VIAL **

N020263 004 Apr 16, 1993

LEVALLORPHAN TARTRATE

INJECTABLE; INJECTION
LORFAN

ROCHE 1MG/ML

N010423 001

LEVAMISOLE HYDROCHLORIDE

TABLET; ORAL

ERGAMISOL

JANSSEN PHARMA EQ 50MG BASE

N020035 001 Jun 18, 1990

LEVETIRACETAM

SOLUTION; ORAL

LEVETIRACETAM

ACI HEALTHCARE LTD 100MG/ML

A078582 001 Jan 15, 2009

APOTEX INC 100MG/ML

A090187 001 Aug 05, 2011

TABLET; ORAL

LEVETIRACETAM

ACTAVIS LABS FL INC 250MG

A077408 001 Mar 02, 2009

500MG

A077408 002 Mar 02, 2009

750MG

A077408 003 Mar 02, 2009

FOSUN PHARMA 250MG

A077324 001 Jan 15, 2009

500MG

A077324 002 Jan 15, 2009

750MG

A077324 003 Jan 15, 2009

MYLAN 250MG

A077324 004 Jan 15, 2009

500MG

A078731 001 Feb 10, 2009

750MG

A078731 002 Feb 10, 2009

1GM

A078731 003 Feb 10, 2009

WATSON LABS INC 250MG

A078731 004 Feb 10, 2009

500MG

A078797 002 Jan 15, 2009

750MG

A078797 003 Jan 15, 2009

1GM

A078797 004 Jan 15, 2009

TABLET, EXTENDED RELEASE; ORAL

LEVETIRACETAM

MYLAN PHARMS INC 500MG

A200475 001 Dec 19, 2011

750MG

A200475 002 Dec 19, 2011

1GM

A200475 003 Dec 07, 2015

SANDOZ 500MG

A091668 001 Nov 01, 2012

750MG

A091668 002 Nov 01, 2012

VIRTUS PHARMS 500MG

A091291 001 Sep 12, 2011

750MG

A091291 002 Sep 12, 2011

LEVOBETAXOLOL HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

BETAXON

ALCON PHARMS LTD EQ 0.5% BASE

N021114 001 Feb 23, 2000

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVOBUNOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

LEVOBUNOLOL HYDROCHLORIDE

| | | | | |
|-----------------|-------|---------|-----|--------------|
| ALCON LABS INC | 0.25% | A074851 | 001 | Oct 28, 1996 |
| APOTEX INC | 0.25% | A075473 | 001 | Aug 03, 2000 |
| | 0.5% | A075475 | 001 | Aug 03, 2000 |
| BAUSCH AND LOMB | 0.25% | A074307 | 001 | Mar 04, 1994 |

LEVOBUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHIROCAINE

| | | | | |
|------------------|------------------|---------|-----|--------------|
| PURDUE PHARMA LP | EQ 2.5MG BASE/ML | N020997 | 001 | Aug 05, 1999 |
| | EQ 5MG BASE/ML | N020997 | 002 | Aug 05, 1999 |
| | EQ 7.5MG BASE/ML | N020997 | 003 | Aug 05, 1999 |

LEVOCABASTINE HYDROCHLORIDE

SUSPENSION/DROPS;OPHTHALMIC

LIVOSTIN

| | | | | |
|----------|---------------|---------|-----|--------------|
| NOVARTIS | EQ 0.05% BASE | N020219 | 001 | Nov 10, 1993 |
|----------|---------------|---------|-----|--------------|

LEVOCARNITINE

INJECTABLE; INJECTION

LEVOCARNITINE

| | | | | |
|-----------------|----------|---------|-----|--------------|
| TEVA PHARMS USA | 200MG/ML | A075881 | 001 | Mar 29, 2001 |
|-----------------|----------|---------|-----|--------------|

SOLUTION; ORAL

CARNITOR

| | | | | |
|---------------------|----------|---------|-----|--------------|
| LEADIANT BIOSCI INC | 1GM/10ML | N018948 | 002 | Apr 27, 1988 |
|---------------------|----------|---------|-----|--------------|

LEVOCETIRIZINE DIHYDROCHLORIDE

TABLET; ORAL

LEVOCETIRIZINE DIHYDROCHLORIDE

| | | | | |
|--------------|-----|---------|-----|--------------|
| FOSUN PHARMA | 5MG | A090486 | 001 | Mar 26, 2013 |
|--------------|-----|---------|-----|--------------|

LEVODOPA

CAPSULE; ORAL

BENDOPA

| | | | | |
|--------------------|-------|---------|-----|--|
| VALEANT PHARM INTL | 100MG | N016948 | 003 | |
| | 250MG | N016948 | 001 | |
| | 500MG | N016948 | 002 | |

DOPAR

| | | | | |
|-------|-------|---------|-----|--|
| SHIRE | 100MG | N016913 | 003 | |
| | 250MG | N016913 | 001 | |
| | 500MG | N016913 | 002 | |

LARODOPA

| | | | | |
|-------|-------|---------|-----|--|
| ROCHE | 100MG | N016912 | 002 | |
| | 250MG | N016912 | 001 | |
| | 500MG | N016912 | 006 | |

TABLET; ORAL

DOPAR

| | | | | |
|-------|-------|---------|-----|--|
| SHIRE | 250MG | N016913 | 004 | |
| | 500MG | N016913 | 005 | |

LARODOPA

| | | | | |
|-------|-------|---------|-----|--|
| ROCHE | 100MG | N016912 | 005 | |
| | 250MG | N016912 | 003 | |
| | 500MG | N016912 | 004 | |

LEVOFLOXACIN

INJECTABLE; INJECTION

LEVAQUIN

| | | | | | |
|---|----------------|----------------------------|---------|-----|--------------|
| + | JANSSEN PHARMS | EQ 500MG/20ML (EQ 25MG/ML) | N020635 | 001 | Dec 20, 1996 |
| + | | EQ 750MG/30ML (EQ 25MG/ML) | N020635 | 004 | Dec 20, 1996 |

LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | |
|---|----------------|-------------------------------|---------|-----|--------------|
| + | JANSSEN PHARMS | EQ 250MG/50ML (EQ 5MG/ML) ** | N020635 | 002 | Dec 20, 1996 |
| + | | EQ 500MG/100ML (EQ 5MG/ML) ** | N020635 | 003 | Dec 20, 1996 |
| + | | EQ 750MG/150ML (EQ 5MG/ML) ** | N020635 | 005 | Dec 20, 1996 |

LEVOFLOXACIN

| | | | | |
|-------------------|----------------------------|---------|-----|--------------|
| AKORN | EQ 500MG/20ML (EQ 25MG/ML) | A091644 | 001 | Jun 20, 2011 |
| | EQ 750MG/30ML (EQ 25MG/ML) | A091644 | 002 | Jun 20, 2011 |
| EMCURE PHARMS LTD | EQ 500MG/20ML (EQ 25MG/ML) | A202590 | 001 | Jan 24, 2013 |
| | EQ 750MG/30ML (EQ 25MG/ML) | A202590 | 002 | Jan 24, 2013 |
| HOSPIRA INC | EQ 500MG/20ML (EQ 25MG/ML) | A078577 | 001 | Aug 12, 2015 |
| | EQ 750MG/30ML (EQ 25MG/ML) | A078577 | 002 | Aug 12, 2015 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVOFLOXACIN

INJECTABLE; INJECTION

LEVOFLOXACIN

| | | | |
|----------------------|----------------------------|-------------|--------------|
| MYLAN ASI | EQ 500MG/20ML (EQ 25MG/ML) | A200560 001 | Jun 20, 2011 |
| | EQ 750MG/30ML (EQ 25MG/ML) | A200560 002 | Jun 20, 2011 |
| ZYDUS PHARMS USA INC | EQ 500MG/20ML (EQ 25MG/ML) | A205968 001 | Jun 01, 2017 |
| | EQ 750MG/30ML (EQ 25MG/ML) | A205968 002 | Jun 01, 2017 |

SOLUTION; ORAL

LEVAQUIN

| | | | |
|------------------|------------|-------------|--------------|
| + JANSSEN PHARMS | 250MG/10ML | N021721 001 | Oct 21, 2004 |
|------------------|------------|-------------|--------------|

SOLUTION/DROPS; OPHTHALMIC

IQUIX

| | | | |
|----------|---------|-------------|--------------|
| + SANTEN | 1.5% ** | N021571 001 | Mar 01, 2004 |
|----------|---------|-------------|--------------|

LEVOFLOXACIN

| | | | |
|------------|------|-------------|--------------|
| APOTEX INC | 0.5% | A078282 001 | Dec 20, 2010 |
|------------|------|-------------|--------------|

QUIXIN

| | | | |
|----------|---------|-------------|--------------|
| + SANTEN | 0.5% ** | N021199 001 | Aug 18, 2000 |
|----------|---------|-------------|--------------|

TABLET; ORAL

LEVAQUIN

| | | | |
|------------------|-------|-------------|--------------|
| + JANSSEN PHARMS | 250MG | N020634 001 | Dec 20, 1996 |
| | 500MG | N020634 002 | Dec 20, 1996 |
| | 750MG | N020634 003 | Sep 08, 2000 |

LEVOFLOXACIN

| | | | |
|-----------------|-------|-------------|--------------|
| MYLAN | 250MG | A076276 001 | Jun 20, 2011 |
| | 500MG | A076276 002 | Jun 20, 2011 |
| | 750MG | A077097 001 | Jun 20, 2011 |
| WATSON LABS INC | 250MG | A201484 001 | Nov 22, 2013 |
| | 500MG | A201484 002 | Nov 22, 2013 |
| | 750MG | A201484 003 | Nov 22, 2013 |

LEVOLEUCOVORIN CALCIUM

SOLUTION; INTRAVENOUS

FUSILEV

| | | | |
|-------------------|-----------------------------------------|-------------|--------------|
| + SPECTRUM PHARMS | EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML) | N020140 002 | Apr 29, 2011 |
| | ** | | |
| | EQ 250MG BASE/25ML (EQ 10MG BASE/ML) ** | N020140 003 | Apr 29, 2011 |

LEVOMEPRIMAZINE

INJECTABLE; INJECTION

LEVOPROME

| | | | |
|---------|---------|-------------|--|
| IMMUNEX | 20MG/ML | N015865 001 | |
|---------|---------|-------------|--|

LEVOMETHADYL ACETATE HYDROCHLORIDE

CONCENTRATE; ORAL

ORLAAM

| | | | |
|----------|------------|-------------|--------------|
| + ROXANE | 10MG/ML ** | N020315 001 | Jul 09, 1993 |
|----------|------------|-------------|--------------|

LEVONORDEFRIN; MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARESTOCAINE HYDROCHLORIDE W/ LEVONORDEFRIN

| | | | |
|--------|---------------|-------------|--|
| SOLVAY | 0.05MG/ML; 2% | A085010 001 | |
|--------|---------------|-------------|--|

CARBOCAINE W/ NEO-COBEFRIN

| | | | |
|---------------|---------------|-------------|--|
| EASTMAN KODAK | 0.05MG/ML; 2% | N012125 002 | |
|---------------|---------------|-------------|--|

ISOCAINE HYDROCHLORIDE W/ LEVONORDEFRIN

| | | | |
|---------------|---------------|-------------|--|
| SEPTODONT INC | 0.05MG/ML; 2% | A084697 001 | |
|---------------|---------------|-------------|--|

MEPIVACAINE HYDROCHLORIDE W/ LEVONORDEFRIN

| | | | |
|-------------|---------------|-------------|--------------|
| BELMORA LLC | 0.05MG/ML; 2% | A084850 002 | Oct 21, 1983 |
|-------------|---------------|-------------|--------------|

POLOCAINE W/ LEVONORDEFRIN

| | | | |
|----------------|---------------|-------------|--------------|
| DENTSPLY PHARM | 0.05MG/ML; 2% | A089517 001 | Apr 14, 1988 |
|----------------|---------------|-------------|--------------|

LEVONORDEFRIN; PROCAINE HYDROCHLORIDE; PROPOXYCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

RAVOCAINE AND NOVOCAIN W/ NEO-COBEFRIN

| | | | |
|---------------|---------------------|-------------|--|
| EASTMAN KODAK | 0.05MG/ML; 2%; 0.4% | N008592 007 | |
|---------------|---------------------|-------------|--|

LEVONORGESTREL

IMPLANT; IMPLANTATION

JADELLE

| | | | |
|----------------------|-----------------|-------------|--------------|
| + POPULATION COUNCIL | 75MG/IMPLANT ** | N020544 001 | Nov 01, 1996 |
|----------------------|-----------------|-------------|--------------|

LEVONORGESTREL

| | | | |
|------------------|--------------|-------------|--------------|
| WYETH PHARMS INC | 75MG/IMPLANT | N020627 001 | Aug 15, 1996 |
|------------------|--------------|-------------|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVONORGESTREL

IMPLANT; IMPLANTATION

NORPLANT

POPULATION COUNCIL 36MG/IMPLANT N019897 001 Dec 10, 1990

NORPLANT SYSTEM IN PLASTIC CONTAINER

WYETH PHARMS INC 36MG/IMPLANT N020088 001 Dec 10, 1990

TABLET; ORAL

LEVONORGESTREL

FDN CONSUMER 0.75MG ** A078665 001 Aug 28, 2009

LUPIN LTD 0.75MG A091328 001 Jan 23, 2013

WATSON LABS 0.75MG A078666 001 Jun 24, 2009

PLAN B

+ FDN CONSUMER 0.75MG ** N021045 001 Jul 28, 1999

+ 0.75MG ** N021045 002 Aug 24, 2006

LEVOPROPOXYPHENE NAPSYLATE ANHYDROUS

CAPSULE; ORAL

NOVRAD

LILLY EQ 50MG BASE N012928 006

EQ 100MG BASE N012928 004

SUSPENSION; ORAL

NOVRAD

LILLY EQ 50MG BASE/5ML N012928 002

LEVORPHANOL TARTRATE

INJECTABLE; INJECTION

LEVO-DROMORAN

VALEANT PHARM INTL 2MG/ML N008719 001 Dec 19, 1991

TABLET; ORAL

LEVO-DROMORAN

+ VALEANT PHARM INTL 2MG ** N008720 001 Dec 19, 1991

LEVOTHYROXINE SODIUM

SOLUTION; ORAL

TIROSINT-SOL

+ INSTITUT BIOCHIMIQUE 13MCG/ML N206977 001 Dec 15, 2016

+ 25MCG/ML N206977 002 Dec 15, 2016

+ 50MCG/ML N206977 003 Dec 15, 2016

+ 75MCG/ML N206977 004 Dec 15, 2016

+ 88MCG/ML N206977 005 Dec 15, 2016

+ 100MCG/ML N206977 006 Dec 15, 2016

+ 112MCG/ML N206977 007 Dec 15, 2016

+ 125MCG/ML N206977 008 Dec 15, 2016

+ 137MCG/ML N206977 009 Dec 15, 2016

+ 150MCG/ML N206977 010 Dec 15, 2016

+ 175MCG/ML N206977 011 Dec 15, 2016

+ 200MCG/ML N206977 012 Dec 15, 2016

TABLET; ORAL

EUTHYROX

PROVELL 0.3MG N021292 012 May 31, 2002

LEVOLET

GENUS LIFESCIENCES 0.025MG N021137 001 Jun 06, 2003

0.05MG N021137 002 Jun 06, 2003

0.075MG N021137 003 Jun 06, 2003

0.088MG N021137 004 Jun 06, 2003

0.1MG N021137 005 Jun 06, 2003

0.112MG N021137 006 Jun 06, 2003

0.125MG N021137 007 Jun 06, 2003

0.137MG N021137 008 Jun 06, 2003

0.15MG N021137 009 Jun 06, 2003

0.175MG N021137 010 Jun 06, 2003

0.2MG N021137 011 Jun 06, 2003

0.3MG N021137 012 Jun 06, 2003

LEVOTHYROXINE SODIUM

MERCK KGAA 0.025MG A076752 001 Jun 16, 2005

0.05MG A076752 002 Jun 16, 2005

0.075MG A076752 003 Jun 16, 2005

0.088MG A076752 004 Jun 16, 2005

0.1MG A076752 005 Jun 16, 2005

0.112MG A076752 006 Jun 16, 2005

0.125MG A076752 007 Jun 16, 2005

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LEVOTHYROXINE SODIUM

TABLET;ORAL

LEVOTHYROXINE SODIUM

| | | |
|---------|-------------|--------------|
| 0.15MG | A076752 008 | Jun 16, 2005 |
| 0.175MG | A076752 009 | Jun 16, 2005 |
| 0.2MG | A076752 010 | Jun 16, 2005 |
| 0.3MG | A076752 011 | Jun 16, 2005 |

LEVOXYL

+ KING PHARMS

| | | |
|----------|-------------|--------------|
| 0.3MG ** | N021301 012 | May 25, 2001 |
|----------|-------------|--------------|

THYRO-TABS

+ LLOYD

| | | |
|------------|-------------|--------------|
| 0.025MG ** | N021116 001 | Oct 24, 2002 |
|------------|-------------|--------------|

+

| | | |
|-----------|-------------|--------------|
| 0.05MG ** | N021116 002 | Oct 24, 2002 |
|-----------|-------------|--------------|

+

| | | |
|------------|-------------|--------------|
| 0.075MG ** | N021116 003 | Oct 24, 2002 |
|------------|-------------|--------------|

+

| | | |
|------------|-------------|--------------|
| 0.088MG ** | N021116 010 | Oct 24, 2002 |
|------------|-------------|--------------|

+

| | | |
|----------|-------------|--------------|
| 0.1MG ** | N021116 004 | Oct 24, 2002 |
|----------|-------------|--------------|

+

| | | |
|------------|-------------|--------------|
| 0.112MG ** | N021116 011 | Oct 24, 2002 |
|------------|-------------|--------------|

+

| | | |
|------------|-------------|--------------|
| 0.125MG ** | N021116 005 | Oct 24, 2002 |
|------------|-------------|--------------|

+

| | | |
|------------|-------------|--------------|
| 0.137MG ** | N021116 012 | Dec 07, 2004 |
|------------|-------------|--------------|

+

| | | |
|-----------|-------------|--------------|
| 0.15MG ** | N021116 006 | Oct 24, 2002 |
|-----------|-------------|--------------|

+

| | | |
|------------|-------------|--------------|
| 0.175MG ** | N021116 007 | Oct 24, 2002 |
|------------|-------------|--------------|

+

| | | |
|----------|-------------|--------------|
| 0.2MG ** | N021116 008 | Oct 24, 2002 |
|----------|-------------|--------------|

+

| | | |
|----------|-------------|--------------|
| 0.3MG ** | N021116 009 | Oct 24, 2002 |
|----------|-------------|--------------|

LIDOCAINE

AEROSOL;ORAL

XYLOCAINE

ASTRAZENECA

10%

N014394 001

FILM, EXTENDED RELEASE;BUCCAL

DENTIPATCH

NOVEN

23MG/PATCH

N020575 001 May 21, 1996

OINTMENT;TOPICAL

ALPHACAINE

CARLISLE

5%

A084944 001

5%

A084946 001

5%

A084947 001

LIDOCAINE

BELMORA LLC

5%

A080210 001

XYLOCAINE

+ ASTRAZENECA

5% **

N008048 001

PATCH;TOPICAL

DENTIPATCH

NOVEN

46.1MG/PATCH

N020575 002 May 21, 1996

SOLUTION;TOPICAL

XYLOCAINE

ASTRAZENECA

5%

N014127 001

SUPPOSITORY;RECTAL

XYLOCAINE

ASTRAZENECA

100MG

N013077 001

LIDOCAINE HYDROCHLORIDE

INJECTABLE;INJECTION

ALPHACAINE HYDROCHLORIDE

CARLISLE

2%

A084721 001

LIDOCAINE HYDROCHLORIDE

ABBOTT

10%

A087980 001 Feb 02, 1983

20%

A089362 001 May 25, 1988

ABRAXIS PHARM

1%

A080420 001

1%

A086761 001

1.5%

A080420 005

2%

A080420 002

2%

A080420 004

2%

A086761 002

2%

N017508 001

4%

N017508 002

20%

N017508 004

AKORN

1%

A085037 001

2%

A085037 002

BEL MAR

1%

A080710 001

2%

A080760 001

BELMORA LLC

2%

A080504 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE

| | | | | |
|-------------------------------------------------------------------|-------------|---------|-----|--------------|
| DELL LABS | 1% | A083387 | 001 | |
| | 2% | A083388 | 001 | |
| ELKINS SINN | 0.5% | A085131 | 001 | |
| | 4% | A084626 | 001 | |
| GD SEARLE LLC | 1% | A083135 | 001 | |
| | 2% | A083135 | 002 | |
| HOSPIRA | 1% | A040013 | 001 | Jun 23, 1995 |
| | 1.5% | A088330 | 001 | May 17, 1984 |
| | 2% | A088331 | 001 | May 17, 1984 |
| INTL MEDICATION | 1% | N017701 | 002 | |
| | 2% | N017701 | 001 | |
| | 1GM/VIAL | N018543 | 001 | |
| | 2GM/VIAL | N018543 | 002 | |
| LUITPOLD | 2% | A083198 | 001 | |
| LYPHOMED | 1% | A080390 | 001 | |
| | 2% | A080390 | 002 | |
| MILES | 1% | A080414 | 001 | |
| | 2% | A080414 | 002 | |
| MYLAN LABS LTD | 0.5% | A091056 | 001 | Dec 08, 2010 |
| | 0.5% | A091058 | 001 | Sep 30, 2010 |
| | 1% | A091056 | 002 | Dec 08, 2010 |
| | 1% | A091058 | 002 | Sep 30, 2010 |
| | 2% | A202242 | 001 | Apr 11, 2014 |
| WATSON LABS | 1% | A080377 | 001 | |
| | 1% | A083627 | 001 | |
| | 2% | A080377 | 002 | |
| | 2% | A083627 | 002 | |
| WEST-WARD PHARMS INT | 1% | A080407 | 001 | |
| | 2% | A080407 | 002 | |
| WYETH AYERST | 1% | A083083 | 001 | |
| | 2% | A083083 | 002 | |
| LIDOCAINE HYDROCHLORIDE 0.1% AND DEXTROSE 5% IN PLASTIC CONTAINER | | | | |
| BAXTER HLTHCARE | 100MG/100ML | N018461 | 001 | |
| LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 200MG/100ML | N018967 | 001 | Mar 30, 1984 |
| LIDOCAINE HYDROCHLORIDE 0.2% IN DEXTROSE 5% | | | | |
| HOSPIRA | 200MG/100ML | A083158 | 005 | |
| LIDOCAINE HYDROCHLORIDE 0.2% IN DEXTROSE 5% IN PLASTIC CONTAINER | | | | |
| ABBOTT | 200MG/100ML | N018954 | 001 | Jul 09, 1985 |
| HOSPIRA | 200MG/100ML | N018388 | 001 | |
| LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 400MG/100ML | N018967 | 002 | Mar 30, 1984 |
| LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5% | | | | |
| HOSPIRA | 400MG/100ML | A083158 | 006 | |
| LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5% IN PLASTIC CONTAINER | | | | |
| HOSPIRA | 400MG/100ML | N018388 | 002 | |
| LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 800MG/100ML | N018967 | 003 | Mar 30, 1984 |
| LIDOCAINE HYDROCHLORIDE 0.8% IN DEXTROSE 5% IN PLASTIC CONTAINER | | | | |
| HOSPIRA | 800MG/100ML | N018388 | 003 | Nov 05, 1982 |
| LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER | | | | |
| HOSPIRA | 1.5% | A088326 | 001 | Jul 31, 1984 |
| | 10% | A088367 | 001 | Jul 31, 1984 |
| | 20% | A088368 | 001 | Jul 31, 1984 |
| LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE | | | | |
| INTL MEDICATION | 4% | N017702 | 002 | |
| | 20% | N017702 | 001 | |
| MYLAN LABS LTD | 2% | A090665 | 001 | Sep 27, 2010 |
| WEST-WARD PHARMS INT | 1% | A084625 | 001 | |
| | 2% | A084625 | 002 | |
| LIDOCATON | | | | |
| PHARMATON | 2% | A084727 | 001 | Aug 17, 1983 |
| LIDOPEN | | | | |
| MERIDIAN MEDCL TECHN | 10% | N017549 | 001 | |
| XYLOCAINE | | | | |
| ASTRAZENECA | 1% | N010418 | 005 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

XYLOCAINE

1.5%

N010418 009

2%

N010418 007

XYLOCAINE 4% PRESERVATIVE FREE

+ FRESENIUS KABI USA 4%

N010417 001

XYLOCAINE DENTAL

DENTSPLY PHARM 2%

N021380 001

XYLOCAINE PRESERVATIVE FREE

FRESENIUS KABI USA 10%

N016801 003

INJECTABLE; SPINAL

XYLOCAINE 1.5% W/ DEXTROSE 7.5%

FRESENIUS KABI USA 1.5%

N016297 001

XYLOCAINE 5% W/ GLUCOSE 7.5%

ASTRAZENECA 5%

N010496 002 Jul 07, 1982

JELLY; TOPICAL

ANESTACON

BLUEPHARMA 2%

A080429 001

LIDOCAINE HYDROCHLORIDE

G AND W LABS INC 2%

A081318 001 Apr 29, 1993

WATSON LABS INC 2%

A040837 001 Mar 23, 2011

SOLUTION; ORAL

LIDOCAINE HYDROCHLORIDE VISCOUS

ACTAVIS MID ATLANTIC 2%

A086578 001

INTL MEDICATION 2%

A086389 001 Feb 02, 1982

XYLOCAINE VISCOUS

FRESENIUS KABI USA 2% **

N009470 001

SOLUTION; TOPICAL

LARYNGOTRACHEAL ANESTHESIA KIT

KENDALL IL 4%

A087931 001 Jun 10, 1983

LIDOCAINE HYDROCHLORIDE

LANNETT CO INC 4%

A040710 001 Feb 27, 2007

PACO 4%

A089688 001 Jun 30, 1989

WOCKHARDT BIO AG 4%

A087881 001 Nov 18, 1982

LTA II KIT

HOSPIRA 4%

A080409 001

4%

A088542 001 Jul 31, 1984

PEDIATRIC LTA KIT

ABBOTT 2%

A088572 001 Jul 31, 1984

HOSPIRA 2%

A085995 001

XYLOCAINE 4% PRESERVATIVE FREE

+ FRESENIUS KABI USA 4%

N010417 002

LIDOCAINE HYDROCHLORIDE; OXYTETRACYCLINE

INJECTABLE; INJECTION

TERRAMYCIN

PFIZER 2%; 50MG/ML

A060567 001

2%; 125MG/ML

A060567 002

LIDOCAINE; PRILOCAINE

DISC; TOPICAL

EMLA

ASTRAZENECA 2.5%; 2.5%

N020962 001 Feb 04, 1998

LINCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

LINCOCIN

PHARMACIA AND UPJOHN EQ 250MG BASE

N050316 001

EQ 500MG BASE

N050316 002

INJECTABLE; INJECTION

LINCOMYCIN HYDROCHLORIDE

WATSON LABS EQ 300MG BASE/ML

A063180 001 Apr 16, 1991

LINDANE

CREAM; TOPICAL

KWELL

REED AND CARNRICK 1%

A084218 001

1%

N006309 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LINDANE

LOTION; TOPICAL

GAMENE

SOLA BARNES HIND 1% A084989 001

KWELL

REED AND CARNRICK 1% A084218 002

1% N006309 003

LINDANE

WOCKHARDT 1% A088190 001 Aug 16, 1984

SCABENE

STIEFEL 1% A086769 001

SHAMPOO; TOPICAL

GAMENE

SOLA BARNES HIND 1% A084988 001

KWELL

REED AND CARNRICK 1% A084219 001

1% N010718 001

SCABENE

STIEFEL 1% A087940 001 Apr 08, 1983

LINEZOLID

SOLUTION; INTRAVENOUS

ZYVOX

+ PHARMACIA AND UPJOHN 400MG/200ML (2MG/ML) N021131 002 Apr 18, 2000

TABLET; ORAL

ZYVOX

+ PHARMACIA AND UPJOHN 400MG ** N021130 001 Apr 18, 2000

LIOTHYRONINE SODIUM

TABLET; ORAL

LIOTHYRONINE SODIUM

WATSON LABS EQ 0.025MG BASE A085755 001 Jan 25, 1982

EQ 0.05MG BASE A085753 001 Feb 03, 1982

LIOTRIX (T4; T3)

TABLET; ORAL

EUTHROID-0.5

PARKE DAVIS 0.03MG; 0.0075MG N016680 001

EUTHROID-1

PARKE DAVIS 0.06MG; 0.015MG N016680 002

EUTHROID-2

PARKE DAVIS 0.12MG; 0.03MG N016680 003

EUTHROID-3

PARKE DAVIS 0.18MG; 0.045MG N016680 004

THYROLAR-0.25

+ ALLERGAN SALES LLC 0.0125MG; 0.0031MG N016807 001

THYROLAR-0.5

+ ALLERGAN SALES LLC 0.025MG; 0.0063MG N016807 005

THYROLAR-1

+ ALLERGAN SALES LLC 0.05MG; 0.0125MG N016807 004

THYROLAR-2

+ ALLERGAN SALES LLC 0.1MG; 0.025MG N016807 002

THYROLAR-3

+ ALLERGAN SALES LLC 0.15MG; 0.0375MG N016807 003

THYROLAR-5

ALLERGAN SALES LLC 0.25MG; 0.0625MG N016807 006

LISINAPRIL

TABLET; ORAL

LISINAPRIL

SANDOZ 2.5MG A075903 001 Jul 01, 2002

2.5MG A075999 001 Jul 01, 2002

5MG A075903 002 Jul 01, 2002

5MG A075999 002 Jul 01, 2002

10MG A075903 003 Jul 01, 2002

10MG A075999 003 Jul 01, 2002

20MG A075903 004 Jul 01, 2002

20MG A075999 004 Jul 01, 2002

30MG A075903 005 Jul 01, 2002

30MG A075999 005 Jul 01, 2002

40MG A075903 006 Jul 01, 2002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LISINAPRILTABLET;ORAL
LISINAPRIL

| | | | |
|----------|-------|-------------|--------------|
| | 40MG | A075999 006 | Jul 01, 2002 |
| TEVA | 2.5MG | A075783 001 | Jul 01, 2002 |
| | 5MG | A075783 002 | Jul 01, 2002 |
| | 10MG | A075783 003 | Jul 01, 2002 |
| | 20MG | A075783 004 | Jul 01, 2002 |
| | 30MG | A075783 005 | Jul 01, 2002 |
| | 40MG | A075783 006 | Jul 01, 2002 |
| PRINIVIL | | | |
| MERCK | 2.5MG | N019558 006 | Jan 28, 1994 |

LITHIUM CARBONATE

CAPSULE;ORAL

| | | | |
|-------------------------------|----------|-------------|--------------|
| ESKALITH | | | |
| NOVEN THERAP | 300MG | N016860 001 | |
| LITHIUM CARBONATE | | | |
| ABLE | 150MG | A076823 001 | Jun 29, 2004 |
| | 300MG | A076121 001 | Sep 27, 2001 |
| | 300MG | A076823 002 | Jun 29, 2004 |
| | 600MG | A076823 003 | Jun 29, 2004 |
| APOTEX INC | 300MG | A076795 001 | Nov 22, 2004 |
| USL PHARMA | 300MG | A072542 001 | Feb 01, 1989 |
| WATSON LABS | 300MG | A070407 001 | Mar 19, 1987 |
| LITHONATE | | | |
| SOLVAY | 300MG | N016782 001 | |
| TABLET;ORAL | | | |
| ESKALITH | | | |
| JDS PHARMS | 300MG | N017971 001 | |
| LITHANE | | | |
| BAYER PHARMS | 300MG | N018833 001 | Jul 18, 1985 |
| LITHIUM CARBONATE | | | |
| HIKMA INTL PHARMS | 300MG | A078715 001 | Dec 28, 2010 |
| PFIZER | 300MG | N016834 001 | |
| LITHOTABS | | | |
| SOLVAY | 300MG | N016980 001 | |
| TABLET, EXTENDED RELEASE;ORAL | | | |
| ESKALITH CR | | | |
| JDS PHARMS | 450MG ** | N018152 001 | Mar 29, 1982 |
| LITHIUM CARBONATE | | | |
| ABLE | 300MG | A076382 001 | Apr 21, 2003 |
| BARR | 300MG | A076170 001 | Jun 10, 2002 |
| | 450MG | A076366 001 | Aug 21, 2003 |
| HIKMA INTL PHARMS | 450MG | A076490 001 | Jun 17, 2003 |

LITHIUM CITRATE

SYRUP;ORAL

| | | | |
|-----------|------------------------|-------------|--|
| LITHONATE | | | |
| SOLVAY | EQ 300MG CARBONATE/5ML | N017672 001 | |

LOMEFLOXACIN HYDROCHLORIDE

TABLET;ORAL

| | | | |
|-----------|---------------|-------------|--------------|
| MAXAQUIN | | | |
| PHARMACIA | EQ 400MG BASE | N020013 001 | Feb 21, 1992 |

LOMUSTINE

CAPSULE;ORAL

| | | | |
|-----------------|-----|-------------|--------------|
| GLEOSTINE | | | |
| + CORDEN PHARMA | 5MG | N017588 004 | Dec 19, 2014 |

LOPERAMIDE HYDROCHLORIDE

CAPSULE;ORAL

| | | | |
|--------------------------|--------|-------------|--------------|
| IMODIUM | | | |
| J AND J CONSUMER INC | 2MG ** | N017690 001 | |
| + | 2MG ** | N017694 001 | |
| LOPERAMIDE HYDROCHLORIDE | | | |
| ROXANE | 2MG | A073080 001 | Nov 27, 1991 |
| TEVA | 2MG | A073122 001 | Aug 30, 1991 |
| YAOPHARMA CO LTD | 2MG | A072993 001 | Aug 28, 1992 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LOPERAMIDE HYDROCHLORIDE

SOLUTION; ORAL

IMODIUM

JANSSEN PHARMS 1MG/5ML N019037 001 Jul 31, 1984

LOPERAMIDE HYDROCHLORIDE

ALLIED 1MG/5ML A073079 001 Apr 30, 1992

ALPHARMA US PHARMS 1MG/5ML A073187 001 Sep 15, 1992

DURAMED PHARMS BARR 1MG/5ML A074991 001 Dec 29, 1997

TEVA 1MG/5ML A073478 001 Jun 23, 1995

WATSON LABS 1MG/5ML A073062 001 May 28, 1993

TABLET; ORAL

LOPERAMIDE HYDROCHLORIDE

ABLE 2MG A073528 001 Nov 30, 1993

CONTRACT PHARMACAL 2MG A073254 001 Jul 30, 1993

PERRIGO 2MG A074194 001 Oct 30, 1992

TABLET, CHEWABLE; ORAL

IMODIUM A-D EZ CHEWS

+ J AND J CONSUMER INC 2MG N020448 001 Jul 24, 1997

LOPERAMIDE HYDROCHLORIDE; SIMETHICONE

TABLET, CHEWABLE; ORAL

IMODIUM MULTI-SYMPTOM RELIEF

+ J AND J CONSUMER INC 2MG;125MG N020606 001 Jun 26, 1996

LOPINAVIR; RITONAVIR

CAPSULE; ORAL

KALETRA

ABBVIE 133.3MG;33.3MG N021226 001 Sep 15, 2000

LORACARBEF

CAPSULE; ORAL

LORABID

KING PHARMS 200MG N050668 001 Dec 31, 1991

400MG N050668 002 Apr 05, 1996

FOR SUSPENSION; ORAL

LORABID

KING PHARMS 100MG/5ML N050667 001 Dec 31, 1991

200MG/5ML N050667 002 Dec 31, 1991

LORATADINE

SYRUP; ORAL

CLARITIN HIVES RELIEF

+ BAYER HEALTHCARE LLC 1MG/ML ** N020641 003 Nov 19, 2003

LORATADINE

APOTEX INC 1MG/ML A075565 001 Oct 05, 2004

RANBAXY LABS LTD 1MG/ML A076529 001 Aug 20, 2004

TABLET; ORAL

LORATADINE

PERRIGO 10MG N021512 001 Jun 24, 2004

LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

LORATADINE AND PSEUDOEPHEDRINE SULFATE

ACTAVIS LABS FL INC 5MG;120MG A076208 001 Jan 28, 2004

LORAZEPAM

INJECTABLE; INJECTION

LORAZEPAM

AKORN 2MG/ML A074974 001 Jul 23, 1998

BEDFORD 2MG/ML A077076 001 Jul 13, 2005

4MG/ML A077076 002 Jul 13, 2005

DAVA PHARMS INC 2MG/ML A074793 001 Mar 16, 2000

4MG/ML A074793 002 Mar 16, 2000

HOSPIRA 2MG/ML A074280 001 May 27, 1994

2MG/ML A074300 001 Apr 12, 1994

4MG/ML A074280 002 May 27, 1994

4MG/ML A074300 003 Mar 19, 1997

MYLAN ASI 2MG/ML A200217 001 Apr 04, 2017

2MG/ML A200542 001 Apr 28, 2017

4MG/ML A200217 002 Apr 04, 2017

4MG/ML A200542 002 Apr 28, 2017

WATSON LABS 2MG/ML A074276 001 Apr 15, 1994

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LORAZEPAM

INJECTABLE; INJECTION

LORAZEPAM

| | | | |
|-----------------------------|-----------|-------------|--------------|
| | 4MG/ML | A074276 002 | Apr 15, 1994 |
| WATSON LABS INC | 1MG/0.5ML | A074551 003 | Sep 12, 1996 |
| | 2MG/ML | A074535 001 | Sep 12, 1996 |
| | 2MG/ML | A074551 001 | Sep 12, 1996 |
| | 4MG/ML | A074535 002 | Sep 12, 1996 |
| | 4MG/ML | A074551 002 | Sep 12, 1996 |
| WEST-WARD PHARMS INT | 2MG/ML | A074496 001 | Sep 28, 1998 |
| | 4MG/ML | A074496 002 | Sep 28, 1998 |
| LORAZEPAM PRESERVATIVE FREE | | | |
| BEDFORD LABS | 2MG/ML | A077074 001 | Jul 13, 2005 |
| | 4MG/ML | A077074 002 | Jul 13, 2005 |

SOLUTION; ORAL

LORAZEPAM

| | | | |
|--------|-----------|-------------|--------------|
| ROXANE | 0.5MG/5ML | A074648 001 | Mar 18, 1997 |
|--------|-----------|-------------|--------------|

TABLET; ORAL

LORAZ

| | | | |
|------------------|-------|-------------|--------------|
| QUANTUM PHARMICS | 0.5MG | A070200 001 | Aug 09, 1985 |
| | 1MG | A070201 001 | Aug 09, 1985 |
| | 2MG | A070202 001 | Aug 09, 1985 |

LORAZEPAM

| | | | |
|--------------------|-------|-------------|--------------|
| AM THERAP | 0.5MG | A070727 001 | Mar 07, 1986 |
| | 1MG | A070728 001 | Mar 07, 1986 |
| | 2MG | A070729 001 | Mar 07, 1986 |
| ANDA REPOSITORY | 0.5MG | A072555 002 | Mar 29, 1991 |
| | 1MG | A072555 003 | Mar 29, 1991 |
| | 2MG | A072555 001 | Mar 29, 1991 |
| HALSEY | 0.5MG | A071434 001 | Sep 01, 1987 |
| | 1MG | A071435 001 | Sep 01, 1987 |
| | 2MG | A071436 001 | Sep 01, 1987 |
| MUTUAL PHARM | 0.5MG | A070472 001 | Dec 10, 1985 |
| | 1MG | A070473 001 | Dec 10, 1985 |
| | 2MG | A070474 001 | Dec 10, 1985 |
| MYLAN | 0.5MG | A071591 002 | Oct 13, 1987 |
| | 1MG | A071591 003 | Oct 13, 1987 |
| | 2MG | A071591 001 | Oct 13, 1987 |
| PAR PHARM | 0.5MG | A070675 001 | Dec 01, 1986 |
| | 1MG | A070676 001 | Dec 01, 1986 |
| | 2MG | A070677 001 | Dec 01, 1986 |
| SANDOZ | 0.5MG | A071193 001 | Apr 15, 1988 |
| | 1MG | A071194 001 | Apr 15, 1988 |
| | 2MG | A071195 001 | Apr 15, 1988 |
| SUN PHARM INDS LTD | 0.5MG | A076045 001 | Aug 29, 2001 |
| | 1MG | A076045 002 | Aug 29, 2001 |
| | 2MG | A076045 003 | Aug 29, 2001 |
| SUPERPHARM | 0.5MG | A071245 001 | Feb 09, 1987 |
| | 1MG | A071246 001 | Feb 09, 1987 |
| | 2MG | A071247 001 | Feb 09, 1987 |
| USL PHARMA | 1MG | A070539 001 | Dec 22, 1986 |
| | 2MG | A070540 001 | Dec 22, 1986 |
| WARNER CHILCOTT | 1MG | A071038 001 | Jan 12, 1988 |
| | 2MG | A071039 001 | Jan 12, 1988 |
| WATSON LABS | 0.5MG | A071086 001 | Mar 23, 1987 |
| | 0.5MG | A071117 001 | Jul 24, 1986 |
| | 1MG | A071087 001 | Mar 23, 1987 |
| | 1MG | A071118 001 | Jul 24, 1986 |
| | 2MG | A071088 001 | Mar 23, 1987 |
| | 2MG | A071110 001 | Jul 24, 1986 |

LOSARTAN POTASSIUM

TABLET; ORAL

COZAAR

| | | | |
|---------------------|------|-------------|--------------|
| + MERCK SHARP DOHME | 25MG | N020386 001 | Apr 14, 1995 |
| + | 50MG | N020386 002 | Apr 14, 1995 |

LOSARTAN POTASSIUM

| | | | |
|-------------|-------|-------------|--------------|
| APOTEX CORP | 25MG | A090790 001 | Oct 06, 2010 |
| | 50MG | A090790 002 | Oct 06, 2010 |
| | 100MG | A090790 003 | Oct 06, 2010 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

LOTEPREDNOL ETABONATE

SUSPENSION/DROPS;OPHTHALMIC

LOTEMAX

PHARMOS

0.5%

N020841 001 Mar 09, 1998

LOVASTATIN

TABLET;ORAL

LOVASTATIN

MYLAN

10MG

A075935 001 Dec 17, 2001

20MG

A075935 002 Dec 17, 2001

40MG

A075935 003 Dec 17, 2001

MEVACOR

+ MERCK

10MG **

N019643 002 Mar 28, 1991

+

20MG **

N019643 003 Aug 31, 1987

+

40MG **

N019643 004 Dec 14, 1988

TABLET, EXTENDED RELEASE;ORAL

ALTOPREV

COVIS PHARMA BV

10MG

N021316 001 Jun 26, 2002

LOXAPINE HYDROCHLORIDE

CONCENTRATE;ORAL

LOXITANE C

ACTAVIS LABS UT INC

EQ 25MG BASE/ML

N017658 001

INJECTABLE;INJECTION

LOXITANE IM

ACTAVIS LABS UT INC

EQ 50MG BASE/ML

N018039 001

LOXAPINE SUCCINATE

CAPSULE;ORAL

LOXITANE

+ ACTAVIS LABS UT INC

EQ 5MG BASE **

N017525 001

+

EQ 10MG BASE **

N017525 002

+

EQ 25MG BASE **

N017525 003

+

EQ 50MG BASE **

N017525 004

TABLET;ORAL

LOXITANE

+ ACTAVIS LABS UT INC

EQ 10MG BASE **

N017525 006

+

EQ 25MG BASE **

N017525 007

+

EQ 50MG BASE **

N017525 008

LUCINACTANT

SUSPENSION;INTRATRACHEAL

SURFAXIN

WINDTREE THERAP

8.5ML

N021746 001 Mar 06, 2012

LUTROPIN ALFA

INJECTABLE;SUBCUTANEOUS

LUVERIS

EMD SERONO

75 IU/VIAL

N021322 001 Oct 08, 2004

LYPRESSIN

SOLUTION;NASAL

DIAPID

NOVARTIS

0.185MG/ML

N016755 001

MAGNESIUM ACETATE TETRAHYDRATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE;INJECTION

PLASMA-LYTE 56 IN PLASTIC CONTAINER

BAXTER HLTHCARE

32MG/100ML;128MG/100ML;234MG/100ML

N019047 001 Jun 15, 1984

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE;INJECTION

ISOLYTE S PH 7.4 IN PLASTIC CONTAINER

B BRAUN

30MG/100ML;37MG/100ML;0.82MG/100ML;370MG/100ML;530MG/100ML;500MG/100ML;12MG/100ML

N019006 001 Apr 04, 1984

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

ISOLYTE S IN PLASTIC CONTAINER

| | | | |
|---------|-------------------------------------------------------------------|-------------|--|
| B BRAUN | 30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG /100ML; 500MG/100ML | N018252 001 | |
|---------|-------------------------------------------------------------------|-------------|--|

SOLUTION; IRRIGATION

PHYSIOSOL IN PLASTIC CONTAINER

| | | | |
|-------------|-------------------------------------------------------------------|-------------|--|
| HOSPIRA INC | 14MG/100ML; 37MG/100ML; 222MG/100ML; 526MG /100ML; 502MG/100ML | N018406 001 | |
|-------------|-------------------------------------------------------------------|-------------|--|

PHYSIOSOL PH 7.4 IN PLASTIC CONTAINER

| | | | |
|-------------|-------------------------------------------------------------------|-------------|--------------|
| HOSPIRA INC | 30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG /100ML; 502MG/100ML | N018406 002 | Jul 08, 1982 |
|-------------|-------------------------------------------------------------------|-------------|--------------|

SYNOVALYTE IN PLASTIC CONTAINER

| | | | |
|-----------------|-------------------------------------------------------------------|-------------|--------------|
| BAXTER HLTHCARE | 30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG /100ML; 502MG/100ML | N019326 001 | Jan 25, 1985 |
|-----------------|-------------------------------------------------------------------|-------------|--------------|

MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE

TABLET; ORAL

MAGNESIUM HYDROXIDE AND OMEPRAZOLE AND SODIUM BICARBONATE

| | | | |
|----------|--------------------|-------------|--------------|
| SANTARUS | 343MG; 20MG; 750MG | N022456 001 | Dec 04, 2009 |
| | 343MG; 40MG; 750MG | N022456 002 | Dec 04, 2009 |

TABLET, CHEWABLE; ORAL

ZEGERID

| | | | |
|----------|--------------------|-------------|--------------|
| SANTARUS | 700MG; 20MG; 600MG | N021850 001 | Mar 24, 2006 |
| | 700MG; 40MG; 600MG | N021850 002 | Mar 24, 2006 |

MAGNESIUM SULFATE; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; POTASSIUM SULFATE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE

SOLUTION; ORAL

SUCLEAR

| | | | |
|------------------|--------------------------------------------------------------------------------------------------------------|-------------|--------------|
| + BRAINTREE LABS | 1.6GM/BOT, 3.13GM/BOT, 17.5GM/BOT, N/A, N/A , N/A, N/A; N/A, N/A, N/A, 210GM, 0.74GM, 2.86GM, 5.6GM ** | N203595 001 | Jan 18, 2013 |
|------------------|--------------------------------------------------------------------------------------------------------------|-------------|--------------|

MALATHION

LOTION; TOPICAL

MALATHION

| | | | |
|------------------|------|-------------|--------------|
| MYLAN PHARMS INC | 0.5% | A078743 001 | Mar 06, 2009 |
|------------------|------|-------------|--------------|

MANGAFODIPIR TRISODIUM

INJECTABLE; INJECTION

TESLASCAN

| | | | |
|------------|-----------|-------------|--------------|
| IC TARGETS | 37.9MG/ML | N020652 001 | Nov 26, 1997 |
|------------|-----------|-------------|--------------|

MANGANESE CHLORIDE TETRAHYDRATE

FOR SOLUTION; ORAL

LUMENHANCE

| | | | |
|--------|-----------|-------------|--------------|
| BRACCO | 3.49MG/GM | N020686 001 | Dec 19, 1997 |
|--------|-----------|-------------|--------------|

MANGANESE SULFATE

INJECTABLE; INJECTION

MANGANESE SULFATE

| | | | |
|---------------|-----------------------|-------------|--------------|
| ABRAXIS PHARM | EQ 0.1MG MANGANESE/ML | N019228 001 | May 05, 1987 |
|---------------|-----------------------|-------------|--------------|

MANNITOL

INJECTABLE; INJECTION

MANNITOL 10%

| | | | |
|---------|------------|-------------|--|
| B BRAUN | 10GM/100ML | N016080 002 | |
| HOSPIRA | 10GM/100ML | N016269 002 | |
| MILES | 10GM/100ML | N016472 002 | |

MANNITOL 10% IN PLASTIC CONTAINER

| | | | |
|-----------------|------------|-------------|--------------|
| ICU MEDICAL INC | 10GM/100ML | N019603 002 | Jan 08, 1987 |
|-----------------|------------|-------------|--------------|

MANNITOL 10% W/ DEXTROSE 5% IN DISTILLED WATER

| | | | |
|---------|------------|-------------|--|
| B BRAUN | 10GM/100ML | N016080 006 | |
|---------|------------|-------------|--|

MANNITOL 15%

| | | | |
|---------|------------|-------------|--|
| B BRAUN | 15GM/100ML | N016080 003 | |
| HOSPIRA | 15GM/100ML | N016269 003 | |
| MILES | 15GM/100ML | N016472 005 | |

MANNITOL 15% IN PLASTIC CONTAINER

| | | | |
|-----------------|------------|-------------|--------------|
| ICU MEDICAL INC | 15GM/100ML | N019603 003 | Jan 08, 1990 |
|-----------------|------------|-------------|--------------|

MANNITOL 15% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.45%

| | | | |
|---------|------------|-------------|--|
| B BRAUN | 15GM/100ML | N016080 005 | |
|---------|------------|-------------|--|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MANNITOL

INJECTABLE; INJECTION

MANNITOL 20%

| | | | |
|---------|------------|---------|-----|
| B BRAUN | 20GM/100ML | N014738 | 001 |
| | 20GM/100ML | N016080 | 004 |
| HOSPIRA | 20GM/100ML | N016269 | 004 |
| MILES | 20GM/100ML | N016472 | 004 |

MANNITOL 25%

| | | | |
|---------------|-------------|---------|------------------|
| ABRAXIS PHARM | 12.5GM/50ML | A086754 | 001 |
| HOSPIRA | 12.5GM/50ML | N016269 | 005 |
| IGI LABS INC | 12.5GM/50ML | A089239 | 001 May 06, 1987 |
| | 12.5GM/50ML | A089240 | 001 May 06, 1987 |
| MERCK | 12.5GM/50ML | N005620 | 001 |
| WATSON LABS | 12.5GM/50ML | A087460 | 001 Jun 27, 1983 |

MANNITOL 5%

| | | | |
|---------|-----------|---------|-----|
| B BRAUN | 5GM/100ML | N016080 | 001 |
| HOSPIRA | 5GM/100ML | N016269 | 001 |

MANNITOL 5% IN PLASTIC CONTAINER

| | | | |
|-----------------|-----------|---------|------------------|
| ICU MEDICAL INC | 5GM/100ML | N019603 | 001 Jan 08, 1987 |
|-----------------|-----------|---------|------------------|

MANNITOL 5% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.12%

| | | | |
|---------|-----------|---------|-----|
| B BRAUN | 5GM/100ML | N016080 | 007 |
|---------|-----------|---------|-----|

SOLUTION; IRRIGATION

RESECTISOL

| | | | |
|---------|-----------|---------|-----|
| B BRAUN | 5GM/100ML | N016704 | 002 |
|---------|-----------|---------|-----|

MANNITOL; SORBITOL

SOLUTION; IRRIGATION

SORBITOL-MANNITOL

| | | | |
|---------|--------------------------|---------|-----|
| HOSPIRA | 540MG/100ML; 2.7GM/100ML | A080224 | 001 |
|---------|--------------------------|---------|-----|

SORBITOL-MANNITOL IN PLASTIC CONTAINER

| | | | |
|---------|--------------------------|---------|-----|
| HOSPIRA | 540MG/100ML; 2.7GM/100ML | N017636 | 001 |
|---------|--------------------------|---------|-----|

MAPROTILINE HYDROCHLORIDE

TABLET; ORAL

LUDIOMIL

| | | | |
|----------|------|---------|------------------|
| NOVARTIS | 25MG | N017543 | 001 |
| | 50MG | N017543 | 002 |
| | 75MG | N017543 | 003 Sep 30, 1982 |

MAPROTILINE HYDROCHLORIDE

| | | | |
|------------------|------|---------|------------------|
| AM THERAP | 25MG | A072129 | 001 Jan 14, 1988 |
| | 50MG | A072130 | 001 Jan 14, 1988 |
| | 75MG | A072131 | 001 Jan 14, 1988 |
| WATSON LABS | 25MG | A071943 | 001 Dec 30, 1987 |
| | 50MG | A071944 | 001 Dec 30, 1987 |
| | 75MG | A071945 | 001 Dec 30, 1987 |
| | 75MG | A072164 | 001 Jun 01, 1988 |
| WATSON LABS TEVA | 25MG | A072162 | 001 Jun 01, 1988 |
| | 50MG | A072163 | 001 Jun 01, 1988 |

MASOPROCOL

CREAM; TOPICAL

ACTINEX

| | | | |
|--------------------|-----|---------|------------------|
| UNIV AZ CANCER CTR | 10% | N019940 | 001 Sep 04, 1992 |
|--------------------|-----|---------|------------------|

MAZINDOL

TABLET; ORAL

MAZANOR

| | | | |
|--------------|-----|---------|-----|
| WYETH AYERST | 1MG | N017980 | 002 |
| | 2MG | N017980 | 001 |

SANOREX

| | | | |
|---------|--------|---------|-----|
| + HEXIM | 1MG ** | N017247 | 001 |
| + | 2MG ** | N017247 | 002 |

MEBENDAZOLE

TABLET, CHEWABLE; ORAL

VERMOX

| | | | |
|------------------|----------|---------|------------------|
| + JANSSEN PHARMS | 100MG ** | N017481 | 001 |
| + | 500MG | N208398 | 001 Oct 19, 2016 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEBUTAMATETABLET; ORAL
DORMATE

MEDPOINTE PHARM HLC 600MG N017374 001

MECAMYLAMINE HYDROCHLORIDETABLET; ORAL
INVERSINE

+ TARGACEPT 2.5MG ** N010251 001

MECASERMIN RINFABATE RECOMBINANTINJECTABLE; SUBCUTANEOUS
IPLEX

INSMED 36MG/0.6ML N021884 001 Dec 12, 2005

MECHLORETHAMINE HYDROCHLORIDEINJECTABLE; INJECTION
MUSTARGEN

+ RECORDATI RARE 10MG/VIAL N006695 001

MECLIZINE HYDROCHLORIDETABLET; ORAL
ANTIVERT

CASPER PHARMA LLC 12.5MG N010721 006

25MG N010721 004

50MG N010721 001 Jan 20, 1982

MECLIZINE HYDROCHLORIDE

ABC HOLDING 12.5MG A085253 001

25MG A085252 001

AMNEAL PHARMS 50MG A201451 003 Feb 23, 2011

ANABOLIC 25MG A085891 001

ANI PHARMS INC 12.5MG A084975 001

25MG A084657 001

BUNDY 12.5MG A084382 001

25MG A084872 001

IVAX SUB TEVA PHARMS 12.5MG A083784 001

KV PHARM 12.5MG A085524 001

25MG A085523 001

MYLAN PHARMS INC 50MG A202640 003 Sep 17, 2012

PAR PHARM 50MG A089674 001 Mar 31, 1988

PLIVA 12.5MG A088732 001 Dec 11, 1985

25MG A088734 001 Dec 11, 1985

RISING PHARMS 12.5MG A040179 001 Jan 30, 1997

25MG A040179 002 Jan 30, 1997

SUPERPHARM 12.5MG A089113 001 Aug 20, 1985

25MG A089114 001 Aug 20, 1985

UDL 12.5MG A088256 001 Jun 13, 1983

25MG A088257 001 Jun 13, 1983

VANGARD 12.5MG A087877 001 Apr 20, 1982

25MG A087620 001 Jan 04, 1982

WATSON LABS 12.5MG A085195 001

12.5MG A085269 001

25MG A085740 001

TABLET, CHEWABLE; ORAL

ANTIVERT

CASPER PHARMA LLC 25MG N010721 005

MECLIZINE HYDROCHLORIDE

IVAX SUB TEVA PHARMS 25MG A084976 001

NEXGEN PHARMA INC 25MG A086392 001

PLIVA 25MG A088733 001 Dec 11, 1985

MECLOCYCLINE SULFOSALICYLATE

CREAM; TOPICAL

MECLAN

JOHNSON AND JOHNSON 1% N050518 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MECLOFENAMATE SODIUM

CAPSULE; ORAL

MECLODIUM

| | | | | |
|------------------|---------------|---------|-----|--------------|
| QUANTUM PHARMICS | EQ 50MG BASE | A071380 | 001 | Jul 14, 1987 |
| | EQ 100MG BASE | A071381 | 001 | Jul 14, 1987 |

MECLOFENAMATE SODIUM

| | | | | |
|------------------|---------------|---------|-----|--------------|
| AM THERAP | EQ 50MG BASE | A071362 | 001 | Feb 10, 1987 |
| | EQ 100MG BASE | A071363 | 001 | Feb 10, 1987 |
| BARR | EQ 50MG BASE | A072848 | 001 | Mar 20, 1989 |
| | EQ 100MG BASE | A072809 | 001 | Mar 20, 1989 |
| FOSUN PHARMA | EQ 50MG BASE | A072262 | 001 | Nov 29, 1988 |
| | EQ 100MG BASE | A072263 | 001 | Nov 29, 1988 |
| PAR PHARM | EQ 50MG BASE | A072077 | 001 | Mar 10, 1988 |
| | EQ 100MG BASE | A072078 | 001 | Mar 10, 1988 |
| USL PHARMA | EQ 50MG BASE | A071007 | 001 | Mar 25, 1988 |
| | EQ 100MG BASE | A071008 | 001 | Mar 25, 1988 |
| VITARINE | EQ 50MG BASE | A071710 | 001 | Jun 15, 1988 |
| | EQ 100MG BASE | A071684 | 001 | Jun 15, 1988 |
| WATSON LABS | EQ 50MG BASE | A070400 | 001 | Nov 25, 1986 |
| | EQ 50MG BASE | A071468 | 001 | Apr 15, 1987 |
| | EQ 50MG BASE | A071640 | 001 | Aug 11, 1987 |
| | EQ 100MG BASE | A070401 | 001 | Nov 25, 1986 |
| | EQ 100MG BASE | A071641 | 001 | Aug 11, 1987 |
| WATSON LABS TEVA | EQ 100MG BASE | A071469 | 001 | Apr 15, 1987 |

MECLOMEN

| | | | | |
|-------------|---------------|---------|-----|--|
| PARKE DAVIS | EQ 50MG BASE | N018006 | 001 | |
| | EQ 100MG BASE | N018006 | 002 | |

MEDROXYPROGESTERONE ACETATE

INJECTABLE; INJECTION

DEPO-PROVERA

| | | | | |
|------------------------|-------------|---------|-----|--|
| + PHARMACIA AND UPJOHN | 100MG/ML ** | N012541 | 002 | |
|------------------------|-------------|---------|-----|--|

MEDROXYPROGESTERONE ACETATE

| | | | | |
|-----------------|----------|---------|-----|--------------|
| SANDOZ INC | 150MG/ML | A078711 | 001 | May 20, 2009 |
| TEVA PHARMS USA | 150MG/ML | A076552 | 001 | Oct 27, 2004 |

TABLET; ORAL

AMEN

| | | | | |
|---------------|-------|---------|-----|--------------|
| AMARIN PHARMS | 10MG | A083242 | 001 | |
| CURRETAB | | | | |
| SOLVAY | 10MG | A085686 | 001 | |
| CYCRIN | | | | |
| ESI | 2.5MG | A081239 | 001 | Oct 30, 1992 |
| | 5MG | A081240 | 001 | Oct 30, 1992 |
| | 10MG | A089386 | 001 | Sep 09, 1987 |

MEDROXYPROGESTERONE ACETATE

| | | | | |
|---------------------|-------|---------|-----|--------------|
| DURAMED PHARMS BARR | 2.5MG | A040311 | 001 | Dec 01, 1999 |
| | 5MG | A040311 | 002 | Dec 01, 1999 |
| | 10MG | A040311 | 003 | Dec 01, 1999 |
| USL PHARMA | 10MG | A088484 | 001 | Jul 26, 1984 |

MEDRYSONE

SUSPENSION; OPHTHALMIC

HMS

| | | | | |
|----------|----|---------|-----|--|
| ALLERGAN | 1% | N016624 | 003 | |
|----------|----|---------|-----|--|

MEFLOQUINE HYDROCHLORIDE

TABLET; ORAL

LARIAM

| | | | | |
|---------|----------|---------|-----|--------------|
| + ROCHE | 250MG ** | N019591 | 001 | May 02, 1989 |
|---------|----------|---------|-----|--------------|

MEFLOQUINE HYDROCHLORIDE

| | | | | |
|---------------------|----------|---------|-----|--------------|
| HIKMA INTL PHARMS | 250MG | A077699 | 001 | Apr 21, 2010 |
| SANDOZ | 250MG | A076175 | 001 | Feb 20, 2002 |
| US ARMY WALTER REED | 250MG ** | N019578 | 001 | May 02, 1989 |

MEGESTROL ACETATE

SUSPENSION; ORAL

MEGACE

| | | | | |
|------------------------|------------|---------|-----|--------------|
| + BRISTOL MYERS SQUIBB | 40MG/ML ** | N020264 | 001 | Sep 10, 1993 |
|------------------------|------------|---------|-----|--------------|

MEGESTROL ACETATE

| | | | | |
|------------|---------|---------|-----|--------------|
| APOTEX INC | 40MG/ML | A077404 | 001 | Feb 16, 2006 |
|------------|---------|---------|-----|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEGESTROL ACETATE

TABLET; ORAL

MEGACE

| | | | | |
|---|----------------------|---------|--|-------------|
| + | BRISTOL MYERS SQUIBB | 20MG ** | | N016979 001 |
| + | | 40MG ** | | N016979 002 |

MEGESTROL ACETATE

| | | | | | |
|--|------------|------|--|-------------|--------------|
| | TEVA | 40MG | | A074745 001 | Feb 27, 1998 |
| | USL PHARMA | 20MG | | A070646 001 | Oct 02, 1987 |
| | | 40MG | | A070647 001 | Oct 02, 1987 |

MELOXICAM

SUSPENSION; ORAL

MOBIC

| | | | | | |
|---|----------------------|--------------|--|-------------|--------------|
| + | BOEHRINGER INGELHEIM | 7.5MG/5ML ** | | N021530 001 | Jun 01, 2004 |
|---|----------------------|--------------|--|-------------|--------------|

TABLET; ORAL

MELOXICAM

| | | | | | |
|--|--------------------|-------|--|-------------|--------------|
| | ANDA REPOSITORY | 7.5MG | | A077935 001 | Jul 19, 2006 |
| | | 15MG | | A077935 002 | Jul 19, 2006 |
| | CR DOUBLE CRANE | 7.5MG | | A078039 001 | Dec 14, 2006 |
| | | 15MG | | A078039 002 | Dec 14, 2006 |
| | IMPAX LABS INC | 7.5MG | | A077930 001 | Jul 19, 2006 |
| | | 15MG | | A077930 002 | Jul 19, 2006 |
| | MYLAN | 7.5MG | | A077923 001 | Jul 19, 2006 |
| | | 7.5MG | | A077934 001 | Jul 20, 2006 |
| | | 15MG | | A077923 002 | Jul 19, 2006 |
| | | 15MG | | A077934 002 | Jul 20, 2006 |
| | ROXANE | 7.5MG | | A077925 001 | Jul 19, 2006 |
| | | 15MG | | A077925 002 | Jul 19, 2006 |
| | SUN PHARM INDS INC | 7.5MG | | A077937 001 | Jul 19, 2006 |
| | | 15MG | | A077937 002 | Jul 19, 2006 |
| | YABAO PHARM | 7.5MG | | A077933 001 | Jul 19, 2006 |
| | | 15MG | | A077933 002 | Jul 19, 2006 |

MELPHALAN HYDROCHLORIDE

INJECTABLE; INJECTION

ALKERAN

| | | | | | |
|---|------------|----------------------|--|-------------|--------------|
| + | APOTEX INC | EQ 50MG BASE/VIAL ** | | N020207 001 | Nov 18, 1992 |
|---|------------|----------------------|--|-------------|--------------|

MELPHALAN HYDROCHLORIDE

| | | | | | |
|--|---------------------|-------------------|--|-------------|--------------|
| | MYLAN INSTITUTIONAL | EQ 50MG BASE/VIAL | | A090299 001 | Oct 27, 2009 |
|--|---------------------|-------------------|--|-------------|--------------|

MEMANTINE HYDROCHLORIDE

SOLUTION; ORAL

NAMENDA

| | | | | | |
|---|--------------------|--------|--|-------------|--------------|
| + | ALLERGAN SALES LLC | 2MG/ML | | N021627 001 | Apr 18, 2005 |
|---|--------------------|--------|--|-------------|--------------|

TABLET; ORAL

MEMANTINE HYDROCHLORIDE

| | | | | | |
|--|-----------------|------|--|-------------|--------------|
| | ORCHID HLTHCARE | 5MG | | A090044 001 | Mar 12, 2012 |
| | | 10MG | | A090044 002 | Mar 12, 2012 |

MENADIOL SODIUM DIPHOSPHATE

INJECTABLE; INJECTION

KAPPADIONE

| | | | | | |
|--|-------|---------|--|-------------|--|
| | LILLY | 10MG/ML | | N005725 001 | |
|--|-------|---------|--|-------------|--|

SYNKAYVITE

| | | | | | |
|--|-------|-----------|--|-------------|--|
| | ROCHE | 5MG/ML | | N003718 004 | |
| | | 10MG/ML | | N003718 006 | |
| | | 37.5MG/ML | | N003718 008 | |

TABLET; ORAL

SYNKAYVITE

| | | | | | |
|--|-------|-----|--|-------------|--|
| | ROCHE | 5MG | | N003718 010 | |
|--|-------|-----|--|-------------|--|

MENADIONE

TABLET; ORAL

MENADIONE

| | | | | | |
|--|-------|-----|--|-------------|--|
| | LILLY | 5MG | | N002139 003 | |
|--|-------|-----|--|-------------|--|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MENOTROPINS (FSH;LH)

INJECTABLE; INJECTION

HUMEGON

| | | | |
|-----------------|-------------------------|-------------|--------------|
| ORGANON USA INC | 75 IU/VIAL;75 IU/VIAL | N020328 001 | Sep 01, 1994 |
| | 150 IU/VIAL;150 IU/VIAL | N020328 002 | Sep 01, 1994 |

MENOTROPINS

| | | | |
|---------|-------------------------|-------------|--------------|
| FERRING | 75 IU/VIAL;75 IU/VIAL | A073598 001 | Jan 30, 1997 |
| | 150 IU/VIAL;150 IU/VIAL | A073599 001 | Jan 30, 1997 |

PERGONAL

| | | | |
|--------|-----------------------|-------------|--------------|
| SERONO | 75 IU/AMP;75 IU/AMP | N017646 001 | |
| | 150 IU/AMP;150 IU/AMP | N017646 002 | May 20, 1985 |

REPRONEX

| | | | |
|---------|-------------------------|-------------|--------------|
| FERRING | 150 IU/VIAL;150 IU/VIAL | N021047 002 | Aug 27, 1999 |
|---------|-------------------------|-------------|--------------|

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

REPRONEX

| | | | |
|---------|-----------------------|-------------|--------------|
| FERRING | 75 IU/VIAL;75 IU/VIAL | N021047 001 | Aug 27, 1999 |
|---------|-----------------------|-------------|--------------|

MEPENZOLATE BROMIDE

SOLUTION; ORAL

CANTIL

| | | | |
|-------------------|----------|-------------|--|
| SANOFI AVENTIS US | 25MG/5ML | N010679 004 | |
|-------------------|----------|-------------|--|

TABLET; ORAL

CANTIL

| | | | |
|---------------------|------|-------------|--|
| + SANOFI AVENTIS US | 25MG | N010679 003 | |
|---------------------|------|-------------|--|

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DEMEROL

| | | | |
|-------------------|----------|-------------|--|
| US PHARM HOLDINGS | 25MG/ML | N005010 007 | |
| | 50MG/ML | N005010 002 | |
| | 75MG/ML | N005010 009 | |
| | 100MG/ML | N005010 003 | |

MEPERIDINE HYDROCHLORIDE

ABBOTT

| | | | |
|--|----------|-------------|--|
| | 25MG/ML | A080388 001 | |
| | 50MG/ML | A080385 001 | |
| | 50MG/ML | A080387 001 | |
| | 75MG/ML | A080389 001 | |
| | 100MG/ML | A080386 001 | |

BAXTER HLTHCARE

| | | | |
|--|----------|-------------|--------------|
| | 25MG/ML | A088279 001 | Jun 15, 1984 |
| | 50MG/ML | A088280 001 | Jun 15, 1984 |
| | 75MG/ML | A088281 001 | Jun 15, 1984 |
| | 100MG/ML | A088282 001 | Jun 15, 1984 |

IGI LABS INC

| | | | |
|--|----------|-------------|--------------|
| | 25MG/ML | A089781 001 | Mar 31, 1989 |
| | 50MG/ML | A089782 001 | Mar 31, 1989 |
| | 50MG/ML | A089783 001 | Mar 31, 1989 |
| | 50MG/ML | A089784 001 | Mar 31, 1989 |
| | 75MG/ML | A089785 001 | Mar 31, 1989 |
| | 100MG/ML | A089786 001 | Mar 31, 1989 |
| | 100MG/ML | A089787 001 | Mar 31, 1989 |
| | 100MG/ML | A089788 001 | Mar 31, 1989 |

INTL MEDICATION

| | | | |
|--|---------|-------------|--|
| | 10MG/ML | A086332 001 | |
|--|---------|-------------|--|

PARKE DAVIS

| | | | |
|--|---------|-------------|--|
| | 50MG/ML | A080364 002 | |
|--|---------|-------------|--|

| | | | |
|--|---------|-------------|--|
| | 75MG/ML | A080364 003 | |
|--|---------|-------------|--|

| | | | |
|--|----------|-------------|--|
| | 100MG/ML | A080364 001 | |
|--|----------|-------------|--|

| | | | |
|-------------|---------|-------------|--------------|
| WATSON LABS | 50MG/ML | A073444 001 | Mar 17, 1992 |
|-------------|---------|-------------|--------------|

| | | | |
|--|----------|-------------|--------------|
| | 100MG/ML | A073445 001 | Mar 17, 1992 |
|--|----------|-------------|--------------|

MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE

| | | | |
|---------|---------|-------------|--------------|
| HOSPIRA | 10MG/ML | A040305 001 | Mar 10, 1999 |
|---------|---------|-------------|--------------|

| | | | |
|-----------------|---------|-------------|--------------|
| ICU MEDICAL INC | 10MG/ML | A088432 001 | Aug 16, 1984 |
|-----------------|---------|-------------|--------------|

| | | | |
|-----------------|---------|-------------|--------------|
| INTL MEDICATION | 10MG/ML | A081309 001 | Aug 30, 1993 |
|-----------------|---------|-------------|--------------|

| | | | |
|------------|---------|-------------|--------------|
| SPECGX LLC | 10MG/ML | A040163 001 | May 12, 1997 |
|------------|---------|-------------|--------------|

| | | | |
|-------------|---------|-------------|--------------|
| WATSON LABS | 10MG/ML | A073443 001 | Mar 17, 1992 |
|-------------|---------|-------------|--------------|

SYRUP; ORAL

DEMEROL

| | | | |
|-------------------|-------------|-------------|--|
| US PHARM HOLDINGS | 50MG/5ML ** | N005010 005 | |
|-------------------|-------------|-------------|--|

TABLET; ORAL

MEPERIDINE HYDROCHLORIDE

| | | | |
|------|------|-------------|--------------|
| BARR | 50MG | A088639 001 | Jul 02, 1984 |
|------|------|-------------|--------------|

| | | | |
|--|-------|-------------|--------------|
| | 100MG | A088640 001 | Sep 19, 1984 |
|--|-------|-------------|--------------|

| | | | |
|---------------------|------|-------------|--------------|
| DURAMED PHARMS BARR | 50MG | A040318 001 | Oct 05, 1999 |
|---------------------|------|-------------|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEPERIDINE HYDROCHLORIDE

TABLET; ORAL

MEPERIDINE HYDROCHLORIDE

| | | | |
|----------------------|-------|-------------|--------------|
| | 100MG | A040318 002 | Oct 05, 1999 |
| SUN PHARM INDUSTRIES | 50MG | A080448 001 | |
| | 100MG | A080448 002 | |
| WATSON LABS | 50MG | A040186 001 | Jun 30, 1997 |
| | 100MG | A040186 002 | Jun 30, 1997 |
| WYETH AYERST | 50MG | A080454 001 | |

MEPERIDINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPERGAN

| | | | |
|----------------------|------------------|-------------|--|
| WEST-WARD PHARMS INT | 25MG/ML; 25MG/ML | N011730 001 | |
|----------------------|------------------|-------------|--|

MEPHENTERMINE SULFATE

INJECTABLE; INJECTION

WYAMINE SULFATE

| | | | |
|----------------------|-----------------|-------------|--|
| BAXTER HLTHCARE CORP | EQ 15MG BASE/ML | N008248 002 | |
| | EQ 30MG BASE/ML | N008248 001 | |

MEPHENYTOIN

TABLET; ORAL

MESANTOIN

| | | | |
|------------|----------|-------------|--|
| + NOVARTIS | 100MG ** | N006008 001 | |
|------------|----------|-------------|--|

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARESTOCAINE HYDROCHLORIDE

| | | | |
|--------|----|-------------|--------------|
| SOLVAY | 3% | A084777 002 | Apr 18, 1982 |
|--------|----|-------------|--------------|

CARBOCAINE

| | | | |
|-----------------|-------|-------------|--|
| + EASTMAN KODAK | 3% ** | N012125 003 | |
|-----------------|-------|-------------|--|

ISOCAINE HYDROCHLORIDE

| | | | |
|---------------|----|-------------|--|
| SEPTODONT INC | 3% | A080925 001 | |
|---------------|----|-------------|--|

MEPIVACAINE HYDROCHLORIDE

| | | | |
|---------------------|----|-------------|--------------|
| BELMORA LLC | 3% | A083559 001 | |
| HOSPIRA INC | 3% | A040806 001 | Apr 28, 2008 |
| INTL MEDICATION SYS | 1% | A087509 001 | Oct 05, 1982 |
| WATSON LABS | 1% | A088769 001 | Nov 20, 1984 |
| | 2% | A088770 001 | Nov 20, 1984 |
| POLOCAINE | | | |
| DENTSPLY PHARM | 3% | A088653 001 | Aug 21, 1984 |

MEPREDNISONE

TABLET; ORAL

BETAPAR

| | | | |
|----------|-----|-------------|--|
| SCHERING | 4MG | N016053 002 | |
|----------|-----|-------------|--|

MEPROBAMATE

CAPSULE; ORAL

EQUANIL

| | | | |
|--------------|-------|-------------|--|
| WYETH AYERST | 400MG | N012455 002 | |
|--------------|-------|-------------|--|

CAPSULE, EXTENDED RELEASE; ORAL

MEPROSPAN

| | | | |
|---------------------|-------|-------------|--|
| MEDPOINTE PHARM HLC | 200MG | N011284 001 | |
| | 400MG | N011284 002 | |

TABLET; ORAL

AMOSENE

| | | | |
|--------------|-------|-------------|--|
| FERNDAL LABS | 400MG | A084030 001 | |
|--------------|-------|-------------|--|

BAMATE

| | | | |
|------|-------|-------------|--|
| ALRA | 200MG | A080380 001 | |
| | 400MG | A080380 002 | |

EQUANIL

| | | | |
|--------------|-------|-------------|--|
| WYETH AYERST | 200MG | N010028 005 | |
| | 400MG | N010028 004 | |

MEPRIAM

| | | | |
|------|-------|-------------|--|
| TEVA | 400MG | N016069 001 | |
|------|-------|-------------|--|

MEPROBAMATE

| | | | |
|---------------------|-------|-------------|--|
| AUROLIFE PHARMA LLC | 400MG | A080655 001 | |
| BARR | 600MG | A084230 001 | |
| ELKINS SINN | 200MG | N015426 002 | |
| | 400MG | N015426 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEPROBAMATE

TABLET; ORAL

MEPROBAMATE

| | | | | | |
|-----------------------|-------|----|---------|-----|--------------|
| HEATHER | 400MG | | N016928 | 003 | |
| | 600MG | | A084329 | 001 | |
| IMPAX LABS | 200MG | | N014322 | 002 | |
| | 400MG | | N014322 | 001 | |
| IVAX SUB TEVA PHARMS | 200MG | | N015438 | 001 | |
| | 400MG | | N015438 | 002 | |
| | 600MG | | A084181 | 001 | |
| IVC INDS | 400MG | | A084153 | 001 | |
| LANNETT | 200MG | | N014882 | 002 | |
| | 400MG | | N014882 | 001 | |
| LEDERLE | 400MG | | A086299 | 001 | |
| LEE KM | 400MG | | A089538 | 001 | Nov 25, 1987 |
| MALLARD | 400MG | | N015072 | 002 | |
| MK LABS | 200MG | | N014368 | 004 | |
| | 400MG | | N014368 | 002 | |
| MYLAN | 400MG | | A083618 | 001 | |
| NEXGEN PHARMA INC | 200MG | | A084220 | 001 | |
| | 400MG | | A084589 | 001 | |
| PARKE DAVIS | 200MG | | A084744 | 001 | |
| | 400MG | | A084744 | 002 | |
| PERRIGO | 200MG | | A084546 | 001 | |
| | 400MG | | A084547 | 001 | |
| PHARMAVITE | 400MG | | A084438 | 001 | |
| PUREPAC PHARM | 200MG | | A084804 | 001 | |
| | 400MG | | A084804 | 002 | |
| PVT FORM | 400MG | | N014601 | 001 | |
| ROXANE | 600MG | | A084332 | 001 | |
| SANDOZ | 200MG | | N014547 | 002 | |
| | 400MG | | N014547 | 001 | |
| SCHERER LABS | 400MG | | A083343 | 001 | |
| SOLVAY | 200MG | | A084435 | 001 | |
| STANLABS PHARM | 200MG | | N014474 | 002 | |
| | 400MG | | N014474 | 004 | |
| SUN PHARM INDUSTRIES | 200MG | | A080699 | 001 | |
| | 400MG | | A080699 | 002 | |
| TABLICAPS | 400MG | | A083494 | 001 | |
| TARO | 200MG | | A200998 | 001 | May 23, 2011 |
| | 400MG | | A200998 | 002 | May 23, 2011 |
| USL PHARMA | 200MG | | A087825 | 001 | Mar 18, 1982 |
| | 400MG | | A087826 | 001 | Mar 18, 1982 |
| VALEANT PHARM INTL | 200MG | | N015139 | 006 | |
| | 400MG | | N015139 | 005 | |
| VANGARD | 400MG | | A088011 | 001 | Jul 14, 1982 |
| WATSON LABS | 200MG | | A085720 | 001 | |
| | 400MG | | A085721 | 001 | |
| | 600MG | | A084274 | 001 | |
| | 600MG | | A085719 | 001 | |
| WEST WARD | 200MG | | N015417 | 003 | |
| | 400MG | | N015417 | 002 | |
| WHITEWORTH TOWN PLSN | 200MG | | A083830 | 001 | |
| | 400MG | | A083442 | 001 | |
| MILTOWN | | | | | |
| + MEDPOINTE PHARM HLC | 200MG | ** | N009698 | 004 | |
| + | 400MG | ** | N009698 | 002 | |
| | 600MG | | A083919 | 001 | |
| NEURAMATE | | | | | |
| HALSEY | 200MG | | N014359 | 002 | |
| | 400MG | | N014359 | 001 | |
| TRANMEP | | | | | |
| SOLVAY | 400MG | | A084369 | 001 | |
| | 400MG | | N016249 | 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MEQUINOL; TRETINOINSOLUTION; TOPICAL
SOLAGE

AQUA PHARMS 2%;0.01% N020922 001 Dec 10, 1999

MEROPENEMINJECTABLE; INJECTION
MEROPENEMSANDOZ 500MG/VIAL A091201 001 Mar 29, 2011
1GM/VIAL A091201 002 Mar 29, 2011MERSALYL SODIUM; THEOPHYLLINEINJECTABLE; INJECTION
MERSALYL-THEOPHYLLINE

WATSON LABS 100MG/ML; 50MG/ML A084875 001

MESALAMINEENEMA; RECTAL
MESALAMINE

G AND W LABS INC 4GM/60ML A076841 001 Sep 30, 2004

SUPPOSITORY; RECTAL

CANASA

ALLERGAN SALES LLC 500MG N021252 001 Jan 05, 2001

ROWASA

+ MEDA PHARMS 500MG ** N019919 001 Dec 18, 1990

TABLET, DELAYED RELEASE; ORAL

ASACOL

APIL 400MG N019651 001 Jan 31, 1992

MESNAINJECTABLE; INTRAVENOUS
MESNA

MYLAN INSTITUTIONAL 100MG/ML A076488 001 Mar 08, 2012

MYLAN LABS LTD 100MG/ML A203364 001 Jul 18, 2014

MESORIDAZINE BESYLATECONCENTRATE; ORAL
SERENTIL

NOVARTIS EQ 25MG BASE/ML N016997 001

INJECTABLE; INJECTION

SERENTIL

NOVARTIS EQ 25MG BASE/ML N016775 001

TABLET; ORAL

SERENTIL

NOVARTIS EQ 10MG BASE N016774 001

EQ 25MG BASE N016774 002

EQ 50MG BASE N016774 003

EQ 100MG BASE N016774 004

MESTRANOL; NORETHINDRONETABLET; ORAL-20
NORINYL

ACTAVIS LABS UT INC 0.1MG; 2MG N013625 004

TABLET; ORAL-21

NORETHIN 1/50M-21

WATSON LABS 0.05MG; 1MG A071539 001 Apr 12, 1988

NORETHINDRONE AND MESTRANOL

WATSON LABS 0.05MG; 1MG A070758 001 Jul 01, 1988

NORINYL 1+50 21-DAY

ACTAVIS LABS UT INC 0.05MG; 1MG N013625 002

NORINYL 1+80 21-DAY

GD SEARLE LLC 0.08MG; 1MG N016724 001

ORTHO-NOVUM 1/50 21

ORTHO MCNEIL PHARM 0.05MG; 1MG N012728 004

ORTHO-NOVUM 1/80 21

ORTHO MCNEIL PHARM 0.08MG; 1MG N016715 001

ORTHO-NOVUM 10-21

ORTHO MCNEIL PHARM 0.06MG; 10MG N012728 001

ORTHO-NOVUM 2-21

ORTHO MCNEIL PHARM 0.1MG; 2MG N012728 005

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MESTRANOL; NORETHINDRONE

TABLET; ORAL-28

NORETHIN 1/50M-28

WATSON LABS

0.05MG;1MG

A071540 001 Apr 12, 1988

NORETHINDRONE AND MESTRANOL

WATSON LABS

0.05MG;1MG

A070759 001 Jul 01, 1988

NORINYL 1+80 28-DAY

GD SEARLE LLC

0.08MG;1MG

N016725 001

ORTHO-NOVUM 1/50 28

ORTHO MCNEIL JANSSEN

0.05MG;1MG

N016709 001

ORTHO-NOVUM 1/80 28

ORTHO MCNEIL PHARM

0.08MG;1MG

N016715 002

MESTRANOL; NORETHYNODREL

TABLET; ORAL

ENOVID

GD SEARLE LLC

0.075MG;5MG

N010976 008

0.15MG;9.85MG

N010976 005

TABLET; ORAL-20

ENOVID

GD SEARLE LLC

0.075MG;5MG

N010976 004

ENOVID-E

GD SEARLE LLC

0.1MG;2.5MG

N010976 006

TABLET; ORAL-21

ENOVID-E 21

GD SEARLE LLC

0.1MG;2.5MG

N010976 007

METAPROTERENOL SULFATE

AEROSOL, METERED; INHALATION

ALUPENT

BOEHRINGER INGELHEIM

0.65MG/INH

N016402 001

SOLUTION; INHALATION

ALUPENT

BOEHRINGER INGELHEIM

0.4%

N018761 002 Oct 10, 1986

0.6%

N018761 001 Jun 30, 1983

5%

N017659 001

METAPROTERENOL SULFATE

APOTEX INC

0.4%

A075402 001 Feb 28, 2001

0.6%

A075403 001 Feb 28, 2001

ASTRAZENECA

0.4%

A071275 001 Jul 27, 1988

0.6%

A071018 001 Jul 27, 1988

DEY

0.33%

A071806 001 Aug 05, 1988

0.5%

A071805 001 Aug 05, 1988

5%

A070805 001 Aug 17, 1987

MYLAN SPECIALITY LP

0.4%

A071786 001 Aug 05, 1988

0.6%

A070804 001 Aug 17, 1987

NEPHRON

0.4%

A071855 001 Jul 14, 1988

0.6%

A071726 001 Jul 14, 1988

WOCKHARDT

0.4%

A075586 001 May 30, 2002

0.6%

A075586 002 May 30, 2002

5%

A072190 001 Jun 07, 1988

PROMETA

MURO

5%

A073340 001 Mar 30, 1992

SYRUP; ORAL

ALUPENT

BOEHRINGER INGELHEIM

10MG/5ML

N017571 001

METAPROTERENOL SULFATE

APOTEX INC

10MG/5ML

A075235 001 Jan 27, 2000

G AND W LABS INC

10MG/5ML

A072761 001 Feb 27, 1992

10MG/5ML

A073034 001 Aug 30, 1991

MORTON GROVE

10MG/5ML

A071656 001 Oct 13, 1987

WOCKHARDT

10MG/5ML

A074702 001 Mar 24, 1997

PROMETA

MURO

10MG/5ML

A072023 001 Sep 15, 1988

TABLET; ORAL

ALUPENT

BOEHRINGER INGELHEIM

10MG

N015874 002

20MG

N015874 001

METAPROTERENOL SULFATE

AM THERAP

10MG

A072054 001 Jun 23, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METAPROTERENOL SULFATE

TABLET; ORAL

METAPROTERENOL SULFATE

| | | | |
|-------------|------|-------------|--------------|
| | 20MG | A072055 001 | Jun 23, 1988 |
| TEVA | 10MG | A072519 001 | Mar 30, 1990 |
| | 20MG | A072520 001 | Mar 30, 1990 |
| USL PHARMA | 10MG | A071013 001 | Jan 25, 1988 |
| | 20MG | A071014 001 | Jan 25, 1988 |
| WATSON LABS | 10MG | A073013 001 | Jan 31, 1991 |
| | 20MG | A072795 001 | Jan 31, 1991 |

METARAMINOL BITARTRATE

INJECTABLE; INJECTION

ARAMINE

| | | | |
|---------|--------------------|-------------|--------------|
| + MERCK | EQ 10MG BASE/ML ** | N009509 002 | Dec 22, 1987 |
|---------|--------------------|-------------|--------------|

METARAMINOL BITARTRATE

| | | | |
|--------------------|-----------------|-------------|--|
| ABRAXIS PHARM | EQ 10MG BASE/ML | A080431 001 | |
| ELKINS SINN | EQ 10MG BASE/ML | A083363 001 | |
| FRESENIUS KABI USA | EQ 10MG BASE/ML | A080722 001 | |
| GD SEARLE LLC | EQ 10MG BASE/ML | A086418 001 | |
| | EQ 20MG BASE/ML | A086418 002 | |

METAXALONE

TABLET; ORAL

METAXALONE

| | | | |
|-----------------|----------|-------------|--------------|
| + PRIMUS PHARMS | 640MG ** | N022503 001 | Jun 01, 2015 |
|-----------------|----------|-------------|--------------|

SKELAXIN

| | | | |
|---------------|----------|-------------|--|
| + KING PHARMS | 400MG ** | N013217 001 | |
|---------------|----------|-------------|--|

METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLUCOPHAGE

| | | | |
|------------------------|----------|-------------|--------------|
| + BRISTOL MYERS SQUIBB | 625MG ** | N020357 003 | Nov 05, 1998 |
|------------------------|----------|-------------|--------------|

| | | | |
|---|----------|-------------|--------------|
| + | 750MG ** | N020357 004 | Nov 05, 1998 |
|---|----------|-------------|--------------|

METFORMIN HYDROCHLORIDE

| | | | |
|----------------------|-------|-------------|--------------|
| BARR | 500MG | A075971 001 | Jan 25, 2002 |
| | 850MG | A075971 002 | Jan 25, 2002 |
| | 1GM | A075971 003 | Jan 25, 2002 |
| IPCA LABS LTD | 500MG | A078422 001 | Aug 06, 2007 |
| | 850MG | A078422 002 | Aug 06, 2007 |
| | 1GM | A078422 003 | Aug 06, 2007 |
| IVAX SUB TEVA PHARMS | 500MG | A075975 001 | Jan 24, 2002 |
| | 625MG | A075975 004 | Jan 24, 2002 |
| | 750MG | A075975 005 | Jan 24, 2002 |
| | 850MG | A075975 002 | Jan 24, 2002 |
| | 1GM | A075975 003 | Jan 24, 2002 |
| MYLAN PHARMS INC | 500MG | A075969 001 | Jan 29, 2002 |
| | 850MG | A075969 002 | Jan 29, 2002 |
| | 1GM | A075969 003 | Jan 29, 2002 |
| PROVIDENT PHARM | 500MG | A077853 001 | Jul 28, 2006 |
| | 850MG | A077853 002 | Jul 28, 2006 |
| | 1GM | A077853 003 | Jul 28, 2006 |
| SANDOZ | 500MG | A075985 001 | Jan 25, 2002 |
| | 850MG | A075985 002 | Jan 25, 2002 |
| | 1GM | A075985 003 | Jan 25, 2002 |
| TEVA | 500MG | A076328 001 | Dec 16, 2002 |
| | 850MG | A076328 002 | Dec 16, 2002 |
| | 1GM | A076328 003 | Dec 16, 2002 |
| WATSON LABS | 500MG | A075979 001 | Jan 24, 2002 |
| | 850MG | A075979 002 | Jan 24, 2002 |
| | 1GM | A075979 003 | Jan 24, 2002 |
| WATSON LABS FLORIDA | 500MG | A075961 001 | Jan 25, 2002 |
| | 850MG | A075961 002 | Jan 25, 2002 |
| | 1GM | A075961 003 | Jan 25, 2002 |

TABLET, EXTENDED RELEASE; ORAL

METFORMIN HYDROCHLORIDE

| | | | |
|-------------------|-------|-------------|--------------|
| ACTAVIS ELIZABETH | 500MG | A076450 001 | Oct 01, 2004 |
| | 750MG | A076878 001 | Apr 13, 2005 |
| BARR | 500MG | A076496 001 | Nov 25, 2005 |
| IMPAX LABS | 500MG | A076249 001 | Jul 30, 2004 |
| | 750MG | A076985 001 | Sep 13, 2005 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

METFORMIN HYDROCHLORIDE

| | | | | |
|----------------------|-------|---------|-----|--------------|
| IVAX SUB TEVA PHARMS | 500MG | A076545 | 001 | Dec 01, 2003 |
| MYLAN | 500MG | A076650 | 001 | Sep 13, 2005 |
| | 750MG | A077113 | 001 | Sep 08, 2005 |
| RANBAXY LABS LTD | 500MG | A076413 | 001 | Jun 18, 2004 |
| | 750MG | A077211 | 001 | Jun 29, 2005 |
| SANDOZ | 500MG | A076223 | 001 | Dec 14, 2004 |
| SUN PHARM INDUSTRIES | 500MG | A077124 | 001 | Dec 21, 2005 |
| TORRENT PHARMS LTD | 750MG | A079226 | 001 | Feb 18, 2010 |
| WATSON LABS INC | 500MG | A076818 | 001 | Dec 14, 2004 |

METFORMIN HYDROCHLORIDE; REPAGLINIDE

TABLET;ORAL

PRANDIMET

| | | | | |
|--------------------|-----------|---------|-----|--------------|
| + NOVO NORDISK INC | 500MG;1MG | N022386 | 001 | Jun 23, 2008 |
| | 500MG;2MG | N022386 | 002 | Jun 23, 2008 |

REPAGLINIDE AND METFORMIN HYDROCHLORIDE

| | | | | |
|-----------|-----------|---------|-----|--------------|
| LUPIN LTD | 500MG;1MG | A200624 | 001 | Jul 15, 2015 |
| | 500MG;2MG | A200624 | 002 | Jul 15, 2015 |

METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE

TABLET;ORAL

AVANDAMET

| | | | | |
|--------------|----------------------|---------|-----|--------------|
| + SB PHARMCO | 500MG;EQ 1MG BASE ** | N021410 | 001 | Oct 10, 2002 |
| | 500MG;EQ 2MG BASE ** | N021410 | 002 | Oct 10, 2002 |
| | 500MG;EQ 4MG BASE ** | N021410 | 003 | Oct 10, 2002 |
| | 1GM;EQ 2MG BASE ** | N021410 | 004 | Aug 25, 2003 |
| | 1GM;EQ 4MG BASE ** | N021410 | 005 | Aug 25, 2003 |

ROSIGLITAZONE MALEATE AND METFORMIN HYDROCHLORIDE

| | | | | |
|------|-------------------|---------|-----|--------------|
| TEVA | 500MG;EQ 2MG BASE | A077337 | 001 | May 07, 2014 |
| | 500MG;EQ 1MG BASE | A077337 | 005 | May 19, 2017 |
| | 500MG;EQ 4MG BASE | A077337 | 002 | May 07, 2014 |
| | 1GM;EQ 4MG BASE | A077337 | 004 | May 07, 2014 |
| | 1GM;EQ 2MG BASE | A077337 | 003 | May 07, 2014 |

METHACHOLINE CHLORIDE

FOR SOLUTION;INHALATION

PROVOCHOLINE

| | | | | |
|--------------|-------------|---------|-----|--------------|
| + METHAPHARM | 1600MG/VIAL | N019193 | 002 | Aug 29, 2016 |
|--------------|-------------|---------|-----|--------------|

METHACYCLINE HYDROCHLORIDE

CAPSULE;ORAL

RONDONMYCIN

| | | | | |
|---------------------|---------------|---------|-----|--|
| MEDPOINTE PHARM HLC | EQ 140MG BASE | A060641 | 001 | |
| | EQ 280MG BASE | A060641 | 002 | |

SYRUP;ORAL

RONDONMYCIN

| | | | | |
|---------------------|------------------|---------|-----|--|
| MEDPOINTE PHARM HLC | EQ 70MG BASE/5ML | A060641 | 003 | |
|---------------------|------------------|---------|-----|--|

METHADONE HYDROCHLORIDE

POWDER;FOR RX COMPOUNDING

METHADONE HYDROCHLORIDE

| | | | | |
|------------------|-----------|---------|-----|--|
| MALLINCKRODT INC | 50GM/BOT | N006383 | 002 | |
| | 100GM/BOT | N006383 | 003 | |
| | 500GM/BOT | N006383 | 004 | |

SYRUP;ORAL

DOLOPHINE HYDROCHLORIDE

| | | | | |
|----------------------|-----------|---------|-----|--|
| WEST-WARD PHARMS INT | 10MG/30ML | N006134 | 004 | |
|----------------------|-----------|---------|-----|--|

TABLET;ORAL

METHADONE HYDROCHLORIDE

| | | | | |
|------------|------|---------|-----|--------------|
| ROXANE | 5MG | A088108 | 001 | Mar 08, 1983 |
| | 10MG | A088109 | 001 | Mar 08, 1983 |
| | 40MG | A074081 | 001 | Apr 28, 1995 |
| VISTAPHARM | 5MG | A040241 | 001 | May 29, 1998 |

TABLET, DISPERSIBLE;ORAL

WESTADONE

| | | | | |
|--------|-------|---------|-----|--|
| SANDOZ | 2.5MG | N017108 | 001 | |
|--------|-------|---------|-----|--|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHADONE HYDROCHLORIDETABLET, EFFERVESCENT;ORAL
WESTADONE

| | | |
|--------|------|-------------|
| SANDOZ | 5MG | N017108 002 |
| | 10MG | N017108 003 |
| | 40MG | N017108 004 |

METHAMPHETAMINE HYDROCHLORIDE

TABLET;ORAL

METHAMPEX

| | | |
|------|------|-------------|
| TEVA | 10MG | A083889 001 |
|------|------|-------------|

METHAMPHETAMINE HYDROCHLORIDE

| | | | |
|-------|------|-------------|--------------|
| ABLE | 5MG | A040529 001 | Feb 25, 2004 |
| REXAR | 5MG | A084931 001 | |
| | 10MG | A084931 002 | |
| TEVA | 5MG | A086359 001 | |

TABLET, EXTENDED RELEASE;ORAL

DESOXYN

| | | |
|----------------|------|-------------|
| RECORDATI RARE | 5MG | N005378 004 |
| | 10MG | N005378 003 |
| | 15MG | N005378 005 |

METHANTHELINE BROMIDE

TABLET;ORAL

BANTHINE

| | | |
|-------|------|-------------|
| SHIRE | 50MG | N007390 001 |
|-------|------|-------------|

METHARBITAL

TABLET;ORAL

GEMONIL

| | | |
|--------|-------|-------------|
| ABBVIE | 100MG | N008322 001 |
|--------|-------|-------------|

METHAZOLAMIDE

TABLET;ORAL

METHAZOLAMIDE

| | | | |
|--------------|------|-------------|--------------|
| APPLIED ANAL | 25MG | A040011 001 | Jul 17, 1997 |
| | 50MG | A040011 002 | Jul 17, 1997 |
| SANDOZ | 25MG | A040102 001 | Aug 28, 1996 |
| | 50MG | A040102 002 | Aug 28, 1996 |

NEPTAZANE

| | | | |
|-----------|---------|-------------|--------------|
| + LEDERLE | 25MG ** | N011721 002 | Nov 25, 1991 |
| + | 50MG ** | N011721 001 | |

METHDILAZINE

TABLET, CHEWABLE;ORAL

TACARYL

| | | |
|-----------------|-------|-------------|
| WESTWOOD SQUIBB | 3.6MG | N011950 009 |
|-----------------|-------|-------------|

METHDILAZINE HYDROCHLORIDE

SYRUP;ORAL

METHDILAZINE HYDROCHLORIDE

| | | |
|--------------------|---------|-------------|
| ALPHARMA US PHARMS | 4MG/5ML | A087122 001 |
|--------------------|---------|-------------|

TACARYL

| | | |
|-----------------|---------|-------------|
| WESTWOOD SQUIBB | 4MG/5ML | N011950 007 |
|-----------------|---------|-------------|

TABLET;ORAL

TACARYL

| | | |
|-----------------|-----|-------------|
| WESTWOOD SQUIBB | 8MG | N011950 006 |
|-----------------|-----|-------------|

METHICILLIN SODIUM

INJECTABLE; INJECTION

STAPHCILLIN

| | | |
|-----------|--------------------|-------------|
| APOTHECON | EQ 900MG BASE/VIAL | A061449 001 |
| | EQ 900MG BASE/VIAL | N050117 001 |
| | EQ 3.6GM BASE/VIAL | A061449 002 |
| | EQ 3.6GM BASE/VIAL | N050117 002 |
| | EQ 5.4GM BASE/VIAL | A061449 003 |
| | EQ 5.4GM BASE/VIAL | N050117 003 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

| | | | |
|----------------|---------|-------------|--------------|
| ECI PHARMS LLC | 15MG | A040619 003 | Jul 12, 2005 |
| | 20MG | A040547 004 | Feb 18, 2005 |
| MYLAN | 20MG | A040350 003 | Jun 07, 2001 |
| TAPAZOLE | | | |
| + KING PHARMS | 5MG ** | N007517 002 | |
| + | 10MG ** | N007517 004 | |

METHIXENE HYDROCHLORIDE

TABLET; ORAL

TREST

| | | | |
|----------|-----|-------------|--|
| NOVARTIS | 1MG | N013420 001 | |
|----------|-----|-------------|--|

METHOCARBAMOL

INJECTABLE; INJECTION

METHOCARBAMOL

| | | | |
|-------------------|----------|-------------|--------------|
| MARSAM PHARMS LLC | 100MG/ML | A089849 001 | Dec 27, 1991 |
| WATSON LABS | 100MG/ML | A086459 001 | |

TABLET; ORAL

DELAXIN

| | | | |
|---------------|-------|-------------|--|
| FERNDALE LABS | 500MG | A085454 001 | |
|---------------|-------|-------------|--|

FORBAXIN

| | | | |
|-------------|-------|-------------|--|
| FOREST LABS | 750MG | A085136 001 | |
|-------------|-------|-------------|--|

METHOCARBAMOL

| | | | |
|----------------------|-------|-------------|--------------|
| ABLE | 500MG | A040413 001 | Mar 17, 2003 |
| | 750MG | A040413 002 | Mar 17, 2003 |
| AM THERAP | 500MG | A089417 001 | Feb 11, 1987 |
| | 750MG | A089418 001 | Feb 11, 1987 |
| ASCOT | 500MG | A087660 001 | Oct 27, 1982 |
| | 750MG | A087661 001 | Oct 27, 1982 |
| CLONMEL HLTHCARE | 500MG | A085961 001 | |
| | 750MG | A085963 001 | |
| FOSUN PHARMA | 500MG | A084616 001 | |
| | 750MG | A084615 001 | |
| HEATHER | 500MG | A084675 001 | |
| | 750MG | A084924 001 | |
| IMPAX LABS | 500MG | A084927 001 | |
| | 750MG | A084928 001 | |
| INWOOD LABS | 500MG | A085137 001 | |
| IVAX SUB TEVA PHARMS | 500MG | A084648 001 | |
| | 750MG | A084649 001 | |
| KV PHARM | 500MG | A085660 001 | |
| | 750MG | A085658 001 | |
| LANNETT CO INC | 500MG | A084756 002 | Mar 31, 2003 |
| | 750MG | A084756 001 | |
| MYLAN | 500MG | A084259 001 | |
| | 750MG | A084323 001 | |
| NYLOS | 750MG | A085033 001 | |
| PIONEER PHARMS | 500MG | A088731 001 | Dec 13, 1985 |
| | 750MG | A089082 001 | Dec 13, 1985 |
| PURACAP PHARM | 500MG | A084231 002 | |
| | 750MG | A084471 001 | |
| PUREPAC PHARM | 500MG | A085718 001 | |
| | 750MG | A085718 002 | |
| ROXANE | 500MG | A088646 001 | Feb 29, 1984 |
| | 750MG | A088647 001 | Feb 29, 1984 |
| SANDOZ | 500MG | A087283 001 | |
| | 750MG | A087282 001 | |
| SOLVAY | 500MG | A084448 001 | |
| | 750MG | A084449 001 | |
| SUN PHARM INDUSTRIES | 500MG | A084488 001 | |
| | 750MG | A084486 001 | |
| SUPERPHARM | 500MG | A087589 001 | Jan 22, 1982 |
| | 750MG | A087590 001 | Jan 22, 1982 |
| TABLICAPS | 500MG | A084846 001 | |
| UPSHER SMITH | 500MG | A087453 001 | |
| | 750MG | A087454 001 | |
| WATSON LABS | 500MG | A083605 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL

| | | |
|-------|---------|-----|
| 500MG | A085180 | 001 |
| 750MG | A083605 | 002 |
| 750MG | A085192 | 001 |

METHOHEXITAL SODIUM

INJECTABLE; INJECTION

BREVITAL SODIUM

PAR STERILE PRODUCTS 200MG/VIAL

N011559 004 Dec 21, 2012

5GM/VIAL

N011559 003

METHOTREXATE

SOLUTION; SUBCUTANEOUS

OTREXUP

+ ANTARES PHARMA INC 7.5MG/0.4ML (7.5MG/0.4ML)

N204824 005 Nov 07, 2014

OTREXUP PFS

+ ANTARES PHARMA INC 10MG/0.4ML (10MG/0.4ML)

N204824 009 May 31, 2017

+ 15MG/0.6ML (15MG/0.6ML)

N204824 010 May 31, 2017

+ 17.5MG/0.7ML (17.5MG/0.7ML)

N204824 011 May 31, 2017

+ 20MG/0.8ML (20MG/0.8ML)

N204824 012 May 31, 2017

+ 22.5MG/0.9ML (22.5MG/0.9ML)

N204824 013 May 31, 2017

+ 25MG/ML (25MG/ML)

N204824 014 May 31, 2017

RASUVO

+ MEDAC PHARMA INC 27.5MG/0.55ML (27.5MG/0.55ML)

N205776 009 Jul 10, 2014

METHOTREXATE SODIUM

INJECTABLE; INJECTION

ABITREXATE

ABIC

EQ 25MG BASE/ML

A089161 001 Mar 10, 1987

EQ 50MG BASE/VIAL

A089354 001 Jul 17, 1987

EQ 100MG BASE/VIAL

A089355 001 Jul 17, 1987

EQ 250MG BASE/VIAL

A089356 001 Jul 17, 1987

FOLEX

PHARMACIA AND UPJOHN EQ 25MG BASE/VIAL

A087695 001 Apr 08, 1983

EQ 50MG BASE/VIAL

A087695 002 Apr 08, 1983

EQ 100MG BASE/VIAL

A087695 003 Apr 08, 1983

EQ 250MG BASE/VIAL

A088954 001 Oct 24, 1985

FOLEX PFS

PHARMACIA AND UPJOHN EQ 25MG BASE/ML

A081242 001 Aug 23, 1991

EQ 25MG BASE/ML

A089180 001 Jan 03, 1986

METHOTREXATE LPF

HOSPIRA EQ 25MG BASE/ML

N011719 007 Mar 31, 1982

METHOTREXATE PRESERVATIVE FREE

HOSPIRA EQ 20MG BASE/2ML (EQ 10MG BASE/ML)

N011719 014 Apr 13, 2005

+ EQ 500MG BASE/20ML (EQ 25MG BASE/ML) **

N011719 013 Apr 13, 2005

EQ 2.5GM BASE/100ML (EQ 25MG BASE/ML)

N011719 011 Apr 13, 2005

METHOTREXATE SODIUM

ABRAXIS PHARM

EQ 2.5MG BASE/ML

A089323 001 Jun 13, 1986

EQ 20MG BASE/VIAL

A088935 001 Oct 11, 1985

EQ 25MG BASE/ML

A089263 001 Jun 13, 1986

EQ 25MG BASE/ML

A089322 001 Jun 13, 1986

EQ 50MG BASE/VIAL

A088936 001 Oct 11, 1985

EQ 100MG BASE/VIAL

A088937 001 Oct 11, 1985

FRESENIUS KABI USA EQ 250MG BASE/10ML (EQ 25MG BASE/ML)

A040263 002 Feb 26, 1999

HOSPIRA

EQ 2.5MG BASE/ML

N011719 004

EQ 20MG BASE/VIAL

N011719 001

EQ 25MG BASE/ML

N011719 005

EQ 50MG BASE/VIAL

N011719 003

EQ 100MG BASE/VIAL

N011719 006

NORBROOK

EQ 25MG BASE/ML

A088648 001 May 09, 1986

PHARMACHEMIE USA

EQ 25MG BASE/ML

A089158 001 Jul 08, 1988

METHOTREXATE SODIUM PRESERVATIVE FREE

HOSPIRA EQ 1GM BASE/VIAL

N011719 009 Apr 07, 1988

MEXATE

BRISTOL

EQ 20MG BASE/VIAL

A086358 001

EQ 50MG BASE/VIAL

A086358 002

EQ 100MG BASE/VIAL

A086358 003

EQ 250MG BASE/VIAL

A086358 004

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHOTREXATE SODIUM

INJECTABLE; INJECTION

MEXATE-AQ

BRISTOL MYERS EQ 25MG BASE/ML

A088760 001 Feb 14, 1985

MEXATE-AQ PRESERVED

BRISTOL MYERS SQUIBB EQ 25MG BASE/ML

A089887 001 Apr 14, 1989

TABLET; ORAL

METHOTREXATE SODIUM

DURAMED PHARMS BARR EQ 2.5MG BASE

A040233 001 Jun 17, 1999

METHOXAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

VASOXYL

GLAXOSMITHKLINE 10MG/ML

N006772 002

20MG/ML

N006772 001

METHOXSALEN

CAPSULE; ORAL

8-MOP

+ VALEANT PHARM INTL 10MG

N009048 001

METHOXSALEN

ANI PHARMS INC 10MG

A087781 001 Jun 08, 1982

LOTION; TOPICAL

OXSORALEN

+ VALEANT PHARM INTL 1%

N009048 002

METHSCOPOLAMINE BROMIDE

TABLET; ORAL

METHSCOPOLAMINE BROMIDE

PVT FORM 2.5MG

A080970 001

PAMINE

FOUGERA PHARMS 2.5MG **

N008848 001

PAMINE FORTE

FOUGERA PHARMS 5MG **

N008848 002 Mar 25, 2003

METHSUXIMIDE

CAPSULE; ORAL

CELONTIN

+ PARKE DAVIS 150MG

N010596 007

METHYCLOTHIAZIDE

TABLET; ORAL

AQUATENSEN

MEDPOINTE PHARM HLC 5MG

N017364 001

ENDURON

+ ABBVIE 2.5MG **

N012524 001

+ 5MG **

N012524 004

METHYCLOTHIAZIDE

IVAX PHARMS 2.5MG

A087913 001 Jun 03, 1982

5MG

A087786 001 May 18, 1982

MYLAN 2.5MG

A087671 001 Aug 17, 1982

PAR PHARM 2.5MG

A089135 001 Feb 12, 1986

5MG

A089136 001 Feb 12, 1986

USL PHARMA 5MG

A088745 001 Mar 21, 1985

WATSON LABS 2.5MG

A085487 001 Mar 11, 1982

2.5MG

A088750 001 Sep 06, 1984

5MG

A085476 001 Mar 11, 1982

5MG

A088724 001 Sep 06, 1984

YAOPHARMA CO LTD 2.5MG

A089835 001 Aug 18, 1988

5MG

A089837 001 Aug 18, 1988

METHYCLOTHIAZIDE; PARGYLINE HYDROCHLORIDE

TABLET; ORAL

EUTRON

ABBOTT 5MG; 25MG

N016047 001

METHYCLOTHIAZIDE; RESERPINE

TABLET; ORAL

DIUTENSEN-R

MEDPOINTE PHARM HLC 2.5MG; 0.1MG

N012708 005

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYL AMINOLEVULINATE HYDROCHLORIDE

CREAM; TOPICAL

METVIXIA

GALDERMA LABS LP

EQ 16.8% BASE

N021415 001 Jul 27, 2004

METHYLDOPA

SUSPENSION; ORAL

ALDOMET

MERCK

250MG/5ML

N018389 001

TABLET; ORAL

ALDOMET

+ MERCK

125MG **

N013400 003

+

250MG **

N013400 001

+

500MG **

N013400 002

METHYLDOPA

ACCORD HLTHCARE

125MG

A070070 003 Oct 15, 1985

DURAMED PHARMS BARR

250MG

A071006 001 Dec 16, 1986

500MG

A071009 001 Dec 16, 1986

HALSEY

125MG

A071751 001 Mar 28, 1988

250MG

A071752 001 Mar 28, 1988

500MG

A071753 001 Mar 28, 1988

PAR PHARM

125MG

A070535 001 Jan 02, 1987

250MG

A070536 001 Jan 02, 1987

500MG

A070537 001 Jan 02, 1987

PARKE DAVIS

125MG

A070331 001 Apr 15, 1986

250MG

A070332 001 Apr 15, 1986

500MG

A070333 001 Apr 15, 1986

PLIVA

125MG

A072126 001 Jul 07, 1988

250MG

A072127 001 Jul 07, 1988

500MG

A072128 001 Jul 07, 1988

PUREPAC PHARM

125MG

A070749 001 Feb 07, 1986

250MG

A070750 001 Feb 07, 1986

500MG

A070452 001 Feb 07, 1986

ROXANE

125MG

A070192 001 Apr 25, 1986

250MG

A070193 001 Apr 25, 1986

500MG

A070194 001 Apr 25, 1986

SUN PHARM INDUSTRIES

125MG

A070073 001 Oct 09, 1986

250MG

A070060 001 Oct 09, 1986

500MG

A070074 001 Oct 09, 1986

SUPERPHARM

250MG

A070669 001 Jun 23, 1989

500MG

A070670 001 Jun 23, 1989

TEVA

125MG

A071105 001 Dec 05, 1986

250MG

A071106 001 Dec 05, 1986

500MG

A071067 001 Dec 05, 1986

WATSON LABS

125MG

A070245 001 Feb 25, 1986

125MG

A070260 001 Jun 24, 1985

250MG

A070246 001 Feb 25, 1986

250MG

A070261 001 Jun 24, 1985

250MG

A070703 001 Jun 06, 1986

500MG

A070247 001 Feb 25, 1986

500MG

A070262 001 Jun 24, 1985

YAOPHARMA CO LTD

125MG

A071700 001 Mar 02, 1988

250MG

N018934 001 Jun 29, 1984

500MG

N018934 002 Jun 29, 1984

METHYLDOPATE HYDROCHLORIDE

INJECTABLE; INJECTION

ALDOMET

+ MERCK

50MG/ML **

N013401 001

METHYLDOPATE HYDROCHLORIDE

ABRAXIS PHARM

50MG/ML

A070652 001 Jun 03, 1986

BAXTER HLTHCARE

50MG/ML

A070291 001 Jul 01, 1986

HOSPIRA

50MG/ML

A070691 001 Jun 19, 1987

50MG/ML

A070698 001 Jun 15, 1987

50MG/ML

A070699 001 Jun 15, 1987

50MG/ML

A070849 001 Jun 19, 1987

MARSAM PHARMS LLC

50MG/ML

A071812 001 Dec 22, 1987

SMITH AND NEPHEW

50MG/ML

A070841 001 Jan 02, 1987

TEVA PARENTERAL

50MG/ML

A072974 001 Nov 22, 1991

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLERGONOVINE MALEATE

TABLET; ORAL

METHERGINE

+ EDISON THERAPS LLC 0.2MG **

N006035 003

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

RITALIN LA

+ NOVARTIS 60MG **

N021284 005 Oct 27, 2014

TABLET; ORAL

METHYLPHENIDATE HYDROCHLORIDE

ABLE 5MG

A040404 001 Mar 29, 2001

10MG

A040404 002 Mar 29, 2001

20MG

A040404 003 Mar 29, 2001

ACTAVIS ELIZABETH 5MG

A040321 001 Feb 05, 2002

10MG

A040321 002 Feb 05, 2002

20MG

A040321 003 Feb 05, 2002

TABLET, CHEWABLE; ORAL

METHYLIN

+ SPECGX LLC 2.5MG **

N021475 001 Apr 15, 2003

+ 5MG **

N021475 002 Apr 15, 2003

+ 10MG **

N021475 003 Apr 15, 2003

TABLET, EXTENDED RELEASE; ORAL

METADATE ER

LANNETT CO INC 10MG

A040306 001 Oct 20, 1999

METHYLPHENIDATE HYDROCHLORIDE

ABLE 20MG

A076032 001 May 09, 2001

ACTAVIS ELIZABETH 20MG

A075450 001 Dec 21, 2001

WATSON LABS 20MG

A040410 001 Feb 09, 2001

RITALIN-SR

+ NOVARTIS 20MG

N018029 001 Mar 30, 1982

METHYLPREDNISOLONE

TABLET; ORAL

MEDROL

PHARMACIA AND UPJOHN 24MG

N011153 005

METHYLPREDNISOLONE

HEATHER 4MG

A085650 001

PAR PHARM 16MG

A089207 001 Apr 25, 1988

24MG

A089208 001 Apr 25, 1988

32MG

A089209 001 Apr 25, 1988

SANDOZ 4MG

A087341 001

TIANJIN TIANYAO 4MG

A204072 001 May 14, 2018

WATSON LABS 4MG

A086161 001 Feb 09, 1982

16MG

A086159 001 Feb 09, 1982

METHYLPREDNISOLONE ACETATE

ENEMA; RECTAL

MEDROL

PHARMACIA AND UPJOHN 40MG/BOT

N018102 001

INJECTABLE; INJECTION

M-PREDROL

BEL MAR 40MG/ML

A086666 001

80MG/ML

A087135 001

METHYLPREDNISOLONE ACETATE

AKORN 40MG/ML

A086903 001 Oct 20, 1982

80MG/ML

A086903 002 Oct 20, 1982

WATSON LABS 20MG/ML

A085597 001

20MG/ML

A087248 001

40MG/ML

A085374 001

40MG/ML

A085600 001

80MG/ML

A085595 001

80MG/ML

A086507 001

OINTMENT; TOPICAL

MEDROL ACETATE

PHARMACIA AND UPJOHN 0.25%

N012421 001

1%

N012421 002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLPREDNISOLONE ACETATE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-MEDROL ACETATE

| | | | | |
|----------------------|------------------------|---------|-----|--|
| PHARMACIA AND UPJOHN | 0.25%;EQ 3.5MG BASE/GM | A060611 | 002 | |
| | 1%;EQ 3.5MG BASE/GM | A060611 | 001 | |

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

A-METHAPRED

| | | | | | |
|-------------------------------------|--------------------|--------------------|---------|--------------|--------------|
| ABBOTT | EQ 40MG BASE/VIAL | A089573 | 001 | Feb 22, 1991 | |
| | EQ 125MG BASE/VIAL | A089574 | 001 | Feb 22, 1991 | |
| | EQ 500MG BASE/VIAL | A089575 | 001 | Feb 22, 1991 | |
| | EQ 1GM BASE/VIAL | A089576 | 001 | Feb 22, 1991 | |
| HOSPIRA | EQ 40MG BASE/VIAL | A085853 | 001 | | |
| | EQ 125MG BASE/VIAL | A085855 | 001 | | |
| | EQ 500MG BASE/VIAL | A085854 | 001 | | |
| | EQ 500MG BASE/VIAL | A089173 | 001 | Aug 18, 1987 | |
| | EQ 1GM BASE/VIAL | A085852 | 001 | | |
| | EQ 1GM BASE/VIAL | A089174 | 001 | Aug 18, 1987 | |
| HOSPIRA INC | EQ 40MG BASE/VIAL | A040793 | 001 | Nov 25, 2008 | |
| | EQ 125MG BASE/VIAL | A040827 | 001 | Nov 25, 2008 | |
| METHYLPREDNISOLONE | | | | | |
| ELKINS SINN | EQ 125MG BASE/VIAL | A086906 | 002 | | |
| | EQ 500MG BASE/VIAL | A086906 | 003 | | |
| | EQ 1GM BASE/VIAL | A086906 | 004 | | |
| ORGANON USA INC | EQ 500MG BASE/VIAL | A087535 | 001 | Jun 25, 1982 | |
| | EQ 1GM BASE/VIAL | A087535 | 002 | Jun 25, 1982 | |
| METHYLPREDNISOLONE SODIUM SUCCINATE | | | | | |
| ABRAXIS PHARM | EQ 40MG BASE/VIAL | A088676 | 001 | Jun 08, 1984 | |
| | EQ 40MG BASE/VIAL | A089143 | 001 | Mar 28, 1986 | |
| | EQ 125MG BASE/VIAL | A088677 | 001 | Jun 08, 1984 | |
| | EQ 125MG BASE/VIAL | A089144 | 001 | Mar 28, 1986 | |
| | EQ 500MG BASE/VIAL | A088678 | 001 | Jun 08, 1984 | |
| | EQ 500MG BASE/VIAL | A089186 | 001 | Mar 28, 1986 | |
| | EQ 500MG BASE/VIAL | A089187 | 001 | Mar 28, 1986 | |
| | EQ 1GM BASE/VIAL | A088679 | 001 | Jun 08, 1984 | |
| | EQ 1GM BASE/VIAL | A089188 | 001 | Mar 28, 1986 | |
| | EQ 1GM BASE/VIAL | A089189 | 001 | Mar 28, 1986 | |
| | BEDFORD LABS | EQ 40MG BASE/VIAL | A040662 | 001 | Feb 21, 2007 |
| | | EQ 125MG BASE/VIAL | A040641 | 002 | Feb 21, 2007 |
| | | EQ 500MG BASE/VIAL | A040641 | 003 | Feb 21, 2007 |
| | | EQ 500MG BASE/VIAL | A040709 | 001 | Feb 21, 2007 |
| EQ 1GM BASE/VIAL | | A040641 | 004 | Feb 21, 2007 | |
| EQ 1GM BASE/VIAL | | A040709 | 002 | Feb 21, 2007 | |
| ELKINS SINN | EQ 40MG BASE/VIAL | A086906 | 001 | | |
| INTL MEDICATION | EQ 40MG BASE/VIAL | A087812 | 001 | Feb 09, 1983 | |
| | EQ 125MG BASE/VIAL | A087813 | 001 | Feb 09, 1983 | |
| | EQ 500MG BASE/VIAL | A087851 | 001 | Feb 09, 1983 | |
| | EQ 1GM BASE/VIAL | A087852 | 001 | Feb 09, 1983 | |
| TEVA PARENTERAL | EQ 125MG BASE/VIAL | A081266 | 001 | Nov 30, 1992 | |
| | EQ 500MG BASE/VIAL | A081267 | 001 | Nov 30, 1992 | |
| | EQ 1GM BASE/VIAL | A081268 | 001 | Nov 30, 1992 | |
| WATSON LABS | EQ 40MG BASE/VIAL | A086953 | 001 | Jul 22, 1982 | |
| | EQ 125MG BASE/VIAL | A087030 | 001 | Jul 22, 1982 | |
| | EQ 500MG BASE/VIAL | A088523 | 001 | Jul 24, 1984 | |
| | EQ 1GM BASE/VIAL | A088524 | 001 | Jul 24, 1984 | |

METHYLPREDNISOLONE; NEOMYCIN SULFATE

OINTMENT; OPHTHALMIC

NEO-MEDROL

| | | | | |
|----------------------|-----------------------|---------|-----|--|
| PHARMACIA AND UPJOHN | 0.1%;EQ 3.5MG BASE/GM | A060645 | 001 | |
|----------------------|-----------------------|---------|-----|--|

METHYLTESTOSTERONE

CAPSULE; ORAL

METHYLTESTOSTERONE

| | | | | |
|----------------|------|---------|-----|--------------|
| HEATHER | 10MG | A084967 | 001 | |
| VIRILON | | | | |
| STAR PHARMS FL | 10MG | A087750 | 001 | Nov 24, 1982 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METHYLTESTOSTERONE

TABLET;BUCCAL

ANDROID 5

VALEANT PHARM INTL 5MG A087222 001

ORETON

SCHERING 10MG A080281 001

TABLET;BUCCAL, SUBLINGUAL

METANDREN

NOVARTIS 5MG N003240 004

10MG N003240 001

10MG N003240 005

25MG N003240 003

METHYLTESTOSTERONE

IMPAX LABS 10MG A084287 001

LILLY 10MG A080256 001

25MG A080256 002

PUREPAC PHARM 10MG A080308 001

10MG A080475 001

10MG A080475 002

25MG A080475 003

PVT FORM 5MG A083836 001

TABLICAPS 10MG A085125 001

USL PHARMA 10MG A080271 001

TABLET;ORAL

ANDROID 10

VALEANT PHARMS NORTH 10MG A086450 001

METHYLTESTOSTERONE

IMPAX LABS 25MG A084310 001

INWOOD LABS 10MG A080839 001

25MG A080973 001

KV PHARM 10MG A084312 001

LANNETT 10MG A087092 001 Nov 05, 1982

25MG A087111 001 Jan 27, 1983

PARKE DAVIS 10MG A084244 001

25MG A084241 001

PUREPAC PHARM 10MG A080309 001

25MG A080310 001

PVT FORM 5MG A080214 001

10MG A080214 002

25MG A080214 003

TABLICAPS 10MG A080313 001

25MG A085270 001

WATSON LABS 10MG A080933 001

25MG A080931 001

WEST WARD 10MG A084331 001

25MG A084331 002

25MG A084642 001

ORETON METHYL

SCHERING 10MG N003158 001

25MG N003158 002

METHYPRYLON

CAPSULE;ORAL

NOLUDAR

ROCHE 300MG N009660 008

ELIXIR;ORAL

NOLUDAR

ROCHE 50MG/5ML N009660 007

TABLET;ORAL

NOLUDAR

ROCHE 50MG N009660 002

200MG N009660 004

METHYSERGIDE MALEATE

TABLET;ORAL

SANSERT

NOVARTIS 2MG N012516 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METOCLOPRAMIDE HYDROCHLORIDE

CONCENTRATE; ORAL

METOCLOPRAMIDE INTENSOL

| | | | |
|--------|-----------------|-------------|--------------|
| ROXANE | EQ 10MG BASE/ML | A072995 001 | Jan 30, 1992 |
|--------|-----------------|-------------|--------------|

INJECTABLE; INJECTION

METOCLOPRAMIDE HYDROCHLORIDE

| | | | |
|------------------|------------------|-------------|--------------|
| BEDFORD | EQ 5MG BASE/ML | A072155 001 | Mar 30, 1992 |
| | EQ 5MG BASE/ML | A072244 001 | Mar 30, 1992 |
| | EQ 5MG BASE/ML | A072247 001 | May 18, 1992 |
| HOSPIRA | EQ 5MG BASE/ML | A070505 001 | Jun 23, 1989 |
| | EQ 5MG BASE/ML | A070506 001 | Jun 22, 1989 |
| | EQ 5MG BASE/ML | A070847 001 | Nov 07, 1988 |
| | EQ 5MG BASE/ML | A071291 001 | Mar 03, 1989 |
| | EQ 5MG BASE/ML | A071990 001 | Jan 18, 1989 |
| | EQ 5MG BASE/ML | A073117 001 | Jan 17, 1991 |
| | EQ 5MG BASE/ML | A074147 001 | Aug 02, 1996 |
| LYPHOMED | EQ 10MG BASE/2ML | A070293 001 | Jan 24, 1986 |
| NORBROOK | EQ 10MG BASE/2ML | A070892 001 | Aug 26, 1988 |
| SMITH AND NEPHEW | EQ 5MG BASE/ML | A070623 001 | Mar 02, 1987 |
| | EQ 10MG BASE/2ML | A070622 001 | Mar 02, 1987 |

REGLAN

| | | | |
|----------------------|-----------------|-------------|--------------|
| WEST-WARD PHARMS INT | EQ 5MG BASE/ML | N017862 001 | |
| | EQ 10MG BASE/ML | N017862 004 | May 28, 1987 |

SOLUTION; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

| | | | |
|----------------------|-----------------|-------------|--------------|
| ACTAVIS MID ATLANTIC | EQ 5MG BASE/5ML | A071340 001 | Aug 18, 1988 |
| LANNETT CO INC | EQ 5MG BASE/5ML | A073680 001 | Oct 27, 1992 |
| MORTON GROVE | EQ 5MG BASE/5ML | A070949 001 | Mar 06, 1987 |
| PACO | EQ 5MG BASE/5ML | A071665 001 | Dec 05, 1988 |
| ROXANE | EQ 5MG BASE/5ML | A072038 001 | Dec 05, 1988 |
| TEVA | EQ 5MG BASE/5ML | A070819 001 | Jul 10, 1987 |
| | EQ 5MG BASE/5ML | A071315 001 | Jun 30, 1993 |

REGLAN

| | | | |
|-------------|--------------------|-------------|--------------|
| + ROBINS AH | EQ 5MG BASE/5ML ** | N018821 001 | Mar 25, 1983 |
|-------------|--------------------|-------------|--------------|

TABLET; ORAL

CLOPRA

| | | | |
|------------------|--------------|-------------|--------------|
| QUANTUM PHARMICS | EQ 5MG BASE | A072384 001 | Jun 02, 1988 |
| | EQ 10MG BASE | A070294 001 | Jul 29, 1985 |

CLOPRA-"YELLOW"

| | | | |
|------------------|--------------|-------------|--------------|
| QUANTUM PHARMICS | EQ 10MG BASE | A070632 001 | Oct 28, 1985 |
|------------------|--------------|-------------|--------------|

MAXOLON

| | | | |
|-------------|--------------|-------------|--------------|
| KING PHARMS | EQ 10MG BASE | A070106 001 | Mar 04, 1986 |
|-------------|--------------|-------------|--------------|

METOCLOPRAMIDE HYDROCHLORIDE

| | | | |
|----------------------|--------------|-------------|--------------|
| CLONMEL | EQ 10MG BASE | A072639 001 | May 09, 1991 |
| HALSEY | EQ 10MG BASE | A070906 001 | Oct 28, 1986 |
| INTERPHARM | EQ 10MG BASE | A071213 001 | Sep 24, 1986 |
| MUTUAL PHARM | EQ 10MG BASE | A070660 001 | Feb 10, 1987 |
| NORTHSTAR HLTHCARE | EQ 5MG BASE | A078374 001 | Nov 30, 2007 |
| | EQ 10MG BASE | A078374 002 | Nov 30, 2007 |
| PAR PHARM | EQ 10MG BASE | A070342 001 | Mar 25, 1986 |
| SANDOZ | EQ 5MG BASE | A072436 001 | Jun 22, 1989 |
| | EQ 10MG BASE | A070850 001 | Feb 03, 1987 |
| SCHERING | EQ 10MG BASE | A070598 001 | Feb 02, 1987 |
| SUN PHARM INDUSTRIES | EQ 5MG BASE | A071536 002 | Jan 16, 1997 |
| | EQ 10MG BASE | A071536 001 | Apr 28, 1993 |
| SUPERPHARM | EQ 10MG BASE | A070926 001 | Jun 26, 1987 |
| USL PHARMA | EQ 10MG BASE | A070339 001 | Jul 29, 1985 |
| WATSON LABS | EQ 10MG BASE | A070363 001 | Mar 02, 1987 |
| | EQ 10MG BASE | A070453 001 | Jun 06, 1986 |
| | EQ 10MG BASE | A070511 001 | Jan 22, 1986 |
| | EQ 10MG BASE | A070645 001 | May 11, 1987 |
| YAOPHARMA CO LTD | EQ 5MG BASE | A074478 001 | Oct 05, 1995 |
| | EQ 10MG BASE | A072215 001 | Jan 30, 1990 |
| | EQ 10MG BASE | A074478 002 | Oct 05, 1995 |

TABLET, ORALLY DISINTEGRATING; ORAL

METOZOLV ODT

| | | | |
|----------------|-----------------|-------------|--------------|
| + SALIX PHARMS | EQ 5MG BASE | N022246 001 | Sep 04, 2009 |
| + | EQ 10MG BASE ** | N022246 002 | Sep 04, 2009 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METOCLOPRAMIDE HYDROCHLORIDE

TABLET, ORALLY DISINTEGRATING;ORAL

REGLAN ODT

MEDA PHARMS

EQ 5MG BASE
EQ 10MG BASEN021793 001 Jun 10, 2005
N021793 002 Jun 10, 2005METOCURINE IODIDE

INJECTABLE; INJECTION

METUBINE IODIDE

LILLY

2MG/ML

N006632 003

METOLAZONE

TABLET;ORAL

DIULO

GD SEARLE LLC

2.5MG
5MG
10MGN018535 001
N018535 002
N018535 003

METOLAZONE

ROXANE

10MG

A076482 002 Apr 29, 2004

TEVA

2.5MG

A076600 001 Jan 06, 2004

5MG

A076833 001 Mar 01, 2004

10MG

A075543 003 Dec 24, 2003

WATSON LABS

10MG

A076891 001 Jul 21, 2004

MYKROX

LANNETT CO INC

0.5MG

N019532 001 Oct 30, 1987

METOPROLOL FUMARATE

TABLET, EXTENDED RELEASE;ORAL

LOPRESSOR

NOVARTIS

EQ 100MG TARTRATE
EQ 200MG TARTRATE
EQ 300MG TARTRATE
EQ 400MG TARTRATEN019786 001 Dec 27, 1989
N019786 002 Dec 27, 1989
N019786 003 Dec 27, 1989
N019786 004 Dec 27, 1989METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE;ORAL

METOPROLOL SUCCINATE

NESHER PHARMS

EQ 25MG TARTRATE
EQ 50MG TARTRATE
EQ 100MG TARTRATE
EQ 200MG TARTRATE
EQ 25MG TARTRATE
EQ 50MG TARTRATE
EQ 100MG TARTRATE
EQ 200MG TARTRATEA077779 001 Mar 20, 2008
A077176 001 May 14, 2008
A076640 002 May 18, 2007
A076640 001 May 18, 2007
A076969 001 Jul 31, 2006
A076969 002 May 18, 2007
A076969 003 Mar 20, 2008
A076969 004 Mar 20, 2008

SANDOZ

METOPROLOL TARTRATE

INJECTABLE; INJECTION

METOPROLOL TARTRATE

WATSON LABS

1MG/ML

A074032 001 Dec 21, 1993

TABLET;ORAL

METOPROLOL TARTRATE

APOTHECON

50MG

A074258 001 Jan 27, 1994

100MG

A074258 002 Jan 27, 1994

MYLAN

50MG

A073666 001 Dec 21, 1993

100MG

A073666 002 Dec 21, 1993

PUREPAC PHARM

50MG

A074380 001 Jul 29, 1994

100MG

A074380 002 Jul 29, 1994

RENATA

50MG

A074453 001 Apr 27, 1995

100MG

A074453 002 Apr 27, 1995

TEVA

50MG

A074143 001 Sep 30, 1994

100MG

A074143 002 Sep 30, 1994

TEVA PHARMS

50MG

A074333 001 Jan 27, 1994

100MG

A074333 002 Jan 27, 1994

YAOPHARMA CO LTD

50MG

A073288 001 Mar 25, 1994

100MG

A073289 001 Mar 25, 1994

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METRIZAMIDE

INJECTABLE; INJECTION

AMIPAQUE

| | | | |
|---------------|-------------|-------------|--------------|
| GE HEALTHCARE | 2.5GM/VIAL | N017982 003 | Sep 12, 1983 |
| | 3.75GM/VIAL | N017982 001 | |
| | 6.75GM/VIAL | N017982 002 | |
| | 13.5GM/VIAL | N017982 004 | Sep 12, 1983 |

METRONIDAZOLE

CAPSULE; ORAL

METRONIDAZOLE

| | | | |
|------|-------|-------------|--------------|
| ABLE | 375MG | A076505 001 | Nov 13, 2003 |
|------|-------|-------------|--------------|

INJECTABLE; INJECTION

FLAGYL I.V. RTU IN PLASTIC CONTAINER

| | | | |
|--------|-------------|-------------|--|
| PFIZER | 500MG/100ML | N018353 002 | |
|--------|-------------|-------------|--|

METRO I.V.

| | | | |
|---------|-------------|-------------|--------------|
| B BRAUN | 500MG/100ML | N018674 001 | Aug 31, 1982 |
|---------|-------------|-------------|--------------|

METRONIDAZOLE

| | | | |
|----------------------|-------------|-------------|--------------|
| ABBOTT | 500MG/100ML | N018889 001 | Nov 18, 1983 |
| ABRAXIS PHARM | 500MG/100ML | A070071 001 | Dec 03, 1984 |
| INTL MEDICATION | 500MG/100ML | A070004 001 | May 08, 1985 |
| WATSON LABS | 500MG/100ML | A070042 001 | Dec 20, 1984 |
| | 500MG/100ML | A070170 001 | Apr 01, 1986 |
| WEST-WARD PHARMS INT | 500MG/100ML | N018907 001 | Mar 30, 1984 |

TABLET; ORAL

METROMIDOL

| | | | |
|---------|-------|-------------|--------------|
| LABS AF | 250MG | A074523 001 | Oct 24, 1996 |
| | 500MG | A074523 002 | Oct 24, 1996 |

METRONIDAZOLE

| | | | |
|----------------------|-------|-------------|--------------|
| ABLE | 250MG | A076519 001 | Jun 27, 2003 |
| | 500MG | A076519 002 | Jun 27, 2003 |
| CHARTWELL MOLECULES | 250MG | N018845 001 | Aug 18, 1983 |
| | 500MG | N018930 001 | Aug 18, 1983 |
| FOSUN PHARMA | 250MG | N018620 001 | Mar 04, 1982 |
| | 250MG | N018740 001 | Oct 22, 1982 |
| | 500MG | N018620 002 | Jun 02, 1983 |
| | 500MG | N018740 002 | Oct 22, 1982 |
| HALSEY | 250MG | A070021 001 | Apr 02, 1985 |
| | 500MG | A070593 001 | Feb 27, 1986 |
| IVAX SUB TEVA PHARMS | 250MG | N018517 001 | |
| | 500MG | N018517 002 | May 05, 1982 |
| LNK | 250MG | N019029 001 | Apr 10, 1984 |
| MUTUAL PHARM | 250MG | N018818 001 | Feb 16, 1983 |
| | 500MG | N018818 002 | Feb 16, 1983 |
| SUPERPHARM | 250MG | A070008 001 | Dec 11, 1984 |
| | 500MG | A070009 001 | Dec 11, 1984 |
| WATSON LABS | 250MG | N018599 001 | Sep 17, 1982 |
| | 250MG | N018764 001 | Sep 17, 1982 |
| | 500MG | N018599 002 | Feb 13, 1984 |
| | 500MG | N018764 002 | Dec 20, 1982 |

PROTOSTAT

| | | | |
|--------------------|-------|-------------|--------------|
| ORTHO MCNEIL PHARM | 250MG | N018871 001 | Mar 02, 1983 |
| | 500MG | N018871 002 | Mar 02, 1983 |

SATRIC

| | | | |
|-------------|-------|-------------|--------------|
| SAVAGE LABS | 250MG | A070029 001 | Mar 19, 1985 |
| | 500MG | A070731 001 | Jun 08, 1987 |

TABLET, EXTENDED RELEASE; ORAL

FLAGYL ER

| | | | |
|-----------------|-------|-------------|--------------|
| + GD SEARLE LLC | 750MG | N020868 001 | Nov 26, 1997 |
|-----------------|-------|-------------|--------------|

METRONIDAZOLE

| | | | |
|--------------------|-------|-------------|--------------|
| ABLE | 750MG | A076462 001 | Jun 25, 2003 |
| ALEMBIC PHARMS LTD | 750MG | A090222 001 | May 05, 2010 |

METRONIDAZOLE HYDROCHLORIDE

INJECTABLE; INJECTION

FLAGYL I.V.

| | | | |
|--------|-----------------------|-------------|--|
| PFIZER | EQ 500MG BASE/VIAL ** | N018353 001 | |
|--------|-----------------------|-------------|--|

METRONIDAZOLE HYDROCHLORIDE

| | | | |
|---------------|--------------------|-------------|--------------|
| ABRAXIS PHARM | EQ 500MG BASE/VIAL | A070295 001 | Oct 15, 1985 |
|---------------|--------------------|-------------|--------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

METYRAPONETABLET; ORAL
METOPIRONE

LABORATORIE HRA 250MG N012911 001

MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

| | | | |
|----------------|-------|-------------|--------------|
| ANI PHARMS INC | 150MG | A074450 001 | May 16, 1996 |
| | 200MG | A074450 002 | May 16, 1996 |
| | 250MG | A074450 003 | May 16, 1996 |
| WATSON LABS | 150MG | A074711 001 | Feb 26, 1997 |
| | 150MG | A074865 001 | Apr 13, 1998 |
| | 200MG | A074711 002 | Feb 26, 1997 |
| | 200MG | A074865 002 | Apr 13, 1998 |
| | 250MG | A074711 003 | Feb 26, 1997 |
| | 250MG | A074865 003 | Apr 13, 1998 |

MEXITIL

| | | | | |
|---|----------------------|-------|-------------|--------------|
| + | BOEHRINGER INGELHEIM | 150MG | N018873 002 | Dec 30, 1985 |
| + | | 200MG | N018873 003 | Dec 30, 1985 |
| + | | 250MG | N018873 004 | Dec 30, 1985 |

MEZLOCILLIN SODIUM MONOHYDRATE

INJECTABLE; INJECTION

MEZLIN

| | | | |
|--------------|-------------------|-------------|--------------|
| BAYER PHARMS | EQ 1GM BASE/VIAL | A062333 001 | |
| | EQ 1GM BASE/VIAL | A062372 005 | Jan 13, 1983 |
| | EQ 1GM BASE/VIAL | N050549 001 | |
| | EQ 2GM BASE/VIAL | A062333 002 | |
| | EQ 2GM BASE/VIAL | A062372 001 | May 13, 1982 |
| | EQ 2GM BASE/VIAL | N050549 002 | |
| | EQ 3GM BASE/VIAL | A062333 003 | |
| | EQ 3GM BASE/VIAL | A062372 002 | May 13, 1982 |
| | EQ 3GM BASE/VIAL | A062697 001 | Jan 22, 1987 |
| | EQ 3GM BASE/VIAL | N050549 003 | |
| | EQ 4GM BASE/VIAL | A062333 004 | |
| | EQ 4GM BASE/VIAL | A062372 003 | May 13, 1982 |
| | EQ 4GM BASE/VIAL | A062697 002 | Jan 22, 1987 |
| | EQ 4GM BASE/VIAL | N050549 004 | |
| | EQ 20GM BASE/VIAL | A062372 004 | Mar 02, 1988 |
| | EQ 20GM BASE/VIAL | N050549 005 | Mar 02, 1988 |

MICONAZOLE

INJECTABLE; INJECTION

MONISTAT

JANSSEN PHARMA 10MG/ML N018040 001

MICONAZOLE NITRATE

CREAM; TOPICAL

MONISTAT-DERM

INSIGHT PHARMS 2% N017494 001

CREAM; VAGINAL

MICONAZOLE NITRATE

| | | | |
|-------------|----|-------------|--------------|
| TEVA | 2% | A074136 001 | Jan 04, 1995 |
| TEVA PHARMS | 2% | A074030 001 | Oct 30, 1992 |

CREAM, SUPPOSITORY; TOPICAL, VAGINAL

M-ZOLE 7 DUAL PACK

ACTAVIS MID ATLANTIC 2%, 100MG A074586 001 Jul 17, 1997

MICONAZOLE 7 COMBINATION PACK

G AND W LABS 2%, 100MG A076585 001 Mar 26, 2004

LOTION; TOPICAL

MONISTAT-DERM

INSIGHT PHARMS 2% N017739 001

TAMPON; VAGINAL

MONISTAT 5

PERSONAL PRODS 100MG N018592 001 Oct 27, 1989

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

| | | | |
|----------------------|-------------------|-------------|--------------|
| AKORN INC | EQ 5MG BASE/ML | A075481 001 | Jun 30, 2000 |
| APOTHECON | EQ 1MG BASE/ML | A075620 001 | Nov 01, 2000 |
| | EQ 5MG BASE/ML | A075620 002 | Nov 01, 2000 |
| | EQ 5MG BASE/ML | A075641 001 | Oct 19, 2000 |
| BAXTER HLTHCARE CORP | EQ 1MG BASE/ML | A075637 001 | Oct 31, 2000 |
| | EQ 5MG BASE/ML | A075637 002 | Oct 31, 2000 |
| BEDFORD | EQ 5MG BASE/ML | A075249 001 | Jun 23, 2000 |
| BEN VENUE | EQ 5MG BASE/ML | A075455 001 | Jun 20, 2000 |
| HOSPIRA | EQ 1MG BASE/ML | A075396 001 | Jun 20, 2000 |
| | EQ 5MG BASE/ML | A075396 002 | Jun 20, 2000 |
| | EQ 5MG BASE/ML | A075484 001 | Jun 20, 2000 |
| HOSPIRA INC | EQ 1MG BASE/ML | A075409 002 | Jun 20, 2000 |
| | EQ 5MG BASE/ML | A075409 001 | Jun 20, 2000 |
| IGI LABS INC | EQ 5MG BASE/ML | A075263 001 | Jun 26, 2000 |
| INTL MEDICATED | EQ 1MG BASE/ML | A076144 001 | Jan 26, 2005 |
| | EQ 5MG BASE/ML | A076144 002 | Jan 26, 2005 |
| INTL MEDICATION | EQ 1MG BASE/ML | A076020 001 | Jul 16, 2004 |
| | EQ 5MG BASE/ML | A076020 002 | Jul 16, 2004 |
| WOCKHARDT | EQ 1MG BASE/ML | A078141 001 | May 30, 2008 |
| | EQ 1MG BASE/ML | A078511 001 | Nov 10, 2008 |
| | EQ 5MG BASE/ML | A078141 002 | May 30, 2008 |
| | EQ 5MG BASE/ML | A078511 002 | Nov 10, 2008 |
| VERSED | | | |
| + HLR | EQ 1MG BASE/ML ** | N018654 002 | May 26, 1987 |
| + | EQ 5MG BASE/ML ** | N018654 001 | Dec 20, 1985 |

SYRUP; ORAL

MIDAZOLAM HYDROCHLORIDE

| | | | |
|--------------------|-------------------|-------------|--------------|
| APOTEX INC | EQ 2MG BASE/ML | A077115 001 | Sep 09, 2005 |
| SUN PHARM INDS LTD | EQ 2MG BASE/ML | A076058 001 | Mar 15, 2002 |
| VERSED | | | |
| + ROCHE | EQ 2MG BASE/ML ** | N020942 001 | Oct 15, 1998 |

MIDODRINE HYDROCHLORIDE

TABLET; ORAL

PROAMATINE

| | | | |
|-------------|-------|-------------|--------------|
| + SHIRE LLC | 2.5MG | N019815 001 | Sep 06, 1996 |
| + | 5MG | N019815 002 | Sep 06, 1996 |
| + | 10MG | N019815 003 | Mar 20, 2002 |

MILNACIPRAN HYDROCHLORIDE

TABLET; ORAL

MILNACIPRAN HYDROCHLORIDE

| | | | |
|------------------|--------|-------------|--------------|
| AMNEAL PHARMS | 12.5MG | A205081 001 | Apr 22, 2016 |
| | 25MG | A205081 002 | Apr 22, 2016 |
| | 50MG | A205081 003 | Apr 22, 2016 |
| | 100MG | A205081 004 | Apr 22, 2016 |
| USPHARMA WINDLAS | 12.5MG | A205071 001 | Jan 27, 2016 |
| | 25MG | A205071 002 | Jan 27, 2016 |
| | 50MG | A205071 003 | Jan 27, 2016 |
| | 100MG | A205071 004 | Jan 27, 2016 |

MILRINONE LACTATE

INJECTABLE; INJECTION

MILRINONE LACTATE

| | | | |
|-------------------------------------------------------|---------------------------------------|-------------|--------------|
| BAXTER HLTHCARE CORP | EQ 1MG BASE/ML | A076427 001 | Sep 21, 2004 |
| HOSPIRA | EQ 1MG BASE/ML | A075830 001 | May 28, 2002 |
| | EQ 1MG BASE/ML | A075884 001 | May 28, 2002 |
| MYLAN INSTITUTIONAL | EQ 1MG BASE/ML | A076428 001 | Jun 16, 2003 |
| WEST-WARD PHARMS INT | EQ 1MG BASE/ML | A075852 001 | May 28, 2002 |
| MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| B BRAUN | EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) | A076414 001 | Aug 18, 2004 |
| BAXTER HLTHCARE | EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) | A076259 001 | Aug 08, 2002 |
| RENAISSANCE SSA LLC | EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) | A077151 001 | Jul 20, 2005 |
| WEST-WARD PHARMS INT | EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) | A075510 001 | May 28, 2002 |
| PRIMACOR | | | |
| + SANOFI AVENTIS US | EQ 1MG BASE/ML ** | N019436 001 | Dec 31, 1987 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MILRINONE LACTATE

INJECTABLE;INJECTION

PRIMACOR IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|---|-------------------|---------------------------------------|-------------|--------------|
| + | SANOFI AVENTIS US | EQ 10MG BASE/100ML ** | N020343 001 | Aug 09, 1994 |
| + | | EQ 15MG BASE/100ML ** | N020343 002 | Aug 09, 1994 |
| + | | EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) | N020343 003 | Aug 09, 1994 |
| | | ** | | |
| + | | EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML) | N020343 004 | Aug 09, 1994 |
| | | ** | | |

MINOCYCLINE HYDROCHLORIDE

CAPSULE;ORAL

MINOCIN

| | | | | |
|---|------------------|-----------------|-------------|--------------|
| + | PRECISION DERMAT | EQ 75MG BASE ** | N050649 003 | Feb 12, 2001 |
| | TRIAx PHARMS | EQ 50MG BASE | N050315 002 | |
| | | EQ 100MG BASE | N050315 001 | |

CAPSULE, EXTENDED RELEASE;ORAL

XIMINO

| | | | | |
|--|--------------------|-----------------|-------------|--------------|
| | SUN PHARM INDS LTD | EQ 67.5MG BASE | N201922 002 | Jul 11, 2012 |
| | | EQ 112.5MG BASE | N201922 004 | Jul 11, 2012 |

INJECTABLE;INJECTION

MINOCIN

| | | | | |
|--|---------|--------------------|-------------|--|
| | LEDERLE | EQ 100MG BASE/VIAL | A062139 001 | |
|--|---------|--------------------|-------------|--|

SUSPENSION;ORAL

MINOCIN

| | | | | |
|--|------------------|------------------|-------------|--|
| | PRECISION DERMAT | EQ 50MG BASE/5ML | N050445 001 | |
|--|------------------|------------------|-------------|--|

TABLET;ORAL

MINOCYCLINE HYDROCHLORIDE

| | | | | |
|---|--------------|------------------|-------------|--------------|
| + | TRIAx PHARMS | EQ 50MG BASE ** | N050451 003 | Aug 10, 1982 |
| + | | EQ 100MG BASE ** | N050451 002 | Aug 10, 1982 |

TABLET, EXTENDED RELEASE;ORAL

MINOCYCLINE HYDROCHLORIDE

| | | | | |
|--|------------------|---------------|-------------|--------------|
| | BARR LABS INC | EQ 45MG BASE | A065485 001 | Mar 17, 2009 |
| | | EQ 80MG BASE | A065485 007 | Apr 26, 2017 |
| | | EQ 90MG BASE | A065485 002 | Mar 17, 2009 |
| | | EQ 105MG BASE | A065485 008 | Apr 26, 2017 |
| | | EQ 135MG BASE | A065485 003 | Mar 17, 2009 |
| | IMPAX LABS INC | EQ 45MG BASE | A090024 001 | Feb 03, 2009 |
| | | EQ 90MG BASE | A090024 002 | Feb 03, 2009 |
| | | EQ 135MG BASE | A090024 003 | Feb 03, 2009 |
| | MYLAN PHARMS INC | EQ 45MG BASE | A090911 001 | Jul 20, 2010 |
| | | EQ 90MG BASE | A090911 002 | Jul 20, 2010 |
| | | EQ 135MG BASE | A090911 003 | Jul 20, 2010 |

SOLODYN

| | | | | |
|---|---------|------------------|-------------|--------------|
| + | MEDICIS | EQ 45MG BASE ** | N050808 001 | May 08, 2006 |
| + | | EQ 90MG BASE ** | N050808 002 | May 08, 2006 |
| + | | EQ 135MG BASE ** | N050808 003 | May 08, 2006 |

MINOXIDIL

SOLUTION;TOPICAL

MINOXIDIL (FOR MEN)

| | | | | |
|--|-----------------|----|-------------|--------------|
| | APOTEX INC | 2% | A074924 001 | Apr 29, 1998 |
| | BAUSCH AND LOMB | 2% | A074643 001 | Apr 09, 1996 |
| | COPELEY PHARM | 2% | A074500 001 | May 23, 1996 |
| | SIGHT PHARMS | 2% | A074743 002 | Oct 18, 1996 |
| | TEVA | 2% | A074589 001 | Apr 05, 1996 |

MINOXIDIL (FOR WOMEN)

| | | | | |
|--|--------------|----|-------------|--------------|
| | APOTEX INC | 2% | A074924 002 | Apr 29, 1998 |
| | SIGHT PHARMS | 2% | A074743 001 | Oct 18, 1996 |

MINOXIDIL EXTRA STRENGTH (FOR MEN)

| | | | | |
|--|------------|----|-------------|--------------|
| | APOTEX INC | 5% | A075839 001 | Oct 01, 2001 |
|--|------------|----|-------------|--------------|

TABLET;ORAL

LONITEN

| | | | | |
|---|----------------------|----------|-------------|--|
| + | PHARMACIA AND UPJOHN | 2.5MG ** | N018154 001 | |
| + | | 10MG ** | N018154 003 | |

MINODYL

| | | | | |
|--|------------------|-------|-------------|--------------|
| | QUANTUM PHARMICS | 2.5MG | A072153 001 | Jul 13, 1988 |
| | | 10MG | A071534 001 | Mar 19, 1987 |

MINOXIDIL

| | | | | |
|--|------------|-------|-------------|--------------|
| | ROYCE LABS | 2.5MG | A071799 001 | Nov 10, 1987 |
| | | 10MG | A071796 001 | Nov 10, 1987 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MINOXIDIL

TABLET; ORAL

MINOXIDIL

USL PHARMA

2.5MG

A071537 001 Dec 16, 1988

MIPOMERSEN SODIUM

SOLUTION; SUBCUTANEOUS

KYNAMRO

+ KASTLE THERAPS LLC

200MG/ML (200MG/ML)

N203568 001 Jan 29, 2013

MIRTAZAPINE

TABLET; ORAL

MIRTAZAPINE

ACTAVIS ELIZABETH

15MG

A076241 001 Jun 25, 2003

15MG

A076308 001 Jun 20, 2003

30MG

A076241 002 Jun 25, 2003

30MG

A076308 002 Jun 20, 2003

45MG

A076241 003 Jun 25, 2003

45MG

A076308 003 Jun 20, 2003

ACTAVIS LABS FL INC

15MG

A076336 001 Jun 20, 2003

30MG

A076336 002 Jun 20, 2003

45MG

A076336 003 Jun 20, 2003

IVAX SUB TEVA PHARMS

15MG

A076244 001 Dec 22, 2003

30MG

A076244 002 Dec 22, 2003

45MG

A076244 003 Dec 22, 2003

MYLAN PHARMS INC

15MG

A076176 001 Jun 19, 2003

30MG

A076176 002 Jun 19, 2003

45MG

A076176 003 Jun 19, 2003

ROXANE

15MG

A076270 001 Jun 19, 2003

30MG

A076270 002 Jun 19, 2003

45MG

A076270 003 Jun 19, 2003

UPSHER SMITH LABS

15MG

A076189 001 Jun 19, 2003

30MG

A076189 002 Jun 19, 2003

45MG

A076189 003 Jun 19, 2003

REMERON

+ ORGANON USA INC

45MG

N020415 003 Mar 17, 1997

TABLET, ORALLY DISINTEGRATING; ORAL

MIRTAZAPINE

ACTAVIS ELIZABETH

15MG

A076689 001 Aug 31, 2005

15MG

A077959 001 Feb 14, 2011

30MG

A076689 002 Aug 31, 2005

30MG

A077959 002 Feb 14, 2011

45MG

A076689 003 Aug 31, 2005

45MG

A077959 003 Feb 14, 2011

ACTAVIS LABS FL INC

15MG

A076307 001 Dec 17, 2003

30MG

A076307 002 Dec 17, 2003

45MG

A076307 003 Feb 28, 2006

MITOMYCIN

INJECTABLE; INJECTION

MITOMYCIN

HOSPIRA

20MG/VIAL

A064106 001 Nov 29, 1995

WEST-WARD PHARMS INT

5MG/VIAL

A064117 001 Apr 19, 1995

20MG/VIAL

A064117 002 Apr 19, 1995

40MG/VIAL

A064117 003 Jun 02, 1999

MITOZYTREX

+ SUPERGEN

5MG/VIAL **

N050763 001 Nov 14, 2002

MUTAMYCIN

+ BRISTOL

5MG/VIAL **

N050450 001

+

20MG/VIAL **

N050450 002

BRISTOL MYERS

5MG/VIAL

A062336 001

20MG/VIAL

A062336 002

40MG/VIAL

A062336 003 Mar 10, 1988

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

MITOXANTRONE HYDROCHLORIDE

FRESENIUS KABI ONCOL EQ 20MG BASE/10ML (EQ 2MG BASE/ML)

A078606 001 May 14, 2008

EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)

A078606 002 May 14, 2008

EQ 30MG BASE/15ML (EQ 2MG BASE/ML)

A078606 003 May 14, 2008

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

NOVANTRONE

EMD SERONO

EQ 20MG BASE/10ML (EQ 2MG BASE/ML)

N019297 001 Dec 23, 1987

+ EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML) **

N019297 002 Dec 23, 1987

+ EQ 30MG BASE/15ML (EQ 2MG BASE/ML) **

N019297 003 Dec 23, 1987

MIVACURIUM CHLORIDE

INJECTABLE; INJECTION

MIVACRON IN DEXTROSE 5% IN PLASTIC CONTAINER

ABBVIE

EQ 0.5MG BASE/ML

N020098 002 Jan 22, 1992

EQ 50MG BASE/100ML

N020098 003 Jan 22, 1992

MIVACURIUM CHLORIDE

MYLAN LABS LTD

EQ 2MG BASE/ML

A078562 001 Apr 30, 2009

SOLUTION; INTRAVENOUS

MIVACRON

+ ABBVIE

EQ 2MG BASE/ML (EQ 2MG BASE/ML) **

N020098 001 Jan 22, 1992

+ EQ 10MG BASE/5ML (EQ 2MG BASE/ML)

N020098 004 Jan 22, 1992

+ EQ 20MG BASE/10ML (EQ 2MG BASE/ML)

N020098 005 Jan 22, 1992

MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

UNIVASC

UCB INC

7.5MG **

N020312 001 Apr 19, 1995

15MG **

N020312 002 Apr 19, 1995

MOLINDONE HYDROCHLORIDE

CAPSULE; ORAL

MOBAN

+ ENDO PHARMS

5MG **

N017111 001

+ 10MG **

N017111 002

+ 25MG **

N017111 003

CONCENTRATE; ORAL

MOBAN

ENDO PHARMS

20MG/ML

N017938 001

TABLET; ORAL

MOBAN

+ ENDO PHARMS

5MG **

N017111 004

+ 10MG **

N017111 005

+ 25MG **

N017111 006

+ 50MG **

N017111 007

+ 100MG **

N017111 008

MOMETASONE FUROATE

CREAM; TOPICAL

ELOCON

MERCK SHARP DOHME

0.1%

N019625 001 May 06, 1987

OINTMENT; TOPICAL

MOMETASONE FUROATE

TARO

0.1%

A076624 001 Dec 03, 2004

MONOBENZONE

CREAM; TOPICAL

BENOQUIN

VALEANT PHARM INTL

20%

N008173 003

MONOCTANOIN

LIQUID; PERFUSION, BILIARY

MOCTANIN

ETHITEK

100%

N019368 001 Oct 29, 1985

MONTELUKAST SODIUM

TABLET; ORAL

MONTELUKAST SODIUM

APOTEX CORP

EQ 10MG BASE

A201294 001 Aug 03, 2012

TABLET, CHEWABLE; ORAL

MONTELUKAST SODIUM

APOTEX INC

EQ 4MG BASE

A201508 001 Aug 03, 2012

EQ 5MG BASE

A201508 002 Aug 03, 2012

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MORICIZINE HYDROCHLORIDE

TABLET; ORAL

ETHMOZINE

| | | | |
|-------|-------|-------------|--------------|
| SHIRE | 200MG | N019753 001 | Jun 19, 1990 |
| | 250MG | N019753 002 | Jun 19, 1990 |
| | 300MG | N019753 003 | Jun 19, 1990 |

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

AVINZA

| | | | |
|-----------------|-------|-------------|--------------|
| KING PHARMS LLC | 30MG | N021260 001 | Mar 20, 2002 |
| | 45MG | N021260 005 | Dec 18, 2008 |
| | 60MG | N021260 002 | Mar 20, 2002 |
| | 75MG | N021260 006 | Dec 18, 2008 |
| | 90MG | N021260 003 | Mar 20, 2002 |
| | 120MG | N021260 004 | Mar 20, 2002 |

INJECTABLE; INJECTION

MORPHINE SULFATE

| | | | |
|-----------------|----------|-------------|--------------|
| + HOSPIRA INC | 15MG/ML | N202515 005 | Nov 14, 2011 |
| ICU MEDICAL INC | 0.5MG/ML | N019917 001 | Oct 30, 1992 |
| SPECGX LLC | 1MG/ML | N020631 001 | Jul 03, 1996 |
| | 2MG/ML | N020631 002 | Jul 03, 1996 |
| WATSON LABS | 0.5MG/ML | A073373 001 | Sep 30, 1991 |
| | 0.5MG/ML | A073375 001 | Sep 30, 1991 |
| | 1MG/ML | A073374 001 | Sep 30, 1991 |
| | 1MG/ML | A073376 001 | Sep 30, 1991 |

INJECTABLE, LIPOSOMAL; EPIDURAL

DEPODUR

| | | | |
|-------------------|----------------------|-------------|--------------|
| PACIRA PHARMS INC | 10MG/ML (10MG/ML) | N021671 001 | May 18, 2004 |
| | 15MG/1.5ML (10MG/ML) | N021671 002 | May 18, 2004 |
| | 20MG/2ML (10MG/ML) | N021671 003 | May 18, 2004 |

TABLET, EXTENDED RELEASE; ORAL

ARYMO ER

| | | | |
|----------|------|-------------|--------------|
| + EGALET | 15MG | N208603 001 | Jan 09, 2017 |
| + | 30MG | N208603 002 | Jan 09, 2017 |
| + | 60MG | N208603 003 | Jan 09, 2017 |

MORPHINE SULFATE

| | | | |
|-----------------|-------|-------------|--------------|
| EPIC PHARMA LLC | 15MG | A091357 001 | Jun 23, 2016 |
| | 30MG | A091357 002 | Jun 23, 2016 |
| | 60MG | A091357 003 | Jun 23, 2016 |
| | 100MG | A091357 004 | Jun 23, 2016 |
| | 200MG | A091357 005 | Jun 23, 2016 |
| WATSON LABS | 100MG | A075656 001 | Jan 30, 2001 |

ORAMORPH SR

| | | | |
|---------------------|-------|-------------|--------------|
| XANODYNE PHARMS INC | 15MG | N019977 004 | Nov 23, 1994 |
| | 30MG | N019977 001 | Aug 15, 1991 |
| | 60MG | N019977 002 | Aug 15, 1991 |
| | 100MG | N019977 003 | Aug 15, 1991 |

MOXALACTAM DISODIUM

INJECTABLE; INJECTION

MOXAM

| | | | |
|-------|--------------------|-------------|--|
| LILLY | EQ 250MG BASE/VIAL | N050550 001 | |
| | EQ 500MG BASE/VIAL | N050550 002 | |
| | EQ 1GM BASE/VIAL | N050550 003 | |
| | EQ 2GM BASE/VIAL | N050550 004 | |
| | EQ 10GM BASE/VIAL | N050550 008 | |

MOXIFLOXACIN HYDROCHLORIDE

SOLUTION; INTRAVENOUS

AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER

| | | | |
|------------------|---------------------------|-------------|--------------|
| + BAYER HLTHCARE | 400MG/250ML (1.6MG/ML) ** | N021277 001 | Nov 30, 2001 |
|------------------|---------------------------|-------------|--------------|

MUPIROCIIN

OINTMENT; TOPICAL

BACTROBAN

| | | | |
|-------------------|-------|-------------|--------------|
| + GLAXOSMITHKLINE | 2% ** | N050591 001 | Dec 31, 1987 |
|-------------------|-------|-------------|--------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

MUPIROICIN CALCIUM

CREAM;TOPICAL

BACTROBAN

+ GLAXOSMITHKLINE EQ 2% BASE

N050746 001 Dec 11, 1997

OINTMENT;NASAL

BACTROBAN

+ GLAXOSMITHKLINE EQ 2% BASE

N050703 001 Sep 18, 1995

MYCOPHENOLATE MOFETIL

CAPSULE;ORAL

MYCOPHENOLATE MOFETIL

APOTEX CORP 250MG

A090419 001 Apr 22, 2009

DR REDDYS LABS LTD 250MG

A091315 001 Oct 27, 2011

JUBILANT CADISTA 250MG

A090762 001 Dec 15, 2014

ZYDUS PHARMS USA INC 250MG

A065433 001 May 04, 2009

TABLET;ORAL

MYCOPHENOLATE MOFETIL

APOTEX 500MG

A090499 001 Apr 22, 2009

DR REDDYS LABS LTD 500MG

A090464 001 Sep 13, 2010

JUBILANT CADISTA 500MG

A090661 001 Dec 15, 2014

ZYDUS PHARMS USA INC 500MG

A065477 001 May 04, 2009

NABUMETONE

TABLET;ORAL

NABUMETONE

COPLY PHARM 750MG

A075179 001 Jun 06, 2000

OXFORD PHARMS 500MG

A079093 001 Feb 27, 2009

750MG

A079093 002 Feb 27, 2009

SANDOZ 500MG

A075590 001 Feb 25, 2002

750MG

A075590 002 Feb 25, 2002

SCIEGEN PHARMS INC 500MG

A078420 001 Sep 24, 2008

750MG

A078420 002 Sep 24, 2008

RELAFEN

+ SMITHKLINE BEECHAM 500MG **

N019583 001 Dec 24, 1991

+ 750MG **

N019583 002 Dec 24, 1991

NADOLOL

TABLET;ORAL

CORGARD

US WORLDMEDS LLC 120MG

N018063 003

160MG

N018063 004

NADOLOL

IVAX SUB TEVA PHARMS 120MG

A074255 002 Jan 24, 1996

160MG

A074255 003 Jan 24, 1996

TEVA PHARMS 80MG

A074368 001 Aug 31, 1994

120MG

A074368 002 Aug 31, 1994

160MG

A074368 003 Aug 31, 1994

NAFCILLIN SODIUM

CAPSULE;ORAL

UNIPEN

WYETH AYERST EQ 250MG BASE

N050111 001

FOR SOLUTION;ORAL

UNIPEN

WYETH AYERST EQ 250MG BASE/5ML

N050199 001

INJECTABLE;INJECTION

NAFCILLIN SODIUM

APOTHECON EQ 500MG BASE/VIAL

A061984 001

EQ 1GM BASE/VIAL

A061984 002

EQ 2GM BASE/VIAL

A061984 003

EQ 4GM BASE/VIAL

A061984 005

SANDOZ EQ 500MG BASE/VIAL

A062527 001 Aug 02, 1984

WATSON LABS INC EQ 500MG BASE/VIAL

A062844 001 Oct 26, 1988

EQ 1GM BASE/VIAL

A062844 002 Oct 26, 1988

EQ 1.5GM BASE/VIAL

A062844 003 Oct 26, 1988

EQ 2GM BASE/VIAL

A062844 004 Oct 26, 1988

EQ 4GM BASE/VIAL

A062844 005 Oct 26, 1988

EQ 10GM BASE/VIAL

A063008 001 Sep 29, 1988

NALLPEN

GLAXOSMITHKLINE EQ 500MG BASE/VIAL

A061999 001

EQ 1GM BASE/VIAL

A061999 002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NAFCILLIN SODIUM

INJECTABLE; INJECTION

NALLPEN

| | | |
|-------------------|-------------|--------------|
| EQ 1GM BASE/VIAL | A062755 001 | Dec 19, 1986 |
| EQ 2GM BASE/VIAL | A061999 003 | |
| EQ 2GM BASE/VIAL | A062755 002 | Dec 19, 1986 |
| EQ 10GM BASE/VIAL | A061999 004 | |

UNIPEN

WYETH AYERST

EQ 500MG BASE/VIAL ** A062717 001 Dec 16, 1986

+

EQ 500MG BASE/VIAL ** N050320 001

EQ 1GM BASE/VIAL ** A062717 002 Dec 16, 1986

EQ 2GM BASE/VIAL ** A062717 004 Dec 16, 1986

+

EQ 2GM BASE/VIAL ** N050320 003

+

EQ 4GM BASE/VIAL ** N050320 004

+

EQ 10GM BASE/VIAL ** N050320 005

+

EQ 20GM BASE/VIAL ** N050320 006

UNIPEN IN PLASTIC CONTAINER

+ WYETH AYERST

EQ 1GM BASE/VIAL ** N050320 002

TABLET; ORAL

UNIPEN

WYETH AYERST

EQ 500MG BASE N050462 001

NAFTIFINE HYDROCHLORIDE

CREAM; TOPICAL

NAFTIN

+ SEBELA IRELAND LTD 1%

N019599 001 Feb 29, 1988

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE

ABRAXIS PHARM

10MG/ML A070751 001 Jul 02, 1986

20MG/ML A070752 001 Sep 24, 1986

NALBUPHINE HYDROCHLORIDE

ABBOTT

20MG/ML A070917 001 Feb 03, 1989

ABBVIE

1.5MG/ML N020200 001 Mar 12, 1993

BARR

10MG/ML A074471 001 Mar 19, 1998

20MG/ML A074471 002 Mar 19, 1998

IGI LABS INC

10MG/ML A072070 001 Apr 10, 1989

10MG/ML A072071 001 Apr 10, 1989

10MG/ML A072072 001 Apr 10, 1989

20MG/ML A072073 001 Apr 10, 1989

20MG/ML A072074 001 Apr 10, 1989

20MG/ML A072075 001 Apr 10, 1989

NUBAIN

+ PAR PHARM INC

10MG/ML ** N018024 001

+

20MG/ML ** N018024 002 May 27, 1982

NALIDIXIC ACID

SUSPENSION; ORAL

NEGGRAM

SANOFI AVENTIS US

250MG/5ML N017430 001

TABLET; ORAL

NALIDIXIC ACID

SUN PHARM INDUSTRIES

250MG A070270 001 Jun 29, 1988

500MG A070271 001 Jun 29, 1988

1GM A070272 001 Jun 29, 1988

WATSON LABS

250MG A071936 001 Jun 29, 1988

500MG A072061 001 Jun 29, 1988

1GM A071919 001 Jun 29, 1988

NEGGRAM

SANOFI AVENTIS US

250MG N014214 002

500MG N014214 004

1GM N014214 005

NALMEFENE HYDROCHLORIDE

INJECTABLE; INJECTION

REVEX

+ EUROHLTH INTL SARL

EQ 0.1MG BASE/ML ** N020459 001 Apr 17, 1995

+

EQ 1MG BASE/ML ** N020459 002 Apr 17, 1995

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

| | | | | |
|----------------------|-----------|---------|-----|--------------|
| WEST-WARD PHARMS INT | 0.4MG/ML | A070298 | 001 | Sep 24, 1986 |
| | 0.4MG/ML | A070496 | 001 | Sep 24, 1986 |
| WYETH AYERST | 0.02MG/ML | A070188 | 001 | Sep 24, 1986 |
| | 0.02MG/ML | A070189 | 001 | Sep 24, 1986 |
| | 0.4MG/ML | A070190 | 001 | Sep 24, 1986 |
| | 0.4MG/ML | A070191 | 001 | Sep 24, 1986 |

NALOXONE HYDROCHLORIDE

| | | | | |
|--------------------|-----------|---------|-----|--------------|
| ABRAXIS PHARM | 0.02MG/ML | A070648 | 001 | Nov 17, 1986 |
| | 0.02MG/ML | A070661 | 001 | Nov 17, 1986 |
| | 0.4MG/ML | A070649 | 001 | Nov 17, 1986 |
| | 1MG/ML | A071604 | 001 | Dec 16, 1988 |
| ASTRAZENECA | 0.02MG/ML | A072081 | 001 | Apr 11, 1989 |
| EUROHLTH INTL SARL | 0.02MG/ML | A071272 | 001 | May 24, 1988 |
| | 1MG/ML | A071273 | 001 | May 24, 1988 |
| | 1MG/ML | A071274 | 001 | May 24, 1988 |
| | 1MG/ML | A071287 | 001 | May 24, 1988 |
| HOSPIRA | 0.02MG/ML | A070171 | 001 | Sep 24, 1986 |
| | 0.02MG/ML | A070252 | 001 | Jan 16, 1987 |
| | 0.02MG/ML | A070253 | 001 | Jan 16, 1987 |
| | 0.4MG/ML | A070255 | 001 | Jan 07, 1987 |
| IGI LABS INC | 0.02MG/ML | A072082 | 001 | Apr 11, 1989 |
| | 0.02MG/ML | A072083 | 001 | Apr 11, 1989 |
| | 0.02MG/ML | A072084 | 001 | Apr 11, 1989 |
| | 0.02MG/ML | A072085 | 001 | Apr 11, 1989 |
| | 0.4MG/ML | A072086 | 001 | Apr 11, 1989 |
| | 0.4MG/ML | A072087 | 001 | Apr 11, 1989 |
| | 0.4MG/ML | A072088 | 001 | Apr 11, 1989 |
| | 0.4MG/ML | A072089 | 001 | Apr 11, 1989 |
| | 0.4MG/ML | A072090 | 001 | Apr 11, 1989 |
| | 1MG/ML | A072091 | 001 | Apr 11, 1989 |
| | 1MG/ML | A072092 | 001 | Apr 11, 1989 |
| | 1MG/ML | A072093 | 001 | Apr 11, 1989 |
| INTL MEDICATION | 0.4MG/ML | A070417 | 001 | Sep 24, 1986 |
| | 1MG/ML | A072115 | 001 | Apr 27, 1988 |
| MARSAM PHARMS LLC | 0.4MG/ML | A071811 | 001 | Jul 19, 1988 |
| SMITH AND NEPHEW | 0.02MG/ML | A071671 | 001 | Nov 17, 1987 |
| | 0.4MG/ML | A071681 | 001 | Nov 17, 1987 |
| | 0.4MG/ML | A071682 | 001 | Nov 17, 1987 |
| SOLOPAK | 0.02MG/ML | A071672 | 001 | Nov 17, 1987 |
| | 0.4MG/ML | A071683 | 001 | Nov 17, 1987 |
| WATSON LABS | 0.4MG/ML | A071339 | 001 | Nov 18, 1987 |

NARCAN

| | | | | |
|---|----------------------|--------------|---------|-----|
| + | ADAPT | 0.02MG/ML ** | N016636 | 002 |
| + | | 0.4MG/ML ** | N016636 | 001 |
| + | | 1MG/ML ** | N016636 | 003 |
| | BRISTOL MYERS SQUIBB | 0.4MG/ML | A071083 | 001 |
| | | 1MG/ML | A071084 | 001 |
| | | 1MG/ML | A071311 | 001 |

SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS

EVZIO

| | | | | |
|---|-----------|---------------------------|---------|-----|
| + | KALEO INC | 0.4MG/0.4ML (0.4MG/0.4ML) | N205787 | 001 |
|---|-----------|---------------------------|---------|-----|

SPRAY, METERED; NASAL

NARCAN

| | | | | |
|---|-------|-----------|---------|-----|
| + | ADAPT | 2MG/SPRAY | N208411 | 002 |
|---|-------|-----------|---------|-----|

NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TARGINIQ

| | | | | |
|---|------------------|------------|---------|-----|
| + | PURDUE PHARMA LP | 5MG; 10MG | N205777 | 001 |
| + | | 10MG; 20MG | N205777 | 002 |
| + | | 20MG; 40MG | N205777 | 003 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

TALWIN NX

SANOFI AVENTIS US EQ 0.5MG BASE; EQ 50MG BASE ** N018733 001 Dec 16, 1982

NALTREXONE HYDROCHLORIDE

TABLET; ORAL

NALTREXONE HYDROCHLORIDE

FOSUN PHARMA 50MG A075434 001 Mar 08, 2000

REVIA

TEVA WOMENS 50MG N018932 001 Nov 20, 1984

NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

TROXYCA ER

PFIZER INC 1.2MG; 10MG N207621 001 Aug 19, 2016

2.4MG; 20MG N207621 002 Aug 19, 2016

3.6MG; 30MG N207621 003 Aug 19, 2016

4.8MG; 40MG N207621 004 Aug 19, 2016

7.2MG; 60MG N207621 005 Aug 19, 2016

9.6MG; 80MG N207621 006 Aug 19, 2016

NANDROLONE DECANOATE

INJECTABLE; INJECTION

DECA-DURABOLIN

ASPEN GLOBAL INC 50MG/ML N013132 001 Jun 12, 1986

100MG/ML N013132 002 Jun 12, 1986

+ 200MG/ML ** N013132 003 Jun 12, 1986

NANDROLONE DECANOATE

ABRAXIS PHARM 100MG/ML A088290 001 Oct 03, 1983

200MG/ML A088317 001 Oct 14, 1983

AKORN 100MG/ML A087519 001 Sep 28, 1983

WATSON LABS 50MG/ML A086385 001 Jan 13, 1984

50MG/ML A087598 001 Oct 06, 1983

50MG/ML A088554 001 Feb 10, 1986

100MG/ML A086598 001 Jan 13, 1984

100MG/ML A087599 001 Oct 06, 1983

200MG/ML A088128 001 Dec 05, 1983

NANDROLONE PHENPROPIONATE

INJECTABLE; INJECTION

DURABOLIN

ORGANON USA INC 25MG/ML N011891 001

50MG/ML N011891 002

NANDROLONE PHENPROPIONATE

WATSON LABS 25MG/ML A086386 001 Jun 17, 1983

50MG/ML A087488 001 Jun 17, 1983

NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

ALBALON

ALLERGAN 0.1% ** A080248 001

NAFAZAIR

BAUSCH AND LOMB 0.1% A040073 001 May 25, 1994

PHARMAFAIR 0.1% A088101 001 Apr 15, 1983

NAPHAZOLINE HYDROCHLORIDE

AKORN INC 0.1% A083590 001

NAPHCON FORTE

ALCON 0.1% A080229 001

OPCON

BAUSCH AND LOMB 0.1% A087506 001

VASOCON

NOVARTIS 0.1% A080235 002 Mar 24, 1983

NAPROXEN

TABLET; ORAL

NAPROSYN

+ ATNAHS PHARMA US 250MG N017581 002

+ 375MG N017581 003

NAPROXEN

CHARTWELL MOLECULES 250MG A074410 001 Apr 28, 1995

375MG A074410 002 Apr 28, 1995

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NAPROXENTABLET; ORAL
NAPROXEN

| | | | |
|----------------------|-------|-------------|--------------|
| | 500MG | A074410 003 | Apr 28, 1995 |
| DAVA PHARMS INC | 250MG | A074105 001 | Dec 21, 1993 |
| | 375MG | A074105 002 | Dec 21, 1993 |
| | 500MG | A074105 003 | Dec 21, 1993 |
| FOSUN PHARMA | 250MG | A074140 001 | Dec 21, 1993 |
| | 375MG | A074140 002 | Dec 21, 1993 |
| | 500MG | A074140 003 | Dec 21, 1993 |
| HAMILTON PHARMS | 250MG | A074110 001 | Oct 30, 1992 |
| | 375MG | A074110 002 | Oct 30, 1992 |
| | 500MG | A074110 003 | Oct 30, 1992 |
| HIKMA INTL PHARMS | 250MG | A076494 001 | Jan 14, 2004 |
| | 375MG | A076494 002 | Jan 14, 2004 |
| | 500MG | A076494 003 | Jan 14, 2004 |
| IVAX SUB TEVA PHARMS | 250MG | A074111 001 | Feb 28, 1995 |
| | 375MG | A074111 002 | Feb 28, 1995 |
| | 500MG | A074111 003 | Feb 28, 1995 |
| PLIVA | 250MG | A074182 001 | Jun 27, 1996 |
| | 375MG | A074182 002 | Jun 27, 1996 |
| | 500MG | A074182 003 | Jun 27, 1996 |
| PUREPAC PHARM | 250MG | A074263 001 | Dec 21, 1993 |
| | 375MG | A074263 002 | Dec 21, 1993 |
| | 500MG | A074263 003 | Dec 21, 1993 |
| ROXANE | 250MG | A074211 001 | Feb 28, 1994 |
| | 375MG | A074211 002 | Feb 28, 1994 |
| | 500MG | A074211 003 | Feb 28, 1994 |
| TEVA | 250MG | A074129 001 | Dec 21, 1993 |
| | 250MG | A074216 001 | Apr 11, 1996 |
| | 375MG | A074129 002 | Dec 21, 1993 |
| | 375MG | A074216 002 | Apr 11, 1996 |
| | 500MG | A074129 003 | Dec 21, 1993 |
| | 500MG | A074216 003 | Apr 11, 1996 |
| TEVA PHARMS | 250MG | A074207 001 | Dec 21, 1993 |
| | 375MG | A074207 002 | Dec 21, 1993 |
| | 500MG | A074207 003 | Dec 21, 1993 |
| WATSON LABS | 250MG | A074457 001 | May 31, 1995 |
| | 375MG | A074457 002 | May 31, 1995 |
| | 500MG | A074457 003 | May 31, 1995 |
| WATSON LABS TEVA | 250MG | A074163 001 | Feb 10, 1995 |
| | 375MG | A074163 002 | Feb 10, 1995 |
| | 500MG | A074163 003 | Feb 10, 1995 |

TABLET, DELAYED RELEASE; ORAL
NAPROXEN

| | | | |
|-------------------|-------|-------------|--------------|
| ACTAVIS ELIZABETH | 375MG | A074936 001 | Feb 24, 1998 |
| | 500MG | A074936 002 | Feb 24, 1998 |
| FOSUN PHARMA | 375MG | A075061 001 | Feb 18, 1998 |
| | 500MG | A075061 002 | Feb 18, 1998 |
| MYLAN PHARMS INC | 375MG | A075390 001 | Apr 19, 2001 |
| | 500MG | A075390 002 | Apr 19, 2001 |

NAPROXEN SODIUM

TABLET; ORAL

ANAPROX

+ ATNAHS PHARMA US

NAPROXEN SODIUM

| | | | |
|----------------------|---------------|-------------|--------------|
| | EQ 250MG BASE | N018164 001 | |
| ABLE | EQ 250MG BASE | A076544 001 | Aug 22, 2003 |
| | EQ 500MG BASE | A076544 002 | Aug 22, 2003 |
| CONTRACT PHARMACAL | EQ 200MG BASE | A074789 001 | Feb 27, 1997 |
| HAMILTON PHARMS | EQ 250MG BASE | A074106 001 | Aug 31, 1993 |
| | EQ 500MG BASE | A074106 002 | Aug 31, 1993 |
| HIKMA | EQ 250MG BASE | A074480 002 | Feb 18, 1998 |
| | EQ 500MG BASE | A074480 001 | May 14, 1996 |
| IVAX SUB TEVA PHARMS | EQ 250MG BASE | A074230 001 | Mar 14, 1995 |
| | EQ 500MG BASE | A074230 002 | Mar 14, 1995 |
| MYLAN | EQ 250MG BASE | A074367 001 | Aug 31, 1994 |
| | EQ 500MG BASE | A074367 002 | Aug 31, 1994 |
| PLD ACQUISITIONS LLC | EQ 200MG BASE | A074646 001 | Jan 13, 1997 |
| PLIVA | EQ 250MG BASE | A074242 001 | Jun 20, 1996 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NAPROXEN SODIUM

TABLET; ORAL

NAPROXEN SODIUM

| | | | |
|---------------|---------------|-------------|--------------|
| | EQ 500MG BASE | A074242 002 | Jun 20, 1996 |
| PUREPAC PHARM | EQ 250MG BASE | A074319 001 | Mar 20, 1995 |
| | EQ 500MG BASE | A074319 002 | Mar 20, 1995 |
| ROXANE | EQ 250MG BASE | A074257 001 | Dec 21, 1993 |
| | EQ 500MG BASE | A074257 002 | Dec 21, 1993 |
| SANDOZ | EQ 250MG BASE | A074162 001 | Dec 21, 1993 |
| | EQ 250MG BASE | A074495 001 | Dec 05, 1994 |
| | EQ 500MG BASE | A074162 002 | Dec 21, 1993 |
| | EQ 500MG BASE | A074495 002 | Dec 05, 1994 |
| TEVA | EQ 250MG BASE | A074142 001 | Dec 21, 1993 |
| | EQ 500MG BASE | A074142 002 | Dec 21, 1993 |
| TEVA PHARMS | EQ 250MG BASE | A074289 001 | Jan 27, 1994 |
| | EQ 500MG BASE | A074289 002 | Jan 27, 1994 |
| WATSON LABS | EQ 250MG BASE | A074195 001 | Dec 21, 1993 |
| | EQ 250MG BASE | A074455 001 | May 31, 1995 |
| | EQ 500MG BASE | A074195 002 | Dec 21, 1993 |
| | EQ 500MG BASE | A074455 002 | May 31, 1995 |

NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE

TABLET; ORAL

TREXIMET

+ PERNIX IRELAND LTD 60MG;EQ 10MG BASE N021926 002 May 14, 2015

NARATRIPTAN HYDROCHLORIDE

TABLET; ORAL

NARATRIPTAN

| | | | |
|-------------|---------------|-------------|--------------|
| APOTEX CORP | EQ 1MG BASE | A091373 001 | Apr 22, 2011 |
| | EQ 2.5MG BASE | A091373 002 | Apr 22, 2011 |

NATEGLINIDE

TABLET; ORAL

NATEGLINIDE

| | | | |
|-------------|-------|-------------|--------------|
| TEVA PHARMS | 60MG | A077467 001 | Sep 09, 2009 |
| | 120MG | A077467 002 | Sep 09, 2009 |

NEBIVOLOL HYDROCHLORIDE

TABLET; ORAL

NEBIVOLOL HYDROCHLORIDE

| | | | |
|---------------------|---------------|-------------|--------------|
| ALKEM LABS LTD | EQ 2.5MG BASE | A203741 001 | Jun 24, 2015 |
| | EQ 5MG BASE | A203741 002 | Jun 24, 2015 |
| | EQ 10MG BASE | A203741 003 | Jun 24, 2015 |
| | EQ 20MG BASE | A203741 004 | Jun 24, 2015 |
| AMERIGEN PHARMS LTD | EQ 2.5MG BASE | A203659 001 | Apr 16, 2015 |
| | EQ 5MG BASE | A203659 002 | Apr 16, 2015 |
| | EQ 10MG BASE | A203659 003 | Apr 16, 2015 |
| | EQ 20MG BASE | A203659 004 | Apr 16, 2015 |
| GLENMARK PHARMS LTD | EQ 2.5MG BASE | A203821 001 | May 25, 2017 |
| | EQ 5MG BASE | A203821 002 | May 25, 2017 |
| | EQ 10MG BASE | A203821 003 | May 25, 2017 |
| | EQ 20MG BASE | A203821 004 | May 25, 2017 |
| INDCHEMIE HEALTH | EQ 2.5MG BASE | A203828 001 | Jul 29, 2015 |
| | EQ 5MG BASE | A203828 002 | Jul 29, 2015 |
| | EQ 10MG BASE | A203828 003 | Jul 29, 2015 |
| | EQ 20MG BASE | A203828 004 | Jul 29, 2015 |
| TORRENT PHARMS LTD | EQ 2.5MG BASE | A203966 001 | Mar 02, 2018 |
| | EQ 5MG BASE | A203966 002 | Mar 02, 2018 |
| | EQ 10MG BASE | A203966 003 | Mar 02, 2018 |
| | EQ 20MG BASE | A203966 004 | Mar 02, 2018 |
| WATSON LABS INC | EQ 2.5MG BASE | A203683 001 | Nov 27, 2015 |
| | EQ 5MG BASE | A203683 002 | Nov 27, 2015 |
| | EQ 10MG BASE | A203683 003 | Nov 27, 2015 |
| | EQ 20MG BASE | A203683 004 | Nov 27, 2015 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NEDOCROMIL SODIUM

AEROSOL, METERED; INHALATION

TILADE

KING PHARMS LLC 1.75MG/INH

N019660 001 Dec 30, 1992

SOLUTION; INHALATION

TILADE

SANOFI AVENTIS US 0.5%

N020750 001 Oct 01, 1997

NEFAZODONE HYDROCHLORIDE

TABLET; ORAL

NEFAZODONE HYDROCHLORIDE

ANI PHARMS INC

50MG

A076072 001 Sep 16, 2003

100MG

A076072 002 Sep 16, 2003

150MG

A076072 003 Sep 16, 2003

200MG

A076072 004 Sep 16, 2003

250MG

A076072 005 Sep 16, 2003

DR REDDYS LABS INC

50MG

A076309 001 Sep 16, 2003

100MG

A076309 002 Sep 16, 2003

150MG

A076309 003 Sep 16, 2003

200MG

A076309 004 Sep 16, 2003

250MG

A076309 005 Sep 16, 2003

FOSUN PHARMA

50MG

A076302 001 Sep 16, 2003

100MG

A076302 002 Sep 16, 2003

150MG

A076302 003 Sep 16, 2003

200MG

A076302 004 Sep 16, 2003

250MG

A076302 005 Sep 16, 2003

IVAX SUB TEVA PHARMS

50MG

A075763 001 Sep 16, 2003

100MG

A075763 002 Sep 16, 2003

150MG

A075763 003 Sep 16, 2003

200MG

A075763 004 Sep 16, 2003

250MG

A075763 005 Sep 16, 2003

MYLAN

100MG

A076129 002 Sep 16, 2003

150MG

A076129 003 Sep 16, 2003

200MG

A076129 004 Sep 16, 2003

250MG

A076129 005 Sep 16, 2003

ROXANE

50MG

A076196 001 Sep 16, 2003

100MG

A076196 002 Sep 16, 2003

150MG

A076196 003 Sep 16, 2003

200MG

A076196 004 Sep 16, 2003

250MG

A076196 005 Sep 16, 2003

SUN PHARM INDS LTD

50MG

A076409 001 Sep 16, 2003

100MG

A076409 002 Sep 16, 2003

150MG

A076409 003 Sep 16, 2003

200MG

A076409 004 Sep 16, 2003

250MG

A076409 005 Sep 16, 2003

WATSON LABS

100MG

A076073 002 Sep 16, 2003

150MG

A076073 003 Sep 16, 2003

200MG

A076073 004 Sep 16, 2003

250MG

A076073 005 Sep 16, 2003

SERZONE

+ BRISTOL MYERS SQUIBB 50MG **

N020152 001 Dec 22, 1994

+ 100MG **

N020152 002 Dec 22, 1994

+ 150MG **

N020152 003 Dec 22, 1994

+ 200MG **

N020152 004 Dec 22, 1994

+ 250MG **

N020152 005 Dec 22, 1994

+ 300MG **

N020152 006 Dec 22, 1994

NELFINAVIR MESYLATE

POWDER; ORAL

VIRACEPT

AGOURON PHARMS EQ 50MG BASE/SCOOPFUL

N020778 001 Mar 14, 1997

NEOMYCIN SULFATE

POWDER; FOR RX COMPOUNDING

NEO-RX

X GEN PHARMS

100%

A061579 001

SOLUTION; ORAL

MYCIFRADIN

PHARMACIA AND UPJOHN EQ 87.5MG BASE/5ML

N050285 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NEOMYCIN SULFATE

SOLUTION; ORAL

NEO-FRADIN

X GEN PHARMS

EQ 87.5MG BASE/5ML

A065010 001 May 23, 2002

TABLET; ORAL

MYCIFRADIN

PHARMACIA AND UPJOHN EQ 350MG BASE

A060520 001

NEOBIOTIC

PFIZER

EQ 350MG BASE

A060475 001

NEOMYCIN SULFATE

BRISTOL MYERS SQUIBB 500MG

A060365 001

LANNETT 500MG

A060607 001

LILLY 500MG

A060385 001

ROXANE 500MG

A062173 001

SANDOZ 500MG

A061586 001

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

CREAM; TOPICAL

NEOSPORIN

GLAXOSMITHKLINE EQ 3.5MG BASE/GM;10,000 UNITS/GM

N050176 002 Jan 14, 1985

OINTMENT; OPHTHALMIC

STATROL

ALCON EQ 3.5MG BASE/GM;10,000 UNITS/GM

N050344 002

SOLUTION/DROPS; OPHTHALMIC

STATROL

ALCON EQ 3.5MG BASE/ML;16,250 UNITS/ML

A062339 001 Nov 30, 1984

EQ 3.5MG BASE/ML;16,250 UNITS/ML

N050456 001

NEOMYCIN SULFATE; POLYMYXIN B SULFATE; PREDNISOLONE ACETATE

SUSPENSION/DROPS; OPHTHALMIC

POLY-PRED

ALLERGAN EQ 0.35% BASE;10,000 UNITS/ML;0.5%

N050081 002

NEOMYCIN SULFATE; PREDNISOLONE ACETATE

OINTMENT; OPHTHALMIC

NEO-DELTA-CORTEF

PHARMACIA AND UPJOHN EQ 3.5MG BASE/GM;0.25%

A061039 002

EQ 3.5MG BASE/GM;0.5%

A061039 001

SUSPENSION/DROPS; OPHTHALMIC

NEO-DELTA-CORTEF

PHARMACIA AND UPJOHN EQ 3.5MG BASE/ML;0.25%

A061037 001

NEOMYCIN SULFATE; PREDNISOLONE SODIUM PHOSPHATE

OINTMENT; OPHTHALMIC

NEO-HYDELTRASOL

MERCCK EQ 3.5MG BASE/GM;EQ 0.25% PHOSPHATE

N050378 001

NEOMYCIN SULFATE; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

MYTRES A

SAVAGE LABS EQ 3.5MG BASE/GM;0.1%

A062598 001 Jul 21, 1986

NEOMYCIN SULFATE-TRIAMCINOLONE ACETONIDE

FOUGERA EQ 3.5MG BASE/GM;0.1%

A062600 001 Jul 21, 1986

PHARMADERM EQ 3.5MG BASE/GM;0.1%

A062595 001 Jul 21, 1986

OINTMENT; TOPICAL

MYTRES A

SAVAGE LABS EQ 3.5MG BASE/GM;0.1%

A062609 001 May 23, 1986

NEOMYCIN SULFATE-TRIAMCINOLONE ACETONIDE

FOUGERA EQ 3.5MG BASE/GM;0.1%

A062608 001 May 23, 1986

PHARMADERM EQ 3.5MG BASE/GM;0.1%

A062607 001 May 23, 1986

NETILMICIN SULFATE

INJECTABLE; INJECTION

NETROMYCIN

SCHERING EQ 10MG BASE/ML

N050544 001 Feb 28, 1983

EQ 25MG BASE/ML

N050544 002 Feb 28, 1983

EQ 100MG BASE/ML

N050544 003 Feb 28, 1983

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NEVIRAPINE

TABLET; ORAL

NEVIRAPINE

| | | | |
|----------------|-------|-------------|--------------|
| APOTEX INC | 200MG | A203021 001 | May 22, 2012 |
| TECH ORGANIZED | 200MG | A203176 001 | May 22, 2012 |

TABLET, EXTENDED RELEASE; ORAL

NEVIRAPINE

| | | | |
|----------------|-------|-------------|--------------|
| APOTEX INC | 400MG | A205258 001 | Apr 03, 2014 |
| TECH ORGANIZED | 100MG | A207467 001 | Jul 31, 2017 |
| | 400MG | A207467 002 | Jul 31, 2017 |

NIACIN

CAPSULE; ORAL

WAMPOCAP

| | | | |
|---------------------|-------|-------------|--|
| MEDPOINTE PHARM HLC | 500MG | N011073 003 | |
|---------------------|-------|-------------|--|

TABLET; ORAL

NIACIN

| | | | |
|----------------------|-------|-------------|--|
| EVERYLIFE | 500MG | A083203 001 | |
| HALSEY | 500MG | A083453 001 | |
| HIKMA PHARMS | 500MG | A083718 001 | |
| IMPAX LABS | 500MG | A083115 001 | |
| IVAX SUB TEVA PHARMS | 500MG | A083180 001 | |
| MK LABS | 500MG | A083525 001 | |
| PUREPAC PHARM | 500MG | A083271 001 | |
| SANDOZ | 500MG | A083306 001 | |
| TABLICAPS | 500MG | A084237 001 | |
| WATSON LABS | 500MG | A083136 001 | |
| | 500MG | A083305 001 | |
| | 500MG | A085172 001 | |

NICOLAR

| | | | |
|-------------------|-------|-------------|--|
| SANOFI AVENTIS US | 500MG | A083823 001 | |
|-------------------|-------|-------------|--|

TABLET, EXTENDED RELEASE; ORAL

NIASPAN

| | | | |
|--------|-------|-------------|--------------|
| ABBVIE | 375MG | N020381 001 | Jul 28, 1997 |
|--------|-------|-------------|--------------|

NIASPAN TITRATION STARTER PACK

| | | | |
|--------|---------------------|-------------|--------------|
| ABBVIE | 375MG; 500MG; 750MG | N020381 005 | Jul 28, 1997 |
|--------|---------------------|-------------|--------------|

NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; TYROSINE

SUSPENSION; ORAL

TPN

| | | | |
|---------------|---------------------------------|-------------|--|
| INTL MINERALS | 15MG/5ML; 3.75MG/5ML; 600MG/5ML | N008378 003 | |
|---------------|---------------------------------|-------------|--|

NICARDIPINE HYDROCHLORIDE

CAPSULE; ORAL

CARDENE

| | | | |
|----------------|---------|-------------|--------------|
| CHIESI USA INC | 20MG ** | N019488 001 | Dec 21, 1988 |
| | 30MG ** | N019488 002 | Dec 21, 1988 |

NICARDIPINE HYDROCHLORIDE

| | | | |
|-------------|------|-------------|--------------|
| WATSON LABS | 20MG | A074670 001 | Oct 28, 1996 |
| | 30MG | A074670 002 | Oct 28, 1996 |

CAPSULE, EXTENDED RELEASE; ORAL

CARDENE SR

| | | | | |
|---|----------------|---------|-------------|--------------|
| + | CHIESI USA INC | 30MG ** | N020005 001 | Feb 21, 1992 |
| + | | 45MG ** | N020005 002 | Feb 21, 1992 |
| + | | 60MG ** | N020005 003 | Feb 21, 1992 |

INJECTABLE; INJECTION

CARDENE

| | | | | |
|---|----------------|----------------------|-------------|--------------|
| + | CHIESI USA INC | 25MG/10ML (2.5MG/ML) | N019734 001 | Jan 30, 1992 |
|---|----------------|----------------------|-------------|--------------|

NICARDIPINE HYDROCHLORIDE

| | | | |
|----------------------|----------------------|-------------|--------------|
| LUITPOLD | 25MG/10ML (2.5MG/ML) | A090534 001 | Nov 17, 2009 |
| MYLAN INSTITUTIONAL | 25MG/10ML (2.5MG/ML) | A090664 001 | Nov 17, 2009 |
| NAVINTA LLC | 25MG/10ML (2.5MG/ML) | A090125 001 | Nov 17, 2009 |
| SUN PHARMA GLOBAL | 25MG/10ML (2.5MG/ML) | N078405 001 | Nov 17, 2009 |
| WEST-WARD PHARMS INT | 25MG/10ML (2.5MG/ML) | A078714 001 | Dec 28, 2009 |
| WOCKHARDT | 25MG/10ML (2.5MG/ML) | A090671 001 | Nov 17, 2009 |

INJECTABLE; INTRAVENOUS

CARDENE IN 5.0% DEXTROSE IN PLASTIC CONTAINER

| | | | | |
|---|----------------|-----------------------|-------------|--------------|
| + | CHIESI USA INC | 40MG/200ML (0.2MG/ML) | N019734 005 | Nov 07, 2008 |
|---|----------------|-----------------------|-------------|--------------|

NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE

| | | | |
|----------------------|-----------------------|-------------|--------------|
| EXELA PHARMA SCIENCE | 20MG/200ML (0.1MG/ML) | N022276 002 | Apr 07, 2016 |
| | 40MG/200ML (0.2MG/ML) | N022276 003 | Apr 07, 2016 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NICLOSAMIDE

TABLET, CHEWABLE;ORAL

NICLOCIDE

BAYER PHARMS

500MG

N018669 001 May 14, 1982

NICOTINE

FILM, EXTENDED RELEASE;TRANSDERMAL

NICOTROL

MCNEIL CONS

15MG/16HR

N020536 001 Jul 03, 1996

PROSTEP

AVEVA

11MG/24HR

N019983 003 Dec 23, 1998

22MG/24HR

N019983 004 Dec 23, 1998

NICOTINE POLACRILEX

GUM, CHEWING;BUCCAL

NICOTINE POLACRILEX

IVAX SUB TEVA PHARMS

EQ 2MG BASE

A076880 001 Feb 18, 2009

EQ 4MG BASE

A077850 001 Feb 18, 2009

THRIVE

GLAXOSMITHKLINE CONS

EQ 2MG BASE

A077658 001 Jun 19, 2007

EQ 4MG BASE

A077656 001 Jun 19, 2007

NIFEDIPINE

CAPSULE;ORAL

ADALAT

BAYER PHARMS

10MG

N019478 001 Nov 27, 1985

20MG

N019478 002 Sep 17, 1986

NIFEDIPINE

CHASE LABS NJ

10MG

A072409 001 Jul 04, 1990

20MG

A073421 001 Jun 19, 1991

TEVA

10MG

A072651 001 Feb 19, 1992

PROCARDIA

+ PFIZER

20MG **

N018482 002 Jul 24, 1986

TABLET, EXTENDED RELEASE;ORAL

AFEDITAB CR

WATSON LABS

60MG

A075659 001 Oct 26, 2001

WATSON LABS TEVA

30MG

A075128 001 Mar 10, 2000

NIFEDIPINE

MARTEC USA LLC

90MG

A075414 003 Mar 23, 2004

MYLAN

30MG

A075108 001 Dec 17, 1999

MYLAN LABS LTD

30MG

A090602 001 Sep 13, 2010

60MG

A090602 002 Sep 13, 2010

90MG

A090602 003 Sep 13, 2010

NILUTAMIDE

TABLET;ORAL

NILANDRON

CONCORDIA PHARMS INC 50MG

N020169 001 Sep 19, 1996

NIMODIPINE

CAPSULE;ORAL

NIMOTOP

+ BAYER PHARMS

30MG **

N018869 001 Dec 28, 1988

NISOLDIPINE

TABLET, EXTENDED RELEASE;ORAL

SULAR

+ COVIS PHARMA BV

10MG **

N020356 001 Feb 02, 1995

+

20MG **

N020356 002 Feb 02, 1995

+

25.5MG **

N020356 006 Jan 02, 2008

+

30MG **

N020356 003 Feb 02, 1995

+

40MG **

N020356 004 Feb 02, 1995

NITRIC OXIDE

GAS;INHALATION

INOMAX

+ MALLINCKRODT HOSP

100PPM **

N020845 002 Dec 23, 1999

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NITROFURANTOIN

CAPSULE; ORAL

NITROFURANTOIN

| | | | |
|-------------|-------|---------|-----|
| WATSON LABS | 50MG | A084326 | 001 |
| | 100MG | A084326 | 002 |

TABLET; ORAL

FURADANTIN

| | | | |
|--------------------|-------|---------|-----|
| PROCTER AND GAMBLE | 50MG | N008693 | 001 |
| | 100MG | N008693 | 002 |

FURALAN

| | | | |
|---------|-------|---------|-----|
| LANNETT | 50MG | A080017 | 001 |
| | 100MG | A080017 | 002 |

NITROFURANTOIN

| | | | |
|----------------------|-------|---------|-----|
| ELKINS SINN | 50MG | A080003 | 001 |
| | 100MG | A080003 | 002 |
| IVAX SUB TEVA PHARMS | 50MG | A080078 | 002 |
| | 100MG | A080078 | 001 |
| SANDOZ | 50MG | A080043 | 001 |
| | 100MG | A080043 | 002 |
| WATSON LABS | 50MG | A080447 | 001 |
| | 50MG | A085797 | 001 |
| | 100MG | A080447 | 002 |
| | 100MG | A085796 | 001 |
| WHITEWORTH TOWN PLSN | 100MG | A084085 | 002 |

NITROFURANTOIN SODIUM

INJECTABLE; INJECTION

IVADANTIN

| | | | |
|--------------------|--------------------|---------|-----|
| PROCTER AND GAMBLE | EQ 180MG BASE/VIAL | N012402 | 001 |
|--------------------|--------------------|---------|-----|

NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

NITROFURANTOIN

| | | | | |
|-------------|-------|---------|-----|--------------|
| MYLAN | 100MG | A074967 | 002 | Jul 09, 1997 |
| SANDOZ | 25MG | A074336 | 001 | Jan 25, 1995 |
| | 50MG | A074336 | 002 | Jan 25, 1995 |
| | 100MG | A074336 | 003 | Jan 25, 1995 |
| WATSON LABS | 25MG | A073696 | 001 | Dec 31, 1992 |
| | 50MG | A073696 | 002 | Dec 31, 1992 |
| | 100MG | A073696 | 003 | Dec 31, 1992 |

NITROFURANTOIN MACROCRYSTALLINE

| | | | | |
|-------------|-------|---------|-----|--------------|
| WATSON LABS | 50MG | A070248 | 001 | Jun 24, 1988 |
| | 100MG | A070249 | 001 | Jun 24, 1988 |

NITROFURANTOIN; NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS)

| | | | | |
|------------------|------------|---------|-----|--------------|
| RANBAXY LABS LTD | 75MG; 25MG | A076951 | 001 | Mar 30, 2005 |
|------------------|------------|---------|-----|--------------|

NITROFURAZONE

CREAM; TOPICAL

FURACIN

| | | | |
|-------|------|---------|-----|
| SHIRE | 0.2% | A083789 | 001 |
|-------|------|---------|-----|

DRESSING; TOPICAL

ACTIN-N

| | | | |
|----------------|------|---------|-----|
| SHERWOOD MEDCL | 0.2% | N017343 | 001 |
|----------------|------|---------|-----|

OINTMENT; TOPICAL

FURACIN

| | | | |
|-------|------|---------|-----|
| SHIRE | 0.2% | N005795 | 001 |
|-------|------|---------|-----|

NITROFURAZONE

| | | | |
|------------------|------|---------|-----|
| AMBIX | 0.2% | A086077 | 001 |
| LANNETT | 0.2% | A084393 | 001 |
| PERRIGO NEW YORK | 0.2% | A084968 | 001 |
| TARO | 0.2% | A086156 | 001 |
| WENDT | 0.2% | A086766 | 001 |

POWDER; TOPICAL

FURACIN

| | | | |
|-------|------|---------|-----|
| SHIRE | 0.2% | A083791 | 001 |
|-------|------|---------|-----|

SOLUTION; TOPICAL

NITROFURAZONE

| | | | |
|------------------|------|---------|-----|
| PERRIGO NEW YORK | 0.2% | A085130 | 001 |
|------------------|------|---------|-----|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NITROFURAZONE

SOLUTION; TOPICAL

NITROFURAZONE

WENDT

0.2%

A087081 001

NITROGLYCERIN

AEROSOL; SUBLINGUAL

NITROLINGUAL

POHL BOSKAMP

0.4MG/SPRAY

N018705 001 Oct 31, 1985

FILM, EXTENDED RELEASE; TRANSDERMAL

NITROGLYCERIN

LANNETT CO INC

0.2MG/HR

A075115 001 Aug 10, 2004

0.4MG/HR

A075115 002 Aug 10, 2004

MYLAN TECHNOLOGIES

0.1MG/HR

A074992 004 Nov 12, 1999

0.2MG/HR

A074992 003 Nov 12, 1999

0.4MG/HR

A074992 002 Nov 12, 1999

0.6MG/HR

A074992 001 Nov 12, 1999

TRANSDERM-NITRO

+ NOVARTIS

0.1MG/HR **

N020144 001 Feb 27, 1996

+

0.2MG/HR **

N020144 002 Feb 27, 1996

+

0.4MG/HR **

N020144 003 Feb 27, 1996

+

0.6MG/HR **

N020144 004 Feb 27, 1996

+

0.8MG/HR **

N020144 005 Feb 27, 1996

INJECTABLE; INJECTION

NITRO IV

POHL BOSKAMP

5MG/ML

N018672 002 Aug 30, 1983

NITRO-BID

SANOFI AVENTIS US

5MG/ML

N018621 001 Jan 05, 1982

10MG/ML

A071159 001 Feb 28, 1990

NITROGLYCERIN

ABRAXIS PHARM

5MG/ML

A070077 001 Dec 13, 1985

5MG/ML

A071203 001 May 08, 1987

+

HOSPIRA

5MG/ML **

N018531 001

INTL MEDICATION

5MG/ML

A070026 001 Sep 10, 1985

LUITPOLD

5MG/ML

A071492 001 May 24, 1988

SMITH AND NEPHEW

5MG/ML

A070633 001 Jun 19, 1986

5MG/ML

A070634 001 Jun 19, 1986

NITROGLYCERIN IN DEXTROSE 5%

HOSPIRA

0.1MG/ML

A074083 001 Oct 26, 1994

10MG/100ML

A071846 001 Aug 31, 1990

20MG/100ML

A071847 001 Aug 31, 1990

40MG/100ML

A071848 001 Aug 31, 1990

NITROL

RORER

0.8MG/ML

N018774 001 Jan 19, 1983

NITRONAL

POHL BOSKAMP

1MG/ML

N018672 001 Aug 30, 1983

NITROSTAT

PARKE DAVIS

0.8MG/ML

N018588 001

5MG/ML

A070863 001 Jan 08, 1987

5MG/ML

N018588 002 Dec 23, 1983

10MG/ML

A070871 001 Jan 08, 1987

10MG/ML

A070872 001 Jan 08, 1987

TRIDIL

HOSPIRA

0.5MG/ML

N018537 002 Jun 16, 1983

5MG/ML

N018537 001

NIZATIDINE

CAPSULE; ORAL

AXID

SMITHKLINE BEECHAM

150MG

N019508 001 Apr 12, 1988

300MG

N019508 002 Apr 12, 1988

NIZATIDINE

ANI PHARMS INC

150MG

A075461 001 Jul 08, 2002

300MG

A075461 002 Jul 08, 2002

APOTEX INC

150MG

A076383 001 Jan 23, 2003

300MG

A076383 002 Jan 23, 2003

MYLAN PHARMS INC

150MG

A075934 001 Jul 09, 2002

300MG

A075934 002 Jul 09, 2002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NIZATIDINE

SOLUTION;ORAL

AXID

+ BRAINTREE

15MG/ML **

N021494 001 May 25, 2004

NONOXYNOL-9

AEROSOL;VAGINAL

DELFIN

PERSONAL PRODS

12.5%

N014349 002

NOREPINEPHRINE BITARTRATE

INJECTABLE;INJECTION

NOREPINEPHRINE BITARTRATE

METRICS PHARM

EQ 1MG BASE/ML

A040522 001 Sep 30, 2004

NOREPINEPHRINE BITARTRATE; PROCAINE HYDROCHLORIDE; PROPOXYCAINE HYDROCHLORIDE

INJECTABLE;INJECTION

RAVOCAINE AND NOVOCAIN W/ LEVOPHED

EASTMAN KODAK

EQ 0.033MG BASE/ML;2%;0.4%

N008592 003

NORETHINDRONE

TABLET;ORAL

NORLUTIN

PARKE DAVIS

5MG

N010895 002

NORETHINDRONE ACETATE

TABLET;ORAL

AYGESTIN

+ DURAMED RES

5MG

N018405 001 Apr 21, 1982

NORLUTATE

PARKE DAVIS

5MG

N012184 002

NORFLOXACIN

SOLUTION/DROPS;OPHTHALMIC

CHIBROXIN

MERCK

0.3%

N019757 001 Jun 17, 1991

TABLET;ORAL

NOROXIN

+ MERCK

400MG **

N019384 002 Oct 31, 1986

NORGESTREL

TABLET;ORAL

OPILL

+ LABORATOIRE HRA

0.075MG

N017031 001

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE;ORAL

AVENTYL HYDROCHLORIDE

LILLY

EQ 10MG BASE

N014684 001

EQ 25MG BASE

N014684 002

NORTRIPTYLINE HYDROCHLORIDE

ANI PHARMS INC

EQ 10MG BASE

A074054 001 Dec 31, 1992

EQ 25MG BASE

A074054 002 Dec 31, 1992

EQ 50MG BASE

A074054 003 Dec 31, 1992

EQ 75MG BASE

A074054 004 Dec 31, 1992

AUROLIFE PHARMA LLC

EQ 10MG BASE

A074835 001 Jun 30, 1997

EQ 25MG BASE

A074835 002 Jun 30, 1997

EQ 50MG BASE

A074835 003 Jun 30, 1997

EQ 75MG BASE

A074835 004 Jun 30, 1997

MYLAN

EQ 10MG BASE

A074234 001 Jul 26, 1993

EQ 25MG BASE

A074234 002 Jul 26, 1993

EQ 50MG BASE

A074234 003 Jul 26, 1993

EQ 75MG BASE

A074234 004 Jul 26, 1993

TEVA

EQ 10MG BASE

A073667 001 Apr 11, 1996

EQ 25MG BASE

A073667 002 Apr 11, 1996

EQ 50MG BASE

A073667 003 Apr 11, 1996

EQ 75MG BASE

A073667 004 Apr 11, 1996

SOLUTION;ORAL

AVENTYL

+ RANBAXY

EQ 10MG BASE/5ML **

N014685 001

PAMELOR

SPECGX LLC

EQ 10MG BASE/5ML

N018012 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NYSTATIN

CREAM; TOPICAL

CANDEX

BAYER PHARMS 100,000 UNITS/GM A061810 001

MYCOSTATIN

DELCOR ASSET CORP 100,000 UNITS/GM ** A060575 001

MYKINAC

ALPHARMA US PHARMS 100,000 UNITS/GM A062387 001 Jul 29, 1982

NILSTAT

LEDERLE 100,000 UNITS/GM A061445 001

NYSTATIN

TARO 100,000 UNITS/GM A062457 001 Jul 28, 1983

LOTION; TOPICAL

CANDEX

BAYER PHARMS 100,000 UNITS/ML N050233 001

OINTMENT; TOPICAL

MYCOSTATIN

DELCOR ASSET CORP 100,000 UNITS/GM ** A060571 001

MYKINAC

ALPHARMA US PHARMS 100,000 UNITS/GM A062731 001 Sep 22, 1986

NILSTAT

LEDERLE 100,000 UNITS/GM A061444 001

PASTILLE; ORAL

MYCOSTATIN

DELCOR ASSET CORP 200,000 UNITS N050619 001 Apr 09, 1987

POWDER; ORAL

BARSTATIN 100

BARLAN 100% A062489 001 Apr 27, 1988

NILSTAT

+ DAVA PHARMS INC 100% ** N050576 001 Dec 22, 1983

NYSTATIN

PADDOCK LLC 100% A062613 001 Nov 26, 1985

POWDER; TOPICAL

MYCOSTATIN

DELCOR ASSET CORP 100,000 UNITS/GM ** A060578 001

NYSTATIN

NESHER PHARMS 100,000 UNITS/GM A065321 001 Aug 18, 2006

SUPPOSITORY; VAGINAL

NYSERT

WARNER CHILCOTT 100,000 UNITS N050478 001

SUSPENSION; ORAL

MYCOSTATIN

DELCOR ASSET CORP 100,000 UNITS/ML A061533 001

NILSTAT

+ GLENMARK GENERICS 100,000 UNITS/ML ** N050299 001

NYSTATIN

ALPHARMA US PHARMS 100,000 UNITS/ML A062571 001 Oct 29, 1985

G AND W LABS INC 100,000 UNITS/ML A062349 001 Jul 14, 1982

100,000 UNITS/ML A062776 001 Dec 17, 1987

MORTON GROVE 100,000 UNITS/ML A062835 001 Nov 19, 1987

PHARMADERM 100,000 UNITS/ML A062518 001 Jul 06, 1984

PHARMAFAIR 100,000 UNITS/ML A062541 001 Jan 16, 1985

SOCORRO 100,000 UNITS/ML A062832 001 Dec 27, 1991

TEVA 100,000 UNITS/ML A062670 001 Jun 18, 1987

NYSTEX

SAVAGE LABS 100,000 UNITS/ML A062519 001 Jul 06, 1984

TABLET; ORAL

MYCOSTATIN

DELCOR ASSET CORP 500,000 UNITS A060574 001

NILSTAT

LEDERLE 500,000 UNITS A061151 001

NYSTATIN

QUANTUM PHARMICS 500,000 UNITS A062525 001 Oct 29, 1984

SANDOZ 500,000 UNITS A062065 001

USL PHARMA 500,000 UNITS A062524 001 Nov 26, 1985

WATSON LABS 500,000 UNITS A062402 001 Dec 16, 1982

TABLET; VAGINAL

KOROSTATIN

HOLLAND RANTOS 100,000 UNITS A061718 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

NYSTATIN

TABLET;VAGINAL

MYCOSTATIN

DELCOR ASSET CORP 100,000 UNITS A060577 001

NILSTAT

LEDERLE 100,000 UNITS A061325 001

NYSTATIN

FOUGERA 100,000 UNITS A062459 001 Nov 09, 1983

ODYSSEY PHARMS 100,000 UNITS A062615 001 Oct 17, 1985

PHARMADERM 100,000 UNITS A062460 001 Nov 09, 1983

QUANTUM PHARMICS 100,000 UNITS A062509 001 Apr 03, 1984

SANDOZ 100,000 UNITS A061965 001

TEVA 100,000 UNITS A062502 001 Dec 23, 1983

WATSON LABS 100,000 UNITS A062176 001

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM;TOPICAL

MYCO-TRIA CET II

TEVA 100,000 UNITS/GM;0.1% A061954 002 Sep 20, 1985

MYCOLOG-II

DELCOR ASSET CORP 100,000 UNITS/GM;0.1% ** A060576 002 May 01, 1985

MYLAN PHARMS INC 100,000 UNITS/GM;0.1% ** A062606 001 May 15, 1985

MYTRES F

SAVAGE LABS 100,000 UNITS/GM;0.1% A062597 001 Oct 08, 1985

NYSTATIN AND TRIAMCINOLONE ACETONIDE

ALPHARMA US PHARMS 100,000 UNITS/GM;0.1% A063010 001 Dec 20, 1988

PERRIGO NEW YORK 100,000 UNITS/GM;0.1% A062186 002 Jun 06, 1985

PHARMAFAIR 100,000 UNITS/GM;0.1% A062657 001 Jul 30, 1986

TARO 100,000 UNITS/GM;0.1% A062347 001 Mar 30, 1987

NYSTATIN TRIAMCINOLONE ACETONIDE

PHARMADERM 100,000 UNITS/GM;0.1% A062596 001 Oct 08, 1985

OINTMENT;TOPICAL

MYCO-TRIA CET II

TEVA 100,000 UNITS/GM;0.1% A062045 002 Nov 26, 1985

MYCOLOG-II

MYLAN PHARMS INC 100,000 UNITS/GM;0.1% ** A060572 001 Jun 28, 1985

MYTRES F

SAVAGE LABS 100,000 UNITS/GM;0.1% A062601 001 Oct 09, 1985

NYSTATIN AND TRIAMCINOLONE ACETONIDE

PERRIGO NEW YORK 100,000 UNITS/GM;0.1% A062280 002 Oct 10, 1985

PHARMAFAIR 100,000 UNITS/GM;0.1% A062656 001 Jul 30, 1986

NYSTATIN-TRIAMCINOLONE ACETONIDE

PHARMADERM 100,000 UNITS/GM;0.1% A062603 001 Oct 09, 1985

OCTREOTIDE ACETATE

INJECTABLE;INJECTION

OCTREOTIDE ACETATE

SUN PHARM INDS EQ 0.05MG BASE/ML A077329 001 Mar 04, 2008

EQ 0.1MG BASE/ML A077329 002 Mar 04, 2008

EQ 0.2MG BASE/ML A077330 001 Mar 04, 2008

EQ 0.5MG BASE/ML A077329 003 Mar 04, 2008

EQ 1MG BASE/ML A077331 001 Mar 04, 2008

WOCKHARDT USA EQ 0.2MG BASE/ML A090986 001 May 11, 2011

EQ 1MG BASE/ML A090986 002 May 11, 2011

OCTREOTIDE ACETATE PRESERVATIVE FREE

WOCKHARDT USA EQ 0.05MG BASE/ML A090985 001 May 11, 2011

EQ 0.1MG BASE/ML A090985 002 May 11, 2011

EQ 0.5MG BASE/ML A090985 003 May 11, 2011

OFLOXACIN

INJECTABLE;INJECTION

FLOXIN

ORTHO MCNEIL PHARM 20MG/ML N020087 002 Mar 31, 1992

40MG/ML N020087 003 Mar 31, 1992

FLOXIN IN DEXTROSE 5%

ORTHO MCNEIL PHARM 400MG/100ML N020087 001 Mar 31, 1992

FLOXIN IN DEXTROSE 5% IN PLASTIC CONTAINER

ORTHO MCNEIL PHARM 4MG/ML N020087 004 Mar 31, 1992

400MG/100ML N020087 005 Mar 31, 1992

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OFLOXACIN

INJECTABLE; INJECTION

OFLOXACIN

BEDFORD

40MG/ML

A075762 001 Jan 16, 2002

SOLUTION/DROPS; OPHTHALMIC

OFLOXACIN

APOTEX INC

0.3%

A076513 001 May 14, 2004

SANDOZ

0.3%

A076848 001 Nov 25, 2008

SOLUTION/DROPS; OTIC

FLOXIN OTIC

+ DAIICHI

0.3% **

N020799 001 Dec 16, 1997

TABLET; ORAL

FLOXIN

JANSSEN PHARMS

200MG **

N019735 001 Dec 28, 1990

300MG **

N019735 002 Dec 28, 1990

400MG **

N019735 003 Dec 28, 1990

OFLOXACIN

LARKEN LABS

200MG

A076093 001 Sep 02, 2003

300MG

A076093 002 Sep 02, 2003

RANBAXY LABS LTD

200MG

A076220 001 Sep 02, 2003

300MG

A076220 002 Sep 02, 2003

400MG

A076220 003 Sep 02, 2003

OLANZAPINE

TABLET; ORAL

OLANZAPINE

AJANTA PHARMA LTD

2.5MG

A206711 001 Aug 30, 2016

5MG

A206711 002 Aug 30, 2016

7.5MG

A206711 003 Aug 30, 2016

10MG

A206711 004 Aug 30, 2016

15MG

A206711 005 Aug 30, 2016

20MG

A206711 006 Aug 30, 2016

MYLAN PHARMS INC

2.5MG

A076866 001 Apr 23, 2012

5MG

A076866 002 Apr 23, 2012

7.5MG

A076866 003 Apr 23, 2012

10MG

A076866 004 Apr 23, 2012

15MG

A076866 005 Apr 23, 2012

20MG

A076866 006 Apr 23, 2012

TABLET, ORALLY DISINTEGRATING; ORAL

OLANZAPINE

AJANTA PHARMA LTD

5MG

A204320 001 May 30, 2017

10MG

A204320 002 May 30, 2017

15MG

A204320 003 May 30, 2017

20MG

A204320 004 May 30, 2017

OLIVE OIL; SOYBEAN OIL

INJECTABLE; INJECTION

CLINOLIPID 20%

+ BAXTER HLTHCARE CORP 16%(160GM/1000ML); 4% (40GM/1000ML)

N204508 001 Oct 03, 2013

OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

OLOPATADINE HYDROCHLORIDE

ZAMBON SPA

EQ 0.1% BASE

A204706 001 Dec 07, 2015

OMEGA-3-ACID ETHYL ESTERS TYPE A

CAPSULE; ORAL

OMTRYG

+ OSMOTICA

1.2GM CONTAINS AT LEAST 900MG OF THE
ETHYL ESTERS OF OMEGA-3 FATTY ACIDS

N204977 001 Apr 23, 2014

OMEGA-3-CARBOXYLIC ACIDS

CAPSULE; ORAL

EPANOVA

+ ASTRAZENECA PHARMS

1GM CONTAINS AT LEAST 850MG OF
POLYUNSATURATED FATTY ACIDS

N205060 001 May 05, 2014

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

PRILOSEC

| | | | | |
|---|--------------------|---------|-------------|--------------|
| + | ASTRAZENECA PHARMS | 10MG ** | N019810 003 | Oct 05, 1995 |
| + | | 20MG ** | N019810 001 | Sep 14, 1989 |
| + | | 40MG ** | N019810 002 | Jan 15, 1998 |

ONDANSETRON

TABLET, ORALLY DISINTEGRATING;ORAL

ONDANSETRON

| | | | | |
|--|---------------------|------|-------------|--------------|
| | CHARTWELL MOLECULES | 4MG | A077406 003 | Dec 26, 2006 |
| | | 8MG | A077406 004 | Dec 26, 2006 |
| | | 16MG | A077406 001 | Dec 26, 2006 |
| | | 24MG | A077406 002 | Dec 26, 2006 |
| | NESHER PHARMS | 4MG | A077717 001 | Jun 25, 2007 |
| | | 8MG | A077717 002 | Jun 25, 2007 |

ONDANSETRON HYDROCHLORIDE

INJECTABLE; INJECTION

ONDANSETRON HYDROCHLORIDE

| | | | | |
|--|---------------------|----------------|-------------|--------------|
| | APOTEX INC | EQ 2MG BASE/ML | A077368 001 | Dec 26, 2006 |
| | HOSPIRA | EQ 2MG BASE/ML | A076695 001 | Dec 26, 2006 |
| | LANNETT CO INC | EQ 2MG BASE/ML | A090116 001 | Apr 14, 2010 |
| | | EQ 2MG BASE/ML | A090883 001 | Aug 05, 2010 |
| | LUITPOLD | EQ 2MG BASE/ML | A077582 001 | Dec 26, 2006 |
| | MYLAN LABS LTD | EQ 2MG BASE/ML | A078257 001 | Apr 23, 2008 |
| | PLIVA HRVATSKA DOO | EQ 2MG BASE/ML | A077544 001 | Dec 26, 2006 |
| | SAGENT PHARMS | EQ 2MG BASE/ML | A078180 001 | Mar 26, 2007 |
| | SUN PHARM INDS (IN) | EQ 2MG BASE/ML | A077172 001 | Dec 26, 2006 |

ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER

| | | | | |
|--|---------|-------------------|-------------|--------------|
| | HOSPIRA | EQ 0.64MG BASE/ML | A076978 001 | Feb 26, 2007 |
|--|---------|-------------------|-------------|--------------|

ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

| | | | | |
|--|---------------------|----------------|-------------|--------------|
| | APOTEX INC | EQ 2MG BASE/ML | A077343 001 | Dec 26, 2006 |
| | HOSPIRA | EQ 2MG BASE/ML | A076696 001 | Dec 26, 2006 |
| | LUITPOLD | EQ 2MG BASE/ML | A077387 001 | Dec 26, 2006 |
| | MYLAN LABS LTD | EQ 2MG BASE/ML | A078244 001 | Apr 23, 2008 |
| | SUN PHARM INDS LTD | EQ 2MG BASE/ML | A077173 001 | Dec 26, 2006 |
| | TARO PHARMS IRELAND | EQ 2MG BASE/ML | A078014 001 | Mar 21, 2008 |

ZOFRAN

| | | | | |
|---|----------------------|-------------------|-------------|--------------|
| + | NOVARTIS PHARMS CORP | EQ 2MG BASE/ML ** | N020007 001 | Jan 04, 1991 |
|---|----------------------|-------------------|-------------|--------------|

ZOFRAN AND DEXTROSE IN PLASTIC CONTAINER

| | | | | |
|---|-----------------|----------------------|-------------|--------------|
| + | GLAXOSMITHKLINE | EQ 0.64MG BASE/ML ** | N020403 001 | Jan 31, 1995 |
|---|-----------------|----------------------|-------------|--------------|

ZOFRAN PRESERVATIVE FREE

| | | | | |
|---|----------------------|-------------------|-------------|--------------|
| + | NOVARTIS PHARMS CORP | EQ 2MG BASE/ML ** | N020007 003 | Dec 10, 1993 |
|---|----------------------|-------------------|-------------|--------------|

TABLET;ORAL

ONDANSETRON HYDROCHLORIDE

| | | | | |
|--|---------------------|--------------|-------------|--------------|
| | CHARTWELL MOLECULES | EQ 4MG BASE | A077303 001 | Jun 25, 2007 |
| | | EQ 8MG BASE | A077303 002 | Jun 25, 2007 |
| | | EQ 24MG BASE | A077303 004 | Jun 25, 2007 |
| | HIKMA INTL PHARMS | EQ 4MG BASE | A077545 001 | Sep 06, 2007 |
| | | EQ 8MG BASE | A077545 002 | Sep 06, 2007 |
| | | EQ 24MG BASE | A077545 003 | Sep 06, 2007 |
| | TARO | EQ 4MG BASE | A077729 001 | Mar 28, 2011 |
| | | EQ 8MG BASE | A077729 002 | Mar 28, 2011 |
| | | EQ 24MG BASE | A077729 003 | Mar 28, 2011 |

ORPHENADRINE CITRATE

INJECTABLE; INJECTION

NORFLEX

| | | | | |
|---|----------|---------|-------------|--|
| + | TELIGENT | 30MG/ML | N013055 001 | |
|---|----------|---------|-------------|--|

ORPHENADRINE CITRATE

| | | | | |
|--|-------------|---------|-------------|--|
| | WATSON LABS | 30MG/ML | A087062 001 | |
|--|-------------|---------|-------------|--|

TABLET, EXTENDED RELEASE;ORAL

NORFLEX

| | | | | |
|---|---------|-------|-------------|--|
| + | MEDICIS | 100MG | N012157 001 | |
|---|---------|-------|-------------|--|

ORPHENADRINE CITRATE

| | | | | |
|--|-------------|-------|-------------|--------------|
| | ASCOT | 100MG | A088067 001 | Apr 06, 1983 |
| | SANDOZ | 100MG | A085046 001 | |
| | WATSON LABS | 100MG | A084303 001 | |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ORPHENADRINE HYDROCHLORIDE

TABLET; ORAL

DISIPAL

3M

50MG

N010653 001

OSELTAMIVIR PHOSPHATE

FOR SUSPENSION; ORAL

TAMIFLU

ROCHE

EQ 12MG BASE/ML

N021246 001 Dec 14, 2000

OXACILLIN SODIUM

CAPSULE; ORAL

BACTOCILL

GLAXOSMITHKLINE

EQ 250MG BASE

A061336 001

EQ 250MG BASE

A062241 001

EQ 500MG BASE

A061336 002

EQ 500MG BASE

A062241 002

OXACILLIN SODIUM

ANI PHARMS INC

EQ 250MG BASE

A062222 001

EQ 500MG BASE

A062222 002

APOTHECON

EQ 250MG BASE

A061450 002

EQ 500MG BASE

A061450 001

PROSTAPHLIN

APOTHECON

EQ 500MG BASE

N050118 002

FOR SOLUTION; ORAL

BACTOCILL

GLAXOSMITHKLINE

EQ 250MG BASE/5ML

A062321 001

OXACILLIN SODIUM

APOTHECON

EQ 250MG BASE/5ML

A061457 001

TEVA

EQ 250MG BASE/5ML

A062252 001

PROSTAPHLIN

APOTHECON

EQ 250MG BASE/5ML

N050194 001

INJECTABLE; INJECTION

BACTOCILL

GLAXOSMITHKLINE

EQ 500MG BASE/VIAL **

A061334 009 Mar 26, 1982

EQ 1GM BASE/VIAL **

A061334 006 Mar 26, 1982

EQ 1GM BASE/VIAL **

A062736 001 Dec 19, 1986

EQ 2GM BASE/VIAL **

A061334 007 Mar 26, 1982

EQ 2GM BASE/VIAL **

A062736 002 Dec 19, 1986

EQ 4GM BASE/VIAL **

A061334 008 Mar 26, 1982

EQ 10GM BASE/VIAL **

A061334 010

OXACILLIN SODIUM

+ APOTHECON

EQ 250MG BASE/VIAL **

N050195 001

+ APOTHECON

EQ 500MG BASE/VIAL **

N050195 002

+ APOTHECON

EQ 1GM BASE/VIAL **

N050195 003

+ APOTHECON

EQ 2GM BASE/VIAL **

N050195 004

+ APOTHECON

EQ 4GM BASE/VIAL **

N050195 005

ELKINS SINN

EQ 250MG BASE/VIAL

A062711 001 Feb 03, 1989

EQ 500MG BASE/VIAL

A062711 002 Feb 03, 1989

EQ 1GM BASE/VIAL

A062711 003 Feb 03, 1989

EQ 2GM BASE/VIAL

A062711 004 Feb 03, 1989

EQ 4GM BASE/VIAL

A062711 005 Feb 03, 1989

EQ 10GM BASE/VIAL

A062711 006 Feb 03, 1989

ISTITUTO BIO ITA SPA

EQ 125MG BASE/VIAL

A062798 003 Dec 11, 1995

EQ 250MG BASE/VIAL

A062798 004 Dec 11, 1995

EQ 500MG BASE/VIAL

A062798 005 Dec 11, 1995

EQ 1GM BASE/VIAL

A062798 001 Dec 11, 1995

EQ 2GM BASE/VIAL

A062798 002 Dec 11, 1995

MYLAN LABS LTD

EQ 1GM BASE/VIAL

A091486 001 Aug 25, 2014

EQ 2GM BASE/VIAL

A091486 002 Aug 25, 2014

SANDOZ

EQ 250MG BASE/VIAL

A061490 001

EQ 500MG BASE/VIAL

A061490 002

EQ 1GM BASE/VIAL

A061490 003

EQ 2GM BASE/VIAL

A061490 004

EQ 10GM BASE/VIAL

A061490 006 May 09, 1991

WATSON LABS INC

EQ 250MG BASE/VIAL

A062856 001 Oct 26, 1988

EQ 500MG BASE/VIAL

A062856 002 Oct 26, 1988

EQ 1GM BASE/VIAL

A062856 003 Oct 26, 1988

EQ 2GM BASE/VIAL

A062856 004 Oct 26, 1988

EQ 4GM BASE/VIAL

A062856 005 Oct 26, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXACILLIN SODIUM

INJECTABLE; INJECTION

OXACILLIN SODIUM

EQ 10GM BASE/VIAL

A062984 001 Sep 29, 1988

POWDER; INTRAVENOUS

OXACILLIN SODIUM

SANDOZ

EQ 1GM BASE/VIAL

A062737 001 Dec 23, 1986

EQ 2GM BASE/VIAL

A062737 002 Dec 23, 1986

OXALIPLATIN

INJECTABLE; IV (INFUSION)

ELOXATIN

+ SANOFI AVENTIS US

50MG/VIAL **

N021492 001 Aug 09, 2002

+

100MG/VIAL **

N021492 002 Aug 09, 2002

+

200MG/40ML (5MG/ML) **

N021759 003 Nov 17, 2006

OXALIPLATIN

FRESENIUS KABI ONCOL

50MG/VIAL

A078810 001 Aug 07, 2009

100MG/VIAL

A078810 002 Aug 07, 2009

SANDOZ

50MG/VIAL

A090849 001 Apr 28, 2011

100MG/VIAL

A090849 002 Apr 28, 2011

SANDOZ INC

50MG/10ML (5MG/ML)

A078812 001 Aug 07, 2009

100MG/20ML (5MG/ML)

A078812 002 Aug 07, 2009

OXAMNIQUINE

CAPSULE; ORAL

VANSIL

PFIZER

250MG

N018069 001

OXANDROLONE

TABLET; ORAL

OXANDRIN

+ GEMINI LABS LLC

2.5MG

N013718 001

+

10MG

N013718 002 Nov 05, 2001

OXANDROLONE

ROXANE

2.5MG

A077249 001 Jul 10, 2007

10MG

A077249 002 Jul 10, 2007

SANDOZ

2.5MG

A076897 001 Dec 01, 2006

10MG

A076897 002 Dec 01, 2006

OXAPROZIN

TABLET; ORAL

OXAPROZIN

ACTAVIS ELIZABETH

600MG

A075843 001 Oct 03, 2001

MYLAN

600MG

A075851 001 Aug 17, 2001

MYLAN PHARMS INC

600MG

A075847 001 Feb 28, 2001

SANDOZ

600MG

A075842 001 Apr 12, 2001

600MG

A075850 001 Apr 27, 2001

WATSON LABS

600MG

A075848 001 Feb 09, 2001

OXAPROZIN POTASSIUM

TABLET; ORAL

DAYPRO ALTA

GD SEARLE

600MG

N020776 001 Oct 17, 2002

OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

AM THERAP

10MG

A071955 001 Mar 03, 1988

15MG

A071956 001 Mar 03, 1988

30MG

A071957 001 Mar 03, 1988

FRONTIDA BIOPHARM

10MG

A071026 002 Aug 10, 1987

15MG

A071026 003 Aug 10, 1987

30MG

A071026 001 Aug 10, 1987

IVAX SUB TEVA PHARMS

10MG

A070943 001 Aug 03, 1987

15MG

A070944 001 Aug 03, 1987

30MG

A070945 001 Aug 03, 1987

MYLAN

10MG

A071713 001 Oct 20, 1987

15MG

A071714 001 Oct 20, 1987

30MG

A071715 001 Oct 20, 1987

WATSON LABS

15MG

A072953 001 Sep 28, 1990

30MG

A072954 001 Sep 28, 1990

WATSON LABS TEVA

10MG

A072952 001 Sep 28, 1990

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXAZEPAM

CAPSULE; ORAL

SERAX

| | | |
|--------------------|---------|-------------|
| ALPHARMA US PHARMS | 10MG ** | N015539 002 |
| | 15MG ** | N015539 004 |
| | 30MG ** | N015539 006 |

ZAXOPAM

| | | | |
|------------------|------|-------------|--------------|
| QUANTUM PHARMICS | 10MG | A070650 001 | Mar 01, 1988 |
| | 15MG | A070640 001 | Mar 01, 1988 |
| | 30MG | A070641 001 | Mar 01, 1988 |

TABLET; ORAL

OXAZEPAM

| | | | |
|----------------------|------|-------------|--------------|
| PARKE DAVIS | 15MG | A071508 001 | Feb 02, 1987 |
| SUN PHARM INDUSTRIES | 15MG | A070683 001 | Jan 16, 1987 |
| WATSON LABS | 15MG | A071494 001 | Apr 21, 1987 |

SERAX

| | | |
|--------------------|---------|-------------|
| ALPHARMA US PHARMS | 15MG ** | N015539 008 |
|--------------------|---------|-------------|

OXCARBAZEPINE

TABLET; ORAL

OXCARBAZEPINE

| | | | |
|------------------|-------|-------------|--------------|
| JUBILANT CADISTA | 150MG | A090239 001 | Jan 25, 2010 |
| | 300MG | A090239 002 | Jan 25, 2010 |
| | 600MG | A090239 003 | Jan 25, 2010 |

OXPRENOLOL HYDROCHLORIDE

CAPSULE; ORAL

TRASICOR

| | | | |
|----------|-------|-------------|--------------|
| NOVARTIS | 20MG | N018166 001 | Dec 28, 1983 |
| | 40MG | N018166 002 | Dec 28, 1983 |
| | 80MG | N018166 003 | Dec 28, 1983 |
| | 160MG | N018166 004 | Dec 28, 1983 |

OXTRIPHYLLINE

SOLUTION; ORAL

CHOLEDYL

| | | | |
|-------------|-----------|-------------|--------------|
| PARKE DAVIS | 100MG/5ML | N009268 012 | Nov 27, 1984 |
|-------------|-----------|-------------|--------------|

OXTRIPHYLLINE

| | | | |
|--------------|-----------|-------------|--------------|
| MORTON GROVE | 100MG/5ML | A088243 001 | Dec 05, 1983 |
|--------------|-----------|-------------|--------------|

SYRUP; ORAL

CHOLEDYL

| | | |
|-------------|----------|-------------|
| PARKE DAVIS | 50MG/5ML | N009268 011 |
|-------------|----------|-------------|

OXTRIPHYLLINE PEDIATRIC

| | | | |
|--------------|----------|-------------|--------------|
| MORTON GROVE | 50MG/5ML | A088242 001 | Dec 05, 1983 |
|--------------|----------|-------------|--------------|

TABLET, DELAYED RELEASE; ORAL

CHOLEDYL

| | | |
|-------------|-------|-------------|
| PARKE DAVIS | 100MG | N009268 003 |
| | 200MG | N009268 007 |

OXTRIPHYLLINE

| | | | |
|-------------|-------|-------------|--------------|
| WATSON LABS | 100MG | A087866 001 | Aug 25, 1983 |
| | 200MG | A087835 001 | Aug 25, 1983 |

TABLET, EXTENDED RELEASE; ORAL

CHOLEDYL SA

| | | | |
|---------------------|-------|-------------|--------------|
| WARNER CHILCOTT LLC | 400MG | A087863 001 | May 24, 1983 |
| | 600MG | A086742 001 | |

OXYBUTYNIN

FILM, EXTENDED RELEASE; TRANSDERMAL

OXYBUTYNIN

| | | | |
|--------------------|------------|-------------|--------------|
| BARR LABS DIV TEVA | 3.9MG/24HR | A090526 001 | Mar 04, 2014 |
|--------------------|------------|-------------|--------------|

GEL, METERED; TRANSDERMAL

GELNIQUE 3%

| | | | |
|----------------------|----|-------------|--------------|
| + ALLERGAN SALES LLC | 3% | N202513 001 | Dec 07, 2011 |
|----------------------|----|-------------|--------------|

OXYBUTYNIN CHLORIDE

SYRUP; ORAL

DITROPAN

| | | |
|------------------------|------------|-------------|
| + ORTHO MCNEIL JANSSEN | 5MG/5ML ** | N018211 001 |
|------------------------|------------|-------------|

OXYBUTYNIN CHLORIDE

| | | | |
|------------|---------|-------------|--------------|
| APOTEX INC | 5MG/5ML | A074997 001 | Oct 15, 1997 |
| MIKART | 5MG/5ML | A075039 001 | Jan 29, 1999 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXYBUTYNIN CHLORIDE

TABLET; ORAL

DITROPAN

+ JANSSEN PHARMS 5MG ** N017577 001

OXYBUTYNIN CHLORIDE

QUANTUM PHARMICS 5MG A072296 001 Dec 08, 1988

USL PHARMA 5MG A070746 001 Mar 10, 1988

WATSON LABS 5MG A072485 001 Apr 19, 1989

TABLET, EXTENDED RELEASE; ORAL

DITROPAN XL

+ JANSSEN PHARMS 15MG ** N020897 003 Jun 22, 1999

OXYCODONE HYDROCHLORIDE

TABLET; ORAL

ROXYBOND

DAIICHI SANKYO INC 5MG N209777 001 Apr 20, 2017

15MG N209777 002 Apr 20, 2017

30MG N209777 003 Apr 20, 2017

TABLET, EXTENDED RELEASE; ORAL

ROXICODONE

ROXANE 10MG N020932 001 Oct 26, 1998

30MG N020932 002 Oct 26, 1998

OXYMETAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

OCUCLEAR

BAYER HEALTHCARE LLC 0.025% N018471 001 May 30, 1986

OXYMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION

OPANA

+ ENDO PHARMS 1MG/ML N011707 002
1.5MG/ML N011707 001

SUPPOSITORY; RECTAL

NUMORPHAN

ENDO PHARMS 5MG N011738 004

TABLET, EXTENDED RELEASE; ORAL

OPANA ER

+ ENDO PHARMS 5MG ** N021610 001 Jun 22, 2006
5MG N201655 001 Dec 09, 2011
7.5MG ** N021610 005 Feb 29, 2008
7.5MG N201655 002 Dec 09, 2011
10MG ** N021610 002 Jun 22, 2006
10MG N201655 003 Dec 09, 2011
15MG ** N021610 006 Feb 29, 2008
15MG N201655 004 Dec 09, 2011
20MG ** N021610 003 Jun 22, 2006
20MG N201655 005 Dec 09, 2011
30MG ** N021610 007 Feb 29, 2008
30MG N201655 006 Dec 09, 2011
40MG ** N021610 004 Jun 22, 2006
40MG N201655 007 Dec 09, 2011

OXYMORPHONE HYDROCHLORIDE

PAR PHARM

5MG A200792 001 Oct 24, 2014

7.5MG A200792 002 Oct 24, 2014

10MG A200792 003 Oct 24, 2014

15MG A200792 004 Oct 24, 2014

20MG A200792 005 Oct 24, 2014

30MG A200792 006 Oct 24, 2014

40MG A200792 007 Oct 24, 2014

SUN PHARM INDS LTD 5MG A203506 001 Apr 24, 2015

7.5MG A203506 002 Apr 24, 2015

10MG A203506 003 Apr 24, 2015

15MG A203506 004 Apr 24, 2015

20MG A203506 005 Apr 24, 2015

30MG A203506 006 Apr 24, 2015

40MG A203506 007 Apr 24, 2015

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

OXYPHENBUTAZONE

TABLET; ORAL

OXYPHENBUTAZONE

WATSON LABS

100MG

A088399 001 Sep 17, 1984

TANDEARIL

NOVARTIS

100MG

N012542 004 Sep 03, 1982

OXYPHENCYCLIMINE HYDROCHLORIDE

TABLET; ORAL

DARICON

PFIZER

10MG

N011612 001

OXYPHENONIUM BROMIDE

TABLET; ORAL

ANTRENYL

NOVARTIS

5MG

N008492 002

OXYTETRACYCLINE

TABLET; ORAL

TERRAMYCIN

PFIZER

250MG

N050287 001

OXYTETRACYCLINE CALCIUM

SYRUP; ORAL

TERRAMYCIN

PFIZER

EQ 125MG BASE/5ML

A060595 001

OXYTETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

OXY-KESSO-TETRA

FERRANTE

EQ 250MG BASE

A060179 001

OXYTETRACYCLINE HYDROCHLORIDE

HIKMA PHARMS

EQ 250MG BASE

A060770 001

IMPAX LABS

EQ 250MG BASE

A060760 001

PROTER

EQ 250MG BASE

A060869 001

PUREPAC PHARM

EQ 250MG BASE

A060634 001

TERRAMYCIN

PFIZER

EQ 125MG BASE

N050286 001

EQ 250MG BASE

N050286 002

INJECTABLE; INJECTION

TERRAMYCIN

PFIZER

EQ 250MG BASE/VIAL

A060586 001

EQ 500MG BASE/VIAL

A060586 002

OXYTETRACYCLINE HYDROCHLORIDE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

TERRAMYCIN W/ POLYMYXIN B SULFATE

CASPER PHARMA LLC

EQ 5MG BASE/GM;10,000 UNITS/GM

N061015 001

OINTMENT; OTIC

TERRAMYCIN W/ POLYMYXIN

PFIZER

EQ 5MG BASE/GM;10,000 UNITS/GM

A061841 001

TABLET; VAGINAL

TERRAMYCIN-POLYMYXIN

PFIZER

EQ 100MG BASE;100,000 UNITS

A061009 001

OXYTOCIN

INJECTABLE; INJECTION

OXYTOCIN

TEVA PHARMS USA

10USP UNITS/ML (10USP UNITS/ML)

A077453 001 Jan 24, 2008

100USP UNITS/10ML (10USP UNITS/ML)

A077453 002 Jan 24, 2008

OXYTOCIN 10 USP UNITS IN DEXTROSE 5%

+ ABBOTT

1USP UNITS/100ML **

N019185 004 Mar 29, 1985

+

2USP UNITS/100ML **

N019185 003 Mar 29, 1985

OXYTOCIN 20 USP UNITS IN DEXTROSE 5%

+ ABBOTT

2USP UNITS/100ML **

N019185 002 Mar 29, 1985

OXYTOCIN 5 USP UNITS IN DEXTROSE 5%

+ ABBOTT

1USP UNITS/100ML **

N019185 001 Mar 29, 1985

SYNTOCINON

NOVARTIS

10USP UNITS/ML

N018245 001

SOLUTION; NASAL

SYNTOCINON

RTRX

40USP UNITS/ML

N012285 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PACLITAXEL

INJECTABLE; INJECTION

PACLITAXEL

| | | | | |
|-----------------|--------|---------|-----|--------------|
| ACCORD HLTHCARE | 6MG/ML | A075436 | 001 | Nov 12, 2004 |
| HOSPIRA | 6MG/ML | A076233 | 001 | Aug 01, 2002 |
| MYLAN | 6MG/ML | A075278 | 001 | Jan 25, 2002 |
| PLIVA LACHEMA | 6MG/ML | A077413 | 001 | Mar 12, 2008 |
| TEVA PHARMS USA | 6MG/ML | A075297 | 001 | Jan 25, 2002 |

PALIPERIDONE

TABLET, EXTENDED RELEASE; ORAL

INVEGA

| | | | | |
|------------------|---------|---------|-----|--------------|
| + JANSSEN PHARMS | 12MG ** | N021999 | 004 | Dec 19, 2006 |
|------------------|---------|---------|-----|--------------|

PALONOSETRON HYDROCHLORIDE

CAPSULE; ORAL

ALOXI

| | | | | |
|--------------------|------------------|---------|-----|--------------|
| + HELSINN HLTHCARE | EQ 0.5MG BASE ** | N022233 | 001 | Aug 22, 2008 |
|--------------------|------------------|---------|-----|--------------|

SOLUTION; INTRAVENOUS

PALONOSETRON HYDROCHLORIDE

| | | | | |
|--------------------|-------------------------------------------|---------|-----|--------------|
| DR REDDYS LABS LTD | EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML) | N203050 | 001 | Mar 01, 2016 |
| | EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML) | N203050 | 002 | Mar 01, 2016 |

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

ARELIA

| | | | | |
|------------|--------------|---------|-----|--------------|
| + NOVARTIS | 30MG/VIAL ** | N020036 | 001 | Oct 31, 1991 |
| | 60MG/VIAL | N020036 | 003 | May 06, 1993 |
| | 90MG/VIAL | N020036 | 004 | May 06, 1993 |

PAMIDRONATE DISODIUM

| | | | | |
|-----------|-----------|---------|-----|--------------|
| AESGEN | 30MG/VIAL | A075594 | 001 | May 06, 2002 |
| | 90MG/VIAL | A075594 | 002 | May 06, 2002 |
| MN PHARMS | 30MG/VIAL | A078300 | 001 | Mar 10, 2009 |
| | 90MG/VIAL | A078300 | 002 | Mar 10, 2009 |

PANCRELIPASE (AMYLASE; LIPASE; PROTEASE)

CAPSULE; ORAL

COTAZYM

| | | | | |
|-----------------|--------------------------------------------------|---------|-----|--------------|
| ORGANON USA INC | 30,000USP UNITS; 8,000USP UNITS; 30,000USP UNITS | N020580 | 001 | Dec 09, 1996 |
|-----------------|--------------------------------------------------|---------|-----|--------------|

CAPSULE, DELAYED RELEASE; ORAL

ULTRESA

| | | | | |
|-------------------|---------------------------------------------------|---------|-----|--------------|
| + FOREST LABS INC | 27,600USP UNITS; 13,800USP UNITS; 27,600USP UNITS | N022222 | 001 | Mar 01, 2012 |
| | 41,400USP UNITS; 20,700USP UNITS; 41,400USP UNITS | N022222 | 002 | Mar 01, 2012 |
| | 46,000USP UNITS; 23,000USP UNITS; 46,000USP UNITS | N022222 | 003 | Mar 01, 2012 |

PANCURONIUM BROMIDE

INJECTABLE; INJECTION

PANCURONIUM BROMIDE

| | | | | |
|--------------|--------|---------|-----|--------------|
| ELKINS SINN | 1MG/ML | A072058 | 001 | Mar 23, 1988 |
| | 2MG/ML | A072059 | 001 | Mar 23, 1988 |
| | 2MG/ML | A072060 | 001 | Mar 23, 1988 |
| HOSPIRA | 2MG/ML | A072321 | 001 | Jan 19, 1989 |
| IGI LABS INC | 1MG/ML | A072210 | 001 | Mar 31, 1988 |
| | 2MG/ML | A072211 | 001 | Mar 31, 1988 |
| | 2MG/ML | A072212 | 001 | Mar 31, 1988 |
| | 2MG/ML | A072213 | 001 | Mar 31, 1988 |

PAVULON

| | | | | |
|-------------------|--------|---------|-----|--|
| + ORGANON USA INC | 1MG/ML | N017015 | 002 | |
| | 2MG/ML | N017015 | 001 | |

PANTOPRAZOLE SODIUM

TABLET, DELAYED RELEASE; ORAL

PANTOPRAZOLE SODIUM

| | | | | |
|--------------------|--------------|---------|-----|--------------|
| SUN PHARM INDS LTD | EQ 20MG BASE | A077058 | 001 | Sep 10, 2007 |
| | EQ 20MG BASE | A200794 | 001 | May 02, 2012 |
| | EQ 40MG BASE | A077058 | 002 | Sep 10, 2007 |
| | EQ 40MG BASE | A200794 | 002 | May 02, 2012 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PARAMETHADIONE

CAPSULE; ORAL

PARADIONE

ABBVIE

150MG

N006800 003

300MG

N006800 001

SOLUTION; ORAL

PARADIONE

ABBVIE

300MG/ML

N006800 002

PARAMETHASONE ACETATE

TABLET; ORAL

HALDRONE

LILLY

1MG

N012772 005

2MG

N012772 006

PARGYLINE HYDROCHLORIDE

TABLET; ORAL

EUTONYL

ABBOTT

10MG

N013448 002

25MG

N013448 003

50MG

N013448 004

PARICALCITOL

CAPSULE; ORAL

ZEMPLAR

+ ABBVIE

4MCG **

N021606 003 May 26, 2005

PAROMOMYCIN SULFATE

CAPSULE; ORAL

HUMATIN

KING PFIZER

EQ 250MG BASE

A062310 001

PARKEDALE

EQ 250MG BASE

A060521 001

SYRUP; ORAL

HUMATIN

PARKE DAVIS

EQ 125MG BASE/5ML

A060522 001

PAROXETINE HYDROCHLORIDE

CAPSULE; ORAL

PAXIL

+ APOTEX TECHNOLOGIES

EQ 10MG BASE **

N020885 001 Oct 09, 1998

+

EQ 20MG BASE **

N020885 002 Oct 09, 1998

+

EQ 30MG BASE **

N020885 003 Oct 09, 1998

+

EQ 40MG BASE **

N020885 004 Oct 09, 1998

SUSPENSION; ORAL

PAROXETINE HYDROCHLORIDE

APOTEX INC

EQ 10MG BASE/5ML

A077395 001 Dec 05, 2006

TABLET; ORAL

PAROXETINE HYDROCHLORIDE

MYLAN PHARMS INC

EQ 10MG BASE

A075716 001 Mar 08, 2004

EQ 20MG BASE

A075716 002 Mar 08, 2004

EQ 30MG BASE

A075716 003 Mar 08, 2004

EQ 40MG BASE

A075716 004 Mar 08, 2004

ROXANE

EQ 10MG BASE

A078026 001 Jun 29, 2007

EQ 20MG BASE

A078026 002 Jun 29, 2007

EQ 30MG BASE

A078026 003 Jun 29, 2007

EQ 40MG BASE

A078026 004 Jun 29, 2007

TEVA PHARMS

EQ 10MG BASE

A077082 001 Jun 29, 2007

EQ 20MG BASE

A077082 002 Jun 29, 2007

EQ 30MG BASE

A077082 003 Jun 29, 2007

EQ 40MG BASE

A077082 004 Jun 29, 2007

UPSHER SMITH LABS

EQ 10MG BASE

A075566 001 Mar 08, 2004

EQ 20MG BASE

A075566 002 Mar 08, 2004

EQ 30MG BASE

A075566 003 Mar 08, 2004

EQ 40MG BASE

A075566 004 Mar 08, 2004

PAXIL

APOTEX TECHNOLOGIES

EQ 50MG BASE

N020031 004 Dec 29, 1992

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PAZOPANIB HYDROCHLORIDE

TABLET;ORAL

VOTRIENT

NOVARTIS PHARMS CORP EQ 400MG BASE

N022465 002 Oct 19, 2009

PEGINESATIDE ACETATE

SOLUTION;INTRAVENOUS, SUBCUTANEOUS

OMONTYS

TAKEDA PHARMS USA EQ 10MG BASE/ML (EQ 10MG BASE/ML)

N202799 007 Mar 27, 2012

EQ 20MG BASE/2ML (EQ 10MG BASE/ML)

N202799 008 Mar 27, 2012

OMONTYS PRESERVATIVE FREE

TAKEDA PHARMS USA EQ 1MG BASE/0.5ML (EQ 1MG BASE/0.5ML)

N202799 001 Mar 27, 2012

EQ 2MG BASE/0.5ML (EQ 2MG BASE/0.5ML)

N202799 002 Mar 27, 2012

EQ 3MG BASE/0.5ML (EQ 3MG BASE/0.5ML)

N202799 003 Mar 27, 2012

EQ 4MG BASE/0.5ML (EQ 4MG BASE/0.5ML)

N202799 004 Mar 27, 2012

EQ 5MG BASE/0.5ML (EQ 5MG BASE/0.5ML)

N202799 005 Mar 27, 2012

EQ 6MG BASE/0.5ML (EQ 6MG BASE/0.5ML)

N202799 006 Mar 27, 2012

PEMIROLAST POTASSIUM

SOLUTION/DROPS;OPHTHALMIC

ALAMAST

SANTEN 0.1%

N021079 001 Sep 24, 1999

PEMOLINE

TABLET;ORAL

CYLERT

ABBOTT 18.75MG

N016832 001

37.5MG

N016832 002

75MG

N016832 003

PEMOLINE

ACTAVIS ELIZABETH 18.75MG

A075595 001 Feb 28, 2000

37.5MG

A075595 002 Feb 28, 2000

75MG

A075595 003 Feb 28, 2000

FOSUN PHARMA 18.75MG

A075286 001 Dec 27, 1999

37.5MG

A075286 002 Jun 30, 1999

75MG

A075286 003 Jun 30, 1999

MALLINCKRODT 18.75MG

A075726 003 Mar 30, 2001

37.5MG

A075726 002 Mar 30, 2001

75MG

A075726 001 Mar 30, 2001

TEVA PHARMS 18.75MG

A075030 003 Feb 22, 2000

37.5MG

A075030 001 Jan 29, 1999

75MG

A075030 002 Jan 29, 1999

VINTAGE PHARMS 18.75MG

A075328 001 Apr 19, 2000

37.5MG

A075328 002 Apr 19, 2000

75MG

A075328 003 Apr 19, 2000

WATSON LABS 18.75MG

A075287 001 Jun 13, 2001

37.5MG

A075287 002 Sep 18, 2000

75MG

A075287 003 Sep 18, 2000

TABLET, CHEWABLE;ORAL

CYLERT

ABBOTT 37.5MG

N017703 001

PEMOLINE

ACTAVIS ELIZABETH 37.5MG

A075678 001 Jul 26, 2000

TEVA PHARMS 37.5MG

A075555 001 Feb 18, 2000

PENBUTOLOL SULFATE

TABLET;ORAL

LEVATOL

+ AUXILIUM PHARMS LLC 10MG **

N018976 001 Dec 30, 1987

+ 20MG **

N018976 004 Jan 05, 1989

PENICILLAMINE

CAPSULE;ORAL

CUPRIMINE

ATON 125MG

N019853 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENICILLIN G BENZATHINE

INJECTABLE; INJECTION

BICILLIN L-A

| | | | | |
|---|-----------------|------------------|---------|-----|
| + | KING PHARMS LLC | 300,000 UNITS/ML | N050141 | 003 |
| | WYETH AYERST | 300,000 UNITS/ML | N050131 | 001 |

PERMAPEN

| | | | | |
|--|-------------------|------------------|---------|-----|
| | CASPER PHARMA LLC | 600,000 UNITS/ML | N060014 | 001 |
|--|-------------------|------------------|---------|-----|

SUSPENSION; ORAL

BICILLIN

| | | | | |
|--|--------------|-------------------|---------|-----|
| | WYETH AYERST | 300,000 UNITS/5ML | N050126 | 002 |
|--|--------------|-------------------|---------|-----|

TABLET; ORAL

BICILLIN

| | | | | |
|--|--------------|---------------|---------|-----|
| | WYETH AYERST | 200,000 UNITS | N050128 | 001 |
|--|--------------|---------------|---------|-----|

PENICILLIN G BENZATHINE; PENICILLIN G PROCAINE

INJECTABLE; INJECTION

BICILLIN C-R

| | | | | |
|---|-----------------|------------------------------------|---------|-----|
| + | KING PHARMS LLC | 150,000 UNITS/ML; 150,000 UNITS/ML | N050138 | 002 |
|---|-----------------|------------------------------------|---------|-----|

PENICILLIN G POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN

| | | | | |
|--|------|-------------------|---------|-----|
| | TEVA | 200,000 UNITS/5ML | A060307 | 002 |
| | | 400,000 UNITS/5ML | A060307 | 004 |

PENICILLIN G POTASSIUM

| | | | | |
|--|---------------|-------------------|---------|-----|
| | MYLAN | 200,000 UNITS/5ML | A060752 | 003 |
| | | 250,000 UNITS/5ML | A060752 | 002 |
| | | 400,000 UNITS/5ML | A060752 | 001 |
| | PUREPAC PHARM | 250,000 UNITS/5ML | A061740 | 001 |
| | | 400,000 UNITS/5ML | A061740 | 002 |

PENICILLIN-2

| | | | | |
|--|------|-------------------|---------|-----|
| | TEVA | 250,000 UNITS/5ML | A060307 | 003 |
|--|------|-------------------|---------|-----|

PENTIDS '200'

| | | | | |
|--|-----------|-------------------|---------|-----|
| | APOTHECON | 200,000 UNITS/5ML | A062149 | 001 |
|--|-----------|-------------------|---------|-----|

PENTIDS '400'

| | | | | |
|--|-----------|-------------------|---------|-----|
| | APOTHECON | 400,000 UNITS/5ML | A062149 | 002 |
|--|-----------|-------------------|---------|-----|

PFIZERPEN G

| | | | | |
|--|--------|-------------------|---------|-----|
| | PFIZER | 400,000 UNITS/5ML | A060587 | 001 |
|--|--------|-------------------|---------|-----|

INJECTABLE; INJECTION

PENICILLIN G POTASSIUM

| | | | | |
|--|--------------------|-------------------------|---------|------------------|
| | APOTHECON | 1,000,000 UNITS/VIAL | A060362 | 001 |
| | | 5,000,000 UNITS/VIAL | A060362 | 003 |
| | | 10,000,000 UNITS/VIAL | A060362 | 004 |
| | | 20,000,000 UNITS/VIAL | A060362 | 002 |
| | CONSOLIDATED PHARM | 500,000 UNITS/VIAL | A060806 | 001 |
| | | 1,000,000 UNITS/VIAL | A060806 | 002 |
| | | 5,000,000 UNITS/VIAL | A060806 | 003 |
| | | 10,000,000 UNITS/VIAL | A060806 | 004 |
| | LILLY | 200,000 UNITS/VIAL | A060384 | 004 |
| | | 500,000 UNITS/VIAL | A060384 | 003 |
| | | 1,000,000 UNITS/VIAL | A060384 | 002 |
| | | 5,000,000 UNITS/VIAL | A060384 | 001 |
| | | 20,000,000 UNITS/VIAL | A060384 | 005 |
| | | 20,000,000 UNITS/VIAL | A060601 | 001 |
| | PARKE DAVIS | 1,000,000 UNITS/VIAL | A062003 | 001 |
| | | 5,000,000 UNITS/VIAL | A062003 | 002 |
| | PFIZER | 20,000,000 UNITS/VIAL | A060074 | 003 |
| | SANDOZ | 1,000,000 UNITS/VIAL ** | A065079 | 001 Aug 30, 2002 |
| | WATSON LABS INC | 1,000,000 UNITS/VIAL | A062991 | 001 Sep 13, 1988 |
| | | 5,000,000 UNITS/VIAL | A062991 | 002 Sep 13, 1988 |
| | | 10,000,000 UNITS/VIAL | A062991 | 003 Sep 13, 1988 |
| | | 20,000,000 UNITS/VIAL | A062991 | 004 Sep 13, 1988 |

PFIZERPEN

| | | | | |
|--|--------|-------------------------|---------|-----|
| | PFIZER | 1,000,000 UNITS/VIAL ** | A060657 | 001 |
|--|--------|-------------------------|---------|-----|

TABLET; ORAL

PENICILLIN G POTASSIUM

| | | | | |
|--|----------------------|---------------|---------|-----|
| | APOTHECON | 250,000 UNITS | A060392 | 003 |
| | IVAX SUB TEVA PHARMS | 400,000 UNITS | A060073 | 004 |
| | LILLY | 250,000 UNITS | A060403 | 001 |
| | MYLAN | 200,000 UNITS | A060781 | 001 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENICILLIN G POTASSIUM

TABLET;ORAL

PENICILLIN G POTASSIUM

| | | |
|---------------|---------------|-------------|
| | 250,000 UNITS | A060781 002 |
| | 400,000 UNITS | A060781 003 |
| | 500,000 UNITS | A060781 005 |
| | 800,000 UNITS | A060781 004 |
| PUREPAC PHARM | 200,000 UNITS | A061588 001 |
| | 250,000 UNITS | A061588 002 |
| | 400,000 UNITS | A061588 003 |
| TEVA | 200,000 UNITS | A060306 001 |
| | 250,000 UNITS | A060306 002 |
| | 400,000 UNITS | A060306 003 |
| | 500,000 UNITS | A060306 004 |
| WYETH AYERST | 200,000 UNITS | A060413 001 |
| | 250,000 UNITS | A060413 002 |
| | 400,000 UNITS | A060413 003 |
| PENTIDS '200' | | |
| APOTHECON | 200,000 UNITS | A062155 001 |
| PENTIDS '250' | | |
| APOTHECON | 250,000 UNITS | A062155 002 |
| PENTIDS '400' | | |
| APOTHECON | 400,000 UNITS | A060392 004 |
| | 400,000 UNITS | A062155 003 |
| PENTIDS '800' | | |
| APOTHECON | 800,000 UNITS | A060392 005 |
| | 800,000 UNITS | A062155 004 |
| PFIZERPEN G | | |
| PFIZER | 50,000 UNITS | A060075 001 |
| | 100,000 UNITS | A060075 002 |
| | 200,000 UNITS | A060075 003 |
| | 250,000 UNITS | A060075 004 |
| | 400,000 UNITS | A060075 005 |
| | 800,000 UNITS | A060075 006 |

PENICILLIN G PROCAINE

INJECTABLE;INJECTION

DURACILLIN A.S.

| | | |
|-----------------------|----------------------|-------------|
| LILLY | 300,000 UNITS/ML | A060093 001 |
| PENICILLIN G PROCAINE | | |
| CONSOLIDATED PHARM | 300,000 UNITS/ML | A060800 001 |
| | 600,000 UNITS/1.2ML | A060800 002 |
| PARKE DAVIS | 300,000 UNITS/ML | A062029 001 |
| PFIZER | 300,000 UNITS/VIAL | A060099 001 |
| | 1,500,000 UNITS/VIAL | A060099 002 |
| PFIZERPEN-AS | | |
| PFIZER | 300,000 UNITS/ML | A060286 001 |
| | 600,000 UNITS/ML | A060286 002 |

PENICILLIN G SODIUM

INJECTABLE;INJECTION

PENICILLIN G SODIUM

| | | |
|---------------------------------------|----------------------|--------------------------|
| BRISTOL MYERS SQUIBB | 5,000,000 UNITS/VIAL | A061935 001 |
| COPANOS | 5,000,000 UNITS/VIAL | A061051 001 |
| PHARMACIA AND UPJOHN | 1,000,000 UNITS/VIAL | A061046 001 |
| INJECTABLE;INTRAMUSCULAR, INTRAVENOUS | | |
| PENICILLIN G SODIUM | | |
| WATSON LABS INC | 5,000,000 UNITS/VIAL | A063014 001 Sep 13, 1988 |

PENICILLIN V

FOR SUSPENSION;ORAL

V-CILLIN

| | | |
|-------|-------------|-------------|
| LILLY | 125MG/0.6ML | A060002 001 |
|-------|-------------|-------------|

PENICILLIN V POTASSIUM

FOR SOLUTION;ORAL

BEEPEN-VK

| | | |
|-----------------|-------------------|-------------|
| GLAXOSMITHKLINE | EQ 125MG BASE/5ML | A062270 001 |
| | EQ 250MG BASE/5ML | A062270 002 |
| BETAPEN-VK | | |
| APOTHECON | EQ 125MG BASE/5ML | A061149 001 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENICILLIN V POTASSIUM

FOR SOLUTION;ORAL

| | | |
|------------------------|-------------------|-------------|
| BETAPEN-VK | EQ 250MG BASE/5ML | A061149 002 |
| LEDERCILLIN VK | | |
| LEDERLE | EQ 125MG BASE/5ML | A060136 001 |
| | EQ 250MG BASE/5ML | A060136 002 |
| PEN-VEE K | | |
| WYETH AYERST | EQ 125MG BASE/5ML | A060007 001 |
| | EQ 250MG BASE/5ML | A060007 002 |
| PENAPAR-VK | | |
| PARKE DAVIS | EQ 125MG BASE/5ML | A062002 001 |
| | EQ 250MG BASE/5ML | A062002 002 |
| PENICILLIN V POTASSIUM | | |
| AM ANTIBIOTICS | EQ 125MG BASE/5ML | A061529 001 |
| | EQ 250MG BASE/5ML | A061529 002 |
| MYLAN | EQ 125MG BASE/5ML | A061624 002 |
| | EQ 250MG BASE/5ML | A061624 001 |
| PUREPAC PHARM | EQ 125MG BASE/5ML | A061758 001 |
| | EQ 250MG BASE/5ML | A061758 002 |
| PFIZERPEN VK | | |
| PFIZER | EQ 125MG BASE/5ML | A061815 001 |
| | EQ 250MG BASE/5ML | A061815 002 |
| V-CILLIN K | | |
| LILLY | EQ 125MG BASE/5ML | A060004 001 |
| | EQ 250MG BASE/5ML | A060004 002 |
| VEETIDS | | |
| APOTHECON | EQ 125MG BASE/5ML | A061410 001 |
| | EQ 250MG BASE/5ML | A061410 002 |
| VEETIDS '125' | | |
| APOTHECON | EQ 125MG BASE/5ML | A061206 001 |
| | EQ 125MG BASE/5ML | A062153 001 |
| VEETIDS '250' | | |
| APOTHECON | EQ 250MG BASE/5ML | A061206 002 |
| | EQ 250MG BASE/5ML | A062153 002 |
| TABLET;ORAL | | |
| BEEPEN-VK | | |
| GLAXOSMITHKLINE | EQ 250MG BASE | A062273 001 |
| | EQ 500MG BASE | A062273 002 |
| BETAPEN-VK | | |
| BRISTOL | EQ 250MG BASE | A061150 001 |
| | EQ 500MG BASE | A061150 002 |
| LEDERCILLIN VK | | |
| LEDERLE | EQ 250MG BASE | A060134 001 |
| | EQ 500MG BASE | A060134 002 |
| PEN-VEE K | | |
| WYETH AYERST | EQ 125MG BASE | A060006 001 |
| | EQ 250MG BASE | A060006 002 |
| | EQ 500MG BASE | A060006 003 |
| PENAPAR-VK | | |
| PARKE DAVIS | EQ 250MG BASE | A062001 001 |
| | EQ 500MG BASE | A062001 002 |
| PENICILLIN V POTASSIUM | | |
| AM ANTIBIOTICS | EQ 250MG BASE | A061528 001 |
| | EQ 500MG BASE | A061528 002 |
| IVAX SUB TEVA PHARMS | EQ 125MG BASE | A060518 001 |
| | EQ 250MG BASE | A060518 002 |
| | EQ 500MG BASE | A060518 003 |
| MYLAN | EQ 250MG BASE | A061530 001 |
| | EQ 500MG BASE | A061530 002 |
| PUREPAC PHARM | EQ 125MG BASE | A061571 001 |
| | EQ 250MG BASE | A061571 002 |
| | EQ 500MG BASE | A061571 003 |
| PFIZERPEN VK | | |
| PFIZER | EQ 250MG BASE | A061836 001 |
| | EQ 500MG BASE | A061836 002 |
| UTICILLIN VK | | |
| PHARMACIA AND UPJOHN | EQ 250MG BASE | A061651 001 |
| | EQ 500MG BASE | A061651 002 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENICILLIN V POTASSIUM

TABLET; ORAL

V-CILLIN K

| | | |
|---------------|------------------|-------------|
| LILLY | EQ 125MG BASE ** | A060003 001 |
| | EQ 250MG BASE ** | A060003 002 |
| | EQ 500MG BASE ** | A060003 003 |
| VEETIDS | | |
| APOTHECON | EQ 250MG BASE | A061411 001 |
| | EQ 500MG BASE | A061411 002 |
| VEETIDS '250' | | |
| APOTHECON | EQ 250MG BASE | A061164 001 |
| | EQ 250MG BASE | A062156 002 |
| VEETIDS '500' | | |
| APOTHECON | EQ 500MG BASE | A061164 002 |
| | EQ 500MG BASE | A062156 001 |

PENTAGASTRIN

INJECTABLE; INJECTION

PEPTAVLON

| | | |
|----------------|--------------|-------------|
| + WYETH AYERST | 0.25MG/ML ** | N017048 001 |
|----------------|--------------|-------------|

PENTAMIDINE ISETHIONATE

FOR SOLUTION; INHALATION

NEBUPENT

| | | | |
|--------------------|------------|-------------|--------------|
| FRESENIUS KABI USA | 600MG/VIAL | N019887 002 | Mar 22, 1996 |
|--------------------|------------|-------------|--------------|

INJECTABLE; INJECTION

PENTACARINAT

| | | | |
|--------------|------------|-------------|--------------|
| ARMOUR PHARM | 300MG/VIAL | A073447 001 | Apr 28, 1994 |
|--------------|------------|-------------|--------------|

PENTAMIDINE ISETHIONATE

| | | | |
|-----------------|------------|-------------|--------------|
| BAXTER HLTHCARE | 300MG/VIAL | A073617 001 | Dec 18, 1995 |
|-----------------|------------|-------------|--------------|

| | | | |
|---------|------------|-------------|--------------|
| HOSPIRA | 300MG/VIAL | A073479 001 | Jun 30, 1992 |
|---------|------------|-------------|--------------|

| | | | |
|-------------|------------|-------------|--------------|
| WATSON LABS | 300MG/VIAL | A074303 001 | Aug 17, 1995 |
|-------------|------------|-------------|--------------|

PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

TALWIN 50

| | | |
|-------------------|--------------|-------------|
| SANOFI AVENTIS US | EQ 50MG BASE | N016732 001 |
|-------------------|--------------|-------------|

PENTAZOCINE LACTATE

INJECTABLE; INJECTION

TALWIN

| | | |
|-----------|-----------------|-------------|
| + HOSPIRA | EQ 30MG BASE/ML | N016194 001 |
|-----------|-----------------|-------------|

PENTETATE CALCIUM TRISODIUM YB-169

INJECTABLE; INJECTION

YTTERBIUM YB 169 DTPA

| | | |
|----|---------|-------------|
| 3M | 2mCi/ML | N017518 001 |
|----|---------|-------------|

PENTOBARBITAL

ELIXIR; ORAL

NEMBUTAL

| | | |
|------------|------------|-------------|
| OAK PHARMS | 18.2MG/5ML | A083244 001 |
|------------|------------|-------------|

PENTOBARBITAL SODIUM

CAPSULE; ORAL

NEMBUTAL SODIUM

| | | |
|------------|------|-------------|
| OAK PHARMS | 30MG | A084095 001 |
|------------|------|-------------|

| | | |
|--|------|-------------|
| | 50MG | A084093 001 |
|--|------|-------------|

| | | |
|--|-------|-------------|
| | 100MG | A083245 001 |
|--|-------|-------------|

PENTOBARBITAL SODIUM

| | | |
|---------|------|-------------|
| LANNETT | 50MG | A085937 001 |
|---------|------|-------------|

| | | |
|--|-------|-------------|
| | 100MG | A085915 001 |
|--|-------|-------------|

| | | |
|----------|-------|-------------|
| VITARINE | 100MG | A083284 001 |
|----------|-------|-------------|

| | | |
|----------------------|-------|-------------|
| WHITEWORTH TOWN PLSN | 100MG | A083338 001 |
|----------------------|-------|-------------|

SODIUM PENTOBARBITAL

| | | |
|----------|-------|-------------|
| ANABOLIC | 100MG | A084590 001 |
|----------|-------|-------------|

| | | |
|-------------|-------|-------------|
| ELKINS SINN | 100MG | A083368 001 |
|-------------|-------|-------------|

| | | |
|-----------|-------|-------------|
| EVERYLIFE | 100MG | A083259 001 |
|-----------|-------|-------------|

| | | |
|--------|-------|-------------|
| HALSEY | 100MG | A084677 001 |
|--------|-------|-------------|

| | | |
|----------------------|------|-------------|
| IVAX SUB TEVA PHARMS | 50MG | A083461 001 |
|----------------------|------|-------------|

| | | |
|--|-------|-------------|
| | 100MG | A083461 002 |
|--|-------|-------------|

| | | |
|-------------|-------|-------------|
| PARKE DAVIS | 100MG | A084156 001 |
|-------------|-------|-------------|

| | | |
|---------|-------|-------------|
| PERRIGO | 100MG | A084560 001 |
|---------|-------|-------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PENTOBARBITAL SODIUM

CAPSULE; ORAL

SODIUM PENTOBARBITAL

| | | | |
|--------------------|-------|---------|-----|
| PUREPAC PHARM | 100MG | A083301 | 001 |
| VALEANT PHARM INTL | 100MG | A083264 | 001 |
| WATSON LABS | 100MG | A085791 | 001 |
| WYETH AYERST | 100MG | A083239 | 001 |

INJECTABLE; INJECTION

PENTOBARBITAL SODIUM

| | | | |
|----------------------|---------|---------|-----|
| ELKINS SINN | 50MG/ML | A083270 | 001 |
| SODIUM PENTOBARBITAL | | | |
| WYETH AYERST | 50MG/ML | A083261 | 001 |

SUPPOSITORY; RECTAL

NEMBUTAL

| | | | | |
|------------|-------|---------|-----|--------------|
| OAK PHARMS | 30MG | A083247 | 001 | Jan 25, 1982 |
| | 60MG | A083247 | 002 | Jan 25, 1982 |
| | 120MG | A083247 | 003 | Jan 25, 1982 |
| | 200MG | A083247 | 004 | Jan 25, 1982 |

TABLET; ORAL

PENTOBARBITAL SODIUM

| | | | |
|----------------------|-------|---------|-----|
| VITARINE | 100MG | A083285 | 001 |
| SODIUM PENTOBARBITAL | | | |
| NEXGEN PHARMA INC | 100MG | A084238 | 001 |

PENTOLINIUM TARTRATE

INJECTABLE; INJECTION

ANSOLYSEN

| | | | |
|--------------|---------|---------|-----|
| WYETH AYERST | 10MG/ML | N009372 | 001 |
|--------------|---------|---------|-----|

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL

PENTOXIFYLLINE

| | | | | |
|---------------------|-------|---------|-----|--------------|
| ACTAVIS ELIZABETH | 400MG | A074878 | 001 | Jul 09, 1997 |
| HERITAGE PHARMS INC | 400MG | A074877 | 001 | Jul 08, 1997 |
| IMPAX LABS | 400MG | A075093 | 001 | Aug 10, 1999 |
| PLIVA | 400MG | A074874 | 001 | May 25, 1999 |
| TEVA | 400MG | A075199 | 001 | Sep 03, 1999 |
| WATSON LABS | 400MG | A075107 | 001 | Sep 04, 1998 |

TRENTAL

| | | | | |
|---------------------|----------|---------|-----|--------------|
| + US PHARM HOLDINGS | 400MG ** | N018631 | 001 | Aug 30, 1984 |
|---------------------|----------|---------|-----|--------------|

PERFLUBRON

LIQUID; ORAL

IMAGENT

| | | | | |
|----------------|------|---------|-----|--------------|
| ALLIANCE PHARM | 100% | N020091 | 001 | Aug 13, 1993 |
|----------------|------|---------|-----|--------------|

PERFLUOROPOLYMETHYLISOPROPYL ETHER; POLYTETRAFLUOROETHYLENE

PASTE; TOPICAL

SKIN EXPOSURE REDUCTION PASTE AGAINST CHEMICAL WARFARE AGENTS

| | | | | |
|---------|---------|---------|-----|--------------|
| US ARMY | 50%;50% | N021084 | 001 | Feb 17, 2000 |
|---------|---------|---------|-----|--------------|

PERGOLIDE MESYLATE

TABLET; ORAL

PERGOLIDE MESYLATE

| | | | | |
|----------------------|----------------|---------|-----|--------------|
| IVAX SUB TEVA PHARMS | EQ 0.05MG BASE | A076094 | 001 | Sep 04, 2003 |
| | EQ 0.25MG BASE | A076094 | 002 | Sep 04, 2003 |
| | EQ 1MG BASE | A076094 | 003 | Sep 04, 2003 |
| PAR PHARM | EQ 0.05MG BASE | A076061 | 001 | Nov 27, 2002 |
| | EQ 0.25MG BASE | A076061 | 002 | Nov 27, 2002 |
| | EQ 1MG BASE | A076061 | 003 | Nov 27, 2002 |

PERMAX

| | | | | |
|--------------------|----------------|---------|-----|--------------|
| VALEANT PHARM INTL | EQ 0.05MG BASE | N019385 | 001 | Dec 30, 1988 |
| | EQ 0.25MG BASE | N019385 | 002 | Dec 30, 1988 |
| | EQ 1MG BASE | N019385 | 003 | Dec 30, 1988 |

PERINDOPRIL ERBUMINE

TABLET; ORAL

ACEON

| | | | | |
|-----------------------|-----|---------|-----|--------------|
| + SYMPLMED PHARMS LLC | 2MG | N020184 | 001 | Dec 30, 1993 |
| + | 4MG | N020184 | 002 | Dec 30, 1993 |
| + | 8MG | N020184 | 003 | Dec 30, 1993 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PERINDOPRIL ERBUMINE

TABLET; ORAL

PERINDOPRIL ERBUMINE

| | | | |
|-----------|-----|-------------|--------------|
| APOTEX | 2MG | A090463 001 | Aug 30, 2010 |
| | 4MG | A090463 002 | Aug 30, 2010 |
| | 8MG | A090463 003 | Aug 30, 2010 |
| LUPIN LTD | 2MG | A078263 001 | Jan 27, 2010 |
| | 4MG | A078263 002 | Jan 27, 2010 |
| | 8MG | A078263 003 | Jan 27, 2010 |

PERMETHRIN

LOTION; TOPICAL

NIX

| | | | |
|-----------------|----|-------------|--------------|
| GLAXOSMITHKLINE | 1% | N019435 001 | Mar 31, 1986 |
|-----------------|----|-------------|--------------|

PERPHENAZINE

CONCENTRATE; ORAL

PERPHENAZINE

| | | | |
|-------------|----------|-------------|--------------|
| PHARM ASSOC | 16MG/5ML | A040360 001 | May 25, 2001 |
|-------------|----------|-------------|--------------|

TRILAFON

| | | | |
|----------|----------|-------------|--|
| SCHERING | 16MG/5ML | N011557 001 | |
|----------|----------|-------------|--|

INJECTABLE; INJECTION

TRILAFON

| | | | |
|----------|--------|-------------|--|
| SCHERING | 5MG/ML | N011213 002 | |
|----------|--------|-------------|--|

SYRUP; ORAL

TRILAFON

| | | | |
|----------|---------|-------------|--|
| SCHERING | 2MG/5ML | N011294 002 | |
|----------|---------|-------------|--|

TABLET; ORAL

PERPHENAZINE

| | | | |
|----------------|-----|-------------|--------------|
| ANI PHARMS INC | 2MG | A089707 001 | Sep 10, 1987 |
|----------------|-----|-------------|--------------|

| | | | |
|--|-----|-------------|--------------|
| | 4MG | A089708 001 | Sep 10, 1987 |
|--|-----|-------------|--------------|

| | | | |
|--|-----|-------------|--------------|
| | 8MG | A089456 001 | Sep 10, 1987 |
|--|-----|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 16MG | A089457 001 | Sep 10, 1987 |
|--|------|-------------|--------------|

TRILAFON

| | | | |
|------------|--------|-------------|--|
| + SCHERING | 2MG ** | N010775 001 | |
|------------|--------|-------------|--|

| | | | |
|---|--------|-------------|--|
| + | 4MG ** | N010775 002 | |
|---|--------|-------------|--|

| | | | |
|---|--------|-------------|--|
| + | 8MG ** | N010775 003 | |
|---|--------|-------------|--|

| | | | |
|---|---------|-------------|--|
| + | 16MG ** | N010775 004 | |
|---|---------|-------------|--|

TABLET, EXTENDED RELEASE; ORAL

TRILAFON

| | | | |
|----------|-----|-------------|--|
| SCHERING | 8MG | N011361 002 | |
|----------|-----|-------------|--|

PHENACEMIDE

TABLET; ORAL

PHENURONE

| | | | |
|----------|----------|-------------|--|
| + ABBVIE | 500MG ** | N007707 001 | |
|----------|----------|-------------|--|

PHENAZOPYRIDINE HYDROCHLORIDE; SULFAMETHOXAZOLE

TABLET; ORAL

AZO GANTANOL

| | | | |
|---------|-----------------|-------------|--------------|
| + ROCHE | 100MG; 500MG ** | N013294 001 | Sep 10, 1987 |
|---------|-----------------|-------------|--------------|

PHENAZOPYRIDINE HYDROCHLORIDE; SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM AND PHENAZOPYRIDINE HYDROCHLORIDE

| | | | |
|------|------------------------------------|-------------|--------------|
| ABLE | 200MG, N/A, N/A; N/A, 800MG, 160MG | N021105 001 | Jun 26, 2001 |
|------|------------------------------------|-------------|--------------|

PHENAZOPYRIDINE HYDROCHLORIDE; SULFISOXAZOLE

TABLET; ORAL

AZO GANTRISIN

| | | | |
|---------|----------------|-------------|--------------|
| + ROCHE | 50MG; 500MG ** | N019358 001 | Aug 31, 1990 |
|---------|----------------|-------------|--------------|

PHENDIMETRAZINE TARTRATE

CAPSULE; ORAL

PHENAZINE

| | | | |
|---------|------|-------------|--|
| MAST MM | 35MG | A086523 001 | |
|---------|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 35MG | A086524 001 | |
|--|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 35MG | A086525 001 | |
|--|------|-------------|--|

PHENDIMETRAZINE TARTRATE

| | | | |
|--------|------|-------------|--|
| SANDOZ | 35MG | A085633 001 | |
|--------|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 35MG | A085694 001 | |
|--|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 35MG | A085702 001 | |
|--|------|-------------|--|

| | | | |
|---------------|------|-------------|--|
| VIRTUS PHARMS | 35MG | A085695 001 | |
|---------------|------|-------------|--|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENDIMETRAZINE TARTRATE

CAPSULE; ORAL

PHENDIMETRAZINE TARTRATE

VITARINE

35MG

A085634 001

35MG

A085645 001

35MG

A085670 001

35MG

A086403 001

35MG

A086408 001

35MG

A086410 001

35MG

A087424 001

SPRX-3

SOLVAY

35MG

A085897 001

STATOBEX

TEVA

35MG

A085507 001

X-TROZINE

SHIRE RICHWOOD

35MG

A087394 001 Sep 22, 1982

CAPSULE, EXTENDED RELEASE; ORAL

BONTRIL

VALEANT

105MG

A088021 001 Sep 21, 1982

MELFIAT-105

NUMARK

105MG

A087487 001 Oct 13, 1982

PHENDIMETRAZINE TARTRATE

GRAHAM DM

105MG

A087214 001 May 26, 1982

105MG

A088020 001 Aug 16, 1982

105MG

A088028 001 Aug 16, 1982

105MG

A088062 001 Sep 13, 1982

105MG

A088063 001 Sep 10, 1982

105MG

A088111 001 Oct 18, 1982

VIRTUS PHARMS

105MG

A087378 001

SPRX-105

NUMARK

105MG

A088024 001 Dec 22, 1982

X-TROZINE L.A.

SHIRE RICHWOOD

105MG

A087371 001 Aug 24, 1982

TABLET; ORAL

ADPHEN

FERNDAL LABS

35MG

A083655 001

ALPHAZINE

SANDOZ

35MG

A085034 001

CAM-METRAZINE

ABC HOLDING

35MG

A085511 001

CAMALL

35MG

A085756 001

CHARTWELL RX

35MG

A083922 001

35MG

A085318 001

35MG

A085320 001

35MG

A085321 001

DI-METREX

PVT FORM

35MG

A085698 001

MELFIAT

NUMARK

35MG

A083790 002

METRA

FOREST PHARMS

35MG

A083754 001

PHENAZINE

MAST MM

35MG

A087305 001

PHENAZINE-35

ABC HOLDING

35MG

A085512 001

PHENDIMETRAZINE TARTRATE

BARR

35MG

A083644 001

35MG

A083684 001

35MG

A083686 001

35MG

A083687 001

35MG

A084831 001

35MG

A084834 001

35MG

A084835 001

CHARTWELL RX

35MG

A085761 001

35MG

A085941 001 Jun 27, 1983

FERNDAL LABS

35MG

A086834 001 Sep 15, 1983

INWOOD LABS

35MG

A084740 001

35MG

A084741 001

35MG

A084742 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENDIMETRAZINE TARTRATE

TABLET;ORAL

PHENDIMETRAZINE TARTRATE

| | | |
|----------------------|---------|-------------|
| | 35MG | A084743 001 |
| IVAX PHARMS | 35MG | A085611 001 |
| | 35MG | A085612 001 |
| IVAX SUB TEVA PHARMS | 35MG | A083682 001 |
| KV PHARM | 35MG | A084138 001 |
| | 35MG | A084141 001 |
| | 35MG | A085525 001 |
| MFG CHEMISTS | 35MG | A085914 001 |
| NEXGEN PHARMA INC | 35MG | A086020 001 |
| NUMARK | 35MG | A083790 001 |
| PVT FORM | 35MG | A085199 001 |
| | 35MG | A085697 001 |
| SANDOZ | 35MG | A085402 001 |
| | 35MG | A085830 001 |
| | 35MG | A086370 001 |
| SOLVAY | 35MG | A083993 001 |
| USL PHARMA | 35MG | A083805 001 |
| | 35MG | A084398 001 |
| | 35MG | A084399 001 |
| VIRTUS PHARMS | 35MG | A085497 001 |
| | 35MG | A086365 001 |
| VITARINE | 35MG | A085519 001 |
| | 35MG | A086005 001 |
| | 35MG | A086106 001 |
| WATSON LABS | 35MG | A085767 001 |
| | 35MG | A085768 001 |
| | 35MG | A085770 001 |
| | 35MG | A085773 001 |
| PLEGINE | | |
| WYETH AYERST | 35MG ** | N012248 001 |
| STATOBEX | | |
| TEVA | 35MG | A086013 001 |
| STATOBEX-G | | |
| TEVA | 35MG | A085095 001 |
| X-TROZINE | | |
| SHIRE RICHWOOD | 35MG | A086550 001 |
| | 35MG | A086551 001 |
| | 35MG | A086552 001 |
| | 35MG | A086553 001 |
| | 35MG | A086554 001 |

PHENINDIONE

TABLET;ORAL

HEDULIN

| | | |
|-------------------|------|-------------|
| SANOFI AVENTIS US | 50MG | N008767 002 |
|-------------------|------|-------------|

PHENMETRAZINE HYDROCHLORIDE

TABLET;ORAL

PRELUDIN

| | | |
|----------------------|------|-------------|
| BOEHRINGER INGELHEIM | 25MG | N010460 005 |
|----------------------|------|-------------|

TABLET, EXTENDED RELEASE;ORAL

PRELUDIN

| | | |
|----------------------|------|-------------|
| BOEHRINGER INGELHEIM | 50MG | N011752 004 |
| | 75MG | N011752 003 |

PHENPROCOUMON

TABLET;ORAL

LIQUAMAR

| | | |
|-----------------|-----|-------------|
| ORGANON USA INC | 3MG | N011228 001 |
|-----------------|-----|-------------|

PHENSUXIMIDE

CAPSULE;ORAL

MILONTIN

| | | |
|-------------|-------|-------------|
| PARKE DAVIS | 500MG | N008855 004 |
|-------------|-------|-------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

FASTIN

| | | | | |
|-----------------|---------|---------|-----|--------------|
| GLAXOSMITHKLINE | 30MG ** | N017352 | 001 | |
| OBESTIN-30 | | | | |
| FERNDALE LABS | 30MG | A087144 | 001 | |
| OBY-TRIM | | | | |
| SHIRE RICHWOOD | 30MG | A087764 | 001 | Mar 18, 1982 |
| ONA-MAST | | | | |
| MAST MM | 30MG | A086511 | 001 | |
| | 30MG | A086516 | 001 | |

PHENTERMINE HYDROCHLORIDE

| | | | | |
|----------------------|---------|---------|-----|--------------|
| ABC HOLDING | 30MG | A085411 | 001 | |
| ABLE | 15MG | A040497 | 001 | Mar 13, 2003 |
| | 30MG | A040403 | 001 | Aug 30, 2001 |
| | 30MG | A040427 | 001 | Aug 30, 2001 |
| CAMALL | 15MG | A086735 | 001 | |
| | 30MG | A087226 | 001 | |
| CHARTWELL RX | 18.75MG | A088576 | 001 | May 23, 1984 |
| | 30MG | A085417 | 001 | |
| | 30MG | A086732 | 002 | |
| | 30MG | A087215 | 001 | |
| | 37.5MG | A087915 | 001 | Dec 22, 1983 |
| | 37.5MG | A087918 | 001 | Dec 22, 1983 |
| | 37.5MG | A087930 | 001 | Oct 14, 1983 |
| | 37.5MG | A088610 | 001 | Jun 04, 1984 |
| | 37.5MG | A088611 | 001 | Jun 04, 1984 |
| | 37.5MG | A088625 | 001 | Aug 23, 1984 |
| DURAMED PHARMS BARR | 30MG | A088948 | 001 | Apr 25, 1986 |
| ELITE LABS INC | 15MG | A040460 | 001 | Jan 14, 2003 |
| | 30MG | A040227 | 001 | Jun 18, 1997 |
| | 30MG | A040448 | 001 | Jan 22, 2003 |
| IVAX PHARMS | 30MG | A086329 | 001 | |
| SANDOZ | 30MG | A087208 | 001 | |
| | 30MG | A087223 | 001 | |
| | 37.5MG | A088414 | 001 | Oct 19, 1983 |
| SUN PHARM INDUSTRIES | 37.5MG | A040527 | 001 | Oct 23, 2003 |
| TEVA | 30MG | A086911 | 001 | |
| | 30MG | A087126 | 001 | |
| | 30MG | A087777 | 001 | Nov 01, 1985 |
| | 30MG | A088612 | 001 | Apr 04, 1984 |
| | 30MG | A088613 | 001 | Apr 09, 1984 |
| | 30MG | A088614 | 001 | Apr 09, 1984 |
| TG UNITED INC | 30MG | A040083 | 001 | Mar 07, 1997 |
| UPSHER SMITH LABS | 30MG | A084487 | 001 | Apr 09, 1982 |
| | 30MG | A088430 | 001 | Mar 27, 1984 |
| USL PHARMA | 30MG | A088797 | 001 | Dec 10, 1984 |
| VITARINE | 30MG | A087202 | 001 | |
| | 30MG | A087235 | 001 | |
| WATSON LABS | 30MG | A086740 | 001 | Mar 21, 1985 |

TABLET; ORAL

ONA-MAST

| | | | | |
|---------|-----|---------|-----|--|
| MAST MM | 8MG | A086260 | 001 | |
|---------|-----|---------|-----|--|

PHENTERMINE HYDROCHLORIDE

| | | | | |
|-------------------|--------|---------|-----|--------------|
| ABLE | 37.5MG | A040402 | 001 | Aug 30, 2001 |
| ACTAVIS ELIZABETH | 37.5MG | A040276 | 001 | Nov 25, 1998 |
| CHARTWELL RX | 8MG | A083923 | 001 | |
| | 8MG | A085319 | 001 | |
| | 37.5MG | A087805 | 001 | Dec 06, 1982 |
| | 37.5MG | A088596 | 001 | Apr 04, 1984 |
| IVAX PHARMS | 8MG | A085553 | 001 | |
| SANDOZ | 8MG | A085671 | 001 | |
| | 8MG | A085689 | 001 | |
| SANDOZ INC | 30MG | A088605 | 001 | Sep 28, 1987 |
| USL PHARMA | 8MG | A083804 | 001 | |
| | 37.5MG | A088910 | 001 | Jul 17, 1985 |
| | 37.5MG | A088917 | 001 | Jul 17, 1985 |
| VITARINE | 8MG | A086453 | 001 | |
| | 8MG | A086456 | 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENTERMINE HYDROCHLORIDE

TABLET;ORAL

| | | | | |
|------------------------------------|-----------|--|---------|------------------|
| PHENTERMINE HYDROCHLORIDE | | | | |
| WATSON LABS | 8MG | | A085739 | 001 |
| TORA | | | | |
| SOLVAY | 8MG | | A084035 | 001 |
| WILPO | | | | |
| + SANDOZ | 8MG ** | | N012737 | 001 |
| TABLET, ORALLY DISINTEGRATING;ORAL | | | | |
| SUPRENZA | | | | |
| CITIUS PHARMS | 15MG ** | | N202088 | 001 Jun 13, 2011 |
| | 30MG ** | | N202088 | 002 Jun 13, 2011 |
| | 37.5MG ** | | N202088 | 003 Mar 27, 2012 |

PHENTERMINE RESIN COMPLEX

CAPSULE, EXTENDED RELEASE;ORAL

| | | | | |
|---------------------------|-----------------|--|---------|------------------|
| IONAMIN | | | | |
| UCB INC | EQ 15MG BASE ** | | N011613 | 004 |
| | EQ 30MG BASE ** | | N011613 | 002 |
| PHENTERMINE RESIN 30 | | | | |
| QUANTUM PHARMICS | EQ 30MG BASE | | A089120 | 001 Feb 04, 1988 |
| PHENTERMINE RESIN COMPLEX | | | | |
| LANNETT CO INC | EQ 15MG BASE | | A040872 | 001 Jul 28, 2011 |
| | EQ 30MG BASE | | A040872 | 002 Jul 28, 2011 |

PHENTOLAMINE MESYLATE

INJECTABLE;INJECTION

| | | | | |
|------------|-------------|--|---------|-----|
| REGITINE | | | | |
| + NOVARTIS | 5MG/VIAL ** | | N008278 | 003 |

PHENYL AMINOSALICYLATE

POWDER;ORAL

| | | | | |
|-------------------|-------|--|---------|-----|
| PHENY-PAS-TEBAMIN | | | | |
| PHARM RES ASSOC | 50% | | N011695 | 002 |
| TABLET;ORAL | | | | |
| PHENY-PAS-TEBAMIN | | | | |
| PHARM RES ASSOC | 500MG | | N011695 | 003 |

PHENYLBUTAZONE

CAPSULE;ORAL

| | | | | |
|----------------------|-------|--|---------|------------------|
| AZOLID | | | | |
| SANOFI AVENTIS US | 100MG | | A087260 | 001 |
| BUTAZOLIDIN | | | | |
| NOVARTIS | 100MG | | N008319 | 009 |
| PHENYLBUTAZONE | | | | |
| FOSUN PHARMA | 100MG | | A087774 | 001 Jun 16, 1982 |
| IVAX PHARMS | 100MG | | A088218 | 001 Jun 24, 1983 |
| SUN PHARM INDUSTRIES | 100MG | | A088994 | 001 Dec 04, 1985 |
| WATSON LABS | 100MG | | A087756 | 001 Dec 17, 1982 |
| TABLET;ORAL | | | | |
| AZOLID | | | | |
| SANOFI AVENTIS US | 100MG | | A087091 | 001 |
| BUTAZOLIDIN | | | | |
| NOVARTIS | 100MG | | N008319 | 008 |
| PHENYLBUTAZONE | | | | |
| FOSUN PHARMA | 100MG | | A084339 | 001 |
| SUN PHARM INDUSTRIES | 100MG | | A088863 | 001 Dec 04, 1985 |
| WATSON LABS | 100MG | | A086151 | 001 |
| | 100MG | | A087674 | 001 Apr 21, 1982 |

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

| | | | | |
|-----------------------|-----------------------|--|---------|------------------|
| PHENERGAN VC | | | | |
| + ANI PHARMS | 5MG/5ML;6.25MG/5ML ** | | N008604 | 003 Apr 02, 1984 |
| PHERAZINE VC | | | | |
| HALSEY | 5MG/5ML;6.25MG/5ML | | A088868 | 001 Mar 02, 1987 |
| PROMETH VC PLAIN | | | | |
| G AND W LABS INC | 5MG/5ML;6.25MG/5ML | | A088761 | 001 Nov 08, 1984 |
| PROMETHAZINE VC PLAIN | | | | |
| CENCI | 5MG/5ML;6.25MG/5ML | | A088815 | 001 Nov 22, 1985 |
| WOCKHARDT | 5MG/5ML;6.25MG/5ML | | A088897 | 001 Jan 04, 1985 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PHENYLEPHRINE HYDROCHLORIDE; PYRILAMINE MALEATE

SOLUTION/DROPS;OPHTHALMIC

PREFRIN-A

ALLERGAN 0.12%;0.1% N007953 001

PHENYTOIN

SUSPENSION;ORAL

DILANTIN-30

PARKE DAVIS 30MG/5ML N008762 002

PHENYTOIN

ACTAVIS MID ATLANTIC 125MG/5ML A089892 001 Sep 25, 1992

PHENYTOIN SODIUM

CAPSULE;ORAL

DIPHENYLAN SODIUM

LANNETT 30MG PROMPT A080857 001

100MG PROMPT A080857 002

EXTENDED PHENYTOIN SODIUM

ANI PHARMS INC 100MG EXTENDED A040435 001 Jun 20, 2003

100MG EXTENDED A089441 001 Dec 18, 1986

WOCKHARDT 30MG EXTENDED A040759 001 Dec 18, 2007

WOCKHARDT USA 100MG EXTENDED A040732 001 Jan 30, 2008

PHENYTEX

WATSON LABS 100MG EXTENDED A088711 001 Dec 21, 1984

PHENYTOIN SODIUM

PHARMERAL 100MG PROMPT A085435 001

WATSON LABS 100MG PROMPT A085894 001

PROMPT PHENYTOIN SODIUM

ANI PHARMS INC 100MG PROMPT A080259 001

WATSON LABS 100MG PROMPT A080905 001

INJECTABLE;INJECTION

DILANTIN

PARKE DAVIS 50MG/ML N010151 001

PHENYTOIN SODIUM

FRESENIUS KABI USA 50MG/ML A089003 001 May 31, 1985

HOSPIRA 50MG/ML A089521 001 Mar 17, 1987

50MG/ML A089744 001 Dec 18, 1987

MARSAM PHARMS LLC 50MG/ML A089501 001 Oct 13, 1987

50MG/ML A089779 001 Nov 27, 1992

SMITH AND NEPHEW 50MG/ML A088519 001 Dec 19, 1984

50MG/ML A088521 001 Dec 18, 1984

SOLOPAK 50MG/ML A088520 001 Dec 17, 1984

WARNER CHILCOTT 50MG/ML A089900 001 Mar 30, 1990

WATSON LABS 50MG/ML A085434 001

PHYTONADIONE

INJECTABLE;INJECTION

AQUAMEPHYTON

+ TELIGENT 1MG/0.5ML ** N012223 002

+ 10MG/ML ** N012223 001

KONAKION

ROCHE 1MG/0.5ML N011745 001

10MG/ML N011745 003

PHYTONADIONE

GLAXOSMITHKLINE 1MG/0.5ML A084060 001

10MG/ML A084060 002

VITAMIN K1

HOSPIRA 10MG/ML A087956 001 Jul 25, 1983

PILOCARPINE

INSERT, EXTENDED RELEASE;OPHTHALMIC

OCUSERT PILO-20

AKORN 5MG N017431 001

OCUSERT PILO-40

AKORN 11MG N017548 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PILOCARPINE HYDROCHLORIDE

GEL;OPHTHALMIC

PILOPINE HS

ALCON

4%

N018796 001 Oct 01, 1984

PINACIDIL

CAPSULE, EXTENDED RELEASE;ORAL

PINDAC

LEO PHARM

12.5MG

N019456 001 Dec 28, 1989

25MG

N019456 002 Dec 28, 1989

PINDOLOL

TABLET;ORAL

PINDOLOL

G AND W LABS INC

5MG

A073661 001 Oct 31, 1993

5MG

A073687 001 Feb 26, 1993

5MG

A074123 001 Apr 17, 1997

10MG

A073661 002 Oct 31, 1993

10MG

A073687 002 Feb 26, 1993

10MG

A074123 002 Apr 17, 1997

MYLAN PHARMS INC

5MG

A074013 001 Sep 24, 1992

10MG

A074018 001 Sep 24, 1992

NOSTRUM LABS

5MG

A074474 001 Oct 28, 1996

10MG

A074474 002 Oct 28, 1996

PUREPAC PHARM

5MG

A074125 001 Apr 28, 1993

10MG

A074125 002 Apr 28, 1993

WATSON LABS

5MG

A074437 001 Feb 27, 1995

10MG

A074437 002 Feb 27, 1995

VISKEN

+ NOVARTIS

5MG **

N018285 001 Sep 03, 1982

+

10MG **

N018285 002 Sep 03, 1982

PIPECURONIUM BROMIDE

INJECTABLE;INJECTION

ARDUAN

ORGANON USA INC

10MG/VIAL

N019638 001 Jun 26, 1990

PIPERACETAZINE

TABLET;ORAL

QUIDE

DOW PHARM

10MG

N013615 001

25MG

N013615 002

PIPERACILLIN SODIUM

INJECTABLE;INJECTION

PIPRACIL

WYETH PHARMS INC

EQ 2GM BASE/VIAL

A062750 001 Oct 13, 1987

+

EQ 2GM BASE/VIAL **

N050545 002

EQ 3GM BASE/VIAL

A062750 002 Oct 13, 1987

+

EQ 3GM BASE/VIAL **

N050545 003

EQ 4GM BASE/VIAL

A062750 003 Oct 13, 1987

+

EQ 4GM BASE/VIAL **

N050545 004

+

EQ 40GM BASE/VIAL **

N050545 006 Sep 30, 1985

PIPERAZINE CITRATE

SYRUP;ORAL

ANTEPAR

GLAXOSMITHKLINE

EQ 500MG BASE/5ML

N009102 001

BRYREL

SANOFI AVENTIS US

EQ 500MG BASE/5ML

N017796 001

MULTIFUGE

BLULINE

EQ 500MG BASE/5ML

N009452 001

PIPERAZINE CITRATE

ALPHARMA US PHARMS

EQ 500MG BASE/5ML

A080774 001

LANNETT

EQ 500MG BASE/5ML

A080963 001

LUITPOLD

EQ 500MG BASE/5ML

A080671 001

VERMIDOL

SOLVAY

EQ 500MG BASE/5ML

A080992 001

TABLET;ORAL

ANTEPAR

GLAXOSMITHKLINE

EQ 500MG BASE

N009102 003

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PIPERAZINE CITRATE

TABLET; ORAL

PIPERAZINE CITRATE

IMPAX LABS

EQ 250MG BASE

A080874 001

PIPERONYL BUTOXIDE; PYRETHRINS

AEROSOL; TOPICAL

RID MOUSSE

BAYER HEALTHCARE LLC 4%;EQ 0.33% BASE

N021043 001 Mar 07, 2000

PIPOBROMAN

TABLET; ORAL

VERCYTE

ABBOTT

10MG

N016245 001

25MG

N016245 002

PIRBUTEROL ACETATE

AEROSOL, METERED; INHALATION

MAXAIR

MEDICIS

EQ 0.2MG BASE/INH

N020014 001 Nov 30, 1992

VALEANT PHARMS

EQ 0.2MG BASE/INH

N019009 001 Dec 30, 1986

PIRENEIDONE

TABLET; ORAL

ESBRIET

+ GENENTECH INC

534MG

N208780 002 Jan 11, 2017

PIROXICAM

CAPSULE; ORAL

PIROXICAM

CYCLE PHARMS LTD

10MG

A073651 001 Feb 26, 1993

20MG

A073651 002 Feb 26, 1993

EGIS

10MG

A074808 001 Jul 08, 1997

20MG

A074808 002 Jul 08, 1997

IVAX SUB TEVA PHARMS

10MG

A074148 001 Jun 03, 1996

20MG

A074148 002 Jun 03, 1996

MYLAN

10MG

A074043 001 Sep 22, 1992

10MG

A074102 001 Jul 31, 1992

20MG

A074043 002 Sep 22, 1992

20MG

A074102 002 Jul 31, 1992

SCS

10MG

A074036 001 May 29, 1992

20MG

A074036 002 May 29, 1992

TEVA

10MG

A073637 001 Jan 28, 1994

20MG

A073638 001 Jan 28, 1994

TEVA PHARMS

10MG

A074103 001 Aug 28, 1992

20MG

A074103 002 Aug 28, 1992

WATSON LABS

10MG

A074287 001 May 16, 1996

10MG

A074460 001 Sep 29, 1995

20MG

A074287 002 May 16, 1996

20MG

A074460 002 Sep 29, 1995

PITAVASTATIN SODIUM

TABLET; ORAL

NIKITA

+ LUPIN LTD

EQ 1MG BASE

N209875 001 Aug 04, 2017

+

EQ 2MG BASE

N209875 002 Aug 04, 2017

+

EQ 4MG BASE

N209875 003 Aug 04, 2017

PLICAMYCIN

INJECTABLE; INJECTION

MITHRACIN

PFIZER

2.5MG/VIAL

N050109 001

PODOFILOX

SOLUTION; TOPICAL

PODOFILOX

BAUSCH AND LOMB INC

0.5%

A090184 001 Jul 21, 2010

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POLYESTRADIOL PHOSPHATEINJECTABLE; INJECTION
ESTRADURIN

WYETH AYERST 40MG/AMP N010753 001

POLYETHYLENE GLYCOL 3350

FOR SOLUTION; ORAL

POLYETHYLENE GLYCOL 3350

BRECKENRIDGE PHARM 17GM/SCOOPFUL A077736 001 May 26, 2006
PADDOCK LLC 17GM/SCOOPFUL A090567 001 Oct 15, 2009
TEVA PHARMS 17GM/SCOOPFUL A077445 001 May 04, 2006POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION; ORAL

PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE

MYLAN 420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/BOT A090409 001 Apr 02, 2010

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE

FOR SOLUTION; ORAL

CLENZ-LYTE

PADDOCK LLC 236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT A090769 001 Jun 07, 2010

SOLUTION; ORAL

OCL

HOSPIRA 6GM/100ML; 75MG/100ML; 168MG/100ML; 146MG/100ML; 1.29GM/100ML N019284 001 Apr 30, 1986

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE ANHYDROUS

FOR SOLUTION; ORAL

COLYTE

MYLAN SPECIALITY LP 120GM/PACKET; 1.49GM/PACKET; 3.36GM/PACKET; 2.92GM/PACKET; 11.36GM/PACKET N018983 005 Oct 26, 1984
227.1GM/PACKET; 2.82GM/PACKET; 6.36GM/PACKET; 5.53GM/PACKET; 21.5GM/PACKET N018983 004 Oct 26, 1984
227.1GM/BOT; 2.82GM/BOT; 6.36GM/BOT; 5.53GM/BOT; 21.5GM/BOT N018983 010 Jan 31, 1989
240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT N018983 007 Jun 12, 1987
360GM/PACKET; 4.47GM/PACKET; 10.08GM/PACKET; 8.76GM/PACKET; 34.08GM/PACKET N018983 006 Oct 26, 1984

COLYTE-FLAVORED

MYLAN SPECIALITY LP 227.1GM/BOT; 2.82GM/BOT; 6.36GM/BOT; 5.53GM/BOT; 21.5GM/BOT N018983 008 Nov 14, 1991
240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT N018983 009 Nov 14, 1991

PEG 3350 AND ELECTROLYTES

MYLAN 236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT A090928 001 Jan 28, 2010

POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES

PADDOCK LLC 240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT A090712 001 Feb 25, 2010

FOR SUSPENSION; ORAL

CO-LAV

VINTAGE PHARMS 240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT A073428 001 Jan 28, 1992

COLOVAGE

DYNAPHARM 227.1GM/PACKET; 2.82GM/PACKET; 6.36GM/PACKET; 5.53GM/PACKET; 21.5GM/PACKET A071320 001 Apr 20, 1988

E-Z-EM PREP LYTE

E Z EM 236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT A071278 001 Nov 21, 1988

GLYCOPREP

GOLDLINE 236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT A072319 001 Dec 23, 1988

GO-EVAC

VINTAGE PHARMS 236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT A073433 001 Apr 28, 1992

PEG-LYTE

SANDOZ 236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT A073098 001 Aug 31, 1993

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POLYMYXIN B SULFATE

INJECTABLE; INJECTION

AEROSPORIN

GLAXOSMITHKLINE EQ 500,000 U BASE/VIAL A062036 001

POWDER; FOR RX COMPOUNDING

POLY-RX

X GEN PHARMS 100,000,000 UNITS/BOT A061578 001

POLYMYXIN B SULFATE

PADDOCK LLC 100,000,000 UNITS/BOT A062455 001 Jul 27, 1983

POLYTHIAZIDE

TABLET; ORAL

RENESE

PFIZER 1MG N012845 001

2MG N012845 002

4MG N012845 003

POLYTHIAZIDE; PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

MINIZIDE

PFIZER 0.5MG;EQ 1MG BASE N017986 001

0.5MG;EQ 2MG BASE N017986 002

0.5MG;EQ 5MG BASE N017986 003

POLYTHIAZIDE; RESERPINE

TABLET; ORAL

RENESE-R

PFIZER 2MG;0.25MG N013636 001

POTASSIUM AMINOSALICYLATE

CAPSULE; ORAL

PASKALIUM

GLENWOOD 500MG N009395 004

POWDER; ORAL

POTASSIUM AMINOSALICYLATE

HEXCEL 100% A080098 001

TABLET; ORAL

PASKALIUM

GLENWOOD 1GM N009395 003

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

K-LEASE

SAVAGE LABS 8MEQ A073398 001 Jan 28, 1992

10MEQ A072427 001 Mar 28, 1990

POTASSIUM CHLORIDE

NESHER PHARMS 10MEQ A070980 001 Feb 17, 1987

TEVA 8MEQ A073531 001 Apr 26, 1996

10MEQ A073532 001 Apr 26, 1996

FOR SUSPENSION, EXTENDED RELEASE; ORAL

MICRO-K LS

KV PHARM 20MEQ/PACKET N019561 003 Aug 26, 1988

INJECTABLE; INJECTION

POTASSIUM CHLORIDE

ABRAXIS PHARM 2MEQ/ML A080204 001

2MEQ/ML A084290 001

2MEQ/ML A086713 001

2MEQ/ML A086714 001

2MEQ/ML A087787 001 Apr 20, 1982

2MEQ/ML A087885 001 Feb 03, 1983

AKORN 2MEQ/ML A088286 001 Sep 05, 1985

BAXTER HLTHCARE 2MEQ/ML A080203 001

2MEQ/ML A085499 001

FRESENIUS KABI USA 2MEQ/ML A087817 001 Oct 20, 1982

GD SEARLE LLC 1MEQ/ML A086219 001

2MEQ/ML A086219 002

2MEQ/ML A086220 002

3MEQ/ML A086219 003

3MEQ/ML A086220 001

4MEQ/ML A086219 004

HOSPIRA 1MEQ/ML A080205 003

1MEQ/ML A083345 003

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POTASSIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE

| | | | | |
|--------------------------------|-------------------|---------|-----|--------------|
| | 1.5MEQ/ML | A083345 | 001 | |
| | 2MEQ/ML | A083345 | 002 | |
| | 2.4MEQ/ML | A080205 | 004 | |
| | 3.2MEQ/ML | A080205 | 005 | |
| INTL MEDICATION | 2MEQ/ML | A083163 | 001 | |
| LILLY | 2MEQ/ML | N007865 | 002 | |
| LUITPOLD | 2MEQ/ML | A080221 | 001 | |
| | 2MEQ/ML | A080736 | 001 | |
| | 2MEQ/ML | A087584 | 001 | |
| | 2MEQ/ML | A087585 | 001 | |
| MILES | 1MEQ/ML | A080195 | 002 | |
| | 2MEQ/ML | A080195 | 001 | |
| | 3MEQ/ML | A080195 | 003 | |
| | 4MEQ/ML | A080195 | 004 | |
| PHARMA SERVE NY | 2MEQ/ML | A086297 | 001 | |
| | 2MEQ/ML | A087362 | 001 | Mar 08, 1983 |
| WATSON LABS | 2MEQ/ML | A086208 | 001 | |
| | 2MEQ/ML | A089163 | 001 | Mar 10, 1988 |
| | 2MEQ/ML | A089421 | 001 | Jan 02, 1987 |
| | 3MEQ/ML | A086210 | 001 | |
| POTASSIUM CHLORIDE 30MEQ IN | PLASTIC CONTAINER | | | |
| + ICU MEDICAL INC | 2.24GM/100ML | N020161 | 003 | Aug 11, 1998 |
| TABLET, EXTENDED RELEASE; ORAL | | | | |
| K+10 | | | | |
| FUTURE PAK | 10MEQ | A070999 | 001 | Oct 22, 1987 |
| K+8 | | | | |
| FUTURE PAK | 8MEQ | A070998 | 001 | Jan 25, 1993 |
| KAON CL | | | | |
| SAVAGE LABS | 6.7MEQ | N017046 | 001 | |
| KAON CL-10 | | | | |
| SAVAGE LABS | 10MEQ | N017046 | 002 | |
| KLOTRIX | | | | |
| APOTHECON | 10MEQ | N017850 | 001 | |
| POTASSIUM CHLORIDE | | | | |
| COPLEY PHARM | 8MEQ | A070618 | 001 | Sep 09, 1987 |
| NESHER PHARMS | 20MEQ | A076044 | 001 | Apr 05, 2002 |
| + SCHERING | 10MEQ ** | N019439 | 002 | Jun 13, 1986 |
| + SCHERING | 20MEQ ** | N019439 | 001 | Jun 13, 1986 |
| SLOW-K | | | | |
| NOVARTIS | 8MEQ | N017476 | 002 | |
| TEN-K | | | | |
| NOVARTIS | 10MEQ | N019381 | 001 | Apr 16, 1986 |

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

| | | | | |
|-------------------------------------------------------------------------|--------------------------|---------|-----|--------------|
| POTASSIUM CHLORIDE 0.037% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 37MG/100ML; 900MG/100ML | N019708 | 001 | Sep 29, 1989 |
| POTASSIUM CHLORIDE 0.075% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 75MG/100ML; 900MG/100ML | N019708 | 002 | Sep 29, 1989 |
| POTASSIUM CHLORIDE 0.11% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 110MG/100ML; 900MG/100ML | N019708 | 003 | Sep 29, 1989 |
| POTASSIUM CHLORIDE 0.22% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 220MG/100ML; 900MG/100ML | N019708 | 005 | Sep 29, 1989 |
| POTASSIUM CHLORIDE 0.224% IN SODIUM CHLORIDE 0.9% | | | | |
| + BAXTER HLTHCARE | 224MG/100ML; 900MG/100ML | N017648 | 003 | |
| POTASSIUM CHLORIDE 0.3% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 300MG/100ML; 900MG/100ML | N019708 | 006 | Sep 29, 1989 |
| SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 75MG/100ML; 900MG/100ML | N018722 | 001 | Nov 09, 1982 |
| BAXTER HLTHCARE | 75MG/100ML; 900MG/100ML | N017648 | 004 | |
| SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 150MG/100ML; 900MG/100ML | N018722 | 002 | Nov 09, 1982 |
| SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 220MG/100ML; 900MG/100ML | N018722 | 003 | Nov 09, 1982 |
| SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER | | | | |
| B BRAUN | 300MG/100ML; 900MG/100ML | N018722 | 004 | Nov 09, 1982 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

POTASSIUM CHLORIDE; SODIUM CHLORIDE; TROMETHAMINE

INJECTABLE; INJECTION

THAM-E

HOSPIRA 370MG/VIAL; 1.75GM/VIAL; 36GM/VIAL N013025 001

POTASSIUM CITRATE

FOR SOLUTION; ORAL

POTASSIUM CITRATE

+ UT SW MEDCTR 10MEQ/PACKET ** N019647 002 Oct 13, 1988

+ 20MEQ/PACKET ** N019647 001 Oct 13, 1988

POTASSIUM IODIDE

SOLUTION; ORAL

POTASSIUM IODIDE

ROXANE 1GM/ML ** N018551 001 Feb 19, 1982

TABLET; ORAL

THYRO-BLOCK

MEDA PHARMS 130MG N018307 001

POTASSIUM PERCHLORATE

CAPSULE; ORAL

PERCHLORACAP

MALLINCKRODT 200MG N017551 001

POVIDONE-IODINE

SOLUTION; TOPICAL

E-Z PREP

CLINIPAD 10% N019382 001 Jul 25, 1989

SPONGE; TOPICAL

E-Z PREP

CLINIPAD 5% N019382 002 Jul 25, 1989

E-Z PREP 220

CLINIPAD 5% N019382 003 Jul 25, 1989

PRALIDOXIME CHLORIDE

INJECTABLE; INJECTION

PRALIDOXIME CHLORIDE

BAXTER HLTHCARE CORP 300MG/ML N018799 001 Dec 13, 1982

TABLET; ORAL

PROTOPAM CHLORIDE

WYETH AYERST 500MG N014122 002

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

MIRAPEX

BOEHRINGER INGELHEIM 1.25MG N020667 004 Jul 01, 1997

PRAMIPEXOLE DIHYDROCHLORIDE

ACTAVIS GRP PTC 0.125MG A091254 001 Nov 30, 2010

0.25MG A091254 002 Nov 30, 2010

0.5MG A091254 003 Nov 30, 2010

0.75MG A091254 004 Nov 30, 2010

1MG A091254 005 Nov 30, 2010

1.5MG A091254 006 Nov 30, 2010

SANDOZ 0.125MG A090190 001 Jul 06, 2010

0.25MG A090190 002 Jul 06, 2010

0.5MG A090190 003 Jul 06, 2010

0.75MG A090190 006 Oct 08, 2010

1MG A090190 004 Jul 06, 2010

1.5MG A090190 005 Jul 06, 2010

WATSON LABS 0.125MG A078551 001 Oct 08, 2010

0.25MG A078551 002 Oct 08, 2010

0.5MG A078551 003 Oct 08, 2010

1MG A078551 004 Oct 08, 2010

1.5MG A078551 005 Oct 08, 2010

PRAMLINTIDE ACETATE

INJECTABLE; SUBCUTANEOUS

SYMLIN

ASTRAZENECA AB EQ 3MG BASE/5ML (EQ 600MCG BASE/ML) N021332 001 Mar 16, 2005

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PRAVASTATIN SODIUM

TABLET; ORAL

PRAVACHOL

+ BRISTOL MYERS SQUIBB 10MG **

N019898 002 Oct 31, 1991

PRAVASTATIN SODIUM

MYLAN

10MG

A077013 001 Oct 23, 2006

20MG

A077013 002 Oct 23, 2006

40MG

A077013 003 Oct 23, 2006

80MG

A077013 004 Dec 28, 2007

PLIVA HRVATSKA DOO

10MG

A077730 001 Nov 21, 2006

20MG

A077730 002 Nov 21, 2006

30MG

A077730 003 Nov 21, 2006

40MG

A077730 005 Nov 21, 2006

RANBAXY LABS LTD

10MG

A076445 001 Apr 23, 2007

20MG

A076445 002 Apr 23, 2007

40MG

A076445 003 Apr 23, 2007

80MG

A076445 004 Apr 23, 2007

PRAZEPAM

CAPSULE; ORAL

CENTRAX

PARKE DAVIS

5MG

N018144 001

10MG

N018144 002

20MG

N018144 003 May 10, 1982

PRAZEPAM

USL PHARMA

5MG

A070427 001 Nov 06, 1987

10MG

A070428 001 Nov 06, 1987

TABLET; ORAL

CENTRAX

PARKE DAVIS

10MG

N017415 001

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

PRAZOSIN HYDROCHLORIDE

AM THERAP

EQ 1MG BASE

A072782 001 May 16, 1989

EQ 2MG BASE

A072783 001 May 16, 1989

EQ 5MG BASE

A072784 001 May 16, 1989

ANI PHARMS INC

EQ 1MG BASE

A072577 002 May 16, 1989

EQ 2MG BASE

A072577 001 May 16, 1989

EQ 5MG BASE

A072577 003 May 16, 1989

DAVA PHARMS INC

EQ 1MG BASE

A072705 001 May 16, 1989

EQ 2MG BASE

A072706 001 May 16, 1989

EQ 5MG BASE

A072707 001 May 16, 1989

PUREPAC PHARM

EQ 1MG BASE

A072991 001 May 16, 1989

EQ 2MG BASE

A072921 001 May 16, 1989

EQ 5MG BASE

A072992 001 May 16, 1989

WATSON LABS

EQ 1MG BASE

A072352 001 May 16, 1989

EQ 2MG BASE

A072333 001 May 16, 1989

EQ 5MG BASE

A072609 001 May 16, 1989

TABLET, EXTENDED RELEASE; ORAL

MINIPRESS XL

PFIZER

2.5MG

N019775 001 Jan 29, 1992

5MG

N019775 002 Jan 29, 1992

PREDNISOLONE

CREAM; TOPICAL

METI-DERM

SCHERING

0.5%

N010209 002

SYRUP; ORAL

PREDNISOLONE

APOTEX INC

5MG/5ML

A040570 001 Aug 25, 2005

15MG/5ML

A040571 001 Aug 25, 2005

IVAX SUB TEVA PHARMS

15MG/5ML

A040287 001 May 28, 1999

NESHER PHARMS

5MG/5ML

A040423 001 Oct 22, 2001

15MG/5ML

A040364 001 Apr 10, 2002

TEVA PHARMS

15MG/5ML

A040322 001 Jan 19, 2000

WE PHARMS

15MG/5ML

A040192 001 May 28, 1998

PRELONE

MURO

5MG/5ML

A089654 001 Jan 17, 1989

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISOLONE

TABLET; ORAL

CORTALONE

| | | |
|--------|-------|-------------|
| HALSEY | 1MG | A080304 003 |
| | 2.5MG | A080304 002 |
| | 5MG | A080304 001 |

DELTA-CORTEF

| | | |
|----------------------|-----|-------------|
| PHARMACIA AND UPJOHN | 5MG | N009987 004 |
|----------------------|-----|-------------|

FERNISOLONE-P

| | | |
|--------------|-----|-------------|
| FERNDAL LABS | 5MG | A083941 001 |
|--------------|-----|-------------|

PREDNISOLONE

| | | |
|---------------------|-----|-------------|
| AUROLIFE PHARMA LLC | 5MG | A084773 001 |
|---------------------|-----|-------------|

| | | |
|------|-----|-------------|
| BARR | 5MG | A084426 002 |
|------|-----|-------------|

| | | |
|-------|-----|-------------|
| BUNDY | 5MG | A083675 001 |
|-------|-----|-------------|

| | | |
|--------------|-----|-------------|
| CHARTWELL RX | 5MG | A084542 001 |
|--------------|-----|-------------|

| | | |
|-------------|-----|-------------|
| ELKINS SINN | 5MG | A080625 001 |
|-------------|-----|-------------|

| | | |
|-----------|-----|-------------|
| EVERYLIFE | 1MG | A084439 001 |
|-----------|-----|-------------|

| | | |
|--|-------|-------------|
| | 2.5MG | A084439 002 |
|--|-------|-------------|

| | | |
|--|-----|-------------|
| | 5MG | A084439 003 |
|--|-----|-------------|

| | | |
|----------|-------|-------------|
| FERRANTE | 2.5MG | A080562 001 |
|----------|-------|-------------|

| | | |
|--|-----|-------------|
| | 5MG | A080562 002 |
|--|-----|-------------|

| | | |
|--------------|-----|-------------|
| FOSUN PHARMA | 5MG | A080339 001 |
|--------------|-----|-------------|

| | | |
|---------|-----|-------------|
| HEATHER | 5MG | A080326 001 |
|---------|-----|-------------|

| | | |
|------------|-----|-------------|
| IMPAX LABS | 5MG | A080780 001 |
|------------|-----|-------------|

| | | |
|-------------|-----|-------------|
| INWOOD LABS | 5MG | A080748 001 |
|-------------|-----|-------------|

| | | |
|----------------------|-----|-------------|
| IVAX SUB TEVA PHARMS | 5MG | A080378 001 |
|----------------------|-----|-------------|

| | | |
|---------|-----|-------------|
| LANNETT | 5MG | A080531 002 |
|---------|-----|-------------|

| | | |
|-----------------|-----|-------------|
| MARSHALL PHARMA | 5MG | A080307 001 |
|-----------------|-----|-------------|

| | | |
|--------|-----|-------------|
| PANRAY | 1MG | A080351 001 |
|--------|-----|-------------|

| | | |
|--|-----|-------------|
| | 5MG | A080351 002 |
|--|-----|-------------|

| | | |
|-----------------|-----|-------------|
| PHOENIX LABS NY | 5MG | A080322 001 |
|-----------------|-----|-------------|

| | | |
|---------------|-----|-------------|
| PUREPAC PHARM | 5MG | A080325 001 |
|---------------|-----|-------------|

| | | |
|----------|-----|-------------|
| PVT FORM | 5MG | A080211 001 |
|----------|-----|-------------|

| | | |
|--------|-----|-------------|
| ROXANE | 5MG | A080327 002 |
|--------|-----|-------------|

| | | |
|--------|-----|-------------|
| SPERTI | 1MG | A080358 001 |
|--------|-----|-------------|

| | | |
|--|-------|-------------|
| | 2.5MG | A080358 002 |
|--|-------|-------------|

| | | |
|--|-----|-------------|
| | 5MG | A080358 003 |
|--|-----|-------------|

| | | |
|------------|-----|-------------|
| SUPERPHARM | 5MG | A088892 001 |
|------------|-----|-------------|

Feb 26, 1985

| | | |
|-----------|-----|-------------|
| TABLICAPS | 5MG | A085170 001 |
|-----------|-----|-------------|

| | | |
|------|-----|-------------|
| TEVA | 5MG | A080398 001 |
|------|-----|-------------|

| | | |
|-----|-----|-------------|
| UDL | 5MG | A087987 001 |
|-----|-----|-------------|

Jan 18, 1983

| | | |
|--------------------|-----|-------------|
| VALEANT PHARM INTL | 5MG | A080236 001 |
|--------------------|-----|-------------|

| | | |
|----------|-----|-------------|
| VITARINE | 5MG | A080534 001 |
|----------|-----|-------------|

| | | |
|-------------|-----|-------------|
| WATSON LABS | 5MG | A085085 002 |
|-------------|-----|-------------|

| | | |
|--|-----|-------------|
| | 5MG | A085415 001 |
|--|-----|-------------|

| | | |
|--|-----|-------------|
| | 5MG | A085416 001 |
|--|-----|-------------|

| | | |
|-----------|-----|-------------|
| WEST WARD | 5MG | A080324 001 |
|-----------|-----|-------------|

| | | |
|----------------------|-----|-------------|
| WHITEWORTH TOWN PLSN | 5MG | A080342 001 |
|----------------------|-----|-------------|

STERANE

| | | |
|--------|-----|-------------|
| PFIZER | 5MG | N009996 001 |
|--------|-----|-------------|

PREDNISOLONE ACETATE

INJECTABLE; INJECTION

METICORTELONGE

| | | |
|----------|---------|-------------|
| SCHERING | 25MG/ML | N010255 002 |
|----------|---------|-------------|

PREDNISOLONE ACETATE

| | | |
|-------|---------|-------------|
| AKORN | 25MG/ML | A083032 001 |
|-------|---------|-------------|

| | | |
|--|---------|-------------|
| | 50MG/ML | A084492 001 |
|--|---------|-------------|

| | | |
|---------|---------|-------------|
| BEL MAR | 25MG/ML | A083738 001 |
|---------|---------|-------------|

| | | |
|--|---------|-------------|
| | 50MG/ML | A083738 002 |
|--|---------|-------------|

| | | |
|-------------|---------|-------------|
| CENT PHARMS | 25MG/ML | A084717 001 |
|-------------|---------|-------------|

| | | |
|--|---------|-------------|
| | 50MG/ML | A084717 002 |
|--|---------|-------------|

| | | |
|-------------|---------|-------------|
| WATSON LABS | 25MG/ML | A083398 001 |
|-------------|---------|-------------|

| | | |
|--|---------|-------------|
| | 25MG/ML | A083654 001 |
|--|---------|-------------|

| | | |
|--|---------|-------------|
| | 40MG/ML | A083767 001 |
|--|---------|-------------|

| | | |
|--|---------|-------------|
| | 50MG/ML | A083764 001 |
|--|---------|-------------|

| | | |
|--|---------|-------------|
| | 50MG/ML | A085781 001 |
|--|---------|-------------|

STERANE

| | | |
|--------|---------|-------------|
| PFIZER | 25MG/ML | N011446 001 |
|--------|---------|-------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISOLONE ACETATE

SUSPENSION; ORAL

FLO-PRED

TARO

EQ 5MG BASE/5ML

N022067 001 Jan 17, 2008

EQ 15MG BASE/5ML

N022067 002 Jan 17, 2008

SUSPENSION/DROPS; OPHTHALMIC

ECONOPRED

ALCON

0.125%

N017468 001

PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC

CETAPRED

ALCON

0.25%;10%

A087771 001 Aug 06, 1993

METIMYD

SCHERING

0.5%;10%

N010210 002 Sep 09, 1984

PREDSULFAIR

PHARMAFAIR

0.5%;10%

A088032 001 Apr 15, 1983

VASOCIDIN

NOVARTIS

0.5%;10%

A088791 001 Oct 05, 1984

SUSPENSION; OPHTHALMIC

ISOPTO CETAPRED

ALCON

0.25%;10%

A087547 001

SUSPENSION/DROPS; OPHTHALMIC

METIMYD

SCHERING

0.5%;10%

N010210 001

PREDAMIDE

AKORN

0.5%;10%

A088059 001 Jul 29, 1983

PREDSULFAIR

PHARMAFAIR

0.5%;10%

A088007 001 Apr 19, 1983

PREDSULFAIR II

PHARMAFAIR

0.2%;10%

A088837 001 Dec 24, 1985

SULPHRIN

BAUSCH AND LOMB

0.5%;10%

A088089 001 Dec 28, 1982

PREDNISOLONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

HYDELTRASOL

MERCK

EQ 20MG PHOSPHATE/ML

N011583 002

PREDNISOLONE SODIUM PHOSPHATE

WATSON LABS

EQ 20MG PHOSPHATE/ML

A080517 001

OINTMENT; OPHTHALMIC, OTIC

HYDELTRASOL

MERCK

EQ 0.25% PHOSPHATE

N011028 001

SOLUTION; ORAL

ORAPRED

CONCORDIA PHARMS INC EQ 15MG BASE/5ML **

A075117 001 Dec 14, 2000

PREDNISOLONE SODIUM PHOSPHATE

AMNEAL PHARMS

EQ 15MG BASE/5ML

A078345 001 Mar 10, 2009

MEDICIS PHARMS

EQ 15MG BASE/5ML

A075250 001 Jul 12, 2002

NESHER PHARMS

EQ 5MG BASE/5ML

A076982 001 May 24, 2005

EQ 15MG BASE/5ML

A076988 001 May 24, 2005

PHARM ASSOC

EQ 5MG BASE/5ML

A076123 001 Dec 23, 2002

VINTAGE PHARMS

EQ 5MG BASE/5ML

A078416 001 Oct 31, 2007

WE PHARMS

EQ 5MG BASE/5ML

A075181 001 Dec 23, 2002

SOLUTION/DROPS; OPHTHALMIC

INFLAMASE FORTE

NOVARTIS

EQ 0.9% PHOSPHATE

A080751 002

INFLAMASE MILD

NOVARTIS

EQ 0.11% PHOSPHATE

A080751 001

METRETON

SCHERING

EQ 0.5% PHOSPHATE

A083834 001

PREDAIR

PHARMAFAIR

EQ 0.11% PHOSPHATE

A088415 001 Feb 29, 1984

PREDAIR FORTE

PHARMAFAIR

EQ 0.9% PHOSPHATE

A088165 001 Mar 28, 1983

PREDNISOLONE SODIUM PHOSPHATE

AKORN

EQ 0.11% PHOSPHATE

A083358 001

EQ 0.9% PHOSPHATE

A083358 002

ALCON PHARMS LTD

EQ 0.11% PHOSPHATE

A081043 001 Oct 24, 1991

EQ 0.9% PHOSPHATE

A081044 001 Oct 24, 1991

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION/DROPS;OPHTHALMIC

PREDNISOLONE SODIUM PHOSPHATE

| | | | |
|------------------|--------------------|-------------|--------------|
| BAUSCH AND LOMB | EQ 0.11% PHOSPHATE | A040065 001 | Jul 29, 1994 |
| SOLA BARNES HIND | EQ 0.11% PHOSPHATE | A084171 001 | |
| | EQ 0.9% PHOSPHATE | A084168 001 | |
| | EQ 0.9% PHOSPHATE | A084169 001 | |
| | EQ 0.9% PHOSPHATE | A084172 001 | |

PREDNISOLONE SODIUM PHOSPHATE; SULFACETAMIDE SODIUM

SOLUTION/DROPS;OPHTHALMIC

SULSTER

| | | | |
|-------|------------------------|-------------|--------------|
| AKORN | EQ 0.23% PHOSPHATE;10% | A074511 001 | Jul 30, 1996 |
|-------|------------------------|-------------|--------------|

VASOCIDIN

| | | | |
|------------|---------------------------|-------------|--------------|
| + NOVARTIS | EQ 0.23% PHOSPHATE;10% ** | N018988 001 | Aug 26, 1988 |
|------------|---------------------------|-------------|--------------|

PREDNISOLONE TEBUTATE

INJECTABLE; INJECTION

HYDELTRA-TBA

| | | | |
|-------|---------|-------------|--|
| MERCK | 20MG/ML | N010562 001 | |
|-------|---------|-------------|--|

PREDNISOLONE TEBUTATE

| | | | |
|-------------|---------|-------------|--------------|
| WATSON LABS | 20MG/ML | A083362 001 | Feb 17, 1984 |
|-------------|---------|-------------|--------------|

PREDNISONE

SOLUTION;ORAL

PREDNISONE

| | | | |
|-----------|---------|-------------|--------------|
| WOCKHARDT | 5MG/5ML | A089726 001 | Aug 02, 1988 |
|-----------|---------|-------------|--------------|

SYRUP;ORAL

LIQUID PRED

| | | | |
|------|---------|-------------|--------------|
| MURO | 5MG/5ML | A087611 002 | Sep 07, 1982 |
|------|---------|-------------|--------------|

TABLET;ORAL

CORTAN

| | | | |
|--------|------|-------------|--|
| HALSEY | 20MG | A087480 001 | |
|--------|------|-------------|--|

DELTA-DOME

| | | | |
|--------------|-----|-------------|--|
| BAYER PHARMS | 5MG | A080293 001 | |
|--------------|-----|-------------|--|

DELTASONE

| | | | |
|------------------------|----------|-------------|--|
| + PHARMACIA AND UPJOHN | 2.5MG ** | N009986 005 | |
|------------------------|----------|-------------|--|

| | | | |
|--|--------|-------------|--|
| | 5MG ** | N009986 002 | |
|--|--------|-------------|--|

| | | | |
|--|---------|-------------|--|
| | 10MG ** | N009986 006 | |
|--|---------|-------------|--|

| | | | |
|--|---------|-------------|--|
| | 20MG ** | N009986 007 | |
|--|---------|-------------|--|

| | | | |
|--|---------|-------------|--|
| | 50MG ** | N009986 008 | |
|--|---------|-------------|--|

FERNISONE

| | | | |
|---------------|-----|-------------|--|
| FERNDALE LABS | 5MG | A083364 001 | |
|---------------|-----|-------------|--|

METICORTEN

| | | | |
|------------|--------|-------------|--|
| + SCHERING | 1MG ** | N009766 002 | |
|------------|--------|-------------|--|

| | | | |
|--|--------|-------------|--|
| | 5MG ** | N009766 001 | |
|--|--------|-------------|--|

ORASONE

| | | | |
|--------|-----|-------------|--|
| SOLVAY | 1MG | A083009 001 | |
|--------|-----|-------------|--|

| | | | |
|--|-----|-------------|--|
| | 5MG | A083009 002 | |
|--|-----|-------------|--|

| | | | |
|--|------|-------------|--|
| | 10MG | A083009 003 | |
|--|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 20MG | A083009 004 | |
|--|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 50MG | A085999 001 | |
|--|------|-------------|--|

PARACORT

| | | | |
|-------------|-----|-------------|--|
| PARKE DAVIS | 5MG | N010962 002 | |
|-------------|-----|-------------|--|

PREDNICEN-M

| | | | |
|----------------|-----|-------------|--|
| SCHWARZ PHARMA | 5MG | A084655 001 | |
|----------------|-----|-------------|--|

PREDNISONE

| | | | |
|-----------|-----|-------------|--------------|
| AM THERAP | 5MG | A089387 001 | Nov 06, 1986 |
|-----------|-----|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 10MG | A089388 001 | Nov 06, 1986 |
|--|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 20MG | A089389 001 | Nov 06, 1986 |
|--|------|-------------|--------------|

| | | | |
|------------------|-----|-------------|--------------|
| AMNEAL PHARMS NY | 5MG | A089597 001 | Oct 05, 1987 |
|------------------|-----|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 10MG | A089598 001 | Oct 05, 1987 |
|--|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 20MG | A089599 001 | Oct 05, 1987 |
|--|------|-------------|--------------|

| | | | |
|---------------------|-----|-------------|--|
| AUROLIFE PHARMA LLC | 5MG | A084774 001 | |
|---------------------|-----|-------------|--|

| | | | |
|--|------|-------------|--------------|
| | 10MG | A089983 001 | Jan 12, 1989 |
|--|------|-------------|--------------|

| | | | |
|--|------|-------------|--|
| | 20MG | A085813 001 | |
|--|------|-------------|--|

| | | | |
|--|------|-------------|--------------|
| | 50MG | A089984 001 | Jan 12, 1989 |
|--|------|-------------|--------------|

| | | | |
|-------|-----|-------------|--|
| BUNDY | 5MG | A083676 001 | |
|-------|-----|-------------|--|

| | | | |
|--------------|-----|-------------|--|
| CHARTWELL RX | 5MG | A083059 001 | |
|--------------|-----|-------------|--|

| | | | |
|--------------------|-----|-------------|--|
| CONTRACT PHARMACAL | 5MG | A080209 001 | |
|--------------------|-----|-------------|--|

| | | | |
|---------------------|-----|-------------|--------------|
| DURAMED PHARMS BARR | 5MG | A088394 001 | Oct 04, 1983 |
|---------------------|-----|-------------|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISONTABLET;ORAL
PREDNISON

| | | | |
|----------------------|-------|-------------|--------------|
| | 10MG | A088395 001 | Oct 04, 1983 |
| | 20MG | A088396 001 | Oct 04, 1983 |
| ELKINS SINN | 5MG | A080491 001 | |
| | 20MG | A085811 001 | |
| EVERYLIFE | 1MG | A084440 001 | |
| | 2.5MG | A084440 002 | |
| | 5MG | A084440 003 | |
| FERRANTE | 2.5MG | A080563 001 | |
| | 5MG | A080563 002 | |
| HALSEY | 5MG | A080300 001 | |
| HEATHER | 5MG | A080320 001 | |
| | 10MG | A084341 001 | |
| | 20MG | A084417 001 | |
| | 20MG | A085543 001 | |
| | 50MG | A086946 001 | |
| HIKMA PHARMS | 1MG | A040890 001 | Nov 01, 2010 |
| | 2.5MG | A040538 001 | Jan 08, 2004 |
| IMPAX LABS | 5MG | A080782 001 | |
| INWOOD LABS | 1MG | A080328 001 | |
| | 2.5MG | A080306 001 | |
| | 5MG | A080279 001 | |
| IVAX SUB TEVA PHARMS | 5MG | A080283 001 | |
| | 10MG | A084133 001 | |
| | 20MG | A084134 001 | |
| KV PHARM | 5MG | A084236 001 | |
| LANNETT | 5MG | A080514 001 | |
| | 20MG | A084275 001 | |
| LEDERLE | 5MG | A086968 001 | |
| MARSHALL PHARMA | 5MG | A080301 001 | |
| MUTUAL PHARM | 5MG | A080701 001 | |
| | 10MG | A086595 001 | |
| | 20MG | A084634 001 | |
| NYLOS | 5MG | A085115 001 | |
| PANRAY | 1MG | A080350 001 | |
| | 2.5MG | A080350 002 | |
| | 5MG | A080350 003 | |
| PHARMAVITE | 5MG | A084662 002 | |
| PHOENIX LABS NY | 5MG | A080321 001 | |
| | 20MG | A083807 001 | |
| PUREPAC PHARM | 5MG | A080353 001 | |
| | 10MG | A086062 001 | |
| | 20MG | A086061 001 | |
| PVT FORM | 20MG | A085151 001 | |
| REXALL | 5MG | A080232 001 | |
| ROXANE | 20MG | N017109 001 | |
| | 25MG | A087833 001 | May 04, 1982 |
| SANDOZ | 5MG | A080336 002 | |
| SCHERER LABS | 5MG | A080371 001 | |
| SPERTI | 1MG | A080359 001 | |
| | 2.5MG | A080359 002 | |
| | 5MG | A080359 003 | |
| SUN PHARM INDUSTRIES | 50MG | A086596 001 | |
| SUPERPHARM | 5MG | A088865 001 | Oct 25, 1984 |
| | 10MG | A088866 001 | Oct 25, 1984 |
| | 20MG | A088867 001 | Oct 25, 1984 |
| TEVA | 5MG | A080397 001 | |
| UDL | 5MG | A087984 001 | Jan 18, 1983 |
| | 10MG | A087985 001 | Jan 18, 1983 |
| | 20MG | A087986 001 | Jan 18, 1983 |
| UPSHER SMITH | 5MG | A087471 001 | |
| | 20MG | A087470 001 | |
| VALEANT PHARM INTL | 5MG | A080237 001 | |
| VANGARD | 5MG | A087682 001 | Jan 15, 1982 |
| | 20MG | A087701 001 | Jan 15, 1982 |
| VITARINE | 5MG | A080334 001 | |
| | 5MG | A080506 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PREDNISONE

TABLET; ORAL

PREDNISONE

| | | | | |
|----------------------|-------|---------|-----|--------------|
| WATSON LABS | 5MG | A085084 | 002 | |
| | 10MG | A087773 | 001 | Jul 13, 1982 |
| | 20MG | A086813 | 001 | |
| | 50MG | A086867 | 001 | |
| | 50MG | A087772 | 001 | Jul 13, 1982 |
| WHITEWORTH TOWN PLSN | 2.5MG | A084913 | 001 | |
| | 5MG | A080343 | 001 | |
| | 10MG | A089028 | 001 | Jul 24, 1986 |
| | 20MG | A084913 | 002 | |
| SERVISONE | | | | |
| LEDERLE | 5MG | A080223 | 001 | |

PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST

| | | | | |
|-----------------------|-------|---------|-----|--|
| + ASTRAZENECA | 1% ** | N014763 | 004 | |
| + | 2% ** | N014763 | 005 | |
| + | 3% ** | N014763 | 003 | |
| CITANEST PLAIN | | | | |
| + ASTRAZENECA | 4% ** | N014763 | 007 | |
| CITANEST PLAIN DENTAL | | | | |
| + DENTSPLY PHARM | 4% | N021382 | 001 | |

PRIMIDONE

SUSPENSION; ORAL

MYSOLINE

| | | | | |
|-------------|-----------|---------|-----|--|
| NURO PHARMA | 250MG/5ML | N010401 | 001 | |
|-------------|-----------|---------|-----|--|

TABLET; ORAL

PRIMIDONE

| | | | | |
|--------------------|-------|---------|-----|--------------|
| DR REDDYS LABS LTD | 50MG | A040862 | 001 | Oct 03, 2008 |
| | 250MG | A040862 | 002 | Oct 03, 2008 |
| HIKMA INTL PHARMS | 50MG | A040667 | 001 | Jul 27, 2006 |
| IMPAX LABS | 50MG | A040717 | 001 | Feb 12, 2008 |
| | 250MG | A040717 | 002 | Feb 12, 2008 |
| WATSON LABS | 250MG | A085052 | 001 | |

PROBENECID

TABLET; ORAL

BENEMID

| | | | | |
|---------|----------|---------|-----|--|
| + MERCK | 500MG ** | N007898 | 004 | |
|---------|----------|---------|-----|--|

PROBENECID

| | | | | |
|----------------------|-------|---------|-----|--------------|
| IVAX SUB TEVA PHARMS | 500MG | A083740 | 001 | May 09, 1984 |
| LEDERLE | 500MG | A086917 | 001 | |
| WATSON LABS | 500MG | A086150 | 002 | Apr 23, 1982 |

PROBUCOL

TABLET; ORAL

LORELCO

| | | | | |
|-------------------|-------|---------|-----|--------------|
| SANOFI AVENTIS US | 250MG | N017535 | 001 | |
| | 500MG | N017535 | 002 | Jul 06, 1988 |

PROCAINAMIDE HYDROCHLORIDE

CAPSULE; ORAL

PROCAINAMIDE HYDROCHLORIDE

| | | | | |
|----------------------|-------|---------|-----|--------------|
| ANI PHARMS INC | 250MG | A089219 | 001 | Jul 01, 1986 |
| | 375MG | A089219 | 002 | Jul 01, 1986 |
| | 500MG | A089219 | 003 | Jul 01, 1986 |
| ASCOT | 250MG | A087542 | 001 | Jan 08, 1982 |
| | 375MG | A087697 | 001 | Mar 01, 1983 |
| | 500MG | A087543 | 001 | Jan 08, 1982 |
| IVAX SUB TEVA PHARMS | 250MG | A084604 | 001 | |
| | 375MG | A084595 | 001 | |
| | 500MG | A084606 | 001 | |
| LANNETT | 250MG | A083693 | 001 | |
| | 500MG | A084696 | 001 | |
| LEDERLE | 250MG | A086942 | 001 | |
| | 375MG | A086952 | 001 | |
| | 500MG | A086943 | 001 | |
| ROXANE | 250MG | A088989 | 001 | Apr 26, 1985 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROCAINAMIDE HYDROCHLORIDE

CAPSULE; ORAL

PROCAINAMIDE HYDROCHLORIDE

| | | | | | |
|-------------|-----------|----------|---------|-----|--------------|
| | 500MG | | A088990 | 001 | Apr 26, 1985 |
| VANGARD | 250MG | | A087643 | 001 | Jun 01, 1982 |
| | 500MG | | A087875 | 001 | Jun 01, 1982 |
| WATSON LABS | 250MG | | A083287 | 001 | |
| | 250MG | | A083795 | 001 | |
| | 250MG | | A085167 | 001 | |
| | 375MG | | A084403 | 001 | |
| | 375MG | | A087020 | 001 | |
| | 500MG | | A084280 | 001 | |
| | 500MG | | A084357 | 001 | |
| | 500MG | | A087021 | 001 | |
| PROCAN | | | | | |
| PARKE DAVIS | 250MG | | A085804 | 001 | |
| | 375MG | | A087502 | 001 | |
| | 500MG | | A085079 | 001 | |
| PROCAPAN | | | | | |
| PANRAY | 250MG | | A083553 | 002 | |
| PRONESTYL | | | | | |
| + | APOTHECON | 250MG ** | N007335 | 001 | |
| + | | 375MG ** | N007335 | 004 | |
| + | | 500MG ** | N007335 | 003 | |

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

| | | | | | |
|----------------------|-----------|-------------|---------|-----|--------------|
| ABRAXIS PHARM | 100MG/ML | | A089415 | 001 | Nov 17, 1986 |
| | 500MG/ML | | A089416 | 001 | Nov 17, 1986 |
| HOSPIRA | 500MG/ML | | A089537 | 001 | Aug 25, 1987 |
| INTL MEDICATION | 500MG/ML | | A088637 | 001 | Jul 31, 1984 |
| PHARMAFAIR | 100MG/ML | | A088824 | 001 | Nov 20, 1985 |
| | 500MG/ML | | A088830 | 001 | Nov 20, 1985 |
| SMITH AND NEPHEW | 100MG/ML | | A088530 | 001 | Mar 04, 1985 |
| | 500MG/ML | | A088531 | 001 | Mar 04, 1985 |
| SOLOPAK | 500MG/ML | | A088532 | 001 | Mar 04, 1985 |
| WARNER CHILCOTT | 100MG/ML | | A089528 | 001 | May 03, 1988 |
| | 500MG/ML | | A089529 | 001 | May 03, 1988 |
| WATSON LABS | 100MG/ML | | A087079 | 001 | |
| | 500MG/ML | | A087080 | 001 | |
| WEST-WARD PHARMS INT | 100MG/ML | | A089029 | 001 | Apr 17, 1986 |
| | 500MG/ML | | A089030 | 001 | Apr 17, 1986 |
| PRONESTYL | | | | | |
| + | APOTHECON | 100MG/ML ** | N007335 | 002 | |
| + | | 500MG/ML ** | N007335 | 005 | |

TABLET; ORAL

PRONESTYL

| | | | | | |
|-----------|-------|--|---------|-----|--|
| APOTHECON | 250MG | | N017371 | 001 | |
| | 375MG | | N017371 | 002 | |
| | 500MG | | N017371 | 003 | |

TABLET, EXTENDED RELEASE; ORAL

PROCAINAMIDE HYDROCHLORIDE

| | | | | | |
|----------------|-------|--|---------|-----|--------------|
| ANI PHARMS INC | 250MG | | A088958 | 001 | Dec 02, 1985 |
| | 250MG | | A089369 | 001 | Aug 14, 1987 |
| | 500MG | | A088959 | 001 | Dec 02, 1985 |
| | 500MG | | A088974 | 001 | Jul 22, 1985 |
| | 500MG | | A089369 | 002 | Jan 09, 1987 |
| | 750MG | | A089369 | 003 | Aug 14, 1987 |
| | 750MG | | A089438 | 001 | Mar 23, 1987 |
| | 1GM | | A040111 | 001 | Dec 13, 1996 |
| INWOOD LABS | 500MG | | A089840 | 001 | Mar 06, 1989 |
| SANDOZ | 500MG | | A089284 | 001 | Jun 23, 1986 |
| WATSON LABS | 250MG | | A088533 | 001 | Dec 03, 1984 |
| | 250MG | | A089026 | 001 | Oct 22, 1985 |
| | 500MG | | A088534 | 001 | Dec 03, 1984 |
| | 500MG | | A089027 | 001 | Oct 22, 1985 |
| | 750MG | | A088535 | 001 | Nov 03, 1984 |
| | 750MG | | A089042 | 001 | Oct 22, 1985 |
| | 1GM | | A089520 | 001 | Jan 15, 1987 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROCAINAMIDE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

PROCAN SR

| | | | | |
|-------------|-------|---------|-----|--------------|
| PARKE DAVIS | 250MG | A086468 | 001 | |
| PARKE DALE | 500MG | A086065 | 001 | |
| | 750MG | A087510 | 001 | Apr 01, 1982 |
| | 1GM | A088489 | 001 | Jan 16, 1985 |

PROCANBID

| | | | | |
|-------------|-------|---------|-----|--------------|
| KING PHARMS | 500MG | N020545 | 001 | Jan 31, 1996 |
| | 1GM | N020545 | 002 | Jan 31, 1996 |

PRONESTYL-SR

| | | | | |
|-----------|-------|---------|-----|--|
| APOTHECON | 500MG | A087361 | 001 | |
|-----------|-------|---------|-----|--|

PROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

NOVOCAIN

| | | | | |
|---------|-----|---------|-----|--|
| HOSPIRA | 1% | A085362 | 003 | |
| | 2% | A085362 | 004 | |
| | 10% | A086797 | 001 | |

PROCAINE HYDROCHLORIDE

| | | | | |
|---------------|----|---------|-----|--|
| ABRAXIS PHARM | 1% | A080384 | 002 | |
| | 1% | A080421 | 001 | |
| | 2% | A080384 | 003 | |
| | 2% | A080421 | 002 | |
| BEL MAR | 1% | A080711 | 001 | |
| | 2% | A080756 | 001 | |
| ELKINS SINN | 1% | A083315 | 001 | |
| | 2% | A083315 | 002 | |
| GD SEARLE LLC | 1% | A086202 | 001 | |
| | 2% | A086202 | 002 | |
| HOSPIRA | 1% | A080416 | 001 | |
| | 2% | A080416 | 002 | |
| MILES | 1% | A080415 | 001 | |
| | 2% | A080415 | 002 | |
| WATSON LABS | 1% | A080658 | 001 | |
| | 1% | A083535 | 001 | |
| | 2% | A080658 | 002 | |
| | 2% | A083535 | 002 | |

PROCAINE HYDROCHLORIDE; TETRACYCLINE HYDROCHLORIDE

INJECTABLE; INJECTION

ACHROMYCIN

| | | | | |
|---------|----------------------|---------|-----|--|
| LEDERLE | 40MG/VIAL;100MG/VIAL | N050276 | 001 | |
| | 40MG/VIAL;250MG/VIAL | N050276 | 003 | |

TETRACYN

| | | | | |
|--------|----------------------|---------|-----|--|
| PFIZER | 40MG/VIAL;100MG/VIAL | A060285 | 002 | |
| | 40MG/VIAL;250MG/VIAL | A060285 | 003 | |

PROCAINE MERETHOXYLLINE; THEOPHYLLINE

INJECTABLE; INJECTION

DICURIN PROCAINE

| | | | | |
|-------|------------------|---------|-----|--|
| LILLY | 100MG/ML;50MG/ML | N008869 | 001 | |
|-------|------------------|---------|-----|--|

PROCHLORPERAZINE

SUPPOSITORY; RECTAL

COMPAZINE

| | | | | |
|-----------------|----------|---------|-----|--|
| GLAXOSMITHKLINE | 2.5MG ** | N011127 | 003 | |
| | 5MG ** | N011127 | 001 | |
| | 25MG ** | N011127 | 002 | |

PROCHLORPERAZINE

| | | | | |
|------|-------|---------|-----|--------------|
| ABLE | 2.5MG | A040407 | 001 | Jul 11, 2001 |
| | 5MG | A040407 | 002 | Jul 11, 2001 |
| | 25MG | A040407 | 003 | Jul 11, 2001 |

PROCHLORPERAZINE EDISYLATE

CONCENTRATE; ORAL

COMPAZINE

| | | | | |
|-----------------|-----------------|---------|-----|--|
| GLAXOSMITHKLINE | EQ 10MG BASE/ML | N011276 | 001 | |
|-----------------|-----------------|---------|-----|--|

PROCHLORPERAZINE

| | | | | |
|--------------------|-----------------|---------|-----|--------------|
| ALPHARMA US PHARMS | EQ 10MG BASE/ML | A087153 | 001 | Jun 08, 1982 |
|--------------------|-----------------|---------|-----|--------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROCHLORPERAZINE EDISYLATE

CONCENTRATE; ORAL

PROCHLORPERAZINE EDISYLATE

MORTON GROVE

EQ 10MG BASE/ML

A088598 001 Oct 25, 1984

INJECTABLE; INJECTION

COMPAZINE

+ GLAXOSMITHKLINE

EQ 5MG BASE/ML **

N010742 002

PROCHLORPERAZINE

BAXTER HLTHCARE

EQ 5MG BASE/ML

A087759 001 Oct 01, 1982

PROCHLORPERAZINE EDISYLATE

HOSPIRA

EQ 5MG BASE/ML

A089703 001 Apr 07, 1988

MARSAM PHARMS LLC

EQ 5MG BASE/ML

A089675 001 Dec 05, 1988

SMITH AND NEPHEW

EQ 5MG BASE/ML

A089251 001 Dec 04, 1986

TEVA PARENTERAL

EQ 5MG BASE/ML

A040505 001 May 30, 2003

WATSON LABS

EQ 5MG BASE/ML

A089530 001 Jul 08, 1987

EQ 5MG BASE/ML

A089605 001 Jul 08, 1987

EQ 5MG BASE/ML

A089606 001 Jul 08, 1987

WEST-WARD PHARMS INT

EQ 5MG BASE/ML

A089523 001 May 03, 1988

WYETH AYERST

EQ 5MG BASE/ML

A086348 001

SYRUP; ORAL

COMPAZINE

GLAXOSMITHKLINE

EQ 5MG BASE/5ML

N011188 001

PROCHLORPERAZINE EDISYLATE

ALPHARMA US PHARMS

EQ 5MG BASE/5ML

A087154 001 Sep 01, 1982

MORTON GROVE

EQ 5MG BASE/5ML

A088597 001 Oct 25, 1984

PROCHLORPERAZINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

COMPAZINE

GLAXOSMITHKLINE

EQ 10MG BASE

N011000 001

EQ 10MG BASE

N021019 001 Oct 06, 1999

EQ 15MG BASE

N011000 002

EQ 15MG BASE

N021019 002 Oct 06, 1999

EQ 30MG BASE

N011000 003

EQ 75MG BASE

N011000 004

TABLET; ORAL

COMPAZINE

GLAXOSMITHKLINE

EQ 5MG BASE **

N010571 001

EQ 10MG BASE **

N010571 002

EQ 25MG BASE **

N010571 003

PROCHLORPERAZINE

WATSON LABS

EQ 5MG BASE

A085580 001

EQ 10MG BASE

A085178 001

EQ 25MG BASE

A085579 001

PROCHLORPERAZINE MALEATE

DURAMED PHARMS BARR

EQ 5MG BASE

A040207 001 May 01, 1997

EQ 5MG BASE

A089484 001 Jan 20, 1987

EQ 10MG BASE

A040207 002 May 01, 1997

EQ 10MG BASE

A089485 001 Jan 20, 1987

EQ 25MG BASE

A089486 001 Jan 20, 1987

IVAX SUB TEVA PHARMS

EQ 5MG BASE

A040162 001 Jan 20, 1998

EQ 10MG BASE

A040162 002 Jan 20, 1998

SANDOZ

EQ 25MG BASE

A040101 003 Jul 19, 1996

PROCYCLIDINE HYDROCHLORIDE

TABLET; ORAL

KEMADRIN

MONARCH PHARMS

2MG

N009818 005

5MG

N009818 003

PROGESTERONE

CAPSULE; ORAL

PROGESTERONE

TEVA PHARMS

100MG

A202121 001 Feb 29, 2012

200MG

A202121 002 Feb 29, 2012

PROMETRIUM

VIRTUS PHARMS

300MG

N019781 003 Oct 15, 1999

INJECTABLE; INJECTION

PROGESTERONE

LILLY

25MG/ML

N009238 002

50MG/ML

N009238 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROGESTERONE

INSERT, EXTENDED RELEASE; INTRAUTERINE

PROGESTASERT

ALZA 38MG N017553 001

PROMAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

SPARINE

WYETH AYERST 30MG/ML N010942 001

100MG/ML N010942 004

INJECTABLE; INJECTION

PROMAZINE HYDROCHLORIDE

WATSON LABS 25MG/ML A084510 001

50MG/ML A084517 001

SPARINE

BAXTER HLTHCARE CORP 25MG/ML N010349 008

50MG/ML N010349 006

SYRUP; ORAL

SPARINE

WYETH AYERST 10MG/5ML N010942 003

TABLET; ORAL

SPARINE

WYETH AYERST 10MG N010348 006

25MG N010348 001

50MG N010348 002

100MG N010348 003

200MG N010348 004

PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PHENERGAN

WYETH AYERST 25MG/ML N008857 002

50MG/ML N008857 003

PROMETHAZINE HYDROCHLORIDE

ABBOTT 25MG/ML A084223 001

50MG/ML A084222 001

AKORN 25MG/ML A083955 002

50MG/ML A083955 001

BEDFORD LABS 25MG/ML A040524 001 Mar 17, 2004

50MG/ML A040524 002 Mar 17, 2004

HOSPIRA 25MG/ML A040372 001 Jun 08, 2000

50MG/ML A040372 002 Jun 08, 2000

50MG/ML A083838 002

LUITPOLD 25MG/ML A040515 001 Mar 19, 2003

MARSAM PHARMS LLC 25MG/ML A089463 001 May 02, 1988

50MG/ML A089477 001 May 02, 1988

MYLAN INSTITUTIONAL 25MG/ML A040471 001 Nov 21, 2002

SANDOZ 25MG/ML A040593 001 Nov 08, 2006

50MG/ML A040593 002 Nov 08, 2006

TEVA PHARMS USA 25MG/ML ** A040454 001 Aug 22, 2002

50MG/ML ** A040454 002 Aug 22, 2002

WATSON LABS 25MG/ML A083532 001

25MG/ML A084591 001

50MG/ML A080629 002

50MG/ML A083532 002

WOCKHARDT 25MG/ML A040785 001 Sep 26, 2008

50MG/ML A040785 002 Sep 26, 2008

ZIPAN-25

ALTANA 25MG/ML A083997 001

ZIPAN-50

ALTANA 50MG/ML A083997 002

SUPPOSITORY; RECTAL

PHENERGAN

+ MYLAN PHARMS INC 12.5MG ** N010926 002

+ 25MG ** N010926 001

+ 50MG ** N011689 001

PROMETHACON

POLYMEDICA 25MG A084901 001

50MG A084902 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROMETHAZINE HYDROCHLORIDE

SUPPOSITORY;RECTAL

PROMETHAZINE HYDROCHLORIDE

| | | | |
|------|--------|-------------|--------------|
| ABLE | 12.5MG | A040504 001 | Apr 11, 2003 |
| | 25MG | A040504 002 | Apr 11, 2003 |
| | 50MG | A040449 001 | Feb 27, 2003 |

SYRUP;ORAL

MYMETHAZINE FORTIS

| | | | |
|------------|----------|-------------|--------------|
| USL PHARMA | 25MG/5ML | A087996 001 | Jan 18, 1983 |
|------------|----------|-------------|--------------|

PROMETH FORTIS

| | | | |
|--------------------|----------|-------------|--|
| ALPHARMA US PHARMS | 25MG/5ML | A084772 001 | |
|--------------------|----------|-------------|--|

PROMETH PLAIN

| | | | |
|----------------------|------------|-------------|--|
| ACTAVIS MID ATLANTIC | 6.25MG/5ML | A085953 001 | |
|----------------------|------------|-------------|--|

PROMETHAZINE

| | | | |
|-------|------------|-------------|--------------|
| CENCI | 6.25MG/5ML | A089013 001 | Sep 20, 1985 |
|-------|------------|-------------|--------------|

PROMETHAZINE HYDROCHLORIDE

| | | | |
|----------|------------|-------------|--|
| KV PHARM | 6.25MG/5ML | A085388 001 | |
|----------|------------|-------------|--|

| | | | |
|--|----------|-------------|--|
| | 25MG/5ML | A085385 001 | |
|--|----------|-------------|--|

| | | | |
|-------------|------------|-------------|--|
| PHARM ASSOC | 6.25MG/5ML | A087518 001 | |
|-------------|------------|-------------|--|

| | | | |
|----------------------|------------|-------------|--|
| WHITEWORTH TOWN PLSN | 6.25MG/5ML | A086395 001 | |
|----------------------|------------|-------------|--|

PROMETHAZINE HYDROCHLORIDE PLAIN

| | | | |
|--------------|---------------|-------------|--------------|
| + ANI PHARMS | 6.25MG/5ML ** | N008381 004 | Apr 18, 1984 |
|--------------|---------------|-------------|--------------|

| | | | |
|---|-------------|-------------|--|
| + | 25MG/5ML ** | N008381 003 | |
|---|-------------|-------------|--|

TABLET;ORAL

PHENERGAN

| | | | |
|---------------------|-----------|-------------|--|
| + DELCOR ASSET CORP | 12.5MG ** | N007935 002 | |
|---------------------|-----------|-------------|--|

| | | | |
|---|---------|-------------|--|
| + | 25MG ** | N007935 003 | |
|---|---------|-------------|--|

| | | | |
|---|---------|-------------|--|
| + | 50MG ** | N007935 004 | |
|---|---------|-------------|--|

PROMETHAZINE HYDROCHLORIDE

| | | | |
|--------|--------|-------------|--|
| ABBOTT | 12.5MG | A084160 001 | |
|--------|--------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 25MG | A084166 001 | |
|--|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 50MG | A084539 001 | |
|--|------|-------------|--|

| | | | |
|------|--------|-------------|--------------|
| ABLE | 12.5MG | A040558 001 | Jul 01, 2004 |
|------|--------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 25MG | A040558 002 | Jul 01, 2004 |
|--|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 50MG | A040558 003 | Jul 01, 2004 |
|--|------|-------------|--------------|

| | | | |
|------------|------|-------------|--------------|
| IMPAX LABS | 25MG | A084214 002 | Jul 07, 1982 |
|------------|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 50MG | A040791 001 | May 20, 2008 |
|--|------|-------------|--------------|

| | | | |
|----------------------|--------|-------------|--|
| IVAX SUB TEVA PHARMS | 12.5MG | A083604 001 | |
|----------------------|--------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 25MG | A083603 001 | |
|--|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 50MG | A083613 001 | |
|--|------|-------------|--|

| | | | |
|---------|--------|-------------|--|
| LANNETT | 12.5MG | A080949 001 | |
|---------|--------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 25MG | A080949 002 | |
|--|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 50MG | A080949 003 | |
|--|------|-------------|--|

| | | | |
|-------|--------|-------------|--------------|
| MYLAN | 12.5MG | A091054 001 | Aug 30, 2011 |
|-------|--------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 25MG | A091054 002 | Aug 30, 2011 |
|--|------|-------------|--------------|

| | | | |
|--|------|-------------|--------------|
| | 50MG | A091054 003 | Aug 30, 2011 |
|--|------|-------------|--------------|

| | | | |
|----------|--------|-------------|--|
| PVT FORM | 12.5MG | A083214 001 | |
|----------|--------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 25MG | A083658 001 | |
|--|------|-------------|--|

| | | | |
|--------|--------|-------------|--------------|
| SANDOZ | 12.5MG | A084176 002 | May 22, 2009 |
|--------|--------|-------------|--------------|

| | | | |
|--|--------|-------------|--|
| | 12.5MG | A084233 001 | |
|--|--------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 25MG | A085146 001 | |
|--|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 50MG | A085146 002 | |
|--|------|-------------|--|

| | | | |
|----------------------|--------|-------------|--|
| SUN PHARM INDUSTRIES | 12.5MG | A084555 001 | |
|----------------------|--------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 25MG | A084554 001 | |
|--|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 50MG | A084557 001 | |
|--|------|-------------|--|

| | | | |
|-----------|--------|-------------|--|
| TABLICAPS | 12.5MG | A084080 001 | |
|-----------|--------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 25MG | A084027 001 | |
|--|------|-------------|--|

| | | | |
|------|------|-------------|--------------|
| TEVA | 25MG | A089109 001 | Sep 10, 1985 |
|------|------|-------------|--------------|

| | | | |
|-------------|--------|-------------|--|
| WATSON LABS | 12.5MG | A083401 001 | |
|-------------|--------|-------------|--|

| | | | |
|--|--------|-------------|--|
| | 12.5MG | A083712 001 | |
|--|--------|-------------|--|

| | | | |
|--|--------|-------------|--|
| | 12.5MG | A085986 001 | |
|--|--------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 25MG | A083204 001 | |
|--|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 25MG | A085684 001 | |
|--|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 50MG | A083403 001 | |
|--|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 50MG | A085664 001 | |
|--|------|-------------|--|

REMSD

| | | | |
|----------------------|------|-------------|--|
| BRISTOL MYERS SQUIBB | 25MG | A083176 002 | |
|----------------------|------|-------------|--|

| | | | |
|--|------|-------------|--|
| | 50MG | A083176 001 | |
|--|------|-------------|--|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPAFENONE HYDROCHLORIDE

TABLET; ORAL

PROPAFENONE HYDROCHLORIDE

| | | | |
|---------------|-------|-------------|--------------|
| NESHER PHARMS | 150MG | A076193 001 | Feb 07, 2002 |
| | 225MG | A076193 002 | Feb 07, 2002 |
| | 300MG | A076193 003 | Feb 07, 2002 |

RHYTHMOL

| | | | |
|-----------------------|-------|-------------|--------------|
| + GLAXOSMITHKLINE LLC | 150MG | N019151 001 | Nov 27, 1989 |
| + | 225MG | N019151 003 | Nov 20, 1992 |
| + | 300MG | N019151 002 | Nov 27, 1989 |

PROPANTHELINE BROMIDE

INJECTABLE; INJECTION

PRO-BANTHINE

| | | | |
|---------------|-----------|-------------|--|
| GD SEARLE LLC | 30MG/VIAL | N008843 001 | |
|---------------|-----------|-------------|--|

TABLET; ORAL

PRO-BANTHINE

| | | | |
|---------|----------|-------------|--|
| + SHIRE | 7.5MG ** | N008732 003 | |
| + | 15MG ** | N008732 002 | |

PROPANTHELINE BROMIDE

| | | | |
|----------------------|-------|-------------|--------------|
| ASCOT | 15MG | A087663 001 | Oct 25, 1982 |
| HEATHER | 15MG | A085780 001 | |
| IMPAX LABS | 15MG | A084541 002 | |
| MYLAN | 15MG | A083706 001 | |
| PAR PHARM | 15MG | A088377 001 | Dec 08, 1983 |
| PVT FORM | 15MG | A080977 001 | |
| SANDOZ | 15MG | A080928 001 | |
| TABLICAPS | 15MG | A084428 001 | |
| WATSON LABS | 15MG | A083029 002 | |
| | 15MG | A083151 001 | |
| WEST-WARD PHARMS INT | 7.5MG | A080927 001 | |

PROPARACAINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

KAINAIR

| | | | |
|------------|------|-------------|--------------|
| PHARMAFAIR | 0.5% | A088087 001 | Jun 07, 1983 |
|------------|------|-------------|--------------|

OPHTHAINE

| | | | |
|-------------|---------|-------------|--|
| + APOTHECON | 0.5% ** | N008883 001 | |
|-------------|---------|-------------|--|

OPHTHETIC

| | | | |
|------------|---------|-------------|--|
| + ALLERGAN | 0.5% ** | N012583 001 | |
|------------|---------|-------------|--|

PARACAINE

| | | | |
|----------|------|-------------|--------------|
| OPTOPICS | 0.5% | A087681 001 | Aug 05, 1982 |
|----------|------|-------------|--------------|

PROPARACAINE HYDROCHLORIDE

| | | | |
|------------------|------|-------------|--|
| SOLA BARNES HIND | 0.5% | A084144 001 | |
| | 0.5% | A084151 001 | |

PROPIOLACTONE

SOLUTION; IRRIGATION

BETAPRONE

| | | | |
|-------------|-----|-------------|--|
| FOREST LABS | N/A | N011657 001 | |
|-------------|-----|-------------|--|

PROPIOMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

LARGON

| | | | |
|----------------------|---------|-------------|--|
| WEST-WARD PHARMS INT | 20MG/ML | N012382 002 | |
|----------------------|---------|-------------|--|

PROPOFOL

INJECTABLE; INJECTION

DIPRIVAN

| | | | |
|--------------------|---------|-------------|--------------|
| FRESENIUS KABI USA | 10MG/ML | N019627 001 | Oct 02, 1989 |
|--------------------|---------|-------------|--------------|

PROPOFOL

| | | | |
|----------------------|---------|-------------|--------------|
| TEVA PARENTERAL | 10MG/ML | A075392 001 | Sep 19, 2000 |
| WEST-WARD PHARMS INT | 10MG/ML | A074848 001 | Apr 19, 2005 |

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

DARVON

| | | | |
|----------------|------|-------------|--|
| XANODYNE PHARM | 32MG | N010997 001 | |
| | 65MG | N010997 003 | |

DOLENE

| | | | |
|---------------------|------|-------------|--|
| HERITAGE PHARMS INC | 65MG | A080530 001 | |
|---------------------|------|-------------|--|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

KESSO-GESIC

MK LABS 65MG A083544 001

PROPHENE 65

HALSEY 65MG A083538 002

PROPOXYPHENE HYDROCHLORIDE

ALRA 65MG A083184 001

IMPAX LABS 65MG A083317 001

IVAX SUB TEVA PHARMS 32MG A083597 001

MUTUAL PHARM 65MG A083186 001

MYLAN 32MG A083528 001

65MG A040569 001 Dec 16, 2004

65MG A083299 001

NEXGEN PHARMA INC 65MG A083185 001

PAR PHARM 65MG A080269 001

PUREPAC PHARM 65MG A083278 001

PVT FORM 32MG A083464 001

65MG A083113 001

ROXANE 32MG A083089 001

65MG A083089 002

SANDOZ 32MG A084014 001

65MG A083125 002

65MG A083688 001

65MG A083870 002

65MG A086495 001

TEVA 65MG A088615 001 Oct 22, 1984

VALEANT PHARM INTL 65MG A080783 001

VINTAGE PHARMS 65MG A040908 001 Jul 17, 2009

WATSON LABS 65MG A080908 002

65MG A085190 001

WEST WARD 65MG A083501 001

WHITEWORTH TOWN PLSN 65MG A084551 001

PROPOXYPHENE HYDROCHLORIDE 65

WARNER CHILCOTT 65MG A083786 001

PROPOXYPHENE NAPSYLATE

SUSPENSION; ORAL

DARVON-N

AAIPHARMA LLC 50MG/5ML N016861 001

TABLET; ORAL

DARVON-N

XANODYNE PHARM 100MG N016862 002

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PROPRANOLOL HYDROCHLORIDE

INWOOD LABS 60MG A072499 001 Apr 11, 1989

80MG A072500 001 Apr 11, 1989

120MG A072501 001 Apr 11, 1989

160MG A072502 001 Apr 11, 1989

MYLAN 60MG A078022 001 Feb 15, 2007

80MG A078022 002 Feb 15, 2007

120MG A078022 003 Feb 15, 2007

160MG A078022 004 Feb 15, 2007

UPSHER SMITH LABS 60MG A078311 001 Mar 06, 2009

80MG A078311 002 Mar 06, 2009

120MG A078311 003 Mar 06, 2009

160MG A078311 004 Mar 06, 2009

CONCENTRATE; ORAL

PROPRANOLOL HYDROCHLORIDE INTENSOL

ROXANE 80MG/ML A071388 001 May 15, 1987

INJECTABLE; INJECTION

PROPRANOLOL HYDROCHLORIDE

+ BAXTER HLTHCARE CORP 1MG/ML N016419 001

FOSUN PHARMA 1MG/ML A076400 001 Feb 26, 2003

SMITH AND NEPHEW 1MG/ML A070135 001 Apr 15, 1986

1MG/ML A070137 001 Apr 15, 1986

SOLOPAK 1MG/ML A070136 001 Apr 15, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPRANOLOL HYDROCHLORIDE

SOLUTION; ORAL

PROPRANOLOL HYDROCHLORIDE

MORTON GROVE

20MG/5ML

A071984 001 Mar 03, 1989

40MG/5ML

A071985 001 Mar 03, 1989

SUSPENSION; ORAL

INDERAL

WYETH AYERST

10MG/ML

N019536 001 Dec 12, 1986

TABLET; ORAL

INDERAL

+ WYETH PHARMS

10MG **

N016418 001

+

20MG **

N016418 003

+

40MG **

N016418 002

+

60MG **

N016418 009 Oct 18, 1982

+

80MG **

N016418 004

+

90MG **

N016418 010 Oct 18, 1982

PROPRANOLOL HYDROCHLORIDE

ANDA REPOSITORY

10MG

A070319 001 Oct 22, 1985

20MG

A070320 001 Oct 22, 1985

40MG

A070103 001 Oct 22, 1985

60MG

A070321 001 Sep 24, 1986

80MG

A070322 001 Aug 04, 1986

ANI PHARMS INC

90MG

A071977 001 Apr 06, 1988

DAVA PHARMS INC

10MG

A070125 001 Jul 30, 1985

20MG

A070126 001 Jul 30, 1985

40MG

A070127 001 Jul 30, 1985

60MG

A071495 001 Dec 31, 1987

80MG

A070128 001 Jul 30, 1985

90MG

A071496 001 Dec 31, 1987

DURAMED PHARMS BARR

10MG

A070306 001 Sep 09, 1985

20MG

A070307 001 Sep 09, 1985

40MG

A070308 001 Sep 09, 1985

60MG

A070309 001 Oct 01, 1986

80MG

A070310 001 Sep 09, 1985

90MG

A071327 001 Oct 01, 1986

INTERPHARM

10MG

A071368 001 May 05, 1987

20MG

A071369 001 May 05, 1987

40MG

A071370 001 May 05, 1987

80MG

A071371 001 May 05, 1987

IVAX SUB TEVA PHARMS

10MG

A072063 001 Jul 29, 1988

20MG

A072066 001 Jul 29, 1988

40MG

A072067 001 Jul 29, 1988

60MG

A072068 001 Jul 29, 1988

80MG

A072069 001 Jul 29, 1988

LEDERLE

10MG

A072117 001 Jun 23, 1988

20MG

A072118 001 Jun 23, 1988

40MG

A072119 001 Jun 23, 1988

80MG

A072120 001 Jun 23, 1988

MYLAN

60MG

A072275 001 Jun 09, 1989

PAR PHARM

90MG

A071288 001 Oct 22, 1986

PUREPAC PHARM

10MG

A070814 001 Nov 03, 1986

20MG

A070815 001 Nov 03, 1986

40MG

A070816 001 Nov 03, 1986

60MG

A070817 001 Nov 03, 1986

80MG

A070757 001 Nov 03, 1986

ROXANE

10MG

A070516 001 Jul 07, 1986

20MG

A070517 001 Jul 07, 1986

40MG

A070518 001 Jul 07, 1986

60MG

A070519 001 Sep 24, 1986

80MG

A070520 001 Jul 07, 1986

90MG

A070521 001 Sep 24, 1986

SANDOZ

10MG

A071658 001 Jul 05, 1988

20MG

A071687 001 Jul 05, 1988

40MG

A071688 001 Jul 05, 1988

60MG

A072197 001 Jul 05, 1988

80MG

A071689 001 Jul 05, 1988

90MG

A072198 001 Jul 05, 1988

SCHERING

10MG

A070120 001 Aug 06, 1985

20MG

A070121 001 Aug 06, 1985

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

| | | | | |
|------------------|------|---------|-----|--------------|
| | 40MG | A070122 | 001 | Aug 06, 1985 |
| | 60MG | A070123 | 001 | Oct 29, 1986 |
| | 80MG | A070124 | 001 | Aug 06, 1985 |
| SUPERPHARM | 10MG | A071515 | 001 | Jun 08, 1988 |
| | 20MG | A071516 | 001 | Jun 08, 1988 |
| | 40MG | A071517 | 001 | Jun 08, 1988 |
| | 80MG | A071518 | 001 | Jun 08, 1988 |
| TEVA | 10MG | A070232 | 001 | Oct 07, 1987 |
| | 20MG | A070233 | 001 | Jun 23, 1986 |
| | 40MG | A070234 | 001 | Jun 23, 1986 |
| WARNER CHILCOTT | 10MG | A070438 | 001 | Sep 15, 1986 |
| | 20MG | A070439 | 001 | Sep 15, 1986 |
| | 40MG | A070440 | 001 | Sep 15, 1986 |
| | 60MG | A070441 | 001 | Sep 24, 1986 |
| | 80MG | A070442 | 001 | Sep 15, 1986 |
| WATSON LABS | 10MG | A070140 | 001 | Jul 30, 1985 |
| | 10MG | A070378 | 001 | Mar 19, 1987 |
| | 20MG | A070141 | 001 | Jul 30, 1985 |
| | 20MG | A070379 | 001 | Mar 19, 1987 |
| | 40MG | A070142 | 001 | Jul 30, 1985 |
| | 40MG | A070380 | 001 | Mar 19, 1987 |
| | 60MG | A070143 | 001 | Jan 15, 1987 |
| | 60MG | A070381 | 001 | Mar 19, 1987 |
| | 60MG | A071098 | 001 | Oct 06, 1986 |
| | 60MG | A071791 | 001 | Jul 15, 1987 |
| | 80MG | A070144 | 001 | Jul 30, 1985 |
| | 80MG | A070382 | 001 | Mar 19, 1987 |
| | 80MG | A070551 | 001 | Jul 10, 1986 |
| | 90MG | A071183 | 001 | Oct 06, 1986 |
| | 90MG | A071792 | 001 | Jul 15, 1987 |
| WATSON LABS TEVA | 10MG | A070548 | 001 | Jul 10, 1986 |
| | 20MG | A070549 | 001 | Apr 11, 1986 |
| | 40MG | A070550 | 001 | Apr 11, 1986 |
| YAOPHARMA CO LTD | 10MG | A070663 | 001 | Jun 13, 1986 |
| | 20MG | A070664 | 001 | Jun 13, 1986 |
| | 40MG | A070665 | 001 | Jun 13, 1986 |
| | 60MG | A070666 | 001 | Oct 10, 1986 |
| | 80MG | A070667 | 001 | Jun 13, 1986 |

PROPYLIDONE

SUSPENSION; INTRATRACHEAL

DIONOSIL AQUEOUS

GLAXOSMITHKLINE 50% N009309 001

DIONOSIL OILY

GLAXOSMITHKLINE 60% N009309 002

PROPYLTHIOURACIL

TABLET; ORAL

PROPYLTHIOURACIL

| | | | |
|----------------------|------|---------|-----|
| ABBOTT | 50MG | A084075 | 001 |
| ANABOLIC | 50MG | A080285 | 001 |
| ANI PHARMS INC | 50MG | A080215 | 001 |
| CHARTWELL RX | 50MG | A084543 | 001 |
| HALSEY | 50MG | A080015 | 001 |
| HIKMA INTL PHARMS | 50MG | A080154 | 001 |
| IMPAX LABS | 50MG | A080159 | 001 |
| LANNETT | 50MG | A080016 | 001 |
| LILLY | 50MG | N006213 | 001 |
| SUN PHARM INDUSTRIES | 50MG | A083982 | 001 |
| TABLICAPS | 50MG | A080840 | 001 |
| WATSON LABS | 50MG | A080932 | 001 |
| | 50MG | A085201 | 001 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PROTAMINE SULFATE

INJECTABLE; INJECTION

PROTAMINE SULFATE

| | | | | | |
|---|----------------------|------------|---------|-----|--------------|
| + | LILLY | 10MG/ML ** | N006460 | 002 | |
| | PHARMACIA AND UPJOHN | 50MG/VIAL | N007413 | 001 | |
| | | 250MG/VIAL | N007413 | 002 | Aug 02, 1984 |
| | WEST-WARD PHARMS INT | 10MG/ML | A089474 | 001 | Nov 05, 1986 |
| | | 10MG/ML | A089475 | 001 | Nov 05, 1986 |

PROTEIN HYDROLYSATE

INJECTABLE; INJECTION

AMINOSOL 5%

| | | | | | |
|--|---------------|----|---------|-----|--------------|
| | ABBVIE | 5% | N005932 | 012 | Jan 31, 1985 |
| | HYPROTIGEN 5% | | | | |
| | B BRAUN | 5% | N006170 | 003 | Jan 10, 1984 |

PROTIRELIN

INJECTABLE; INJECTION

THYPINONE

| | | | | | |
|--|------------|----------|---------|-----|--|
| | ABBOTT | 0.5MG/ML | N017638 | 001 | |
| | THYREL TRH | | | | |
| | FERRING | 0.5MG/ML | N018087 | 001 | |

PROTOKYLOL HYDROCHLORIDE

TABLET; ORAL

VENTAIRE

| | | | | | |
|--|-------------------|-----|---------|-----|--|
| | SANOFI AVENTIS US | 2MG | A083459 | 001 | |
|--|-------------------|-----|---------|-----|--|

PROTRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

VIVACTIL

| | | | | | |
|--|-------------|---------|---------|-----|--|
| | TEVA WOMENS | 5MG ** | N016012 | 001 | |
| | | 10MG ** | N016012 | 002 | |

PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

NOVAFED

| | | | | | |
|---|-------------------|----------|---------|-----|--|
| | SANOFI AVENTIS US | 120MG | N017603 | 001 | |
| | SUDAFED 12 HOUR | | | | |
| + | GLAXOSMITHKLINE | 120MG ** | N017941 | 002 | |

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

ACTIFED

| | | | | | |
|--|-------------------------------------------------|------------|---------|-----|--------------|
| | GLAXOSMITHKLINE | 120MG; 5MG | N018996 | 001 | Jun 17, 1985 |
| | TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES | | | | |
| | KV PHARM | 120MG; 5MG | A071798 | 001 | Mar 16, 1989 |

SYRUP; ORAL

ACTAHIST

| | | | | | |
|--|-------|----------------------|---------|-----|--------------|
| | CENCI | 30MG/5ML; 1.25MG/5ML | A088344 | 001 | Feb 09, 1984 |
|--|-------|----------------------|---------|-----|--------------|

HISTAFED

| | | | | | |
|--|-------|----------------------|---------|-----|--------------|
| | CENCI | 30MG/5ML; 1.25MG/5ML | A088283 | 001 | Apr 20, 1984 |
|--|-------|----------------------|---------|-----|--------------|

MYFED

| | | | | | |
|--|------------|----------------------|---------|-----|--------------|
| | USL PHARMA | 30MG/5ML; 1.25MG/5ML | A088116 | 001 | Mar 04, 1983 |
|--|------------|----------------------|---------|-----|--------------|

TRILITRON

| | | | | | |
|--|----------------|----------------------|---------|-----|--------------|
| | NEWTRON PHARMS | 30MG/5ML; 1.25MG/5ML | A088474 | 001 | Feb 12, 1985 |
|--|----------------|----------------------|---------|-----|--------------|

TABLET; ORAL

ALLERFED

| | | | | | |
|--|----------|-------------|---------|-----|--------------|
| | PVT FORM | 60MG; 2.5MG | A088860 | 001 | Jan 31, 1985 |
|--|----------|-------------|---------|-----|--------------|

CORPHED

| | | | | | |
|--|--------------|-------------|---------|-----|--------------|
| | FOSUN PHARMA | 60MG; 2.5MG | A088602 | 001 | Apr 11, 1985 |
|--|--------------|-------------|---------|-----|--------------|

PSEUDOEPHEDRINE HYDROCHLORIDE AND TRIPROLIDINE HYDROCHLORIDE

| | | | | | |
|--|--------|-------------|---------|-----|--------------|
| | SANDOZ | 60MG; 2.5MG | A088193 | 001 | May 17, 1983 |
|--|--------|-------------|---------|-----|--------------|

TRILITRON

| | | | | | |
|--|----------------|-------------|---------|-----|--------------|
| | NEWTRON PHARMS | 60MG; 2.5MG | A088515 | 001 | Jan 09, 1985 |
|--|----------------|-------------|---------|-----|--------------|

TRIPHED

| | | | | | |
|--|------|-------------|---------|-----|--------------|
| | TEVA | 60MG; 2.5MG | A088630 | 001 | May 17, 1984 |
|--|------|-------------|---------|-----|--------------|

TRIPROLIDINE AND PSEUDOEPHEDRINE

| | | | | | |
|--|-------------|-------------|---------|-----|--------------|
| | WATSON LABS | 60MG; 2.5MG | A088318 | 002 | Jan 13, 1984 |
|--|-------------|-------------|---------|-----|--------------|

| | | | | | |
|--|-----------|-------------|---------|-----|--------------|
| | WEST WARD | 60MG; 2.5MG | A088117 | 001 | Apr 19, 1983 |
|--|-----------|-------------|---------|-----|--------------|

TRIPROLIDINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

| | | | | | |
|--|----------------------|-------------|---------|-----|--------------|
| | IVAX SUB TEVA PHARMS | 60MG; 2.5MG | A085273 | 001 | Dec 12, 1984 |
|--|----------------------|-------------|---------|-----|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

TABLET; ORAL

TRIPROLIDINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE
SUPERPHARM 60MG; 2.5MG

A088578 001 Feb 21, 1985

TABLET, EXTENDED RELEASE; ORAL

TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES
KV PHARM 120MG; 5MG

A072758 001 Nov 25, 1991

PSEUDOEPHEDRINE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL

PSEUDO-12

UCB INC EQ 60MG HYDROCHLORIDE/5ML

N019401 001 Jun 19, 1987

PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

AFRINOL

+ SCHERING PLOUGH 120MG

N018191 001

PYRIDOSTIGMINE BROMIDE

TABLET; ORAL

PYRIDOSTIGMINE BROMIDE

ANI PHARMS INC 30MG

A040512 002 Jul 20, 2005

60MG

A040512 001 Oct 08, 2003

IMPAX LABS INC 60MG

A040457 001 Dec 26, 2002

SOLVAY 30MG

A089572 001 Nov 27, 1990

US ARMY 30MG

N020414 001 Feb 05, 2003

PYRIDOXINE HYDROCHLORIDE

INJECTABLE; INJECTION

HEXA-BETALIN

LILLY 100MG/ML

A080854 001

PYRIDOXINE HYDROCHLORIDE

AKORN 100MG/ML

A087967 001 Oct 01, 1982

BEL MAR 100MG/ML

A080761 001

DELL LABS 50MG/ML

A083771 001

100MG/ML

A083772 001

ELKINS SINN 100MG/ML

A080581 001

LUITPOLD 100MG/ML

A080669 001

MYLAN INSTITUTIONAL 100MG/ML

A204879 001 Jul 14, 2016

WATSON LABS 100MG/ML

A080572 001

100MG/ML

A083760 001

PYRILAMINE MALEATE

TABLET; ORAL

PYRILAMINE MALEATE

IMPAX LABS 25MG

A080808 001

WATSON LABS 25MG

A085231 001

PYRIMETHAMINE; SULFADOXINE

TABLET; ORAL

FANSIDAR

ROCHE 25MG; 500MG

N018557 001

PYRITHIONE ZINC

LOTION; TOPICAL

HEAD & SHOULDERS CONDITIONER

WARNER CHILCOTT 0.3%

N019412 002 Mar 10, 1986

PYRVINIUM PAMOATE

SUSPENSION; ORAL

POVAN

PARKE DAVIS EQ 50MG BASE/5ML

N011964 001

TABLET; ORAL

POVAN

PARKE DAVIS EQ 50MG BASE

N012485 002

QUAZEPAM

TABLET; ORAL

DORAL

GALT PHARMS 7.5MG

N018708 003 Feb 26, 1987

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

QUETIAPINE FUMARATE

TABLET; ORAL

QUETIAPINE FUMARATE

ACTAVIS GRP PTC

EQ 25MG BASE

A201762 001 Feb 27, 2013

EQ 50MG BASE

A201762 002 Feb 27, 2013

EQ 100MG BASE

A201762 003 Feb 27, 2013

EQ 150MG BASE

A201762 004 Feb 27, 2013

EQ 200MG BASE

A201762 005 Feb 27, 2013

EQ 300MG BASE

A201762 006 Feb 27, 2013

EQ 400MG BASE

A201762 007 Feb 27, 2013

MYLAN PHARMS INC

EQ 25MG BASE

A090323 001 Mar 27, 2012

SEROQUEL

+ ASTRAZENECA PHARMS

EQ 150MG BASE **

N020639 004 Dec 20, 1998

QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

QUINAPRIL HYDROCHLORIDE

ACTAVIS ELIZABETH

EQ 5MG BASE

A076459 001 Dec 22, 2004

EQ 10MG BASE

A076459 002 Dec 22, 2004

EQ 20MG BASE

A076459 003 Dec 22, 2004

EQ 40MG BASE

A076459 004 Dec 22, 2004

ACTAVIS LABS FL INC

EQ 5MG BASE

A076049 001 Jan 14, 2005

EQ 10MG BASE

A076049 002 Jan 14, 2005

EQ 20MG BASE

A076049 003 Jan 14, 2005

EQ 40MG BASE

A076049 004 Jan 14, 2005

APOTEX INC

EQ 5MG BASE

A076240 001 Jan 26, 2006

EQ 10MG BASE

A076240 002 Jan 26, 2006

EQ 20MG BASE

A076240 003 Jan 26, 2006

EQ 40MG BASE

A076240 004 Jan 26, 2006

MYLAN

EQ 5MG BASE

A076036 001 Jan 28, 2005

EQ 10MG BASE

A076036 002 Jan 28, 2005

EQ 20MG BASE

A076036 003 Jan 28, 2005

EQ 40MG BASE

A076036 004 Jan 28, 2005

SUN PHARM INDS LTD

EQ 5MG BASE

A076607 001 Dec 15, 2004

EQ 5MG BASE

A090800 001 Jun 18, 2009

EQ 10MG BASE

A076607 002 Dec 15, 2004

EQ 10MG BASE

A090800 002 Jun 18, 2009

EQ 20MG BASE

A076607 003 Dec 15, 2004

EQ 20MG BASE

A090800 003 Jun 18, 2009

EQ 40MG BASE

A076607 004 Dec 15, 2004

EQ 40MG BASE

A090800 004 Jun 18, 2009

YAOPHARMA CO LTD

EQ 5MG BASE

A076803 001 Mar 02, 2005

EQ 10MG BASE

A076803 002 Mar 02, 2005

EQ 20MG BASE

A076803 003 Mar 02, 2005

EQ 40MG BASE

A076803 004 Mar 02, 2005

QUINESTROL

TABLET; ORAL

ESTROVIS

PARKE DAVIS

0.1MG

N016768 002

0.2MG

N016768 003

QUINETHAZONE

TABLET; ORAL

HYDROMOX

LEDERLE

50MG

N013264 001

QUINETHAZONE; RESERPINE

TABLET; ORAL

HYDROMOX R

LEDERLE

50MG; 0.125MG

N013927 001

QUINIDINE GLUCONATE

INJECTABLE; INJECTION

QUINIDINE GLUCONATE

+ LILLY

80MG/ML

N007529 002 Feb 10, 1989

TABLET; ORAL

QUINACT

BAYER HLTHCARE

266MG

A085978 001

400MG

A086099 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

QUINIDINE GLUCONATE

TABLET, EXTENDED RELEASE;ORAL

DURAQUIN

WARNER CHILCOTT 330MG N017917 001

QUINAGLUTE

BAYER HLTHCARE 324MG N016647 001

QUINALAN

LANNETT 324MG A088081 001 Feb 10, 1986

QUINATIME

WATSON LABS 324MG A087448 001

QUINIDINE GLUCONATE

ASCOT 324MG A088582 001 Jun 17, 1985

AUROLIFE PHARMA LLC 324MG A089894 001 Dec 15, 1988

CYCLE PHARMS LTD 324MG A088431 001 Jan 06, 1984

HALSEY 324MG A089476 001 Apr 10, 1987

SUPERPHARM 324MG A089164 001 Nov 21, 1985

WATSON LABS 324MG A087785 001 Jan 24, 1983

324MG A087810 001 Sep 29, 1982

QUINIDINE POLYGALACTURONATE

TABLET;ORAL

CARDIOQUIN

PHARM RES ASSOC 275MG N011642 002

QUINIDINE SULFATE

CAPSULE;ORAL

CIN-QUIN

SOLVAY 200MG A085296 001

300MG A085297 001

QUINIDINE SULFATE

LILLY 200MG A085103 001

TABLET;ORAL

CIN-QUIN

SOLVAY 100MG A085299 001

200MG A084932 001

300MG A085298 001

QUINIDINE SULFATE

BARR 200MG A084177 001

CONTRACT PHARMACAL 200MG A083808 001

CYCLE PHARMS LTD 200MG A083640 001

300MG A085632 001

DAVA PHARMS INC 200MG A087011 001

ELKINS SINN 200MG A083622 001

EVERYLIFE 200MG A083439 001

HALSEY 200MG A083583 001

HIKMA PHARMS 200MG A083862 001

IMPAX LABS 200MG A083347 001

IVAX SUB TEVA PHARMS 200MG A084549 001

KING PHARMS 200MG A085175 001

KV PHARM 200MG A085276 001

LANNETT 200MG A083743 001

LEDERLE 200MG A086176 001

LILLY 200MG A085038 001

PERRIGO 200MG A085322 001

PHARMAVITE 200MG A084627 001

PUREPAC PHARM 200MG A084003 001

SANDOZ 200MG A084631 001

200MG A084914 001

300MG A089839 001 Sep 29, 1988

SCHERER LABS 200MG A085068 001

SUN PHARM INDUSTRIES 100MG A081029 001 Apr 14, 1989

SUPERPHARM 200MG A088973 001 Apr 10, 1985

USL PHARMA 200MG A087837 001 Apr 14, 1982

VALEANT PHARM INTL 200MG A083393 001

VANGARD 200MG A087909 001 Jul 13, 1982

VINTAGE PHARMS 200MG A083963 001

WARNER CHILCOTT 200MG A083879 001

WATSON LABS 100MG A085584 001

200MG A085140 002

WHITEWORTH TOWN PLSN 200MG A085444 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

QUINIDINE SULFATE

TABLET;ORAL

QUINORA

| | | | | |
|------------|-------|---------|-----|--|
| KEY PHARMS | 200MG | A083576 | 001 | |
| SCHERING | 300MG | A085222 | 001 | |

TABLET, EXTENDED RELEASE;ORAL

QUINIDEX

| | | | | |
|------------------|-------|---------|-----|--|
| WYETH PHARMS INC | 300MG | N012796 | 002 | |
|------------------|-------|---------|-----|--|

QUINIDINE SULFATE

| | | | | |
|------------------|-------|---------|-----|--------------|
| G AND W LABS INC | 300MG | A040045 | 001 | Jun 30, 1994 |
|------------------|-------|---------|-----|--------------|

RABEPRAZOLE SODIUM

TABLET, DELAYED RELEASE;ORAL

ACIPHEX

| | | | | |
|-------------|---------|---------|-----|--------------|
| + EISAI INC | 10MG ** | N020973 | 001 | May 29, 2002 |
|-------------|---------|---------|-----|--------------|

RAMIPRIL

CAPSULE;ORAL

RAMIPRIL

| | | | | |
|-------------------|--------|---------|-----|--------------|
| ACTAVIS ELIZABETH | 1.25MG | A077513 | 001 | Jun 18, 2008 |
| | 2.5MG | A077513 | 002 | Jun 18, 2008 |
| | 5MG | A077513 | 003 | Jun 18, 2008 |
| | 10MG | A077513 | 004 | Jun 18, 2008 |
| CIPLA | 1.25MG | A077004 | 001 | Aug 07, 2008 |
| | 2.5MG | A077004 | 002 | Aug 07, 2008 |
| | 5MG | A077004 | 003 | Aug 07, 2008 |
| RANBAXY LABS LTD | 10MG | A077004 | 004 | Aug 07, 2008 |
| | 5MG | A078849 | 001 | Mar 06, 2009 |
| | 10MG | A078849 | 002 | Mar 06, 2009 |
| WATSON LABS | 5MG | A076549 | 003 | Oct 24, 2005 |
| | 1.25MG | A077514 | 001 | Jun 18, 2008 |
| YAOPHARMA CO LTD | 2.5MG | A077514 | 002 | Jun 18, 2008 |
| | 5MG | A077514 | 003 | Jun 18, 2008 |
| | 10MG | A077514 | 004 | Jun 18, 2008 |

TABLET;ORAL

ALTACE

| | | | | |
|---------------|-----------|---------|-----|--------------|
| + KING PFIZER | 1.25MG ** | N022021 | 001 | Feb 27, 2007 |
| + | 2.5MG ** | N022021 | 002 | Feb 27, 2007 |
| + | 5MG ** | N022021 | 003 | Feb 27, 2007 |
| + | 10MG ** | N022021 | 004 | Feb 27, 2007 |

RAMIPRIL

| | | | | |
|----------------------|--------|---------|-----|--------------|
| APOTEX INC | 1.25MG | A091069 | 001 | Dec 02, 2015 |
| | 2.5MG | A091069 | 002 | Dec 02, 2015 |
| | 5MG | A091069 | 003 | Dec 02, 2015 |
| | 10MG | A091069 | 004 | Dec 02, 2015 |
| MYLAN PHARMS INC | 1.25MG | A090650 | 001 | Jun 30, 2011 |
| | 2.5MG | A090650 | 002 | Jun 30, 2011 |
| | 5MG | A090650 | 003 | Jun 30, 2011 |
| | 10MG | A090650 | 004 | Jun 30, 2011 |
| ZYDUS PHARMS USA INC | 1.25MG | A090697 | 001 | Sep 24, 2009 |
| | 2.5MG | A090697 | 002 | Sep 24, 2009 |
| | 5MG | A090697 | 003 | Sep 24, 2009 |
| | 10MG | A090697 | 004 | Sep 24, 2009 |

RANITIDINE BISMUTH CITRATE

TABLET;ORAL

TRITEC

| | | | | |
|-----------------|-------|---------|-----|--------------|
| GLAXOSMITHKLINE | 400MG | N020559 | 001 | Aug 08, 1996 |
|-----------------|-------|---------|-----|--------------|

RANITIDINE HYDROCHLORIDE

CAPSULE;ORAL

RANITIDINE HYDROCHLORIDE

| | | | | |
|-------|---------------|---------|-----|--------------|
| MYLAN | EQ 150MG BASE | A075564 | 001 | Oct 27, 2000 |
| | EQ 300MG BASE | A075564 | 002 | Oct 27, 2000 |
| TEVA | EQ 150MG BASE | A075557 | 001 | Oct 31, 2003 |
| | EQ 300MG BASE | A075557 | 002 | Oct 31, 2003 |

ZANTAC 150

| | | | | |
|-------------------|------------------|---------|-----|--------------|
| + GLAXOSMITHKLINE | EQ 150MG BASE ** | N020095 | 001 | Mar 08, 1994 |
|-------------------|------------------|---------|-----|--------------|

ZANTAC 300

| | | | | |
|-------------------|------------------|---------|-----|--------------|
| + GLAXOSMITHKLINE | EQ 300MG BASE ** | N020095 | 002 | Mar 08, 1994 |
|-------------------|------------------|---------|-----|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RANITIDINE HYDROCHLORIDE

GRANULE, EFFERVESCENT;ORAL

ZANTAC 150

GLAXO GRP LTD EQ 150MG BASE/PACKET N020251 002 Mar 31, 1994

INJECTABLE;INJECTION

RANITIDINE HYDROCHLORIDE

BEDFORD EQ 25MG BASE/ML A074764 001 Nov 19, 2004

ZANTAC IN PLASTIC CONTAINER

TELIGENT EQ 1MG BASE/ML N019593 002 Sep 27, 1991

EQ 50MG BASE/100ML N019593 001 Dec 17, 1986

SYRUP;ORAL

RANITIDINE HYDROCHLORIDE

APOTEX INC EQ 15MG BASE/ML A077602 001 Sep 17, 2007

RANBAXY EQ 15MG BASE/ML A078448 001 Dec 13, 2007

WOCKHARDT EQ 15MG BASE/ML A079211 001 May 26, 2009

EQ 15MG BASE/ML A079212 001 Feb 23, 2009

ZANTAC

+ GLAXO GRP LTD EQ 15MG BASE/ML N019675 001 Dec 30, 1988

TABLET;ORAL

RANITIDINE HYDROCHLORIDE

BOEHRINGER INGELHEIM EQ 150MG BASE A074662 001 Aug 29, 1997

EQ 300MG BASE A074662 002 Aug 29, 1997

CONTRACT PHARMACAL EQ 75MG BASE A075094 001 Jun 21, 1999

MYLAN EQ 150MG BASE A074023 001 Aug 22, 1997

EQ 150MG BASE A074552 001 Jul 30, 1998

EQ 300MG BASE A074023 002 Aug 22, 1997

EQ 300MG BASE A074552 002 Jul 30, 1998

RANBAXY EQ 75MG BASE A075254 001 Jan 14, 2000

EQ 150MG BASE A075000 001 Jan 30, 1998

EQ 300MG BASE A075000 002 Jan 30, 1998

SANDOZ EQ 75MG BASE A075519 001 Sep 26, 2002

SUN PHARM INDS LTD EQ 75MG BASE A075132 001 Jan 14, 2000

EQ 150MG BASE A075439 001 Apr 19, 2000

EQ 300MG BASE A075439 002 Apr 19, 2000

WATSON LABS EQ 75MG BASE A075212 001 Jan 14, 2000

EQ 150MG BASE A074864 001 Oct 20, 1997

EQ 300MG BASE A074864 002 Oct 20, 1997

WATSON LABS INC EQ 150MG BASE A077426 001 Dec 19, 2005

EQ 300MG BASE A077426 002 Dec 19, 2005

WOCKHARDT EQ 75MG BASE A078884 001 Jul 31, 2008

EQ 150MG BASE A078653 001 Nov 26, 2007

EQ 150MG BASE A078701 001 Nov 12, 2009

EQ 300MG BASE A078701 002 Dec 11, 2009

ZANTAC 150

+ GLAXO GRP LTD EQ 150MG BASE N018703 001 Jun 09, 1983

ZANTAC 300

+ GLAXO GRP LTD EQ 300MG BASE N018703 002 Dec 09, 1985

TABLET, EFFERVESCENT;ORAL

ZANTAC 150

GLAXO GRP LTD EQ 150MG BASE N020251 001 Mar 31, 1994

ZANTAC 25

GLAXO GRP LTD EQ 25MG BASE N020251 003 Apr 01, 2004

ZANTAC 75

+ SANOFI US EQ 75MG BASE ** N020745 001 Feb 26, 1998

RANOLAZINE

TABLET, EXTENDED RELEASE;ORAL

RANOLAZINE

LUPIN LTD 500MG A201046 001 Jul 29, 2013

1GM A201046 002 Jul 29, 2013

RAPACURONIUM BROMIDE

INJECTABLE;INJECTION

RAPLON

ORGANON USA INC 100MG/VIAL N020984 001 Aug 18, 1999

200MG/VIAL N020984 002 Aug 18, 1999

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RASAGILINE MESYLATE

TABLET; ORAL

RASAGILINE MESYLATE

APOTEX INC

EQ 0.5MG BASE

A201950 001 Sep 12, 2013

EQ 1MG BASE

A201950 002 Sep 12, 2013

RAUWOLFIA SERPENTINA ROOT

TABLET; ORAL

HIWOLFIA

BOWMAN PHARMS

50MG

N009276 003

50MG

N009276 005

100MG

N009276 004

HYSERPIN

PHYS PRODS VA

50MG

N010581 001

KOGLUCOID

PANRAY

50MG

N009278 001

100MG

N009278 002

RAUDIXIN

APOTHECON

50MG

N008842 001

100MG

N008842 002

RAUSERPIN

FERNDAL LABS

50MG

N009926 002

100MG

N009926 004

RAUVAL

PAL PAK

50MG

N009108 002

100MG

N009108 004

RAUWOLFIA SERPENTINA

BUNDY

50MG

N009477 001

100MG

N009477 002

HALSEY

50MG

A080498 001

100MG

A080498 002

IMPAX LABS

50MG

N009273 001

100MG

N009273 002

IVAX SUB TEVA PHARMS

50MG

N011521 001

100MG

N011521 002

PUREPAC PHARM

50MG

A080842 001

100MG

A080842 002

PVT FORM

50MG

A080583 001

100MG

A080583 002

SOLVAY

50MG

A080500 001

100MG

A080500 002

TABLICAPS

50MG

A083867 001

100MG

A083444 001

VALEANT PHARM INTL

50MG

N009668 001

100MG

N009668 002

WATSON LABS

50MG

A080907 001

100MG

A080914 001

WOLFINA

FOREST PHARMS

50MG

N009255 008

100MG

N009255 006

RESCINNAMINE

CAPSULE; ORAL

CINNASIL

PANRAY

0.5MG

A084736 001

TABLET; ORAL

MODERIL

PFIZER

0.25MG

N010686 003

0.5MG

N010686 006

RESERPINE

ELIXIR; ORAL

SERPASIL

NOVARTIS

0.2MG/4ML

N009115 005

INJECTABLE; INJECTION

SANDRIL

LILLY

2.5MG/ML

N010012 001

SERPASIL

NOVARTIS

2.5MG/ML

N009434 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RESERPINE

TABLET; ORAL

HISERPIA

| | | |
|---------------|--------|-------------|
| BOWMAN PHARMS | 0.1MG | N009631 002 |
| | 0.25MG | N009631 004 |

RAU-SED

| | | |
|----------------------|--------|-------------|
| BRISTOL MYERS SQUIBB | 0.1MG | N009357 001 |
| | 0.25MG | N009357 004 |
| | 0.5MG | N009357 006 |
| | 1MG | N009357 008 |

RESERPINE

| | | | |
|----------------------|--------|-------------|--------------|
| BARR | 0.25MG | A080721 002 | |
| BELL PHARMA | 0.1MG | A083058 001 | |
| | 0.25MG | A083058 002 | |
| BUNDY | 0.1MG | N009663 001 | |
| | 0.25MG | N009663 003 | |
| CYCLE PHARMS LTD | 0.1MG | N009859 001 | |
| | 0.25MG | N009859 002 | |
| ELKINS SINN | 0.1MG | A083145 001 | |
| | 0.25MG | A083145 002 | |
| EVERYLIFE | 0.1MG | N010441 001 | |
| | 0.25MG | N010441 002 | |
| | 0.5MG | N010441 003 | |
| | 1MG | N010441 004 | |
| HALSEY | 0.1MG | A080457 002 | |
| | 0.25MG | A080457 001 | |
| | 1MG | A080457 003 | |
| HIKMA INTL PHARMS | 0.1MG | A080975 001 | |
| | 0.25MG | A080975 002 | |
| | 1MG | A080975 003 | |
| IMPAX LABS | 0.1MG | N009627 001 | |
| | 0.25MG | N009627 002 | |
| IVAX SUB TEVA PHARMS | 0.1MG | N011185 001 | |
| | 0.25MG | N011185 002 | |
| MARSHALL PHARMA | 0.1MG | A080492 001 | |
| | 0.25MG | A080492 002 | |
| MK LABS | 0.1MG | A080525 002 | |
| | 0.25MG | A080525 001 | |
| MYLAN | 1MG | A084974 001 | |
| PHARMAVITE | 0.25MG | A084663 001 | |
| PUREPAC PHARM | 0.1MG | A080753 002 | |
| | 0.25MG | A080753 001 | |
| PVT FORM | 0.1MG | A086117 001 | |
| | 0.25MG | A080582 001 | |
| | 0.25MG | A085775 001 | |
| | 1MG | A080582 002 | |
| REXALL | 0.25MG | A080637 001 | |
| + SANDOZ | 0.1MG | N009838 001 | |
| + | 0.25MG | N009838 002 | |
| SOLVAY | 0.25MG | A080446 001 | |
| TABLICAPS | 0.25MG | A085207 001 | |
| TEVA | 0.1MG | A089020 001 | Mar 07, 1985 |
| | 0.25MG | A089019 001 | Mar 07, 1985 |
| VALEANT PHARM INTL | 0.1MG | N009667 001 | |
| | 0.25MG | N009667 002 | |
| WATSON LABS | 0.1MG | A080679 001 | |
| | 0.25MG | A080393 001 | |
| | 0.25MG | A085401 001 | |
| | 1MG | A080749 001 | |
| WHITEWORTH TOWN PLSN | 0.1MG | A080723 001 | |
| | 0.25MG | A080723 002 | |
| | 1MG | A080723 003 | |
| SANDRIL | | | |
| LILLY | 0.1MG | N009376 004 | |
| | 0.25MG | N009376 001 | |
| SERPALAN | | | |
| LANNETT | 0.1MG | N010124 001 | |
| | 0.25MG | N010124 002 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RESERPINE

TABLET; ORAL

SERPANRAY

| | | |
|--------|--------|-------------|
| PANRAY | 0.1MG | N009391 001 |
| | 0.25MG | N009391 002 |
| | 1MG | N009391 004 |

SERPASIL

| | | |
|----------|--------|-------------|
| NOVARTIS | 0.1MG | N009115 001 |
| | 0.25MG | N009115 003 |
| | 1MG | N009115 004 |

SERPATE

| | | |
|------|--------|-------------|
| VALE | 0.1MG | N009453 001 |
| | 0.25MG | N009453 002 |

SERPIVITE

| | | |
|----------|--------|-------------|
| VITARINE | 0.25MG | N009645 002 |
|----------|--------|-------------|

RESERPINE; TRICHLORMETHIAZIDE

TABLET; ORAL

METATENSIN #2

| | | |
|-------------------|------------|-------------|
| SANOFI AVENTIS US | 0.1MG; 2MG | N012972 001 |
|-------------------|------------|-------------|

METATENSIN #4

| | | |
|-------------------|------------|-------------|
| SANOFI AVENTIS US | 0.1MG; 4MG | N012972 002 |
|-------------------|------------|-------------|

NAQUIVAL

| | | |
|----------|------------|-------------|
| SCHERING | 0.1MG; 4MG | N012265 003 |
|----------|------------|-------------|

TRICHLORMETHIAZIDE W/ RESERPINE

| | | |
|-------------|------------|-------------|
| WATSON LABS | 0.1MG; 4MG | A085248 001 |
|-------------|------------|-------------|

RIBAVIRIN

CAPSULE; ORAL

REBETOL

| | | | |
|-------------------|----------------------------------------------------------------------------------------------------------------------------|-------------|--------------|
| MERCK SHARP DOHME | 200MG**Indicated for use and comarketed with Interferon ALFA-2B, Recombinant (INTRON A), as Rebetron Combination Therapy** | N020903 001 | Jun 03, 1998 |
|-------------------|----------------------------------------------------------------------------------------------------------------------------|-------------|--------------|

TABLET; ORAL

COPEGUS

| | | | |
|-------|-------|-------------|--------------|
| ROCHE | 200MG | N021511 001 | Dec 03, 2002 |
| | 400MG | N021511 002 | Jun 21, 2005 |

RIMANTADINE HYDROCHLORIDE

SYRUP; ORAL

FLUMADINE

| | | | |
|-------------|----------|-------------|--------------|
| FOREST LABS | 50MG/5ML | N019650 001 | Sep 17, 1993 |
|-------------|----------|-------------|--------------|

TABLET; ORAL

RIMANTADINE HYDROCHLORIDE

| | | | |
|-------------------|-------|-------------|--------------|
| ACTAVIS ELIZABETH | 100MG | A076375 001 | Jan 14, 2003 |
|-------------------|-------|-------------|--------------|

| | | | |
|----------------|-------|-------------|--------------|
| IMPAX LABS INC | 100MG | A075916 001 | Nov 02, 2001 |
|----------------|-------|-------------|--------------|

RIMEXOLONE

SUSPENSION/DROPS; OPHTHALMIC

VEXOL

| | | | |
|-------|----|-------------|--------------|
| ALCON | 1% | N020474 001 | Dec 30, 1994 |
|-------|----|-------------|--------------|

RISEDRONATE SODIUM

TABLET; ORAL

ACTONEL

+ APIL 75MG **

| | |
|-------------|--------------|
| N020835 004 | Apr 16, 2007 |
|-------------|--------------|

TABLET, DELAYED RELEASE; ORAL

RISEDRONATE SODIUM

| | | | |
|----------------|------|-------------|--------------|
| IMPAX LABS INC | 35MG | A205066 001 | Jun 29, 2018 |
|----------------|------|-------------|--------------|

RISPERIDONE

SOLUTION; ORAL

RISPERIDONE

| | | | |
|----------------|--------|-------------|--------------|
| LANNETT CO INC | 1MG/ML | A202386 001 | Jan 12, 2015 |
|----------------|--------|-------------|--------------|

| | | | |
|----------------|--------|-------------|--------------|
| PRECISION DOSE | 1MG/ML | A076797 001 | Jun 28, 2010 |
|----------------|--------|-------------|--------------|

| | | | |
|-----------|--------|-------------|--------------|
| WOCKHARDT | 1MG/ML | A078744 001 | Oct 08, 2009 |
|-----------|--------|-------------|--------------|

TABLET; ORAL

RISPERDAL

| | | | |
|----------------|-----|-------------|--------------|
| JANSSEN PHARMS | 5MG | N020272 005 | Dec 29, 1993 |
|----------------|-----|-------------|--------------|

RISPERIDONE

| | | | |
|------------------|--------|-------------|--------------|
| JUBILANT CADISTA | 0.25MG | A078828 001 | Mar 23, 2009 |
|------------------|--------|-------------|--------------|

| | | | |
|--|-------|-------------|--------------|
| | 0.5MG | A078828 002 | Mar 23, 2009 |
|--|-------|-------------|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RISPERIDONE

TABLET; ORAL

RISPERIDONE

| | | | |
|-------------------------------------|--------|-------------|--------------|
| | 1MG | A078828 003 | Mar 23, 2009 |
| | 2MG | A078828 004 | Mar 23, 2009 |
| | 3MG | A078828 005 | Mar 23, 2009 |
| | 4MG | A078828 006 | Mar 23, 2009 |
| RATIOPHARM | 0.25MG | A077784 001 | Jun 08, 2010 |
| | 0.5MG | A077784 002 | Jun 08, 2010 |
| | 1MG | A077784 003 | Jun 08, 2010 |
| | 2MG | A077784 004 | Jun 08, 2010 |
| | 3MG | A077784 005 | Jun 08, 2010 |
| | 4MG | A077784 006 | Jun 08, 2010 |
| SYNTHON PHARMS | 0.25MG | A078187 001 | Oct 22, 2009 |
| | 0.5MG | A078187 002 | Oct 22, 2009 |
| | 1MG | A078187 003 | Oct 22, 2009 |
| | 2MG | A078187 004 | Oct 22, 2009 |
| | 3MG | A078187 005 | Oct 22, 2009 |
| | 4MG | A078187 006 | Oct 22, 2009 |
| WATSON LABS | 0.25MG | A077860 001 | Dec 05, 2008 |
| | 0.5MG | A077860 002 | Dec 05, 2008 |
| | 1MG | A077860 003 | Dec 05, 2008 |
| | 2MG | A077860 004 | Dec 05, 2008 |
| | 3MG | A077860 005 | Dec 05, 2008 |
| | 4MG | A077860 006 | Dec 05, 2008 |
| WEST WARD PHARMS | 0.25MG | A078740 001 | May 29, 2009 |
| | 0.5MG | A078740 002 | May 29, 2009 |
| | 1MG | A078740 003 | May 29, 2009 |
| | 2MG | A078740 004 | May 29, 2009 |
| | 3MG | A078740 005 | May 29, 2009 |
| | 4MG | A078740 006 | May 29, 2009 |
| TABLET, ORALLY DISINTEGRATING; ORAL | | | |
| RISPERIDONE | | | |
| MYLAN PHARMS INC | 0.25MG | A091537 006 | Feb 12, 2013 |
| | 0.5MG | A091537 001 | Mar 30, 2011 |
| | 1MG | A091537 002 | Mar 30, 2011 |
| | 2MG | A091537 003 | Mar 30, 2011 |
| | 3MG | A091537 004 | Mar 30, 2011 |
| | 4MG | A091537 005 | Mar 30, 2011 |

RITODRINE HYDROCHLORIDE

INJECTABLE; INJECTION

RITODRINE HYDROCHLORIDE

| | | | |
|-------------------------------------------------------------|------------|-------------|--------------|
| ABRAXIS PHARM | 10MG/ML | A071188 001 | Jul 23, 1987 |
| | 15MG/ML | A071189 001 | Jul 23, 1987 |
| HOSPIRA | 10MG/ML | A071618 001 | Feb 28, 1991 |
| | 15MG/ML | A071619 001 | Feb 28, 1991 |
| RITODRINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER | | | |
| HOSPIRA | 30MG/100ML | A071438 001 | Jan 22, 1991 |
| YUTOPAR | | | |
| ASTRAZENECA | 10MG/ML | N018580 001 | |
| | 15MG/ML | N018580 002 | |
| TABLET; ORAL | | | |
| YUTOPAR | | | |
| ASTRAZENECA | 10MG | N018555 001 | |

RITONAVIR

CAPSULE; ORAL

NORVIR

| | | | |
|----------|-------|-------------|--------------|
| ABBOTT | 100MG | N020680 001 | Mar 01, 1996 |
| + ABBVIE | 100MG | N020945 001 | Jun 29, 1999 |

RIVASTIGMINE TARTRATE

SOLUTION; ORAL

EXELON

| | | | |
|----------|----------------|-------------|--------------|
| NOVARTIS | EQ 2MG BASE/ML | N021025 001 | Apr 21, 2000 |
|----------|----------------|-------------|--------------|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

RIZATRIPTAN BENZOATE

TABLET; ORAL

MAXALT

+ MERCK EQ 5MG BASE N020864 001 Jun 29, 1998

RIZATRIPTAN BENZOATE

APOTEX INC EQ 5MG BASE A202244 001 Dec 31, 2012

EQ 10MG BASE A202244 002 Dec 31, 2012

TABLET, ORALLY DISINTEGRATING; ORAL

MAXALT-MLT

+ MERCK EQ 5MG BASE N020865 001 Jun 29, 1998

RIZATRIPTAN BENZOATE

APOTEX INC EQ 5MG BASE A202477 001 Jul 01, 2013

EQ 10MG BASE A202477 002 Jul 01, 2013

ROCURONIUM BROMIDE

INJECTABLE; INJECTION

ZEMURON

+ ORGANON USA INC 50MG/5ML (10MG/ML) ** N020214 001 Mar 17, 1994

+ 10MG/ML (10MG/ML) ** N020214 002 Mar 17, 1994

+ 100MG/10ML (10MG/ML) ** N020214 003 Mar 17, 1994

ROFECOXIB

SUSPENSION; ORAL

VIOXX

MERCK 12.5MG/5ML N021052 001 May 20, 1999

25MG/5ML N021052 002 May 20, 1999

TABLET; ORAL

VIOXX

MERCK 12.5MG N021042 001 May 20, 1999

25MG N021042 002 May 20, 1999

50MG N021042 003 Feb 25, 2000

ROFLUMILAST

TABLET; ORAL

ROFLUMILAST

MYLAN PHARMS INC 500MCG A208257 001 Jul 13, 2018

ROLAPITANT HYDROCHLORIDE

EMULSION; INTRAVENOUS

VARUBI

+ TERSERA THERAPS LLC EQ 166.5MG BASE/92.5ML (EQ 1.8MG N208399 001 Oct 25, 2017

BASE/ML)

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

ROPINIROLE HYDROCHLORIDE

EPIC PHARMA LLC EQ 0.25MG BASE A078230 001 May 20, 2008

EQ 0.5MG BASE A078230 002 May 20, 2008

EQ 1MG BASE A078230 003 May 20, 2008

EQ 2MG BASE A078230 004 May 20, 2008

EQ 3MG BASE A078230 005 May 20, 2008

EQ 4MG BASE A078230 006 May 20, 2008

EQ 5MG BASE A078230 007 May 20, 2008

G AND W LABS INC EQ 0.25MG BASE A077460 001 May 05, 2008

EQ 0.5MG BASE A077460 002 May 05, 2008

EQ 1MG BASE A077460 003 May 05, 2008

EQ 2MG BASE A077460 004 May 05, 2008

EQ 3MG BASE A077460 005 May 05, 2008

EQ 4MG BASE A077460 006 May 05, 2008

EQ 5MG BASE A077460 007 May 19, 2008

TABLET, EXTENDED RELEASE; ORAL

REQUIP XL

+ GLAXOSMITHKLINE LLC EQ 3MG BASE ** N022008 002 Jun 13, 2008

ROPINIROLE HYDROCHLORIDE

MYLAN PHARMS INC EQ 2MG BASE A200462 001 Oct 15, 2012

EQ 3MG BASE A200462 002 Oct 15, 2012

EQ 4MG BASE A200462 003 Oct 15, 2012

EQ 6MG BASE A200462 004 Oct 15, 2012

EQ 8MG BASE A200462 005 Oct 15, 2012

EQ 12MG BASE A200462 006 Oct 15, 2012

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ROPIVACAINE HYDROCHLORIDE

SOLUTION; INJECTION

NAROPIN

| | | | |
|--------------------|----------------------|-------------|--------------|
| FRESENIUS KABI USA | 50MG/10ML (5MG/ML) | N020533 013 | May 01, 1998 |
| | 75MG/10ML (7.5MG/ML) | N020533 012 | Sep 24, 1996 |

ROSE BENGAL SODIUM I-131

INJECTABLE; INJECTION

ROBENGATOPE

| | | | |
|--------|-------------|-------------|--|
| BRACCO | 0.5mCi/VIAL | N016224 001 | |
| | 1mCi/VIAL | N016224 002 | |
| | 2mCi/VIAL | N016224 003 | |

SODIUM ROSE BENGAL I 131

| | | | |
|-------|-----------|-------------|--|
| SORIN | 0.5mCi/ML | N017318 001 | |
|-------|-----------|-------------|--|

ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDIA

+ SB PHARMCO

EQ 8MG BASE N021071 004 May 25, 1999

ROSIGLITAZONE MALEATE

TEVA

| | | |
|-------------|-------------|--------------|
| EQ 2MG BASE | A076747 001 | Jan 25, 2013 |
| EQ 4MG BASE | A076747 002 | Jan 25, 2013 |
| EQ 8MG BASE | A076747 003 | Jan 25, 2013 |

RUFINAMIDE

TABLET; ORAL

BANZEL

+ EISAI INC

100MG ** N021911 001 Nov 14, 2008

SAFFLOWER OIL

INJECTABLE; INJECTION

LIPOSYN 10%

ABBOTT

10% (10GM/100ML) N018203 001

LIPOSYN 20%

ABBOTT

20% (20GM/100ML) N018614 001

SAFFLOWER OIL; SOYBEAN OIL

INJECTABLE; INJECTION

LIPOSYN II 10%

HOSPIRA

5%;5% (5GM/100ML) N018997 001 Aug 27, 1984

LIPOSYN II 20%

HOSPIRA

10%;10% (10GM/100ML) N018991 001 Aug 27, 1984

SALMETEROL XINAFOATE

AEROSOL, METERED; INHALATION

SEREVENT

GLAXOSMITHKLINE

EQ 0.021MG BASE/INH N020236 001 Feb 04, 1994

SAQUINAVIR

CAPSULE; ORAL

FORTOVASE

+ HOFFMANN LA ROCHE

200MG ** N020828 001 Nov 07, 1997

SAQUINAVIR MESYLATE

CAPSULE; ORAL

INVIRASE

+ HOFFMANN LA ROCHE

EQ 200MG BASE N020628 001 Dec 06, 1995

SARALASIN ACETATE

INJECTABLE; INJECTION

SARENIN

PROCTER AND GAMBLE

EQ 0.6MG BASE/ML N018009 001

SECOBARBITAL SODIUM

CAPSULE; ORAL

SECOBARBITAL SODIUM

ANABOLIC

100MG A084422 001

BARR

100MG A084225 001

EVERYLIFE

100MG A085895 001

HALSEY

100MG A084676 001

IVAX PHARMS

100MG A085869 001

KV PHARM

100MG A085285 001

LANNETT

50MG A085909 001

100MG A085903 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SECOBARBITAL SODIUM

CAPSULE; ORAL

SECOBARBITAL SODIUM

| | | | |
|----------------------|-------|---------|-----|
| PARKE DAVIS | 100MG | A084762 | 001 |
| PERRIGO | 100MG | A084561 | 001 |
| PUREPAC PHARM | 100MG | A085867 | 001 |
| VALEANT PHARM INTL | 100MG | A085477 | 001 |
| VITARINE | 100MG | A085898 | 001 |
| | 100MG | A086273 | 001 |
| WATSON LABS | 100MG | A085792 | 001 |
| WEST WARD | 100MG | A084926 | 001 |
| WHITEWORTH TOWN PLSN | 100MG | A085798 | 001 |
| WYETH AYERST | 100MG | A086390 | 001 |

INJECTABLE; INJECTION

SECOBARBITAL SODIUM

| | | | |
|--------------|------------|---------|-----|
| ELKINS SINN | 100MG/VIAL | A083281 | 001 |
| WYETH AYERST | 50MG/ML | A083262 | 001 |

SECONAL SODIUM

| | | | |
|-------|---------|---------|-----|
| LILLY | 50MG/ML | N007392 | 002 |
|-------|---------|---------|-----|

SUPPOSITORY; RECTAL

SECONAL SODIUM

| | | | |
|-------|-------|---------|-----|
| LILLY | 30MG | A086530 | 001 |
| | 60MG | A086530 | 002 |
| | 120MG | A086530 | 003 |
| | 200MG | A086530 | 004 |

SECRETIN

INJECTABLE; INJECTION

SECRETIN-FERRING

| | | | |
|---------|-----------|---------|-----|
| FERRING | 75CU/VIAL | N018290 | 001 |
|---------|-----------|---------|-----|

SECRETIN SYNTHETIC PORCINE

FOR SOLUTION; INTRAVENOUS

SECFEFLO

| | | | | |
|------------|------------|---------|-----|--------------|
| CHIRHOCLIN | 16MCG/VIAL | N021136 | 001 | Apr 04, 2002 |
|------------|------------|---------|-----|--------------|

SELEGILINE HYDROCHLORIDE

CAPSULE; ORAL

ELDEPRYL

| | | | | |
|------------|-----|---------|-----|--------------|
| + SOMERSET | 5MG | N020647 | 001 | May 15, 1996 |
|------------|-----|---------|-----|--------------|

SELEGILINE HYDROCHLORIDE

| | | | | |
|----------------|-----|---------|-----|--------------|
| LANNETT CO INC | 5MG | A075145 | 001 | Sep 15, 2003 |
|----------------|-----|---------|-----|--------------|

TABLET; ORAL

SELEGILINE HYDROCHLORIDE

| | | | | |
|---------------------|--------|---------|-----|--------------|
| CHARTWELL MOLECULES | 5MG | A074565 | 001 | Aug 02, 1996 |
| | 5MG | A074641 | 001 | Aug 02, 1996 |
| G AND W LABS INC | 5MG | A074537 | 001 | Aug 02, 1996 |
| | 5MG | A074744 | 001 | Jan 27, 1997 |
| | 5MG | A074756 | 001 | Nov 25, 1998 |
| SIEGFRIED | 5MG | A074672 | 001 | Apr 01, 1997 |
| + SOMERSET | 5MG ** | N019334 | 001 | Jun 05, 1989 |

SELENIUM SULFIDE

LOTION/SHAMPOO; TOPICAL

EXSEL

| | | | |
|------------------|------|---------|-----|
| ALLERGAN HERBERT | 2.5% | A083892 | 001 |
|------------------|------|---------|-----|

SELENIUM SULFIDE

| | | | |
|----------------------|------|---------|-----|
| ACTAVIS MID ATLANTIC | 2.5% | A084394 | 001 |
| G AND W LABS INC | 2.5% | A086209 | 001 |
| IVAX PHARMS | 2.5% | A085777 | 001 |

SELSUN

| | | | |
|-----------|------|---------|-----|
| + CHATTEM | 2.5% | N007936 | 001 |
|-----------|------|---------|-----|

SELENOMETHIONINE SE-75

INJECTABLE; INJECTION

SELENOMETHIONINE SE 75

| | | | |
|---------------|-----------|---------|-----|
| GE HEALTHCARE | 250uCi/ML | N017257 | 001 |
| MALLINCKRODT | 100uCi/ML | N017098 | 001 |
| PHARMALUCENCE | 500uCi/ML | N017322 | 001 |

SETHOTOPE

| | | | |
|--------|--------------|---------|-----|
| BRACCO | 85-550uCi/ML | N017047 | 001 |
|--------|--------------|---------|-----|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SERMORELIN ACETATE

INJECTABLE; INJECTION

GEREF

| | | | | | |
|---|----------------|-----------------------|---------|-----|--------------|
| + | EMD SERONO | EQ 0.05MG BASE/AMP ** | N019863 | 001 | Dec 28, 1990 |
| + | EMD SERONO INC | EQ 0.5MG BASE/VIAL ** | N020443 | 001 | Sep 26, 1997 |
| + | | EQ 1MG BASE/VIAL ** | N020443 | 002 | Sep 26, 1997 |

SERTRALINE HYDROCHLORIDE

CONCENTRATE; ORAL

SERTRALINE HYDROCHLORIDE

| | | | | | |
|--|--------------------|-----------------|---------|-----|--------------|
| | ACI HEALTHCARE LTD | EQ 20MG BASE/ML | A076934 | 001 | Jun 30, 2006 |
| | RANBAXY LABS LTD | EQ 20MG BASE/ML | A078053 | 001 | Feb 05, 2007 |

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

| | | | | | |
|--|----------------------|---------------|---------|-----|--------------|
| | ACI HEALTHCARE LTD | EQ 25MG BASE | A076881 | 001 | Feb 06, 2007 |
| | | EQ 50MG BASE | A076881 | 002 | Feb 06, 2007 |
| | | EQ 100MG BASE | A076881 | 003 | Feb 06, 2007 |
| | ACTAVIS ELIZABETH | EQ 25MG BASE | A077345 | 001 | Feb 06, 2007 |
| | | EQ 50MG BASE | A077345 | 002 | Feb 06, 2007 |
| | | EQ 100MG BASE | A077345 | 003 | Feb 06, 2007 |
| | ANDA REPOSITORY | EQ 25MG BASE | A077818 | 001 | Feb 06, 2007 |
| | | EQ 50MG BASE | A077818 | 002 | Feb 06, 2007 |
| | | EQ 100MG BASE | A077818 | 003 | Feb 06, 2007 |
| | CHARTWELL MOLECULAR | EQ 25MG BASE | A077162 | 001 | Feb 06, 2007 |
| | | EQ 50MG BASE | A077162 | 002 | Feb 06, 2007 |
| | | EQ 100MG BASE | A077162 | 003 | Feb 06, 2007 |
| | FOSUN PHARMA | EQ 25MG BASE | A077713 | 001 | Feb 06, 2007 |
| | | EQ 50MG BASE | A077713 | 002 | Feb 06, 2007 |
| | | EQ 100MG BASE | A077713 | 003 | Feb 06, 2007 |
| | HIKMA PHARMS | EQ 25MG BASE | A077864 | 001 | Aug 10, 2009 |
| | | EQ 50MG BASE | A077864 | 002 | Aug 10, 2009 |
| | | EQ 100MG BASE | A077864 | 003 | Aug 10, 2009 |
| | IVAX SUB TEVA PHARMS | EQ 25MG BASE | A075719 | 003 | Jun 30, 2006 |
| | | EQ 50MG BASE | A075719 | 001 | Jun 30, 2006 |
| | | EQ 100MG BASE | A075719 | 002 | Jun 30, 2006 |
| | MYLAN | EQ 25MG BASE | A076671 | 001 | Feb 06, 2007 |
| | | EQ 50MG BASE | A076671 | 002 | Feb 06, 2007 |
| | | EQ 100MG BASE | A076671 | 003 | Feb 06, 2007 |
| | MYLAN PHARMS INC | EQ 25MG BASE | A076540 | 001 | Mar 20, 2007 |
| | | EQ 50MG BASE | A076540 | 002 | Mar 20, 2007 |
| | | EQ 100MG BASE | A076540 | 003 | Mar 20, 2007 |
| | | EQ 100MG BASE | A078626 | 003 | Jan 31, 2008 |
| | PLIVA HRVATSKA DOO | EQ 25MG BASE | A077299 | 001 | Feb 06, 2007 |
| | | EQ 50MG BASE | A077299 | 002 | Feb 06, 2007 |
| | | EQ 100MG BASE | A077299 | 003 | Feb 06, 2007 |
| | SCIEGEN PHARMS INC | EQ 25MG BASE | A076442 | 001 | Apr 30, 2007 |
| | | EQ 50MG BASE | A076442 | 002 | Apr 30, 2007 |
| | | EQ 100MG BASE | A076442 | 003 | Apr 30, 2007 |
| | SUN PHARM INDS (IN) | EQ 25MG BASE | A078108 | 001 | Feb 06, 2007 |
| | | EQ 50MG BASE | A078108 | 002 | Feb 06, 2007 |
| | | EQ 100MG BASE | A078108 | 003 | Feb 06, 2007 |
| | WATSON LABS TEVA | EQ 25MG BASE | A077663 | 001 | Feb 06, 2007 |
| | | EQ 50MG BASE | A077663 | 002 | Feb 06, 2007 |
| | | EQ 100MG BASE | A077663 | 003 | Feb 06, 2007 |

ZOLOFT

| | | | | | |
|---|--------|------------------|---------|-----|--------------|
| + | PFIZER | EQ 150MG BASE ** | N019839 | 003 | Dec 30, 1991 |
| + | | EQ 200MG BASE ** | N019839 | 004 | Dec 30, 1991 |

SEVELAMER HYDROCHLORIDE

CAPSULE; ORAL

RENAGEL

| | | | | | |
|--|---------|-------|---------|-----|--------------|
| | GENZYME | 403MG | N020926 | 001 | Oct 30, 1998 |
|--|---------|-------|---------|-----|--------------|

SIBUTRAMINE HYDROCHLORIDE

CAPSULE; ORAL

MERIDIA

| | | | | | |
|--|--------|------|---------|-----|--------------|
| | ABBOTT | 5MG | N020632 | 001 | Nov 22, 1997 |
| | | 10MG | N020632 | 002 | Nov 22, 1997 |
| | | 15MG | N020632 | 003 | Nov 22, 1997 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SILDENAFIL CITRATE

TABLET;ORAL

SILDENAFIL CITRATE

ACTAVIS GRP PTC

EQ 20MG BASE

A200149 001 Feb 25, 2013

APOTEX CORP

EQ 20MG BASE

A091379 001 Nov 06, 2012

SILVER SULFADIAZINE

CREAM;TOPICAL

SSD AF

DR REDDYS LA

1%

N018578 003 Jul 11, 1990

DRESSING;TOPICAL

SILDAFLO

FRANKLIN PHARMS

1%

N019608 001 Nov 30, 1989

SIMEPREVIR SODIUM

CAPSULE;ORAL

OLYSIO

+ JANSSEN PRODS

EQ 150MG BASE

N205123 001 Nov 22, 2013

SIMETHICONE-CELLULOSE

SUSPENSION;ORAL

SONORX

BRACCO

7.5MG/ML

N020773 001 Oct 29, 1998

SIMVASTATIN

TABLET;ORAL

SIMVASTATIN

HISUN PHARM HANGZHOU

10MG

A206557 001 Nov 13, 2017

20MG

A206557 002 Nov 13, 2017

40MG

A206557 003 Nov 13, 2017

80MG

A206557 004 Nov 13, 2017

IVAX SUB TEVA PHARMS

5MG

A076052 001 Jun 23, 2006

10MG

A076052 002 Jun 23, 2006

20MG

A076052 003 Jun 23, 2006

40MG

A076052 004 Jun 23, 2006

80MG

A076052 005 Dec 20, 2006

MYLAN PHARMS INC

5MG

A090868 001 Jun 08, 2010

10MG

A090868 002 Jun 08, 2010

20MG

A090868 003 Jun 08, 2010

40MG

A090868 004 Jun 08, 2010

80MG

A090868 005 Jun 08, 2010

SUN PHARM INDS LTD

5MG

A076285 001 Dec 20, 2006

10MG

A076285 002 Dec 20, 2006

20MG

A076285 003 Dec 20, 2006

40MG

A076285 004 Dec 20, 2006

80MG

A076285 005 Jun 23, 2006

YAOPHARMA CO LTD

5MG

A077766 001 Dec 20, 2006

10MG

A077766 002 Dec 20, 2006

20MG

A077766 003 Dec 20, 2006

40MG

A077766 004 Dec 20, 2006

80MG

A077766 005 Dec 20, 2006

TABLET, ORALLY DISINTEGRATING;ORAL

SIMVASTATIN

SYNTHON PHARMS

10MG

N021961 001 Oct 09, 2007

20MG

N021961 002 Oct 09, 2007

40MG

N021961 003 Oct 09, 2007

80MG

N021961 004 Oct 09, 2007

SIMVASTATIN; SITAGLIPTIN PHOSPHATE

TABLET;ORAL

JUVISYNC

+ MERCK SHARP DOHME

10MG;EQ 50MG BASE **

N202343 004 Sep 18, 2012

+

10MG;EQ 100MG BASE **

N202343 001 Oct 07, 2011

+

20MG;EQ 50MG BASE **

N202343 005 Sep 18, 2012

+

20MG;EQ 100MG BASE **

N202343 002 Oct 07, 2011

+

40MG;EQ 50MG BASE **

N202343 006 Sep 18, 2012

+

40MG;EQ 100MG BASE **

N202343 003 Oct 07, 2011

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SIROLIMUS

TABLET; ORAL

RAPAMUNE

+ PF PRISM CV

5MG **

N021110 003 Feb 23, 2004

SODIUM BENZOATE; SODIUM PHENYLACETATE

SOLUTION; ORAL

UCEPHAN

B BRAUN

100MG/ML; 100MG/ML

N019530 001 Dec 23, 1987

SODIUM BICARBONATE

INJECTABLE; INJECTION

SODIUM BICARBONATE

HOSPIRA INC

0.5MEQ/ML

A202679 001 Mar 07, 2017

0.5MEQ/ML

A202981 001 Mar 04, 2016

SODIUM BICARBONATE IN PLASTIC CONTAINER

+ ABBOTT

0.9MEQ/ML **

N019443 001 Jun 03, 1986

+

1MEQ/ML **

N019443 002 Jun 03, 1986

SODIUM BICARBONATE; TARTARIC ACID

GRANULE, EFFERVESCENT; ORAL

BAROS

MALLINCKRODT INC

460MG/GM; 420MG/GM

N018509 001 Aug 07, 1985

SODIUM CHLORIDE

INJECTABLE; INJECTION

BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

ABRAXIS PHARM

9MG/ML

A088909 001 Feb 07, 1985

SODIUM CHLORIDE

ABBOTT

20GM/100ML

N017013 001

B BRAUN

20GM/100ML

N017038 001

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN

450MG/100ML

N018184 001

MILES

450MG/100ML

N018503 001

SODIUM CHLORIDE 0.9%

+ MEDEFIL INC

18MG/2ML (9MG/ML)

N202832 002 Jan 06, 2012

+

22.5MG/2.5ML (9MG/ML)

N202832 003 Jan 06, 2012

+

27MG/3ML (9MG/ML)

N202832 004 Jan 06, 2012

+

45MG/5ML (9MG/ML)

N202832 005 Jan 06, 2012

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

ABBOTT

9MG/ML

N019218 001 Jul 13, 1984

+ ICU MEDICAL INC

9MG/ML

N019217 001 Jul 13, 1984

+ MEDEFIL INC

9MG/ML (9MG/ML)

N202832 001 Jan 06, 2012

MILES

900MG/100ML

N018502 001

SODIUM CHLORIDE 23.4% IN PLASTIC CONTAINER

+ ABRAXIS PHARM

234MG/ML **

N019329 001 Apr 22, 1987

SOLUTION; IRRIGATION

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

BAXTER HLTHCARE

450MG/100ML

N017864 001

450MG/100ML

N018497 001 Feb 19, 1982

HOSPIRA

450MG/100ML

N017670 001

450MG/100ML

N018380 001

SODIUM CHLORIDE IN PLASTIC CONTAINER

MILES

900MG/100ML

N018247 001

SODIUM CHROMATE CR-51

INJECTABLE; INJECTION

CHROMITOPE SODIUM

BRACCO

2mCi/VIAL

N013993 002

200uCi/ML

N013993 001

SODIUM CHROMATE CR 51

MALLINKRODT NUCLEAR

100uCi/ML

N016708 001

SODIUM FLUORIDE F-18

INJECTABLE; INTRAVENOUS

FLUORINE F-18

+ GE HEALTHCARE

2mCi/ML **

N017042 001

SODIUM FLUORIDE F 18

NIH NCI DCTD

10-200mCi/ML **

N022494 001 Jan 26, 2011

SODIUM FLUORIDE F-18

UIHC PET IMAGING

10-200mCi/ML

A204462 001 Nov 17, 2015

UNIV TX MD ANDERSON

10-200mCi/ML

A203247 001 Dec 23, 2013

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SODIUM IODIDE I-123

CAPSULE;ORAL

SODIUM IODIDE I 123

| | | | |
|---------------------|--------|-------------|--------------|
| CARDINAL HEALTH 418 | 400uCi | N018671 003 | May 27, 1982 |
| GE HEALTHCARE | 100uCi | N017630 001 | |

SOLUTION;ORAL

SODIUM IODIDE I 123

| | | | |
|---------------|------------|-------------|--|
| GE HEALTHCARE | 2mCi/ML ** | N017630 002 | |
|---------------|------------|-------------|--|

SODIUM IODIDE I-131

CAPSULE;ORAL

IODOTOPE

| | | | |
|--------|----------|-------------|--|
| BRACCO | 1-130mCi | N010929 001 | |
| | 1-150mCi | N010929 003 | |

SODIUM IODIDE I 131

| | | | |
|-----|--------|-------------|--|
| CIS | 50uCi | N017316 001 | |
| | 100uCi | N017316 002 | |

| | | | |
|--------------------|----------|-------------|--------------|
| JUBILANT DRAXIMAGE | 2-200mCi | N021305 004 | Nov 18, 2004 |
|--------------------|----------|-------------|--------------|

| | | | |
|---------------------|------------|-------------|--|
| MALLINKRODT NUCLEAR | 0.8-100mCi | N016515 002 | |
|---------------------|------------|-------------|--|

| | | | |
|---|------------|-------------|--|
| + | 0.8-100mCi | N016517 001 | |
|---|------------|-------------|--|

| | | | |
|--|-----------|-------------|--|
| | 15-100uCi | N016517 002 | |
|--|-----------|-------------|--|

SOLUTION;ORAL

HICON

| | | | |
|--------------------|-----------------|-------------|--------------|
| JUBILANT DRAXIMAGE | 1-250mCi/0.25ML | N021305 002 | Jan 24, 2003 |
| | 1-500mCi/0.5ML | N021305 003 | Jan 24, 2003 |
| | 1-1000mCi/ML | N021305 005 | Apr 04, 2006 |

IODOTOPE

| | | | |
|--------|--------------|-------------|--|
| BRACCO | 7-106mCi/BOT | N010929 002 | |
|--------|--------------|-------------|--|

SODIUM IODIDE I 131

| | | | |
|-----|----------|-------------|--|
| CIS | 50mCi/ML | N017315 001 | |
|-----|----------|-------------|--|

| | | | |
|---|---------------------|-----------------|-------------|
| + | MALLINKRODT NUCLEAR | 3.5-150mCi/VIAL | N016515 001 |
|---|---------------------|-----------------|-------------|

SODIUM LACTATE

INJECTABLE;INJECTION

SODIUM LACTATE 0.167 MOLAR IN PLASTIC CONTAINER

| | | | |
|---------|--------------|-------------|--|
| B BRAUN | 1.87GM/100ML | N018186 001 | |
|---------|--------------|-------------|--|

| | | | |
|-----------------|--------------|-------------|--|
| BAXTER HLTHCARE | 1.87GM/100ML | N016692 001 | |
|-----------------|--------------|-------------|--|

| | | | |
|---------|--------------|-------------|--|
| HOSPIRA | 1.87GM/100ML | N018249 001 | |
|---------|--------------|-------------|--|

SODIUM LACTATE 1/6 MOLAR IN PLASTIC CONTAINER

| | | | |
|---------|--------------|-------------|--------------|
| B BRAUN | 1.87GM/100ML | N020004 001 | Apr 21, 1992 |
|---------|--------------|-------------|--------------|

SODIUM MONOFLUOROPHOSPHATE

GEL;DENTAL

EXTRA-STRENGTH AIM

| | | | |
|-------------------|------|-------------|--------------|
| CHESEBROUGH PONDS | 1.2% | N019518 002 | Aug 06, 1986 |
|-------------------|------|-------------|--------------|

PASTE;DENTAL

EXTRA-STRENGTH AIM

| | | | |
|-------------------|------|-------------|--------------|
| CHESEBROUGH PONDS | 1.2% | N019518 001 | Jun 03, 1987 |
|-------------------|------|-------------|--------------|

SODIUM NITROPRUSSIDE

INJECTABLE;INJECTION

NIPRIDE

| | | | |
|-------|-----------|-------------|--|
| ROCHE | 50MG/VIAL | N017546 001 | |
|-------|-----------|-------------|--|

NITROPRESS

| | | | |
|--------|-----------|-------------|--------------|
| ABBOTT | 50MG/VIAL | A071555 001 | Nov 16, 1987 |
|--------|-----------|-------------|--------------|

| | | | |
|---|--------|--------------|-------------|
| + | ABBVIE | 50MG/VIAL ** | N018450 001 |
|---|--------|--------------|-------------|

| | | | |
|---------|-----------|-------------|--------------|
| HOSPIRA | 50MG/VIAL | A070566 001 | Jun 09, 1986 |
|---------|-----------|-------------|--------------|

SODIUM NITROPRUSSIDE

| | | | |
|---------------|-----------|-------------|--------------|
| ABRAXIS PHARM | 50MG/VIAL | A070031 001 | Jan 17, 1985 |
|---------------|-----------|-------------|--------------|

| | | | |
|---|-----------------|--------------|-------------|
| + | BAXTER HLTHCARE | 50MG/VIAL ** | N018581 001 |
|---|-----------------|--------------|-------------|

| | | | |
|-----------------|---------|-------------|--------------|
| TEVA PARENTERAL | 25MG/ML | A073465 001 | Mar 30, 1992 |
|-----------------|---------|-------------|--------------|

SODIUM PHOSPHATE P-32

SOLUTION;INJECTION, ORAL

PHOSPHOTOPE

| | | | |
|--------|-------------|-------------|--|
| BRACCO | 1-8mCi/VIAL | N010927 001 | |
|--------|-------------|-------------|--|

SODIUM PHOSPHATE P 32

| | | | |
|--------------|------------|-------------|--|
| MALLINCKRODT | 0.67mCi/ML | N011777 001 | |
|--------------|------------|-------------|--|

| | | | |
|--|-------------|-------------|--|
| | 1.5mCi/VIAL | N011777 002 | |
|--|-------------|-------------|--|

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SODIUM PHOSPHATE, DIBASIC ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL

VISICOL

SALIX PHARMS 0.398GM;1.102GM N021097 001 Sep 21, 2000

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL

MONOBASIC SODIUM PHOSPHATE AND DIBASIC SODIUM PHOSPHATE

NOVEL LABS INC 0.398GM;1.102GM A079247 001 Dec 30, 2011

SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL

KAYEXALATE

+ CONCORDIA PHARMS INC 453.6GM/BOT ** N011287 001

SODIUM POLYSTYRENE SULFONATE

CITRUSPHARMA 454GM/BOT A040909 001 Dec 03, 2008

WOCKHARDT 453.6GM/BOT A088786 001 Sep 11, 1984

SUSPENSION; ORAL, RECTAL

SODIUM POLYSTYRENE SULFONATE

MORTON GROVE 15GM/60ML A088717 001 Sep 11, 1984

ROXANE 15GM/60ML A088453 001 Nov 17, 1983

SODIUM SUCCINATE

INJECTABLE; INJECTION

SODIUM SUCCINATE

ELKINS SINN 30% A080516 001

SODIUM TETRADECYL SULFATE

INJECTABLE; INJECTION

SOTRADECOL

+ ELKINS SINN 1% ** N005970 004

+ 3% ** N005970 005

SODIUM THIOSULFATE

INJECTABLE; INJECTION

SODIUM THIOSULFATE

US ARMY 250MG/ML N020166 001 Feb 14, 1992

SOMATREM

INJECTABLE; INJECTION

PROTROPIN

GENENTECH 5MG/VIAL N019107 001 Oct 17, 1985

10MG/VIAL N019107 002 Oct 24, 1989

SOMATROPIN

INJECTABLE; INJECTION

ASELLACRIN 10

SERONO 10 IU/VIAL N017726 001

ASELLACRIN 2

SERONO 2 IU/VIAL N017726 002 Jul 21, 1983

BIO-TROPIN

FERRING 4.8MG/VIAL N019774 001 May 25, 1995

CRESCORMON

GENENTECH 4 IU/VIAL N017992 001

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION

ACCRETROPIN

EMERGENT 5MG/ML (5MG/ML) N021538 001 Jan 23, 2008

GENOTROPIN PRESERVATIVE FREE

+ PHARMACIA AND UPJOHN 1.5MG/VIAL N020280 004 Aug 24, 1995

HUMATROPE

LILLY 2MG/VIAL N019640 001 Jun 23, 1987

NORDITROPIN

NOVO NORDISK INC 4MG/VIAL N019721 001 May 08, 1995

5MG/1.5ML N021148 001 Jun 20, 2000

8MG/VIAL N019721 002 May 08, 1995

10MG/1.5ML N021148 002 Jun 20, 2000

15MG/1.5ML N021148 003 Jun 20, 2000

NORDITROPIN NORDIFLEX

NOVO NORDISK INC 5MG/1.5ML N021148 004 Oct 01, 2004

10MG/1.5ML N021148 005 Oct 01, 2004

15MG/1.5ML N021148 006 Oct 01, 2004

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION

NORDITROPIN NORDIFLEX

30MG/3ML

N021148 007 Mar 10, 2009

NUTROPIN

GENENTECH

5MG/VIAL

N020168 001 Nov 17, 1993

10MG/VIAL

N020168 002 Nov 17, 1993

NUTROPIN AQ

GENENTECH

10MG/2ML (5MG/ML)

N020522 001 Dec 29, 1995

NUTROPIN AQ PEN

+ GENENTECH

10MG/2ML (5MG/ML)

N020522 002 Apr 22, 2002

+ GENENTECH

20MG/2ML (10MG/ML)

N020522 006 Jan 03, 2008

NUTROPIN DEPOT

GENENTECH

13.5MG/VIAL

N021075 001 Dec 22, 1999

18MG/VIAL

N021075 002 Dec 22, 1999

22.5MG/VIAL

N021075 003 Dec 22, 1999

SAIZEN

EMD SERONO

4MG/VIAL

N019764 005 Jan 16, 2007

6MG/VIAL

N019764 001 Oct 08, 1996

SEROSTIM

EMD SERONO

8.8MG/VIAL

N020604 004 Sep 06, 2001

VALTROPIN

LG CHEM LTD

5MG/VIAL

N021905 001 Apr 19, 2007

ZORBITIVE

EMD SERONO

4MG/VIAL

N021597 001 Dec 01, 2003

5MG/VIAL

N021597 002 Dec 01, 2003

6MG/VIAL

N021597 003 Dec 01, 2003

INJECTABLE; SUBCUTANEOUS

SEROSTIM LQ

EMD SERONO

6MG/0.5ML (6MG/0.5ML)

N020604 005 Feb 11, 2005

SORBITOL

SOLUTION; IRRIGATION

SORBITOL 3% IN PLASTIC CONTAINER

BAXTER HLTHCARE

3GM/100ML

N018512 001 May 27, 1982

SOTALOL HYDROCHLORIDE

TABLET; ORAL

BETAPACE

COVIS PHARMA BV

320MG

N019865 004 Oct 30, 1992

BETAPACE AF

COVIS PHARMA BV

40MG

N021151 006 Apr 02, 2003

60MG

N021151 007 Apr 02, 2003

100MG

N021151 005 Mar 14, 2003

SOTALOL HYDROCHLORIDE

IMPAX PHARMS

80MG

A075663 001 Nov 07, 2000

120MG

A075663 002 Nov 07, 2000

160MG

A075663 003 Nov 07, 2000

240MG

A075663 004 Nov 07, 2000

MYLAN

80MG

A075237 001 May 01, 2000

80MG

A075725 001 Dec 19, 2000

120MG

A075237 002 May 01, 2000

120MG

A075725 002 Dec 19, 2000

160MG

A075237 003 May 01, 2000

160MG

A075725 003 Dec 19, 2000

240MG

A075237 004 May 01, 2000

240MG

A075725 004 Dec 19, 2000

SUN PHARM INDUSTRIES

80MG

A075515 001 Oct 15, 2001

80MG

A076576 001 Apr 08, 2004

120MG

A075515 004 Oct 15, 2001

120MG

A076576 002 Apr 08, 2004

160MG

A075515 002 Oct 15, 2001

160MG

A076576 003 Apr 08, 2004

240MG

A075515 003 Oct 15, 2001

TEVA

80MG

A076883 001 Jul 26, 2004

120MG

A076883 002 Jul 26, 2004

160MG

A076883 003 Jul 26, 2004

WATSON LABS

80MG

A075238 001 Jul 13, 2000

120MG

A075238 002 Jul 13, 2000

160MG

A075238 003 Jul 13, 2000

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SOTALOL HYDROCHLORIDE

TABLET; ORAL

SOTALOL HYDROCHLORIDE

240MG

A075238 004 Jul 13, 2000

SOYBEAN OIL

INJECTABLE; INJECTION

LIPOSYN III 10%

HOSPIRA

10%

N018969 001 Sep 24, 1984

LIPOSYN III 20%

HOSPIRA

20%

N018970 001 Sep 25, 1984

LIPOSYN III 30%

HOSPIRA

30%

N020181 001 Jan 13, 1998

SOYACAL 10%

ALPHA THERA

10%

N018465 001 Jun 29, 1983

SOYACAL 20%

ALPHA THERA

20%

N018786 001 Jun 29, 1983

TRAVAMULSION 10%

BAXTER HLTHCARE

10%

N018660 001 Feb 26, 1982

TRAVAMULSION 20%

BAXTER HLTHCARE

20%

N018758 001 Feb 15, 1983

SPARFLOXACIN

TABLET; ORAL

ZAGAM

MYLAN

200MG

N020677 001 Dec 19, 1996

SPECTINOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

TROBICIN

PFIZER

EQ 2GM BASE/VIAL

N050347 001

EQ 4GM BASE/VIAL

N050347 002

SPIRAPRIL HYDROCHLORIDE

TABLET; ORAL

RENORMAX

SCHERING

3MG

N020240 001 Dec 29, 1994

6MG

N020240 002 Dec 29, 1994

12MG

N020240 003 Dec 29, 1994

24MG

N020240 004 Dec 29, 1994

SPIRONOLACTONE

TABLET; ORAL

SPIRONOLACTONE

ASCOT

25MG

A087687 001 Oct 20, 1982

AUROBINDO PHARMA LTD

25MG

A202187 001 Mar 06, 2014

50MG

A202187 002 Mar 06, 2014

100MG

A202187 003 Mar 06, 2014

IVAX PHARMS

25MG

A087108 001

LEDERLE

25MG

A087634 001

MUTUAL PHARM

25MG

A087265 001

MYLAN

25MG

A087086 001

PUREPAC PHARM

25MG

A087998 001 Oct 14, 1983

25MG

A088053 001 Aug 25, 1983

SUPERPHARM

25MG

A089364 001 Nov 07, 1986

UPSHER SMITH

25MG

A087554 001

VANGARD

25MG

A087648 001 Feb 01, 1982

WARNER CHILCOTT

25MG

A087952 001 Nov 18, 1982

WATSON LABS

25MG

A086898 002 Mar 02, 1982

25MG

A087078 001

STANOZOLOL

TABLET; ORAL

WINSTROL

+ LUNDBECK INC

2MG

N012885 001 May 14, 1984

STAVUDINE

CAPSULE; ORAL

STAVUDINE

MYLAN LABS LTD

30MG

A078775 001 Jan 05, 2009

40MG

A078775 002 Jan 05, 2009

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

STAVUDINECAPSULE; ORAL
ZERIT

BRISTOL MYERS SQUIBB 5MG N020412 001 Jun 24, 1994

CAPSULE, EXTENDED RELEASE; ORAL
ZERIT XRBRISTOL MYERS SQUIBB 37.5MG N021453 001 Dec 31, 2002
50MG N021453 002 Dec 31, 2002
75MG N021453 003 Dec 31, 2002
100MG N021453 004 Dec 31, 2002FOR SOLUTION; ORAL
STAVUDINEAUROBINDO PHARMA 1MG/ML A077774 001 Dec 29, 2008
CIPLA LTD 1MG/ML A078030 001 Mar 20, 2009STERILE WATER FOR INJECTION

LIQUID; N/A

BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER

ABRAXIS PHARM 100% A089099 001 Dec 29, 1987
100% A089100 001 Dec 29, 1987

STERILE WATER FOR INJECTION IN PLASTIC CONTAINER

B BRAUN 100% N019077 001 Mar 02, 1984

STERILE WATER FOR IRRIGATION

LIQUID; IRRIGATION

STERILE WATER IN PLASTIC CONTAINER

MILES 100% N018246 001

STREPTOMYCIN SULFATE

INJECTABLE; INJECTION

STREPTOMYCIN SULFATE

COPANOS EQ 500MG BASE/ML A060684 001
LILLY EQ 1GM BASE/VIAL A060107 001
EQ 1GM BASE/2ML A060404 001
EQ 5GM BASE/VIAL A060107 002
PFIZER EQ 1GM BASE/VIAL ** A060076 001
EQ 1GM BASE/2.5ML A060111 001
EQ 5GM BASE/VIAL ** A060076 002SUCCINYLCHOLINE CHLORIDE

INJECTABLE; INJECTION

ANECTINE

SANDOZ INC 50MG/ML N008453 003
500MG/VIAL N008453 001
1GM/VIAL N008453 004

QUELICIN PRESERVATIVE FREE

+ HOSPIRA 20MG/ML N008845 001
50MG/ML N008845 002
100MG/ML N008845 004

SUCCINYLCHOLINE CHLORIDE

INTL MEDICATION 100MG/VIAL A085400 001 Feb 04, 1982
ORGANON USA INC 20MG/ML A080997 001

SUCOSTRIN

APOTHECON 20MG/ML N008847 001
100MG/ML N008847 003SUCRALFATE

TABLET; ORAL

SUCRALFATE

MYLAN IRELAND LTD 1GM A074415 001 Jun 08, 1998

SUFENTANIL CITRATE

INJECTABLE; INJECTION

SUFENTANIL CITRATE

WATSON LABS EQ 0.05MG BASE/ML A074406 001 Dec 15, 1995

SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC

BLEPH-10

ALLERGAN 10% A084015 001

CETAMIDE

ALCON 10% A080021 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFACETAMIDE SODIUM

OINTMENT;OPHTHALMIC

SODIUM SULAMYD

+ SCHERING 10% ** N005963 002

SULFAIR 10

PHARMAFAIR 10% A088000 001 Dec 22, 1982

SOLUTION/DROPS;OPHTHALMIC

BLEPH-30

ALLERGAN 30% A080028 002

ISOPTO CETAMIDE

ALCON 15% A080020 002

OCUSULF-10

MIZA PHARMS USA 10% A080660 001

OCUSULF-30

MIZA PHARMS USA 30% A080660 002

SODIUM SULAMYD

+ SCHERING 10% ** N005963 001

+ 30% ** N005963 003

SODIUM SULFACETAMIDE

AKORN 10% A083021 001

15% A083021 002

30% A083021 003

SOLA BARNES HIND 10% A084143 001

10% A084145 001

30% A084146 001

30% A084147 001

SULF-10

NOVARTIS 10% A080025 001

SULF-15

NOVARTIS 15% A089047 001 Oct 31, 1995

SULFACEL-15

OPTOPICS 15% A080024 001

SULFACETAMIDE SODIUM

AKORN 30% A040216 001 May 25, 1999

ALCON PHARMS LTD 30% A089068 001 May 05, 1987

PHARMAFAIR 10% A088947 001 May 17, 1985

SULFAIR 10

PHARMAFAIR 10% A087949 001 Dec 13, 1982

SULFAIR FORTE

PHARMAFAIR 30% A088385 001 Oct 13, 1983

SULFAIR-15

PHARMAFAIR 15% A088186 001 May 25, 1983

SULTEN-10

BAUSCH AND LOMB 10% A087818 001 Feb 03, 1983

SULFACYTINE

TABLET;ORAL

RENOQUID

GLENWOOD 250MG N017569 001

SULFADIAZINE

TABLET;ORAL

SULFADIAZINE

ABBVIE 300MG N004125 005

EVERYLIFE 500MG A080088 001

IMPAX LABS 500MG A080081 001

LANNETT 500MG A080084 001

LEDERLE 500MG N004054 001

+ LILLY 500MG N004122 002

SULFADIAZINE SODIUM

INJECTABLE; INJECTION

SULFADIAZINE SODIUM

LEDERLE 250MG/ML N004054 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFADIAZINE; SULFAMERAZINE

| | | | |
|---------------------|----------------------|--|-------------|
| SUSPENSION; ORAL | | | |
| SULFONAMIDES DUPLEX | | | |
| LILLY | 250MG/5ML; 250MG/5ML | | N006317 007 |

SULFAMETER

| | | | |
|----------------|-------|--|-------------|
| TABLET; ORAL | | | |
| SULLA | | | |
| BAYER HLTHCARE | 500MG | | N016000 002 |

SULFAMETHIZOLE

| | | | |
|---------------|-------|--|-------------|
| TABLET; ORAL | | | |
| MICROSUL | | | |
| FOREST PHARMS | 1GM | | A086012 001 |
| PROKLAR | | | |
| FOREST PHARMS | 500MG | | A080273 001 |
| THIOSULFIL | | | |
| WYETH AYERST | 250MG | | N008565 001 |
| | 500MG | | N008565 004 |

SULFAMETHOXAZOLE

| | | | |
|---------------------|-----------|--|--------------------------|
| SUSPENSION; ORAL | | | |
| GANTANOL | | | |
| ROCHE | 500MG/5ML | | N013664 002 |
| TABLET; ORAL | | | |
| GANTANOL | | | |
| ROCHE | 500MG | | N012715 002 |
| GANTANOL-DS | | | |
| ROCHE | 1GM | | N012715 003 |
| SULFAMETHOXAZOLE | | | |
| ASCOT | 500MG | | A087662 001 Oct 20, 1982 |
| AUROLIFE PHARMA LLC | 500MG | | A085844 001 |
| BARR | 500MG | | A087189 001 Jul 25, 1983 |
| HEATHER | 500MG | | A086163 001 |
| WATSON LABS | 500MG | | A085053 001 |
| | 1GM | | A086000 001 |
| UROBAK | | | |
| SHIONOGI | 500MG | | A087307 001 |

SULFAMETHOXAZOLE; TRIMETHOPRIM

| | | | |
|-----------------------------------|------------------------|--|--------------------------|
| INJECTABLE; INJECTION | | | |
| BACTRIM | | | |
| + SUN PHARM INDS INC | 80MG/ML; 16MG/ML ** | | N018374 001 |
| SEPTRA | | | |
| MONARCH PHARMS | 80MG/ML; 16MG/ML | | N018452 001 |
| SULFAMETHOXAZOLE AND TRIMETHOPRIM | | | |
| ABRAXIS PHARM | 80MG/ML; 16MG/ML | | A070223 001 Dec 29, 1987 |
| BEDFORD | 80MG/ML; 16MG/ML | | A072383 001 Apr 29, 1992 |
| HOSPIRA | 80MG/ML; 16MG/ML | | A073199 001 Sep 11, 1992 |
| WATSON LABS | 80MG/ML; 16MG/ML | | A071556 001 Dec 29, 1987 |
| WEST-WARD PHARMS INT | 80MG/ML; 16MG/ML | | A070627 001 Dec 29, 1987 |
| | 80MG/ML; 16MG/ML | | A070628 001 Dec 29, 1987 |
| SUSPENSION; ORAL | | | |
| BACTRIM | | | |
| + SUN PHARM INDUSTRIES | 200MG/5ML; 40MG/5ML ** | | N017560 001 |
| BACTRIM PEDIATRIC | | | |
| SUN PHARM INDUSTRIES | 200MG/5ML; 40MG/5ML ** | | N017560 002 |
| SEPTRA | | | |
| MONARCH PHARMS | 200MG/5ML; 40MG/5ML ** | | N017598 001 |
| SEPTRA GRAPE | | | |
| MONARCH PHARMS | 200MG/5ML; 40MG/5ML ** | | N017598 002 Feb 12, 1986 |
| SULFAMETHOXAZOLE AND TRIMETHOPRIM | | | |
| ANI PHARMS INC | 200MG/5ML; 40MG/5ML | | A070028 001 Jun 02, 1987 |
| | 200MG/5ML; 40MG/5ML ** | | A077612 001 Nov 13, 2006 |
| TEVA | 200MG/5ML; 40MG/5ML | | N018812 001 Jan 28, 1983 |
| | 200MG/5ML; 40MG/5ML | | N018812 002 Jun 10, 1983 |
| SULFATRIM | | | |
| PHARM ASSOC | 200MG/5ML; 40MG/5ML | | N018615 002 Jan 07, 1983 |
| SULMEPRIM | | | |
| USL PHARMA | 200MG/5ML; 40MG/5ML | | A070063 001 Aug 01, 1986 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFAMETHOXAZOLE; TRIMETHOPRIM

SUSPENSION; ORAL

SULMEPRIM PEDIATRIC

USL PHARMA

200MG/5ML; 40MG/5ML

A070064 001 Aug 01, 1986

TRIMETH/SULFA

ALPHARMA US PHARMS

200MG/5ML; 40MG/5ML

A072289 001 May 23, 1988

200MG/5ML; 40MG/5ML

A072398 001 May 23, 1988

NASKA

200MG/5ML; 40MG/5ML

A072399 001 May 23, 1988

TABLET; ORAL

COTRIM

TEVA

400MG; 80MG

A070034 001 May 16, 1985

COTRIM D.S.

TEVA

800MG; 160MG

A070048 001 Mar 18, 1985

SULFAMETHOPRIM

NOVEL LABS INC

400MG; 80MG

A070022 001 Feb 15, 1985

SULFAMETHOPRIM-DS

NOVEL LABS INC

800MG; 160MG

A070032 001 Feb 15, 1985

SULFAMETHOXAZOLE AND TRIMETHOPRIM

FOSUN PHARMA

400MG; 80MG

A070889 001 Nov 13, 1986

400MG; 80MG

N018598 003 May 19, 1982

800MG; 160MG

A070890 001 Nov 13, 1986

HEATHER

400MG; 80MG

N018946 001 Aug 10, 1984

800MG; 160MG

N018946 002 Aug 10, 1984

INTERPHARM

400MG; 80MG

A071299 001 Oct 27, 1987

800MG; 160MG

A071300 001 Oct 27, 1987

MARTEC USA LLC

400MG; 80MG

A072408 001 Dec 07, 1988

MUTUAL PHARM

400MG; 80MG

A070006 001 Nov 14, 1984

PLIVA

400MG; 80MG

A070215 001 Sep 10, 1985

800MG; 160MG

A070216 001 Sep 10, 1985

ROXANE

400MG; 80MG

A072768 001 Aug 30, 1991

TEVA

400MG; 80MG

N018242 001

800MG; 160MG

N018242 002

USL PHARMA

400MG; 80MG

A070203 001 Nov 08, 1985

800MG; 160MG

A070204 001 Nov 08, 1985

WATSON LABS

400MG; 80MG

A070002 001 Nov 07, 1984

400MG; 80MG

N018852 001 May 09, 1983

800MG; 160MG

A070000 001 Nov 07, 1984

SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH

FOSUN PHARMA

800MG; 160MG

N018598 004 May 19, 1982

MARTEC USA LLC

800MG; 160MG

A072417 001 Dec 07, 1988

MUTUAL PHARM

800MG; 160MG

A070007 001 Nov 14, 1984

ROXANE

800MG; 160MG

A072769 001 Aug 30, 1991

WATSON LABS

800MG; 160MG

N018854 001 May 09, 1983

SULFATRIM-DS

SUPERPHARM

800MG; 160MG

A070066 001 Jun 24, 1985

SULFATRIM-SS

SUPERPHARM

400MG; 80MG

A070065 002 Jun 24, 1985

UROPLUS DS

SHIONOGI

800MG; 160MG

A071816 001 Sep 28, 1987

UROPLUS SS

SHIONOGI

400MG; 80MG

A071815 001 Sep 28, 1987

SULFANILAMIDE

CREAM; VAGINAL

SULFANILAMIDE

G AND W LABS INC

15%

A088718 001 Sep 19, 1985

SUPPOSITORY; VAGINAL

AVC

MYLAN SPECIALITY LP

1.05GM

N006530 004 Jan 27, 1987

SULFAPHENAZOLE

SUSPENSION; ORAL

SULFABID

PHARM RES ASSOC

500MG/5ML

N013093 001

TABLET; ORAL

SULFABID

PURDUE FREDERICK

500MG

N013092 002

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFAPYRIDINE

TABLET; ORAL

SULFAPYRIDINE

LILLY

500MG

N000159 001

SULFASALAZINE

SUSPENSION; ORAL

AZULFIDINE

PHARMACIA AND UPJOHN

250MG/5ML

N018605 001

TABLET; ORAL

S.A.S.-500

SOLVAY

500MG

A083450 001

SULFASALAZINE

HERITAGE PHARMS INC

500MG

A080197 001

SANDOZ

500MG

A086184 001

SUN PHARM INDUSTRIES

500MG

A089590 001 Oct 19, 1987

SUPERPHARM

500MG

A089339 001 Oct 26, 1987

WATSON LABS

500MG

A084964 001

500MG

A087197 001

TABLET, DELAYED RELEASE; ORAL

SULFASALAZINE

WATSON LABS

500MG

A088052 001 May 24, 1983

SULFINPYRAZONE

CAPSULE; ORAL

ANTURANE

+ NOVARTIS

200MG **

N011556 004

SULFINPYRAZONE

BARR

200MG

A087666 001 Sep 17, 1982

IVAX PHARMS

200MG

A087770 001 Nov 19, 1982

PAR PHARM

200MG

A088934 001 Sep 06, 1985

VANGARD

200MG

A088666 001 Feb 17, 1984

TABLET; ORAL

ANTURANE

NOVARTIS

100MG **

N011556 003

SULFINPYRAZONE

BARR

100MG

A087665 001 Sep 17, 1982

IVAX PHARMS

100MG

A087769 001 Jun 01, 1982

PAR PHARM

100MG

A088933 001 Sep 06, 1985

WATSON LABS

100MG

A087667 001 May 26, 1982

SULFISOXAZOLE

TABLET; ORAL

GANTRISIN

ROCHE

500MG

N006525 001

SOSOL

MK LABS

500MG

A080036 001

SOXAZOLE

ALRA

500MG

A080366 001

SULFALAR

PARKE DAVIS

500MG

A084955 001

SULFISOXAZOLE

ANI PHARMS INC

500MG

A080142 001

AUROLIFE PHARMA LLC

500MG

A085628 001

BARR

500MG

A084031 001

HEATHER

500MG

A080189 001

IMPAX LABS

500MG

A080109 001

LANNETT

500MG

A080085 001

LEDERLE

500MG

A087649 001

PHARMERAL

500MG

A084385 001

PUREPAC PHARM

500MG

A080087 001

ROXANE

500MG

A080082 001

VALEANT PHARM INTL

500MG

A080268 002

VITARINE

500MG

A087332 001

WATSON LABS

500MG

A085534 001

WEST WARD

500MG

A080379 001

SULSOXIN

SOLVAY

500MG

A080040 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SULFISOXAZOLE ACETYL

| | | | |
|---------------------|-------------------|--|-------------|
| EMULSION; ORAL | | | |
| LIPO GANTRISIN | | | |
| ROCHE | EQ 1GM BASE/5ML | | N009182 009 |
| SUSPENSION; ORAL | | | |
| GANTRISIN PEDIATRIC | | | |
| ROCHE | EQ 500MG BASE/5ML | | N009182 004 |
| SYRUP; ORAL | | | |
| GANTRISIN | | | |
| ROCHE | EQ 500MG BASE/5ML | | N009182 002 |

SULFISOXAZOLE DIOLAMINE

| | | | |
|----------------------------|------------------|--|-------------|
| INJECTABLE; INJECTION | | | |
| GANTRISIN | | | |
| ROCHE | EQ 400MG BASE/ML | | N006917 001 |
| OINTMENT; OPHTHALMIC | | | |
| GANTRISIN | | | |
| ROCHE | EQ 4% BASE | | N008414 002 |
| SOLUTION/DROPS; OPHTHALMIC | | | |
| GANTRISIN | | | |
| ROCHE | EQ 4% BASE | | N007757 002 |
| SULFISOXAZOLE DIOLAMINE | | | |
| SOLA BARNES HIND | EQ 4% BASE | | A084148 001 |

SULFOXONE SODIUM

| | | | |
|-------------------------------|-------|--|-------------|
| TABLET, DELAYED RELEASE; ORAL | | | |
| DIASONE SODIUM | | | |
| ABBVIE | 165MG | | N006044 003 |

SULFUR

| | | | |
|-----------------|--------|--|-------------|
| POWDER; TOPICAL | | | |
| BENSULFOID | | | |
| POYTHRESS | 33.32% | | N002918 001 |

SULINDAC

| | | | |
|-----------------|----------|-------------|--------------|
| TABLET; ORAL | | | |
| CLINORIL | | | |
| + MERCK | 150MG ** | | N017911 001 |
| + | 200MG ** | | N017911 002 |
| SULINDAC | | | |
| ANI PHARMS INC | 150MG | A072972 001 | Feb 28, 1992 |
| | 200MG | A072973 001 | Feb 28, 1992 |
| EPIC PHARMA LLC | 150MG | A073262 002 | Sep 06, 1991 |
| | 200MG | A073262 001 | Sep 06, 1991 |
| FOSUN PHARMA | 150MG | A072712 001 | Aug 30, 1991 |
| | 200MG | A072713 001 | Aug 30, 1991 |

SUMATRIPTAN

| | | | |
|-----------------|------------|-------------|--------------|
| SPRAY; NASAL | | | |
| IMITREX | | | |
| GLAXOSMITHKLINE | 10MG/SPRAY | N020626 002 | Aug 26, 1997 |

SUMATRIPTAN SUCCINATE

| | | | |
|--------------------------|-------------------------------------|-------------|--------------|
| INJECTABLE; SUBCUTANEOUS | | | |
| ALSUMA | | | |
| MERIDIAN MEDCL | EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML) | N022377 001 | Jun 29, 2010 |
| SUMATRIPTAN SUCCINATE | | | |
| FRESENIUS KABI USA | EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML) | A079240 002 | Sep 18, 2009 |
| | EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML) | A079240 001 | Sep 18, 2009 |
| INJECTALIA | EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML) | A090310 001 | Aug 11, 2010 |
| SANDOZ INC | EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML) | A078067 002 | Feb 06, 2009 |
| | EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML) | A078067 001 | Feb 06, 2009 |
| TEVA PARENTERAL | EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML) | A078318 001 | Feb 06, 2009 |
| | EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML) | A078318 002 | Feb 06, 2009 |
| SUMAVEL DOSEPRO | | | |
| + ENDO VENTURES LTD | EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML) | N022239 002 | Nov 26, 2013 |
| + | EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML) | N022239 001 | Jul 15, 2009 |
| SYSTEM; IONTOPHORESIS | | | |
| ZECURITY | | | |
| + TEVA BRANDED PHARM | EQ 6.5MG BASE/4HR | N202278 001 | Jan 17, 2013 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

SUMATRIPTAN SUCCINATE

TABLET; ORAL

SUMATRIPTAN SUCCINATE

| | | | |
|--------------|---------------|-------------|--------------|
| FOSUN PHARMA | EQ 25MG BASE | A076976 001 | Aug 10, 2009 |
| | EQ 50MG BASE | A076976 002 | Aug 10, 2009 |
| | EQ 100MG BASE | A076976 003 | Aug 10, 2009 |
| HIKMA PHARMS | EQ 25MG BASE | A078298 001 | May 21, 2013 |
| | EQ 50MG BASE | A078298 002 | May 21, 2013 |
| | EQ 100MG BASE | A078298 003 | May 21, 2013 |
| MYLAN | EQ 25MG BASE | A077163 001 | Nov 02, 2009 |
| | EQ 50MG BASE | A077163 002 | Nov 02, 2009 |
| | EQ 100MG BASE | A077163 003 | Nov 02, 2009 |
| ROXANE | EQ 25MG BASE | A078241 001 | Aug 10, 2009 |
| | EQ 50MG BASE | A078241 002 | Aug 10, 2009 |
| | EQ 100MG BASE | A078241 003 | Aug 10, 2009 |
| TEVA | EQ 25MG BASE | A076840 001 | Feb 09, 2009 |
| | EQ 50MG BASE | A076840 002 | Feb 09, 2009 |
| | EQ 100MG BASE | A076840 003 | Feb 09, 2009 |

SUPROFEN

SOLUTION/DROPS; OPHTHALMIC

PROFENAL

| | | | |
|-------|----|-------------|--------------|
| ALCON | 1% | N019387 001 | Dec 23, 1988 |
|-------|----|-------------|--------------|

SUTILAINS

OINTMENT; TOPICAL

TRAVASE

| | | | |
|----------|--------------------|-------------|--|
| + ABBOTT | 82,000 UNITS/GM ** | N012828 001 | |
|----------|--------------------|-------------|--|

TACRINE HYDROCHLORIDE

CAPSULE; ORAL

COGNEX

| | | | |
|--------------|--------------|-------------|--------------|
| SHIONOGI INC | EQ 10MG BASE | N020070 001 | Sep 09, 1993 |
| | EQ 20MG BASE | N020070 002 | Sep 09, 1993 |
| | EQ 30MG BASE | N020070 003 | Sep 09, 1993 |
| | EQ 40MG BASE | N020070 004 | Sep 09, 1993 |

TACROLIMUS

CAPSULE; ORAL

TACROLIMUS

| | | | |
|-------------|-------------|-------------|--------------|
| WATSON LABS | EQ 5MG BASE | A090402 001 | Jul 01, 2010 |
|-------------|-------------|-------------|--------------|

TALBUTAL

TABLET; ORAL

LOTUSATE

| | | | |
|-------------------|-------|-------------|--|
| SANOFI AVENTIS US | 120MG | N009410 005 | |
|-------------------|-------|-------------|--|

TAMOXIFEN CITRATE

TABLET; ORAL

NOLVADEX

| | | | |
|---------------|-----------------|-------------|--------------|
| + ASTRAZENECA | EQ 10MG BASE ** | N017970 001 | |
| + | EQ 20MG BASE ** | N017970 002 | Mar 21, 1994 |

TAMOXIFEN CITRATE

| | | | |
|----------------------|--------------|-------------|--------------|
| ACTAVIS LABS FL INC | EQ 10MG BASE | A076179 001 | Feb 20, 2003 |
| | EQ 20MG BASE | A076179 002 | Feb 20, 2003 |
| AEGIS PHARMS | EQ 10MG BASE | A076398 001 | Mar 31, 2003 |
| | EQ 20MG BASE | A076398 002 | Mar 31, 2003 |
| IVAX SUB TEVA PHARMS | EQ 10MG BASE | A075740 001 | Feb 20, 2003 |
| | EQ 20MG BASE | A075740 002 | Feb 20, 2003 |
| PHARMACHEMIE | EQ 10MG BASE | A074539 001 | Mar 31, 2003 |
| ROXANE | EQ 10MG BASE | A076027 001 | Feb 20, 2003 |
| | EQ 20MG BASE | A076027 002 | Feb 20, 2003 |
| TEVA | EQ 10MG BASE | A074504 001 | Apr 28, 2003 |
| | EQ 20MG BASE | A074504 002 | Apr 28, 2003 |

TAPENTADOL HYDROCHLORIDE

SOLUTION; ORAL

NUCYNTA

| | | | |
|------------|-----------------|-------------|--------------|
| + ASSERTIO | EQ 20MG BASE/ML | N203794 001 | Oct 15, 2012 |
|------------|-----------------|-------------|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TECHNETIUM TC-99M ALBUMIN AGGREGATED

| | | | |
|-----------------------|---------|--|-------------|
| INJECTABLE; INJECTION | | | |
| TC 99M-LUNGAGGREGATE | | | |
| GE HEALTHCARE | 5mCi/ML | | N017848 001 |

TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

| | | | |
|---------------------------------|-----|--|-------------|
| INJECTABLE; INJECTION | | | |
| A-N STANNOUS AGGREGATED ALBUMIN | | | |
| SYNCOR PHARMS | N/A | | N017916 001 |
| AN-MAA | | | |
| PHARMALUCENCE | N/A | | N017792 001 |
| LUNGAGGREGATE REAGENT | | | |
| GE HEALTHCARE | N/A | | N017838 001 |
| MACROTEC | | | |
| BRACCO | N/A | | N017833 001 |
| PULMOLITE | | | |
| + JUBILANT DRAXIMAGE | N/A | | N017776 001 |
| TECHNESCAN MAA | | | |
| MALLINCKRODT | N/A | | N017842 001 |
| TECHNETIUM TC 99M MAA | | | |
| GE HEALTHCARE | N/A | | N017773 001 |

TECHNETIUM TC-99M ALBUMIN COLLOID KIT

| | | | |
|-----------------------|-----|--|--------------------------|
| INJECTABLE; INJECTION | | | |
| MICROLITE | | | |
| PHARMALUCENCE | N/A | | N018263 001 Mar 25, 1983 |

TECHNETIUM TC-99M ALBUMIN KIT

| | | | |
|-----------------------|-----|--|-------------|
| INJECTABLE; INJECTION | | | |
| TECHNETIUM TC 99M HSA | | | |
| GE HEALTHCARE | N/A | | N017775 001 |

TECHNETIUM TC-99M ALBUMIN MICROSPHERES KIT

| | | | |
|-----------------------|-----|--|-------------|
| INJECTABLE; INJECTION | | | |
| INSTANT MICROSPHERES | | | |
| 3M | N/A | | N017832 001 |

TECHNETIUM TC-99M APCITIDE

| | | | |
|-----------------------|-----|--|--------------------------|
| INJECTABLE; INJECTION | | | |
| ACUTECT | | | |
| CIS BIO INTL SA | N/A | | N020887 001 Sep 14, 1998 |

TECHNETIUM TC-99M DEPREOTIDE

| | | | |
|-----------------------|-----|--|--------------------------|
| INJECTABLE; INJECTION | | | |
| NEO TECT KIT | | | |
| CIS BIO INTL SA | N/A | | N021012 001 Aug 03, 1999 |

TECHNETIUM TC-99M ETIDRONATE KIT

| | | | |
|-----------------------------------------|-----|--|-------------|
| INJECTABLE; INJECTION | | | |
| CINTICHEM TECHNETIUM 99M HEDSPA | | | |
| GE HEALTHCARE | N/A | | N017653 001 |
| MPI STANNOUS DIPHOSPHONATE | | | |
| GE HEALTHCARE | N/A | | N017667 001 |
| OSTEOSCAN | | | |
| MALLINCKRODT | N/A | | N017454 001 |
| TECHNETIUM TC 99M DIPHOSPHONATE-TIN KIT | | | |
| GE HEALTHCARE | N/A | | N017562 001 |

TECHNETIUM TC-99M FERRENTEATE KIT

| | | | |
|-----------------------|-----|--|-------------|
| INJECTABLE; INJECTION | | | |
| RENOTEC | | | |
| BRACCO | N/A | | N017045 001 |

TECHNETIUM TC-99M GLUCEPTATE KIT

| | | | |
|-----------------------|-----|--|--------------------------|
| INJECTABLE; INJECTION | | | |
| GLUCOSCAN | | | |
| BRISTOL MYERS SQUIBB | N/A | | N017907 001 |
| TECHNESCAN GLUCEPTATE | | | |
| DRAXIMAGE | N/A | | N018272 001 Jan 27, 1982 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TECHNETIUM TC-99M LIDOFENIN KIT

INJECTABLE; INJECTION

TECHNESCAN HIDA

DRAXIMAGE

N/A

N018489 001 Oct 31, 1986

TECHNETIUM TC-99M MEDRONATE

INJECTABLE; INJECTION

DRAXIMAGE MDP-10

JUBILANT DRAXIMAGE

N/A

N018035 001

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION

AMERSCAN MDP KIT

GE HEALTHCARE

N/A

N018335 001 Aug 05, 1982

MDP-BRACCO

CARDINAL HEALTH 414

N/A

N018107 001

OSTEOLITE

PHARMALUCENCE

N/A

N017972 001

TECHNETIUM TC 99M MPI MDP

GE HEALTHCARE

N/A

N018141 001

N/A

N018141 002 Jun 12, 1989

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION

AN-DTPA

JUBILANT DRAXIMAGE

N/A

N017714 001

MPI DTPA KIT - CHELATE

GE HEALTHCARE

N/A

N017255 001

TECHNETIUM TC-99M PENTETATE KIT

GE HEALTHCARE

N/A

N017264 002

TECHNETIUM TC-99M POLYPHOSPHATE KIT

INJECTABLE; INJECTION

SODIUM POLYPHOSPHATE-TIN KIT

GE HEALTHCARE

N/A

N017664 001

TECHNETIUM TC-99M PYRO/TRIMETA PHOSPHATES KIT

INJECTABLE; INJECTION

PYROLITE

PHARMALUCENCE

N/A

N017684 001

TECHNETIUM TC-99M PYROPHOSPHATE KIT

INJECTABLE; INJECTION

PHOSPHOTEC

BRACCO

N/A

N017680 001

TECHNETIUM TC-99M RED BLOOD CELL KIT

INJECTABLE; INJECTION

RBC-SCAN

CADEMA

N/A

N020063 001 Jun 11, 1992

TECHNETIUM TC-99M SESTAMIBI KIT

INJECTABLE; INJECTION

MIRALUMA

LANTHEUS MEDCL

N/A

N019785 003 May 23, 1997

TECHNETIUM TC-99M SODIUM PERTECHNETATE

SOLUTION; INJECTION, ORAL

SODIUM PERTECHNETATE TC 99M

+ GE HEALTHCARE

2-100mCi/ML **

N017471 001

+ MALLINCKRODT

10-60mCi/ML **

N017725 001

PHARMALUCENCE

12mCi/ML

N017321 001

24mCi/ML

N017321 002

48mCi/ML

N017321 003

TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION; INJECTION, ORAL

MINITEC

BRACCO

0.22-2.22 CI/GENERATOR

N017339 001

SOLUTION; INTRAVENOUS

TECHNELITE

LANTHEUS MEDCL

0.0083-2.7 CI/GENERATOR

N017771 001

ULTRA-TECHNEKOW FM

MALLINCKRODT NUCLEAR

0.25-3 CI/GENERATOR

N017243 002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION;INTRAVENOUS, ORAL
 TECHNETIUM TC 99M GENERATOR
 GE HEALTHCARE 830-16600mCi/GENERATOR N017693 001

TECHNETIUM TC-99M SUCCIMER KIT

INJECTABLE;INJECTION
 MPI DMSA KIDNEY REAGENT
 GE HEALTHCARE N/A N017944 001 May 18, 1982

TECHNETIUM TC-99M SULFUR COLLOID

SOLUTION;INJECTION, ORAL
 TECHNETIUM TC 99M SULFUR COLLOID
 GE HEALTHCARE 4mCi/ML N017456 001

SOLUTION;ORAL
 TECHNETIUM TC 99M SULFUR COLLOID
 MALLINCKRODT 3mCi/ML N017724 001

TECHNETIUM TC-99M SULFUR COLLOID KIT

SOLUTION;INJECTION, ORAL
 TECHNOCOLL
 MALLINCKRODT N/A N017059 001
 TECHNETIUM TC 99M TSC
 GE HEALTHCARE N/A N017784 001
 TESULOID
 BRACCO N/A N016923 001

TECHNETIUM TC-99M TEBOROXIME KIT

INJECTABLE;INJECTION
 CARDIOTEC
 BRACCO N/A N019928 001 Dec 19, 1990

TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE;INJECTION
 MYOVUE
 + GE HEALTHCARE N/A N020372 001 Feb 09, 1996

TEGASEROD MALEATE

TABLET;ORAL
 ZELNORM
 US WORLDMEDS LLC EQ 2MG BASE N021200 001 Jul 24, 2002
 EQ 6MG BASE N021200 002 Jul 24, 2002

TELAPREVIR

TABLET;ORAL
 INCIVEK
 VERTEX PHARMS 375MG N201917 001 May 23, 2011

TELAVANCIN HYDROCHLORIDE

POWDER;INTRAVENOUS
 VIBATIV
 + CUMBERLAND PHARMS EQ 250MG BASE/VIAL N022110 001 Sep 11, 2009

TELBIVUDINE

SOLUTION;ORAL
 TYZEKA
 NOVARTIS 100MG/5ML N022154 001 Apr 28, 2009

TABLET;ORAL
 TYZEKA
 + NOVARTIS 600MG N022011 001 Oct 25, 2006

TELITHROMYCIN

TABLET;ORAL
 KETEK
 SANOFI AVENTIS US 300MG N021144 002 Feb 09, 2005
 400MG N021144 001 Apr 01, 2004

TELMISARTAN

TABLET;ORAL
 TELMISARTAN
 WATSON LABS 40MG A078710 002 Jan 08, 2014

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TEMAZEPAM

CAPSULE; ORAL

TEMAZ

| | | | | |
|------------------|------|---------|-----|--------------|
| QUANTUM PHARMICS | 15MG | A070564 | 001 | Oct 15, 1985 |
| | 30MG | A070547 | 001 | Oct 15, 1985 |

TEMAZEPAM

| | | | | |
|----------------------|------|---------|-----|--------------|
| DURAMED PHARMS BARR | 15MG | A071708 | 001 | Sep 29, 1988 |
| | 30MG | A071709 | 001 | Sep 29, 1988 |
| SUN PHARM INDUSTRIES | 15MG | A071174 | 001 | Jul 10, 1986 |
| | 30MG | A071175 | 001 | Jul 10, 1986 |
| USL PHARMA | 15MG | A070489 | 001 | Jul 07, 1986 |
| | 30MG | A070490 | 001 | Jul 07, 1986 |
| WATSON LABS | 15MG | A070383 | 001 | Mar 23, 1987 |
| | 15MG | A071446 | 001 | May 21, 1993 |
| | 30MG | A070384 | 001 | Mar 23, 1987 |
| | 30MG | A071447 | 001 | May 21, 1993 |

TEMOZOLOMIDE

CAPSULE; ORAL

TEMOZOLOMIDE

| | | | | |
|------------------|-------|---------|-----|--------------|
| MYLAN PHARMS INC | 5MG | A205227 | 001 | Jun 29, 2016 |
| | 20MG | A205227 | 002 | Jun 29, 2016 |
| | 100MG | A205227 | 003 | Jun 29, 2016 |
| | 140MG | A205227 | 004 | Jun 29, 2016 |
| | 180MG | A205227 | 005 | Jun 29, 2016 |
| | 250MG | A205227 | 006 | Jun 29, 2016 |

TENIPOSIDE

INJECTABLE; INJECTION

VUMON

| | | | | |
|--------------------|---------|---------|-----|--------------|
| + HQ SPECLT PHARMA | 10MG/ML | N020119 | 001 | Jul 14, 1992 |
|--------------------|---------|---------|-----|--------------|

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

HYTRIN

| | | | | |
|----------|-----------------|---------|-----|--------------|
| + ABBOTT | EQ 1MG BASE ** | N020347 | 001 | Dec 14, 1994 |
| | EQ 2MG BASE ** | N020347 | 002 | Dec 14, 1994 |
| | EQ 5MG BASE ** | N020347 | 003 | Dec 14, 1994 |
| | EQ 10MG BASE ** | N020347 | 004 | Dec 14, 1994 |

TERAZOSIN HYDROCHLORIDE

| | | | | |
|--------------------|--------------|---------|-----|--------------|
| MYLAN TECHNOLOGIES | EQ 1MG BASE | A075384 | 001 | Dec 01, 2000 |
| | EQ 2MG BASE | A075384 | 002 | Dec 01, 2000 |
| | EQ 5MG BASE | A075384 | 003 | Dec 01, 2000 |
| | EQ 10MG BASE | A075384 | 004 | Dec 01, 2000 |
| RANBAXY LABS LTD | EQ 1MG BASE | A076021 | 001 | Aug 22, 2002 |
| | EQ 2MG BASE | A076021 | 002 | Aug 22, 2002 |
| | EQ 5MG BASE | A076021 | 003 | Aug 22, 2002 |
| | EQ 10MG BASE | A076021 | 004 | Aug 22, 2002 |
| SANDOZ | EQ 1MG BASE | A075667 | 001 | Jul 28, 2000 |
| | EQ 2MG BASE | A075667 | 002 | Jul 28, 2000 |
| | EQ 5MG BASE | A075667 | 003 | Jul 28, 2000 |
| | EQ 10MG BASE | A075667 | 004 | Jul 28, 2000 |

TABLET; ORAL

HYTRIN

| | | | | |
|--------|--------------|---------|-----|--------------|
| ABBOTT | EQ 1MG BASE | N019057 | 001 | Aug 07, 1987 |
| | EQ 2MG BASE | N019057 | 002 | Aug 07, 1987 |
| | EQ 5MG BASE | N019057 | 003 | Aug 07, 1987 |
| | EQ 10MG BASE | N019057 | 004 | Aug 07, 1987 |

TERAZOSIN HYDROCHLORIDE

| | | | | |
|----------------------|--------------|---------|-----|--------------|
| IVAX SUB TEVA PHARMS | EQ 1MG BASE | A074530 | 001 | Apr 21, 2000 |
| | EQ 2MG BASE | A074530 | 002 | Apr 21, 2000 |
| | EQ 5MG BASE | A074530 | 003 | Apr 21, 2000 |
| | EQ 10MG BASE | A074530 | 004 | Apr 21, 2000 |
| SANDOZ | EQ 1MG BASE | A074315 | 001 | Dec 31, 1998 |
| | EQ 1MG BASE | A074657 | 001 | Apr 28, 2000 |
| | EQ 2MG BASE | A074315 | 002 | Dec 31, 1998 |
| | EQ 2MG BASE | A074657 | 002 | Apr 28, 2000 |
| | EQ 5MG BASE | A074315 | 003 | Dec 31, 1998 |
| | EQ 5MG BASE | A074657 | 003 | Apr 28, 2000 |
| | EQ 10MG BASE | A074315 | 004 | Dec 31, 1998 |
| | EQ 10MG BASE | A074657 | 004 | Apr 28, 2000 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TERAZOSIN HYDROCHLORIDE

TABLET; ORAL

TERAZOSIN HYDROCHLORIDE

TEVA

EQ 1MG BASE
EQ 2MG BASE
EQ 5MG BASE
EQ 10MG BASEA074446 001 May 18, 2000
A074446 002 May 18, 2000
A074446 003 May 18, 2000
A074446 004 May 18, 2000TERBINAFINE

GEL; TOPICAL

LAMISIL

GLAXOSMITHKLINE CONS 1%

N020846 001 Apr 29, 1998

TERBINAFINE HYDROCHLORIDE

CREAM; TOPICAL

LAMISIL

NOVARTIS

1%

N020192 001 Dec 30, 1992

GRANULE; ORAL

LAMISIL

+ NOVARTIS

EQ 125MG BASE/PACKET

N022071 001 Sep 28, 2007

+

EQ 187.5MG BASE/PACKET

N022071 002 Sep 28, 2007

SOLUTION; TOPICAL

LAMISIL

GLAXOSMITHKLINE CONS 1%

N020749 001 Oct 17, 1997

TABLET; ORAL

TERBINAFINE HYDROCHLORIDE

APOTEX

EQ 250MG BASE

A078199 001 Jul 02, 2007

GEDEON RICHTER USA

EQ 250MG BASE

A077065 001 Jul 02, 2007

MYLAN

EQ 250MG BASE

A077136 001 Jul 02, 2007

EQ 250MG BASE

A077195 001 Jul 02, 2007

ROXANE

EQ 250MG BASE

A077223 001 Jul 02, 2007

WOCKHARDT

EQ 250MG BASE

A078229 001 Jul 02, 2007

TERBUTALINE SULFATE

AEROSOL, METERED; INHALATION

BRETHAIRE

NOVARTIS

0.2MG/INH

N018762 001 Aug 17, 1984

BRICANYL

SANOFI AVENTIS US

0.2MG/INH

N018000 001 Mar 19, 1985

INJECTABLE; INJECTION

BRETHINE

+ PHARMACARE

1MG/ML **

N018571 001

BRICANYL

SANOFI AVENTIS US

1MG/ML

N017466 001

TERBUTALINE SULFATE

TEVA PHARMS USA

1MG/ML

A076853 001 Jul 20, 2004

TABLET; ORAL

BRICANYL

SANOFI AVENTIS US

2.5MG

N017618 001

5MG

N017618 002

TERCONAZOLE

CREAM; VAGINAL

TERAZOL 3

+ JANSSEN PHARMS

0.8%

N019964 001 Feb 21, 1991

TERAZOL 7

+ JANSSEN PHARMS

0.4%

N019579 001 Dec 31, 1987

SUPPOSITORY; VAGINAL

TERAZOL 3

+ JANSSEN PHARMS

80MG

N019641 001 May 24, 1988

TERCONAZOLE

FOUGERA PHARMS

80MG

A076850 001 Jul 12, 2006

TERIPARATIDE ACETATE

INJECTABLE; INJECTION

PARATHAR

SANOFI AVENTIS US

200 UNITS/VIAL

N019498 001 Dec 23, 1987

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TERIPARATIDE RECOMBINANT HUMAN

INJECTABLE; SUBCUTANEOUS

FORTEO

LILLY 0.75MG/3ML (0.25MG/ML) N021318 001 Nov 26, 2002

TESTOLACTONE

INJECTABLE; INJECTION

TESLAC

BRISTOL MYERS SQUIBB 100MG/ML N016119 001

TABLET; ORAL

TESLAC

BRISTOL MYERS SQUIBB 50MG N016118 001

250MG N016118 002

TESTOSTERONE

FILM, EXTENDED RELEASE; TRANSDERMAL

ANDRODERM

ALLERGAN SALES LLC 2.5MG/24HR N020489 001 Sep 29, 1995

5MG/24HR N020489 002 May 02, 1997

TESTODERM

ALZA 4MG/24HR N019762 001 Oct 12, 1993

6MG/24HR N019762 002 Oct 12, 1993

TESTODERM TTS

ALZA 5MG/24HR N020791 001 Dec 18, 1997

GEL; TRANSDERMAL

TESTOSTERONE

ANI PHARMS INC 25MG/2.5GM PACKET N202763 001 Feb 14, 2012

50MG/5GM PACKET N202763 002 Feb 14, 2012

PERRIGO ISRAEL 25MG/2.5GM PACKET N203098 002 Jan 31, 2013

50MG/5GM PACKET N203098 003 Jan 31, 2013

GEL, METERED; TRANSDERMAL

TESTOSTERONE

PERRIGO ISRAEL 12.5MG/1.25GM ACTUATION N203098 001 Jan 31, 2013

INJECTABLE; INJECTION

TESTOSTERONE

WATSON LABS 25MG/ML A086420 001 May 10, 1983

50MG/ML A086419 001 Aug 23, 1983

100MG/ML A086417 001 Jul 07, 1983

SOLUTION, METERED; TRANSDERMAL

AXIRON

+ ELI LILLY AND CO 30MG/1.5ML ACTUATION ** N022504 001 Nov 23, 2010

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

DEPO-TESTOSTERONE

PHARMACIA AND UPJOHN 50MG/ML A085635 001

TESTOSTERONE CYPIONATE

WATSON LABS 100MG/ML A084401 001

100MG/ML A086029 001

200MG/ML A084401 002

TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

DELATESTRYL

ENDO PHARMS 200MG/ML N009165 001

+ 200MG/ML N009165 003

TESTOSTERONE ENANTHATE

MYLAN INSTITUTIONAL 200MG/ML A040647 001 Oct 05, 2009

WATSON LABS 100MG/ML A083667 001

100MG/ML A085599 001

200MG/ML A083667 002

TESTOSTERONE PROPIONATE

INJECTABLE; INJECTION

TESTOSTERONE PROPIONATE

BEL MAR 25MG/ML A080741 001

50MG/ML A080742 001

100MG/ML A080743 001

ELKINS SINN 25MG/ML A080276 001

LILLY 50MG/ML A080254 002

WATSON LABS 25MG/ML A080188 001

25MG/ML A085490 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TESTOSTERONE PROPIONATE

INJECTABLE; INJECTION

TESTOSTERONE PROPIONATE

| | |
|----------|-------------|
| 50MG/ML | A080188 002 |
| 50MG/ML | A085490 002 |
| 100MG/ML | A080188 003 |
| 100MG/ML | A083595 003 |

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

BRISTACYCLINE

BRISTOL

| | |
|-------|-------------|
| 250MG | A061658 001 |
| 250MG | A061888 001 |
| 500MG | A061658 002 |
| 500MG | A061888 002 |

CYCLOPAR

WARNER CHILCOTT

| | |
|-------|-------------|
| 250MG | A061725 001 |
| 250MG | A062175 001 |
| 250MG | A062332 001 |
| 500MG | A061725 002 |
| 500MG | A062332 002 |

PANMYCIN

PHARMACIA AND UPJOHN

| | |
|-------|-------------|
| 250MG | A060347 001 |
|-------|-------------|

RETET

SOLVAY

| | |
|-------|-------------|
| 250MG | A061443 001 |
| 500MG | A061443 002 |

ROBITET

WYETH AYERST

| | |
|-------|-------------|
| 250MG | A061734 001 |
| 500MG | A061734 002 |

SUMYCIN

APOTHECON

| | |
|-------|-------------|
| 100MG | A060429 002 |
| 125MG | A060429 004 |
| 250MG | A060429 001 |
| 500MG | A060429 003 |

TETRACHEL

ANGUS

| | |
|-------|-------------|
| 250MG | A060343 001 |
| 500MG | A060343 003 |

TETRACYCLINE HYDROCHLORIDE

ABBOTT

| | |
|-------|-------------|
| 250MG | A061802 001 |
| 500MG | A061802 002 |

ANI PHARMS INC

| | |
|-------|-------------|
| 250MG | A061471 001 |
|-------|-------------|

ELKINS SINN

| | |
|-------|-------------|
| 250MG | A060059 001 |
|-------|-------------|

FERRANTE

| | |
|-------|-------------|
| 125MG | A060173 001 |
| 250MG | A060173 002 |

HEATHER

| | |
|-------|-------------|
| 250MG | A061148 001 |
| 500MG | A061148 002 |

HIKMA PHARMS

| | |
|-------|-------------|
| 250MG | A060768 001 |
| 500MG | A060768 002 |

IMPAX LABS

| | |
|-------|-------------|
| 100MG | A060469 002 |
| 250MG | A060469 001 |

IVAX SUB TEVA PHARMS

| | |
|-------|-------------|
| 500MG | A060469 003 |
| 250MG | A060704 001 |

MAST MM

| | |
|-------|-------------|
| 500MG | A060704 002 |
| 250MG | A062085 001 |

MYLAN

| | |
|-------|-------------|
| 250MG | A060783 001 |
| 500MG | A060783 002 |

PUREPAC PHARM

| | |
|-------|-------------|
| 250MG | A060290 001 |
| 500MG | A060290 002 |

PVT FORM

| | | |
|-------|-------------|--------------|
| 250MG | A062686 001 | Jul 24, 1986 |
| 500MG | A062686 002 | Jul 24, 1986 |

ROXANE

| | |
|-------|-------------|
| 500MG | A061214 002 |
|-------|-------------|

SUN PHARM INDUSTRIES

| | |
|-------|-------------|
| 250MG | A060736 001 |
| 500MG | A060736 002 |

SUPERPHARM

| | | |
|-------|-------------|--------------|
| 250MG | A062540 001 | Mar 21, 1985 |
| 500MG | A062540 002 | Mar 21, 1985 |

VALEANT PHARM INTL

| | |
|-------|-------------|
| 250MG | A060471 001 |
| 500MG | A060471 002 |

WARNER CHILCOTT

| | |
|-------|-------------|
| 250MG | A062300 001 |
| 500MG | A062300 002 |

WATSON LABS

| | |
|-------|-------------|
| 250MG | A062103 001 |
|-------|-------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TETRACYCLINE HYDROCHLORIDE

CAPSULE;ORAL

TETRACYCLINE HYDROCHLORIDE

| | |
|--------------------|-------------|
| 250MG | A062343 001 |
| 500MG | A062103 002 |
| 500MG | A062343 002 |
| WYETH AYERST 250MG | A061685 001 |
| 500MG | A061685 002 |

TETRACYN

| | |
|-------------------|-------------|
| PFIPHARMECS 250MG | A060082 003 |
| 500MG | A060082 004 |

FIBER, EXTENDED RELEASE;PERIODONTAL

ACTISITE

| | | |
|----------------------------|-------------|--------------|
| SCHIFF AND CO 12.7MG/FIBER | N050653 001 | Mar 25, 1994 |
|----------------------------|-------------|--------------|

FOR SOLUTION;TOPICAL

TOPICYCLINE

| | |
|----------------|-------------|
| SHIRE 2.2MG/ML | N050493 001 |
|----------------|-------------|

INJECTABLE;INJECTION

ACHROMYCIN

| | |
|--------------------|-------------|
| LEDERLE 250MG/VIAL | N050273 002 |
| 500MG/VIAL | N050273 003 |

TETRACYN

| | |
|-------------------|-------------|
| PFIZER 250MG/VIAL | A060096 001 |
| 500MG/VIAL | A060096 002 |

OINTMENT;OPHTHALMIC

ACHROMYCIN

| | |
|---------------|-------------|
| STORZ 10MG/GM | N050266 001 |
|---------------|-------------|

SUSPENSION;ORAL

ACHROMYCIN V

| | |
|-------------------|-------------|
| LEDERLE 125MG/5ML | N050263 002 |
|-------------------|-------------|

SUMYCIN

| | |
|---------------------|-------------|
| PAR PHARM 125MG/5ML | A060400 001 |
|---------------------|-------------|

TETRACYCLINE HYDROCHLORIDE

| | |
|------------------------------|-------------|
| ALPHARMA US PHARMS 125MG/5ML | A060633 001 |
| FERRANTE 125MG/5ML | A060174 001 |
| PROTER 125MG/5ML | A060446 001 |
| PUREPAC PHARM 125MG/5ML | A060291 001 |

TETRACYN

| | |
|-----------------------|-------------|
| PFIPHARMECS 125MG/5ML | A060095 001 |
|-----------------------|-------------|

TETRAMED

| | |
|--------------------------------|-------------|
| IVAX SUB TEVA PHARMS 125MG/5ML | A061468 001 |
|--------------------------------|-------------|

SUSPENSION/DROPS;OPHTHALMIC

ACHROMYCIN

| | |
|----------|-------------|
| STORZ 1% | N050268 001 |
|----------|-------------|

TABLET;ORAL

PANMYCIN

| | |
|----------------------------|-------------|
| PHARMACIA AND UPJOHN 250MG | A061705 001 |
| 500MG | A061705 002 |

SUMYCIN

| | |
|----------------|-------------|
| PAR PHARM 50MG | A061147 003 |
| 100MG | A061147 002 |
| 250MG | A061147 001 |
| 500MG | A061147 004 |

TETRACYCLINE PHOSPHATE COMPLEX

CAPSULE;ORAL

TETREX

| | |
|--------------------------------|-------------|
| BRISTOL EQ 100MG HYDROCHLORIDE | A061653 001 |
| EQ 250MG HYDROCHLORIDE | A061653 002 |
| EQ 250MG HYDROCHLORIDE | A061889 002 |
| EQ 250MG HYDROCHLORIDE | N050212 002 |
| EQ 500MG HYDROCHLORIDE | A061653 003 |
| EQ 500MG HYDROCHLORIDE | A061889 001 |
| EQ 500MG HYDROCHLORIDE | N050212 003 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THALLOUS CHLORIDE TL-201

INJECTABLE; INJECTION

THALLOUS CHLORIDE TL 201

BRACCO 1mCi/ML

N018548 001 Dec 30, 1982

TRACE LIFE 1mCi/ML

A075569 001 Nov 21, 2001

INJECTABLE; INTRAVENOUS

THALLOUS CHLORIDE TL 201

+ LANTHEUS MEDCL 2mCi/ML

N017806 002 Oct 09, 1998

MALLINKRODT NUCLEAR 2mCi/ML

A077698 001 Nov 09, 2006

THEOPHYLLINE

CAPSULE; ORAL

BRONKODYL

SANOFI AVENTIS US 100MG

A085264 001

200MG

A085264 002

ELIXOPHYLLIN

FOREST LABS 100MG

A085545 001 Jul 31, 1984

200MG

A083921 001 Jul 31, 1984

SOMOPHYLLIN-T

FISONS 100MG

A087155 001 Feb 25, 1985

200MG

A087155 002 Feb 25, 1985

250MG

A087155 003 Feb 25, 1985

THEOPHYLLINE

KV PHARM 100MG

A085263 001

200MG

A085263 002

SCHERER RP 100MG

A084731 002 Nov 07, 1986

200MG

A084731 001 Nov 07, 1986

250MG

A084731 003 Nov 07, 1986

CAPSULE, EXTENDED RELEASE; ORAL

AEROLATE III

FLEMING PHARMS 65MG

A085075 003 Nov 24, 1986

AEROLATE JR

FLEMING PHARMS 130MG

A085075 002 Nov 24, 1986

AEROLATE SR

FLEMING PHARMS 260MG

A085075 001 Nov 24, 1986

ELIXOPHYLLIN SR

FOREST LABS 125MG

A086826 001 Jan 29, 1985

250MG

A086826 002 Jan 29, 1985

SLO-BID

SANOFI AVENTIS US 50MG

A088269 001 Jan 31, 1985

75MG

A089539 001 May 10, 1989

100MG

A087892 001 Jan 31, 1985

125MG

A089540 001 May 10, 1989

200MG

A087893 001 Jan 31, 1985

300MG

A087894 001 Jan 31, 1985

SLO-PHYLLIN

SANOFI AVENTIS US 60MG

A085206 001 May 24, 1982

125MG

A085203 001 May 24, 1982

250MG

A085205 001 May 24, 1982

SOMOPHYLLIN-CRT

GRAHAM DM 50MG

A087763 001 Feb 27, 1985

100MG

A087194 001

200MG

A088382 001 Feb 27, 1985

250MG

A087193 001

300MG

A088383 001 Feb 27, 1985

THEO-DUR

SCHERING 50MG

A088022 001 Sep 10, 1985

75MG

A088015 001 Sep 10, 1985

125MG

A088016 001 Sep 10, 1985

200MG

A087995 001 Sep 10, 1985

THEOBID

WHITBY 260MG

A085983 001 Mar 20, 1985

THEOBID JR.

WHITBY 130MG

A087854 001 Mar 20, 1985

THEOCLEAR L.A.-130

SCHWARZ PHARMA 130MG

A086569 001 May 27, 1982

THEOCLEAR L.A.-260

SCHWARZ PHARMA 260MG

A086569 002 May 27, 1982

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THEOPHYLLINE

CAPSULE, EXTENDED RELEASE;ORAL

THEOPHYL-SR

| | | | | |
|--------------------|-------|---------|-----|--------------|
| ORTHO MCNEIL PHARM | 125MG | A086480 | 001 | Feb 08, 1985 |
| | 250MG | A086471 | 001 | Feb 08, 1985 |

THEOPHYLLINE

| | | | | |
|-------------|-------|---------|-----|--------------|
| CENT PHARMS | 125MG | A088654 | 001 | Feb 12, 1985 |
| | 250MG | A088689 | 001 | Feb 12, 1985 |
| HOSPIRA | 100MG | A089976 | 001 | Jan 04, 1995 |
| | 200MG | A089977 | 001 | Jan 04, 1995 |
| | 300MG | A089932 | 001 | Jan 04, 1995 |
| INWOOD LABS | 100MG | A040052 | 001 | Feb 14, 1994 |
| | 125MG | A040052 | 002 | Feb 14, 1994 |
| | 200MG | A040052 | 003 | Feb 14, 1994 |
| | 300MG | A040052 | 004 | Feb 14, 1994 |
| SANDOZ | 260MG | A087462 | 001 | May 11, 1982 |

THEOPHYLLINE-SR

| | | | | |
|------------|-------|---------|-----|--------------|
| SCHERER RP | 300MG | A088255 | 001 | Jun 12, 1986 |
|------------|-------|---------|-----|--------------|

THEOVENT

| | | | | |
|----------|-------|---------|-----|--------------|
| SCHERING | 125MG | A087010 | 001 | Jan 31, 1985 |
| | 250MG | A087910 | 001 | Jan 31, 1985 |

ELIXIR;ORAL

ELIXOMIN

| | | | | |
|-------|-----------|---------|-----|--------------|
| CENCI | 80MG/15ML | A088303 | 001 | Jan 25, 1984 |
|-------|-----------|---------|-----|--------------|

LANOPHYLLIN

| | | | | |
|---------|-----------|---------|-----|--|
| LANNETT | 80MG/15ML | A084578 | 001 | |
|---------|-----------|---------|-----|--|

THEOLIXIR

| | | | | |
|--------|-----------|---------|-----|--|
| PANRAY | 80MG/15ML | A084559 | 001 | |
|--------|-----------|---------|-----|--|

THEOPHYL-225

| | | | | |
|--------------------|--------------|---------|-----|--|
| ORTHO MCNEIL PHARM | 112.5MG/15ML | A086485 | 001 | |
|--------------------|--------------|---------|-----|--|

THEOPHYLLINE

| | | | | |
|--------------------|-----------|---------|-----|--------------|
| ALPHARMA US PHARMS | 80MG/15ML | A089223 | 001 | May 27, 1988 |
| CENCI | 80MG/15ML | A087679 | 001 | Apr 15, 1982 |
| CHARTWELL RX | 80MG/15ML | A085952 | 001 | |
| HALSEY | 80MG/15ML | A085169 | 001 | |
| PHARM ASSOC | 80MG/15ML | A086720 | 001 | |
| PRECISION DOSE | 80MG/15ML | A085863 | 001 | |
| ROXANE | 80MG/15ML | A084739 | 001 | |
| TARO | 80MG/15ML | A089626 | 001 | Oct 28, 1988 |
| WOCKHARDT | 80MG/15ML | A086748 | 001 | |

INJECTABLE;INJECTION

THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|---------|------------|---------|-----|--------------|
| B BRAUN | 40MG/100ML | N019083 | 001 | Nov 07, 1984 |
|---------|------------|---------|-----|--------------|

THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|---------|------------|---------|-----|--------------|
| B BRAUN | 80MG/100ML | N019083 | 002 | Nov 07, 1984 |
|---------|------------|---------|-----|--------------|

THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|---------|-------------|---------|-----|--------------|
| B BRAUN | 160MG/100ML | N019083 | 003 | Nov 07, 1984 |
|---------|-------------|---------|-----|--------------|

THEOPHYLLINE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|---------|-------------|---------|-----|--------------|
| B BRAUN | 200MG/100ML | N019212 | 001 | Nov 07, 1984 |
| | 200MG/100ML | N019826 | 004 | Aug 14, 1992 |

THEOPHYLLINE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|---------|-------------|---------|-----|--------------|
| B BRAUN | 4MG/ML | N019212 | 003 | Nov 07, 1984 |
| | 400MG/100ML | N019212 | 002 | Nov 07, 1984 |
| | 400MG/100ML | N019826 | 005 | Aug 14, 1992 |

THEOPHYLLINE AND DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | |
|-----------------|-------------|---------|-----|--------------|
| BAXTER HLTHCARE | 4MG/ML | N018649 | 007 | Jul 26, 1982 |
| | 40MG/100ML | N018649 | 001 | Jul 26, 1982 |
| | 80MG/100ML | N018649 | 002 | Jul 26, 1982 |
| | 160MG/100ML | N018649 | 003 | Jul 26, 1982 |
| | 200MG/100ML | N018649 | 004 | Jul 26, 1982 |
| | 320MG/100ML | N018649 | 006 | Nov 13, 1985 |
| | 400MG/100ML | N018649 | 005 | Jul 26, 1982 |

THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER

| | | | | | |
|---|-------------|-------------|---------|-----|--------------|
| + | HOSPIRA INC | 4MG/ML | N019211 | 007 | Dec 14, 1984 |
| + | | 40MG/100ML | N019211 | 001 | Dec 14, 1984 |
| | | 80MG/100ML | N019211 | 002 | Dec 14, 1984 |
| + | | 160MG/100ML | N019211 | 003 | Dec 14, 1984 |
| | | 200MG/100ML | N019211 | 004 | Dec 14, 1984 |
| + | | 320MG/100ML | N019211 | 006 | Jan 20, 1988 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THEOPHYLLINE

INJECTABLE; INJECTION

THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER
400MG/100ML

N019211 005 Dec 14, 1984

SOLUTION; ORAL

AEROLATE

FLEMING PHARMS

150MG/15ML

A089141 001 Dec 03, 1986

THEOLAIR

3M

80MG/15ML

A086107 001

THEOPHYLLINE

ROXANE

80MG/15ML

A087449 001 Sep 15, 1983

SUSPENSION; ORAL

ELIXICON

FOREST LABS

100MG/5ML

A085502 001

SYRUP; ORAL

ACCURBRON

SANOFI AVENTIS US

150MG/15ML

A088746 001 Nov 22, 1985

AQUAPHYLLIN

FERNDAL LABS

80MG/15ML

A087917 001 Jan 18, 1983

SLO-PHYLLIN

SANOFI AVENTIS US

80MG/15ML

A085187 001

THEOCLEAR-80

CENT PHARMS

80MG/15ML

A087095 001 Mar 01, 1982

THEOPHYLLINE

ALPHARMA US PHARMS

80MG/15ML

A086001 001

150MG/15ML

A086545 001

TABLET; ORAL

QUIBRON-T

MONARCH PHARMS

300MG

A088656 001 Aug 22, 1985

SLO-PHYLLIN

SANOFI AVENTIS US

100MG

A085202 001

200MG

A085204 001

THEOCLEAR-100

CENT PHARMS

100MG

A085353 002

THEOCLEAR-200

CENT PHARMS

200MG

A085353 001

THEOLAIR

MEDICIS

125MG

A086399 001

250MG

A086399 002

THEOPHYL-225

ORTHO MCNEIL PHARM

225MG

A084726 001

TABLET, CHEWABLE; ORAL

THEOPHYL

ORTHO MCNEIL PHARM

100MG

A086506 001 Sep 12, 1985

TABLET, EXTENDED RELEASE; ORAL

DURAPHYL

FOREST LABS

100MG

A088503 001 Apr 03, 1985

200MG

A088504 001 Apr 03, 1985

300MG

A088505 001 Apr 03, 1985

LABID

WARNER CHILCOTT

250MG

A087225 001

QUIBRON-T/SR

MONARCH PHARMS

300MG

A087563 001 Jun 21, 1983

SUSTAIRE

ROERIG

100MG

A085665 001

300MG

A085665 002

T-PHYL

PHARM RES ASSOC

200MG

A088253 001 Aug 17, 1983

THEO-DUR

SCHERING

100MG

A085328 001

200MG

A086998 001

300MG

A085328 002

450MG

A089131 001 Jun 25, 1986

THEOCHRON

NOSTRUM PHARMS LLC

300MG

A087400 002 Jan 11, 1983

THEOLAIR-SR

3M

200MG

A088369 001 Jul 16, 1987

250MG

A086363 002 Jul 16, 1987

300MG

A088364 001 Jul 16, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THEOPHYLLINE

TABLET, EXTENDED RELEASE;ORAL

THEOLAIR-SR

500MG

A089132 001 Jul 16, 1987

THEOPHYLLINE

ABLE

300MG

A040548 001 Apr 30, 2004

400MG

A040543 001 Apr 27, 2004

450MG

A040546 001 Apr 30, 2004

600MG

A040539 001 Apr 27, 2004

INWOOD LABS

450MG

A040034 001 Apr 28, 1995

TEVA PHARMS

450MG

A081236 001 Nov 09, 1992

UNI-DUR

SCHERING

400MG

A089822 001 Jan 04, 1995

600MG

A089823 001 Jan 04, 1995

THEOPHYLLINE SODIUM GLYCINATE

ELIXIR;ORAL

SYNOHYLATE

CENT PHARMS

EQ 165MG BASE/15ML

N006333 008

TABLET;ORAL

ASBRON

NOVARTIS

EQ 150MG BASE

A085148 001

THIABENDAZOLE

SUSPENSION;ORAL

MINTEZOL

MERCK SHARP DOHME

500MG/5ML

N016097 001

TABLET, CHEWABLE;ORAL

MINTEZOL

MERCK SHARP DOHME

500MG

N016096 001

THIAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

BETALIN S

LILLY

100MG/ML

A080853 001

THIAMINE HYDROCHLORIDE

ABRAXIS PHARM

100MG/ML

A080509 001

AKORN

100MG/ML

A087968 001 Oct 01, 1982

BEL MAR

100MG/ML

A080718 001

200MG/ML

A080712 001

DELL LABS

100MG/ML

A083775 001

HOSPIRA

100MG/ML

A040079 001 May 03, 1996

LUITPOLD

100MG/ML

A080667 001

PARKE DAVIS

100MG/ML

A080770 001

WATSON LABS

100MG/ML

A080571 001

100MG/ML

A083534 001

200MG/ML

A080571 002

200MG/ML

A083534 002

WEST-WARD PHARMS INT

100MG/ML

A080575 001

WYETH AYERST

100MG/ML

A080553 001

THIAMYLAL SODIUM

INJECTABLE; INJECTION

SURITAL

PARKEDALE

1GM/VIAL

N007600 003

5GM/VIAL

N007600 005

10GM/VIAL

N007600 009

THIETHYLPERAZINE MALATE

INJECTABLE; INJECTION

TORECAN

NOVARTIS

5MG/ML

N012754 002

THIETHYLPERAZINE MALEATE

SUPPOSITORY; RECTAL

TORECAN

NOVARTIS

10MG

N013247 001

TABLET; ORAL

TORECAN

NOVARTIS

10MG

N012753 001

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THIOPENTAL SODIUM

SUSPENSION;RECTAL

PENTOTHAL

ABBOTT

400MG/GM

N011679 001

THIORIDAZINE

SUSPENSION;ORAL

MELLARIL-S

NOVARTIS

EQ 25MG HYDROCHLORIDE/5ML **

N017923 001

EQ 100MG HYDROCHLORIDE/5ML **

N017923 002

THIORIDAZINE HYDROCHLORIDE

CONCENTRATE;ORAL

MELLARIL

NOVARTIS

30MG/ML **

N011808 012

100MG/ML **

N011808 018

THIORIDAZINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC

100MG/ML

A088229 001 Aug 23, 1983

ALPHARMA US PHARMS

30MG/ML

A087766 001 Apr 26, 1983

ANI PHARMS INC

30MG/ML

A089602 001 Nov 09, 1987

100MG/ML

A089603 001 Nov 09, 1987

HI TECH PHARMA

30MG/ML

A040125 001 Aug 16, 1996

100MG/ML

A040126 001 Aug 16, 1996

PHARM ASSOC

30MG/ML

A040187 001 Aug 28, 1997

100MG/ML

A040213 001 May 29, 1998

SANDOZ

30MG/ML

A088307 001 Nov 23, 1983

100MG/ML

A088308 001 Nov 23, 1983

WOCKHARDT

30MG/ML

A088258 001 Jul 25, 1983

100MG/ML

A088227 001 Jul 05, 1983

THIORIDAZINE HYDROCHLORIDE INTENSOL

ROXANE

30MG/ML

A088941 001 Dec 16, 1985

100MG/ML

A088942 001 Dec 16, 1985

TABLET;ORAL

MELLARIL

+ NOVARTIS

10MG **

N011808 003

+

15MG **

N011808 016

+

25MG **

N011808 006

+

50MG **

N011808 011

+

100MG **

N011808 009

+

150MG **

N011808 017

+

200MG **

N011808 015

THIORIDAZINE HYDROCHLORIDE

ANI PHARMS INC

10MG

A088270 001 Apr 14, 1983

10MG

A088493 001 May 17, 1985

15MG

A088271 001 Apr 14, 1983

25MG

A088272 001 Apr 14, 1983

50MG

A088194 001 Apr 14, 1983

100MG

A088273 001 Oct 03, 1983

100MG

A088456 001 May 17, 1985

FOSUN PHARMA

10MG

A088131 001 Aug 30, 1983

15MG

A088132 001 Aug 30, 1983

25MG

A088133 001 Aug 30, 1983

50MG

A088134 001 Aug 30, 1983

100MG

A088135 001 Nov 20, 1984

150MG

A088136 001 Sep 17, 1986

200MG

A088137 001 Sep 17, 1986

MUTUAL PHARM

10MG

A088375 001 Nov 18, 1983

25MG

A087264 001 Nov 18, 1983

50MG

A088370 001 Nov 18, 1983

100MG

A088379 001 Nov 16, 1983

MYLAN

10MG

A088332 001 Jun 27, 1983

25MG

A088333 001 Jun 27, 1983

50MG

A088334 001 Jun 27, 1983

100MG

A088335 001 Nov 18, 1983

PAR PHARM

10MG

A088351 001 Dec 05, 1983

15MG

A088352 001 Dec 05, 1983

25MG

A088336 001 Dec 05, 1983

50MG

A088322 001 Dec 05, 1983

100MG

A088480 001 Dec 29, 1983

150MG

A089764 001 Feb 09, 1988

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THIORIDAZINE HYDROCHLORIDE

TABLET; ORAL

THIORIDAZINE HYDROCHLORIDE

| | | | |
|----------------------|-------|-------------|--------------|
| | 200MG | A089765 001 | Feb 09, 1988 |
| ROXANE | 10MG | A088663 001 | Mar 15, 1984 |
| | 25MG | A088664 001 | Mar 15, 1984 |
| | 50MG | A088665 001 | Mar 15, 1984 |
| | 100MG | A089048 001 | Feb 26, 1985 |
| SUN PHARM INDUSTRIES | 15MG | A088461 001 | Nov 18, 1983 |
| | 150MG | A088737 001 | Sep 26, 1984 |
| | 200MG | A088738 001 | Oct 16, 1984 |
| SUPERPHARM | 10MG | A089103 001 | Jul 02, 1985 |
| | 25MG | A089104 001 | Jul 02, 1985 |
| | 50MG | A089105 001 | Jul 02, 1985 |
| WATSON LABS | 10MG | A088412 001 | Sep 12, 1983 |
| | 10MG | A088476 001 | Nov 08, 1983 |
| | 10MG | A088561 001 | May 11, 1984 |
| | 15MG | A088345 001 | Jul 28, 1983 |
| | 15MG | A088562 001 | May 11, 1984 |
| | 25MG | A088296 001 | Jul 28, 1983 |
| | 25MG | A088478 001 | Nov 08, 1983 |
| | 25MG | A088755 001 | Jul 24, 1984 |
| | 50MG | A088323 001 | Jul 28, 1983 |
| | 50MG | A088479 001 | Nov 08, 1983 |
| | 50MG | A088563 001 | May 11, 1984 |
| | 100MG | A088284 001 | Aug 25, 1983 |
| | 100MG | A088564 001 | May 11, 1984 |
| | 100MG | A088736 001 | Jul 24, 1984 |
| | 150MG | A088410 001 | Mar 05, 1984 |
| | 150MG | A088869 001 | Jun 28, 1985 |
| | 200MG | A088381 001 | Mar 14, 1984 |
| WATSON LABS TEVA | 15MG | A088477 001 | Nov 08, 1983 |
| | 25MG | A088567 001 | May 11, 1984 |
| | 200MG | A088872 001 | Apr 26, 1985 |
| WEST WARD | 10MG | A088658 001 | Mar 26, 1984 |
| | 15MG | A088659 001 | Mar 26, 1984 |
| | 25MG | A088660 001 | Mar 26, 1984 |
| | 50MG | A088661 001 | Mar 26, 1984 |

THIOTEPA

INJECTABLE; INJECTION

THIOPLEX

+ IMMUNEX 15MG/VIAL ** N020058 001 Dec 22, 1994

THIOTEPA

FRESENIUS KABI USA 15MG/VIAL A075698 001 Sep 20, 2001

IMMUNEX 15MG/VIAL N011683 001

TEVA PARENTERAL 15MG/VIAL ** A075730 001 Apr 20, 2001

30MG/VIAL ** A075730 002 Apr 20, 2001

THIOTHIXENE

CAPSULE; ORAL

NAVANE

+ PFIZER 1MG ** N016584 001

+ 2MG ** N016584 002

+ 5MG ** N016584 003

+ 10MG ** N016584 004

+ 20MG ** N016584 005

THIOTHIXENE

AM THERAP 1MG A071884 001 Aug 12, 1987

2MG A071885 001 Aug 12, 1987

5MG A071886 001 Aug 12, 1987

10MG A071887 001 Aug 12, 1987

20MG A072200 001 Dec 17, 1987

SANDOZ 1MG A071529 002 Jun 24, 1987

2MG A071529 003 Jun 24, 1987

5MG A071529 001 Jun 24, 1987

10MG A071529 004 Jun 24, 1987

WATSON LABS 1MG A070600 001 Jun 05, 1987

2MG A070601 001 Jun 05, 1987

2MG A071626 001 Jun 25, 1987

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

THIOTHIXENE

CAPSULE; ORAL

THIOTHIXENE

| | | |
|------|-------------|--------------|
| 5MG | A070602 001 | Jun 05, 1987 |
| 5MG | A071627 001 | Jun 25, 1987 |
| 10MG | A070603 001 | Jun 05, 1987 |
| 10MG | A071628 001 | Jun 25, 1987 |

THIOTHIXENE HYDROCHLORIDE

CONCENTRATE; ORAL

NAVANE

| | | |
|------------------------------------|-------------------|--------------------------|
| PFIZER | EQ 5MG BASE/ML | N016758 001 |
| THIOTHIXENE HYDROCHLORIDE | | |
| ALPHARMA US PHARMS | EQ 5MG BASE/ML | A070969 001 Oct 16, 1987 |
| PACO | EQ 1MG BASE/ML | A071917 001 Sep 20, 1989 |
| | EQ 5MG BASE/ML | A071939 001 Dec 16, 1988 |
| TEVA | EQ 5MG BASE/ML | A071184 001 Jun 22, 1987 |
| TEVA PHARMS | EQ 5MG BASE/ML | A071554 001 Oct 16, 1987 |
| THIOTHIXENE HYDROCHLORIDE INTENSOL | | |
| CYCLE PHARMS LTD | EQ 5MG BASE/ML | A073494 001 Jun 30, 1992 |
| INJECTABLE; INJECTION | | |
| NAVANE | | |
| PFIZER | EQ 2MG BASE/ML | N016904 001 |
| | EQ 10MG BASE/VIAL | N016904 002 |

THYROGLOBULIN

TABLET; ORAL

PROLOID

| | | |
|-------------|-------|-------------|
| PARKE DAVIS | 16MG | N002245 009 |
| | 32MG | N002245 005 |
| | 65MG | N002245 002 |
| | 100MG | N002245 008 |
| | 130MG | N002245 010 |
| | 200MG | N002245 007 |
| | 325MG | N002245 004 |

THYROGLOBULIN

| | | |
|------------|--------|-------------|
| IMPAX LABS | 64.8MG | A080151 001 |
|------------|--------|-------------|

THYROTROPIN

INJECTABLE; INJECTION

THYTROPAR

| | | |
|-------------------|------------|-------------|
| SANOFI AVENTIS US | 10 IU/VIAL | N008682 001 |
|-------------------|------------|-------------|

TIAGABINE HYDROCHLORIDE

TABLET; ORAL

GABITRIL

| | | |
|----------|------|--------------------------|
| CEPHALON | 6MG | N020646 006 Nov 29, 2005 |
| | 8MG | N020646 007 Nov 29, 2005 |
| | 10MG | N020646 008 Nov 29, 2005 |
| | 20MG | N020646 004 Sep 30, 1997 |

TICARCILLIN DISODIUM

INJECTABLE; INJECTION

TICAR

| | | |
|-----------------|-------------------|--------------------------|
| GLAXOSMITHKLINE | EQ 1GM BASE/VIAL | N050497 001 |
| | EQ 3GM BASE/VIAL | A062690 001 Dec 19, 1986 |
| | EQ 3GM BASE/VIAL | N050497 002 |
| | EQ 6GM BASE/VIAL | N050497 003 |
| | EQ 20GM BASE/VIAL | N050497 004 |
| | EQ 30GM BASE/VIAL | N050497 005 Apr 04, 1984 |

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLID

| | | |
|---------------------------|-------|--------------------------|
| ROCHE PALO | 125MG | N019979 001 Mar 24, 1993 |
| | 250MG | N019979 002 Oct 31, 1991 |
| TICLOPIDINE HYDROCHLORIDE | | |
| ACTAVIS ELIZABETH | 250MG | A075253 001 Aug 20, 1999 |
| MYLAN | 250MG | A075161 001 Sep 13, 1999 |
| | 250MG | A075316 001 Nov 02, 1999 |
| WATSON LABS | 250MG | A075309 001 Apr 26, 2000 |
| YAOPHARMA CO LTD | 250MG | A075318 001 Aug 20, 1999 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TICLOPIDINE HYDROCHLORIDE

TABLET;ORAL

TICLOPIDINE HYDROCHLORIDE

250MG

A075326 001 Aug 20, 1999

TILUDRONATE DISODIUM

TABLET;ORAL

SKELID

+ SANOFI AVENTIS US

EQ 200MG BASE **

N020707 001 Mar 07, 1997

TIMOLOL MALEATE

SOLUTION/DROPS;OPHTHALMIC

TIMOLOL MALEATE

AKORN

EQ 0.25% BASE

A074465 001 Mar 25, 1997

EQ 0.25% BASE

A074515 001 Mar 25, 1997

APOTEX INC

EQ 0.25% BASE

A075411 001 Sep 08, 2000

EQ 0.5% BASE

A075412 001 Sep 08, 2000

FOUGERA

EQ 0.25% BASE

A074667 001 Mar 25, 1997

EQ 0.5% BASE

A074668 001 Mar 25, 1997

TABLET;ORAL

BLOCADREN

MERCK

5MG **

N018017 001

10MG **

N018017 002

20MG **

N018017 004

TIMOLOL MALEATE

QUANTUM PHARMICS

5MG

A072466 001 May 19, 1989

10MG

A072467 001 May 19, 1989

20MG

A072468 001 May 19, 1989

TEVA

5MG

A072648 001 Jun 16, 1993

10MG

A072649 001 Jun 16, 1993

20MG

A072650 001 Jun 16, 1993

USL PHARMA

5MG

A072001 001 Apr 11, 1989

10MG

A072002 001 Apr 11, 1989

20MG

A072003 001 Apr 11, 1989

WATSON LABS

5MG

A072269 001 Apr 11, 1989

5MG

A072917 001 Jul 31, 1991

10MG

A072270 001 Apr 11, 1989

10MG

A072918 001 Jul 31, 1991

20MG

A072271 001 Apr 11, 1989

20MG

A072919 001 Jul 31, 1991

YAOPHARMA CO LTD

5MG

A072550 001 Apr 13, 1989

10MG

A072551 001 Apr 13, 1989

20MG

A072552 001 Apr 13, 1989

TINZAPARIN SODIUM

INJECTABLE;INJECTION

INNOHEP

LEO PHARMA AS

20,000 IU/ML

N020484 001 Jul 14, 2000

TIOCONAZOLE

CREAM;TOPICAL

TZ-3

PFIZER

1%

N018682 001 Feb 18, 1983

TIROFIBAN HYDROCHLORIDE

INJECTABLE;INJECTION

AGGRASTAT

MEDICURE

EQ 12.5MG BASE/50ML (EQ 0.25MG BASE/ML)

N020912 001 May 14, 1998

EQ 25MG BASE/500ML (EQ 0.05MG BASE/ML)

N020913 001 May 14, 1998

TIZANIDINE HYDROCHLORIDE

TABLET;ORAL

TIZANIDINE HYDROCHLORIDE

ACTAVIS ELIZABETH

EQ 2MG BASE

A076283 001 Jul 12, 2002

EQ 4MG BASE

A076283 002 Jul 12, 2002

BARR

EQ 2MG BASE

A076371 001 Apr 09, 2003

EQ 4MG BASE

A076371 002 Apr 09, 2003

IVAX SUB TEVA PHARMS

EQ 2MG BASE

A076321 001 Sep 30, 2004

EQ 4MG BASE

A076321 002 Sep 30, 2004

MYLAN PHARMS INC

EQ 2MG BASE

A076282 001 Dec 16, 2003

EQ 4MG BASE

A076282 002 Dec 16, 2003

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TIZANIDINE HYDROCHLORIDE

TABLET; ORAL

ZANAFLEX

+ COVIS PHARMA BV EQ 2MG BASE ** N020397 002 Feb 04, 2000

TOBRAMYCIN

SOLUTION/DROPS; OPHTHALMIC

TOBRAMYCIN

ALCON PHARMS LTD 0.3% A063176 001 May 25, 1994

APOTEX INC 0.3% A065087 001 Feb 25, 2002

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

NEBCIN

LILLY EQ 10MG BASE/ML A062008 004

EQ 10MG BASE/ML A062707 001 Apr 29, 1987

+ EQ 10MG BASE/ML ** N050477 005

EQ 40MG BASE/ML A062008 001

+ EQ 1.2GM BASE/VIAL ** N050519 001

TOBRAMYCIN SULFATE

APOTHECON EQ 10MG BASE/ML A064021 001 May 31, 1994

EQ 40MG BASE/ML A064021 002 May 31, 1994

EQ 40MG BASE/ML A064026 001 May 31, 1994

HOSPIRA EQ 10MG BASE/ML A063080 001 Apr 30, 1991

EQ 40MG BASE/ML A063161 001 May 29, 1991

IGI LABS INC EQ 10MG BASE/ML A063119 001 Oct 31, 1994

EQ 40MG BASE/ML A063120 001 Oct 31, 1994

EQ 40MG BASE/ML A063121 001 Oct 31, 1994

EQ 40MG BASE/ML A063122 001 Oct 31, 1994

WATSON LABS INC EQ 10MG BASE/ML A062945 001 Aug 09, 1989

EQ 40MG BASE/ML A062945 002 Aug 09, 1989

WEST-WARD PHARMS INT EQ 10MG BASE/ML A063113 001 Apr 26, 1991

EQ 10MG BASE/ML A063128 001 Nov 27, 1991

EQ 40MG BASE/ML A063118 001 Jul 29, 1991

EQ 40MG BASE/ML A063127 001 Nov 27, 1991

TOBRAMYCIN SULFATE (PHARMACY BULK)

HOSPIRA EQ 40MG BASE/ML ** A063116 001 May 18, 1992

TOCAINIDE HYDROCHLORIDE

TABLET; ORAL

TONOCARD

ASTRAZENECA 400MG N018257 001 Nov 09, 1984

600MG N018257 002 Nov 09, 1984

TOLAZAMIDE

TABLET; ORAL

TOLAZAMIDE

BARR 100MG A070162 001 Jan 14, 1986

250MG A070163 001 Jan 14, 1986

500MG A070164 001 Jan 14, 1986

DURAMED PHARMS BARR 100MG A070165 001 Jan 10, 1986

250MG A070166 001 Jan 10, 1986

500MG A070167 001 Jan 10, 1986

G AND W LABS INC 100MG N018894 001 Nov 02, 1984

250MG N018894 002 Nov 02, 1984

500MG N018894 003 Nov 02, 1984

INTERPHARM 250MG A071270 001 Sep 23, 1986

500MG A071271 001 Sep 23, 1986

PAR PHARM 100MG A070159 001 Jan 06, 1986

250MG A070160 001 Jan 06, 1986

500MG A070161 001 Jan 06, 1986

SUN PHARM INDUSTRIES 100MG A071357 001 Jul 16, 1987

250MG A071358 001 Jul 16, 1987

500MG A071359 001 Jul 16, 1987

SUPERPHARM 250MG A070763 001 Jun 16, 1986

500MG A070764 001 Jun 16, 1986

USL PHARMA 100MG A071355 001 Jan 11, 1988

250MG A070168 001 Apr 02, 1986

500MG A070169 001 Apr 02, 1986

WATSON LABS 100MG A070242 001 Aug 01, 1986

100MG A070513 001 Jan 09, 1986

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TOLAZAMIDETABLET; ORAL
TOLAZAMIDE

| | | | | |
|------------------|----------------------|---------|-----|--------------|
| | 250MG | A070243 | 001 | Aug 01, 1986 |
| | 250MG | A070514 | 001 | Jan 09, 1986 |
| | 500MG | A070244 | 001 | Aug 01, 1986 |
| | 500MG | A070515 | 001 | Jan 09, 1986 |
| YAOPHARMA CO LTD | 100MG | A071633 | 001 | Dec 09, 1987 |
| | 250MG | A070289 | 001 | Mar 13, 1986 |
| | 500MG | A070290 | 001 | Mar 13, 1986 |
| TOLINASE | | | | |
| + | PHARMACIA AND UPJOHN | 100MG | ** | N015500 002 |
| + | | 250MG | ** | N015500 004 |
| + | | 500MG | ** | N015500 005 |

TOLAZOLINE HYDROCHLORIDE

INJECTABLE; INJECTION

PRISCOLINE

| | | | | |
|----------|---------|---------|-----|--------------|
| NOVARTIS | 25MG/ML | N006403 | 005 | Feb 22, 1985 |
|----------|---------|---------|-----|--------------|

TOLBUTAMIDE

TABLET; ORAL

ORINASE

| | | | | |
|----------------------|-------|----|---------|-----|
| PHARMACIA AND UPJOHN | 250MG | ** | N010670 | 002 |
| | 500MG | ** | N010670 | 001 |

TOLBUTAMIDE

| | | | | |
|------------------|-------|---------|-----|--------------|
| ALRA | 500MG | A086141 | 001 | |
| ASCOT | 500MG | A087541 | 001 | Mar 01, 1983 |
| BARR | 500MG | A087121 | 001 | |
| DAVA PHARMS INC | 500MG | A086926 | 001 | |
| IVAX PHARMS | 500MG | A087093 | 001 | |
| PARKE DAVIS | 500MG | A086047 | 001 | |
| PUREPAC PHARM | 500MG | A088950 | 001 | Jun 17, 1985 |
| SANDOZ | 500MG | N012678 | 001 | |
| SUPERPHARM | 500MG | A088893 | 001 | Nov 19, 1984 |
| VANGARD | 500MG | A087876 | 001 | Apr 20, 1982 |
| WATSON LABS | 250MG | A089110 | 001 | May 29, 1987 |
| | 500MG | A086109 | 001 | |
| | 500MG | A087318 | 001 | |
| | 500MG | A089111 | 001 | May 29, 1987 |
| YAOPHARMA CO LTD | 500MG | A086574 | 001 | |

TOLBUTAMIDE SODIUM

INJECTABLE; INJECTION

ORINASE DIAGNOSTIC

| | | | | |
|----------------------|------------------|---------|-----|--|
| PHARMACIA AND UPJOHN | EQ 1GM BASE/VIAL | N012095 | 001 | |
|----------------------|------------------|---------|-----|--|

TOLCAPONE

TABLET; ORAL

TASMAR

| | | | | |
|--------------------|-------|---------|-----|--------------|
| VALEANT PHARMS LLC | 200MG | N020697 | 002 | Jan 29, 1998 |
|--------------------|-------|---------|-----|--------------|

TOLMETIN SODIUM

CAPSULE; ORAL

TOLECTIN DS

| | | | | |
|----------------------|---------------|---------|-----|--|
| ORTHO MCNEIL JANSSEN | EQ 400MG BASE | N018084 | 001 | |
|----------------------|---------------|---------|-----|--|

TOLMETIN SODIUM

| | | | | |
|----------------------|---------------|---------|-----|--------------|
| ACTAVIS ELIZABETH | EQ 400MG BASE | A073308 | 001 | Jan 24, 1992 |
| FOSUN PHARMA | EQ 400MG BASE | A073462 | 001 | Apr 30, 1992 |
| IVAX SUB TEVA PHARMS | EQ 400MG BASE | A073392 | 001 | Jan 24, 1992 |
| SUN PHARM INDUSTRIES | EQ 400MG BASE | A073311 | 001 | Nov 27, 1991 |
| TEVA | EQ 400MG BASE | A073519 | 001 | May 29, 1992 |

TABLET; ORAL

TOLECTIN

| | | | | |
|----------------------|---------------|---------|-----|--|
| ORTHO MCNEIL JANSSEN | EQ 200MG BASE | N017628 | 001 | |
|----------------------|---------------|---------|-----|--|

TOLECTIN 600

| | | | | |
|----------------------|---------------|---------|-----|--------------|
| ORTHO MCNEIL JANSSEN | EQ 600MG BASE | N017628 | 002 | Mar 08, 1989 |
|----------------------|---------------|---------|-----|--------------|

TOLMETIN SODIUM

| | | | | |
|-------------------|---------------|---------|-----|--------------|
| ACTAVIS ELIZABETH | EQ 600MG BASE | A073527 | 001 | Jun 30, 1992 |
| FOSUN PHARMA | EQ 200MG BASE | A073588 | 001 | Jul 31, 1992 |
| | EQ 600MG BASE | A074002 | 001 | Sep 27, 1993 |
| G AND W LABS INC | EQ 600MG BASE | A074399 | 001 | Mar 28, 1996 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TOLMETIN SODIUM

TABLET; ORAL

TOLMETIN SODIUM

| | | | | |
|----------------------|---------------|---------|-----|--------------|
| | EQ 600MG BASE | A074729 | 001 | Feb 27, 1997 |
| SUN PHARM INDUSTRIES | EQ 200MG BASE | A073310 | 001 | Nov 27, 1991 |

TOLTERODINE TARTRATE

TABLET; ORAL

TOLTERODINE TARTRATE

| | | | | |
|-------------|-----|---------|-----|--------------|
| APOTEX CORP | 1MG | A200164 | 001 | Sep 25, 2012 |
| | 2MG | A200164 | 002 | Sep 25, 2012 |

TOLVAPTAN

TABLET; ORAL

SAMSCA

| | | | | |
|------------------------|---------|---------|-----|--------------|
| + OTSUKA AMERICA PHARM | 60MG ** | N022275 | 003 | May 19, 2009 |
|------------------------|---------|---------|-----|--------------|

TOPIRAMATE

CAPSULE; ORAL

TOPAMAX SPRINKLE

| | | | | |
|----------------|------|---------|-----|--------------|
| JANSSEN PHARMS | 50MG | N020844 | 003 | Oct 26, 1998 |
|----------------|------|---------|-----|--------------|

TOPIRAMATE

| | | | | |
|--------------|------|---------|-----|--------------|
| BARR | 15MG | A076448 | 001 | Apr 15, 2009 |
| | 25MG | A076448 | 002 | Apr 15, 2009 |
| FOSUN PHARMA | 15MG | A079206 | 001 | Oct 14, 2009 |
| | 25MG | A079206 | 002 | Oct 14, 2009 |
| MYLAN | 15MG | A078418 | 001 | Oct 14, 2009 |
| | 25MG | A078418 | 002 | Oct 14, 2009 |

TABLET; ORAL

TOPAMAX

| | | | | |
|----------------|-------|---------|-----|--------------|
| JANSSEN PHARMS | 300MG | N020505 | 003 | Dec 24, 1996 |
| | 400MG | N020505 | 006 | Dec 24, 1996 |

TOPIRAMATE

| | | | | |
|--------------------|-------|---------|-----|--------------|
| ACTAVIS TOTOWA | 25MG | A078637 | 001 | Feb 27, 2013 |
| | 50MG | A078637 | 002 | Feb 27, 2013 |
| | 100MG | A078637 | 003 | Feb 27, 2013 |
| | 200MG | A078637 | 004 | Feb 27, 2013 |
| BARR | 25MG | A076315 | 001 | Mar 27, 2009 |
| | 100MG | A076315 | 002 | Mar 27, 2009 |
| | 200MG | A076315 | 003 | Mar 27, 2009 |
| HIKMA PHARMS | 25MG | A091185 | 001 | Nov 25, 2013 |
| | 50MG | A091185 | 002 | Nov 25, 2013 |
| | 100MG | A091185 | 003 | Nov 25, 2013 |
| | 200MG | A091185 | 004 | Nov 25, 2013 |
| MYLAN | 25MG | A076314 | 001 | Mar 27, 2009 |
| | 50MG | A076314 | 002 | Mar 27, 2009 |
| | 100MG | A076314 | 003 | Mar 27, 2009 |
| | 200MG | A076314 | 004 | Mar 27, 2009 |
| PLIVA HRVATSKA DOO | 25MG | A077905 | 001 | Mar 30, 2009 |
| | 50MG | A077905 | 002 | Mar 30, 2009 |
| | 100MG | A077905 | 003 | Mar 30, 2009 |
| | 200MG | A077905 | 004 | Mar 30, 2009 |
| ROXANE | 25MG | A076306 | 001 | Mar 27, 2009 |
| | 50MG | A076306 | 002 | Mar 27, 2009 |
| | 100MG | A076306 | 003 | Mar 27, 2009 |
| | 200MG | A076306 | 004 | Mar 27, 2009 |
| WATSON LABS | 25MG | A077643 | 001 | Mar 27, 2009 |
| | 50MG | A077643 | 002 | Mar 27, 2009 |
| | 100MG | A077643 | 003 | Mar 27, 2009 |
| | 200MG | A077643 | 004 | Mar 27, 2009 |
| WOCKHARDT USA | 25MG | A090353 | 001 | Sep 01, 2010 |
| | 50MG | A090353 | 002 | Sep 01, 2010 |
| | 100MG | A090353 | 003 | Sep 01, 2010 |
| | 200MG | A090353 | 004 | Sep 01, 2010 |

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TOPOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

TOPOTECAN HYDROCHLORIDE

| | | | | |
|----------------------|------------------|---------|-----|--------------|
| FRESENIUS KABI ONCOL | EQ 4MG BASE/VIAL | A091376 | 001 | Nov 29, 2010 |
| SUN PHARM INDS LTD | EQ 4MG BASE/VIAL | A202203 | 001 | Aug 29, 2013 |

SOLUTION; INTRAVENOUS

TOPOTECAN

| | | | | | |
|---|------------|-------------------------------------|---------|-----|--------------|
| + | SANDOZ INC | EQ 1MG BASE/ML (EQ 1MG BASE/ML) ** | N200199 | 001 | Feb 25, 2011 |
| + | | EQ 3MG BASE/3ML (EQ 1MG BASE/ML) ** | N200199 | 002 | Feb 25, 2011 |
| + | | EQ 4MG BASE/4ML (EQ 1MG BASE/ML) ** | N200199 | 003 | Feb 25, 2011 |

TORSEMIDE

INJECTABLE; INJECTION

DEMADEX

| | | | | | |
|---|-------|-----------------------|---------|-----|--------------|
| + | ROCHE | 50MG/5ML (10MG/ML) ** | N020137 | 002 | Aug 23, 1993 |
| + | | 20MG/2ML (10MG/ML) ** | N020137 | 001 | Aug 23, 1993 |

TORSEMIDE

| | | | | |
|----------------------|--------------------|---------|-----|--------------|
| LUITPOLD | 20MG/2ML (10MG/ML) | A090656 | 001 | Apr 21, 2010 |
| | 50MG/5ML (10MG/ML) | A090656 | 002 | Apr 21, 2010 |
| WEST-WARD PHARMS INT | 20MG/2ML (10MG/ML) | A078007 | 001 | Jun 11, 2008 |
| | 50MG/5ML (10MG/ML) | A078007 | 002 | Jun 11, 2008 |

TABLET; ORAL

TORSEMIDE

| | | | | |
|----------------|-------|---------|-----|--------------|
| SUN PHARM INDS | 5MG | A078478 | 001 | Feb 26, 2008 |
| | 10MG | A078478 | 002 | Feb 26, 2008 |
| | 20MG | A078478 | 003 | Feb 26, 2008 |
| | 100MG | A078478 | 004 | Feb 26, 2008 |

TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE

| | | | | |
|----------------------|------|---------|-----|--------------|
| ACCORD HLTHCARE | 50MG | A202390 | 001 | May 16, 2013 |
| ACTAVIS ELIZABETH | 50MG | A075960 | 001 | Jun 19, 2002 |
| ASTA | 50MG | A075974 | 001 | Jul 12, 2002 |
| FOSUN PHARMA | 50MG | A075968 | 001 | Jun 25, 2002 |
| IVAX SUB TEVA PHARMS | 50MG | A075963 | 001 | Jul 03, 2002 |
| MYLAN PHARMS INC | 50MG | A075980 | 001 | Nov 21, 2002 |
| NORTHSTAR HLTHCARE | 50MG | A078935 | 001 | May 26, 2010 |
| WATSON LABS | 50MG | A075962 | 001 | Jun 24, 2002 |

ULTRAM

| | | | | |
|----------------|-------|---------|-----|--------------|
| JANSSEN PHARMS | 100MG | N020281 | 001 | Mar 03, 1995 |
|----------------|-------|---------|-----|--------------|

TABLET, EXTENDED RELEASE; ORAL

RYZOLT

| | | | | | |
|---|---------------|----------|---------|-----|--------------|
| + | PURDUE PHARMA | 100MG ** | N021745 | 001 | Dec 30, 2008 |
| + | | 200MG ** | N021745 | 002 | Dec 30, 2008 |
| + | | 300MG ** | N021745 | 003 | Dec 30, 2008 |

ULTRAM ER

| | | | | | |
|---|----------------|-------|---------|-----|--------------|
| + | VALEANT PHARMS | 100MG | N021692 | 001 | Sep 08, 2005 |
| + | | 200MG | N021692 | 002 | Sep 08, 2005 |
| + | | 300MG | N021692 | 003 | Sep 08, 2005 |

TABLET, ORALLY DISINTEGRATING; ORAL

RYBIX ODT

| | | | | |
|--------------|------|---------|-----|--------------|
| SHIONOGI INC | 50MG | N021693 | 001 | May 05, 2005 |
|--------------|------|---------|-----|--------------|

TRAMETINIB DIMETHYL SULFOXIDE

TABLET; ORAL

MEKINIST

| | | | | | |
|---|----------------------|--------|---------|-----|--------------|
| + | NOVARTIS PHARMS CORP | EQ 1MG | N204114 | 002 | May 29, 2013 |
|---|----------------------|--------|---------|-----|--------------|

TRANDOLAPRIL

TABLET; ORAL

MAVIK

| | | | | | |
|---|--------|-----|---------|-----|--------------|
| + | ABBVIE | 1MG | N020528 | 001 | Apr 26, 1996 |
| + | | 2MG | N020528 | 002 | Apr 26, 1996 |
| + | | 4MG | N020528 | 003 | Apr 26, 1996 |

TRANDOLAPRIL

| | | | | |
|--------------------|-----|---------|-----|--------------|
| CIPLA | 1MG | A077307 | 002 | Jun 12, 2007 |
| | 2MG | A077307 | 001 | Jun 12, 2007 |
| | 4MG | A077307 | 003 | Jun 12, 2007 |
| DR REDDYS LABS LTD | 1MG | A078493 | 001 | Aug 25, 2008 |
| | 2MG | A078493 | 002 | Aug 25, 2008 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRANDOLAPRIL

TABLET;ORAL

TRANDOLAPRIL

| | | | |
|-----------------|-----|-------------|--------------|
| | 4MG | A078493 003 | Aug 25, 2008 |
| EPIC PHARMA LLC | 1MG | A077256 001 | Jun 12, 2007 |
| | 2MG | A077256 002 | Jun 12, 2007 |
| | 4MG | A077256 003 | Jun 12, 2007 |
| INVAGEN PHARMS | 1MG | A078320 001 | Jun 12, 2007 |
| | 2MG | A078320 002 | Jun 12, 2007 |
| | 4MG | A078320 003 | Jun 12, 2007 |
| MYLAN | 1MG | A078346 001 | Apr 28, 2008 |
| | 2MG | A078346 002 | Apr 28, 2008 |
| | 4MG | A078346 003 | Apr 28, 2008 |
| WATSON LABS | 1MG | A077805 001 | Jun 12, 2007 |
| | 2MG | A077805 002 | Jun 12, 2007 |
| | 4MG | A077805 003 | Jun 12, 2007 |

TRANEXAMIC ACID

TABLET;ORAL

CYKLOKAPRON

| | | | |
|----------------------|-------|-------------|--------------|
| PHARMACIA AND UPJOHN | 500MG | N019280 001 | Dec 30, 1986 |
| TRANEXAMIC ACID | | | |
| AMERIGEN PHARMS LTD | 650MG | A203256 001 | Jul 25, 2016 |

TRAVOPROST

SOLUTION/DROPS;OPHTHALMIC

IZBA

| | | | |
|------------------------|-----------|-------------|--------------|
| + NOVARTIS PHARMS CORP | 0.003% ** | N204822 001 | May 15, 2014 |
| TRAVATAN | | | |
| + ALCON PHARMS LTD | 0.004% ** | N021257 001 | Mar 16, 2001 |

TRAZODONE HYDROCHLORIDE

TABLET;ORAL

DESYREL

| | | | |
|----------|----------|-------------|--------------|
| + PRAGMA | 50MG ** | N018207 001 | |
| + | 100MG ** | N018207 002 | |
| + | 150MG ** | N018207 003 | Mar 25, 1985 |
| + | 300MG ** | N018207 004 | Nov 07, 1988 |

TRAZODONE HYDROCHLORIDE

| | | | |
|---------------------|-------|-------------|--------------|
| AM THERAP | 50MG | A071139 001 | Oct 29, 1986 |
| | 100MG | A071140 001 | Oct 29, 1986 |
| AUROLIFE PHARMA LLC | 50MG | A072484 001 | Apr 30, 1990 |
| FOSUN PHARMA | 100MG | A072483 001 | Apr 30, 1990 |
| MYLAN | 50MG | A071405 001 | Feb 27, 1991 |
| | 100MG | A071406 001 | Feb 27, 1991 |
| MYLAN PHARMS INC | 50MG | A090514 001 | Jun 02, 2009 |
| | 100MG | A090514 002 | Jun 02, 2009 |
| | 150MG | A090514 003 | Jun 02, 2009 |
| | 300MG | A090514 004 | Jun 02, 2009 |
| QUANTUM PHARMICS | 100MG | A070921 001 | Dec 01, 1986 |
| TEVA | 150MG | A074357 001 | Apr 30, 1997 |
| USL PHARMA | 50MG | A070491 001 | Apr 29, 1987 |
| | 100MG | A070492 001 | Apr 29, 1987 |
| WATSON LABS | 50MG | A070857 001 | Oct 10, 1986 |
| | 50MG | A071112 001 | Nov 17, 1986 |
| | 100MG | A070858 001 | Oct 10, 1986 |
| | 100MG | A071113 001 | Nov 17, 1986 |

TRIALODINE

| | | | |
|------------------|------|-------------|--------------|
| QUANTUM PHARMICS | 50MG | A070942 001 | Dec 01, 1986 |
|------------------|------|-------------|--------------|

TABLET, EXTENDED RELEASE;ORAL

OLEPTRO

| | | | |
|-------------------|----------|-------------|--------------|
| + ANGELINI PHARMA | 150MG ** | N022411 001 | Feb 02, 2010 |
| + | 300MG ** | N022411 002 | Feb 02, 2010 |

TRETINOIN

CAPSULE;ORAL

VESANOID

| | | | |
|---------------|---------|-------------|--------------|
| + CHEPLAPHARM | 10MG ** | N020438 001 | Nov 22, 1995 |
|---------------|---------|-------------|--------------|

CREAM;TOPICAL

TRETINOIN

| | | | |
|--------------------|---------|-------------|--------------|
| ALLERGAN SALES LLC | 0.0375% | A090098 001 | Mar 22, 2010 |
|--------------------|---------|-------------|--------------|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRETINOINCREAM;TOPICAL
TRETINOIN

0.075%

A202209 001 Oct 11, 2012

SOLUTION;TOPICAL

RETIN-A

+ VALEANT INTL

0.05%

N016921 001

TRETINOIN

TEVA PHARMS

0.05%

A074873 001 Jun 19, 1998

WOCKHARDT

0.05%

A075260 001 Jan 25, 1999

SWAB;TOPICAL

RETIN-A

VALEANT INTL

0.05%

N016921 002

TRIAMCINOLONE

TABLET;ORAL

ARISTOCORT

ASTELLAS

1MG

N011161 009

2MG

N011161 004

4MG

N011161 007

8MG

N011161 011

16MG

N011161 010

KENACORT

DELCOR ASSET CORP

1MG

N011283 003

2MG

N011283 008

4MG

N011283 006

8MG

N011283 010

TRIAMCINOLONE

BARR

2MG

A084286 001

2MG

A084318 001

4MG

A084267 001

4MG

A084319 001

8MG

A084268 001

8MG

A084320 001

IMPAX LABS

4MG

A084340 001

IVAX SUB TEVA PHARMS

4MG

A083750 001

MYLAN

2MG

A084406 001

PUREPAC PHARM

2MG

A084020 002

4MG

A084020 003

ROXANE

2MG

A084708 001

4MG

A084709 001

8MG

A084707 001

SANDOZ

4MG

A085601 001

TEVA

4MG

A084775 001

WATSON LABS

4MG

A084270 001

4MG

A085834 001

TRIAMCINOLONE ACETONIDE

AEROSOL, METERED;INHALATION

AZMACORT

ABBVIE

0.1MG/INH

N018117 001 Apr 23, 1982

AEROSOL, METERED;NASAL

NASACORT

SANOFI AVENTIS US

0.055MG/INH

N019798 001 Jul 11, 1991

CREAM;TOPICAL

ARISTOCORT

ASTELLAS

0.025%

A083017 003

0.1%

A083016 004

0.5%

A083015 002

ARISTOCORT A

ASTELLAS

0.025%

A083017 004

0.025%

A088818 001 Oct 16, 1984

0.1%

A083016 005

0.1%

A088819 001 Oct 16, 1984

0.5%

A083015 003

0.5%

A088820 001 Oct 16, 1984

FLUTEX

IVAX PHARMS

0.025%

A085539 001

0.1%

A085539 002

0.5%

A085539 003

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIAMCINOLONE ACETONIDE

CREAM;TOPICAL

| | | | |
|----------------------------------------------------------|--------------------------|-------------|--------------|
| KENALOG | | | |
| DELCOR ASSET CORP | 0.5% | A083943 001 | |
| KENALOG-H | | | |
| DELCOR ASSET CORP | 0.1% | A086240 001 | |
| TRIA CET | | | |
| TEVA | 0.025% | A084908 001 | |
| | 0.1% | A084908 002 | |
| | 0.5% | A084908 003 | |
| TRIA CORT | | | |
| SOLVAY | 0.1% | A087113 001 | |
| TRIAMCINOLONE ACETONIDE | | | |
| ACTAVIS MID ATLANTIC | 0.1% | A087798 001 | Jun 04, 1982 |
| ALPHARMA US PHARMS | 0.025% | A087797 001 | Jun 07, 1982 |
| AMBIX | 0.025% | A087932 001 | May 09, 1983 |
| MORTON GROVE | 0.025% | A088094 001 | Sep 01, 1983 |
| | 0.1% | A088095 001 | Sep 01, 1983 |
| | 0.5% | A088096 001 | Sep 01, 1983 |
| PHARMADERM | 0.025% | A087990 001 | Jul 07, 1983 |
| | 0.1% | A087991 001 | Jul 07, 1983 |
| | 0.5% | A087992 001 | Jul 07, 1983 |
| PHARMAFAIR | 0.025% | A087921 001 | Aug 10, 1982 |
| | 0.1% | A087912 001 | Aug 10, 1982 |
| | 0.5% | A087922 001 | Aug 10, 1982 |
| TARO | 0.025% | A040038 001 | Oct 26, 1994 |
| | 0.025% | A086277 001 | |
| | 0.1% | A086276 001 | |
| | 0.5% | A086275 001 | |
| TOPIDERM | 0.025% | A089274 001 | Feb 21, 1989 |
| | 0.1% | A089275 001 | Feb 21, 1989 |
| | 0.5% | A089276 001 | Feb 21, 1989 |
| TRIA TEX | | | |
| IVAX PHARMS | 0.025% | A087430 001 | Nov 01, 1988 |
| | 0.1% | A087429 001 | Nov 01, 1988 |
| | 0.5% | A087428 001 | Nov 01, 1988 |
| TRYMEX | | | |
| SAVAGE LABS | 0.025% | A088196 001 | Mar 25, 1983 |
| | 0.1% | A088197 001 | Mar 25, 1983 |
| | 0.5% | A088198 001 | Mar 25, 1983 |
| GEL;TOPICAL | | | |
| ARISTOGEL | | | |
| ASTELLAS | 0.1% | A083380 001 | |
| INJECTABLE; INJECTION | | | |
| TRIAMCINOLONE ACETONIDE | | | |
| PARNELL | 3MG/ML | N019503 001 | Oct 16, 1987 |
| SAN DOZ INC | 10MG/ML | A090166 001 | May 27, 2009 |
| | 40MG/ML | A090164 001 | Jun 01, 2009 |
| WATSON LABS | 40MG/ML | A085825 001 | |
| INJECTABLE; INTRA-ARTICULAR, INTRAMUSCULAR, INTRAVITREAL | | | |
| TRIVARIS | | | |
| + ALLERGAN | 8MG/0.1ML (8MG/0.1ML) ** | N022220 001 | Jun 16, 2008 |
| LOTION;TOPICAL | | | |
| KENALOG | | | |
| DELCOR ASSET CORP | 0.025% ** | A084343 001 | |
| + | 0.025% ** | N011602 003 | |
| | 0.1% ** | A084343 002 | |
| + | 0.1% ** | N011602 001 | |
| TRIAMCINOLONE ACETONIDE | | | |
| ALPHARMA US PHARMS | 0.025% | A087191 001 | Sep 08, 1982 |
| | 0.1% | A087192 001 | Sep 08, 1982 |
| OINTMENT;TOPICAL | | | |
| ARISTOCORT | | | |
| ASTELLAS | 0.1% | A080750 004 | |
| | 0.5% ** | A080745 002 | |
| ARISTOCORT A | | | |
| ASTELLAS | 0.1% | A080750 003 | |
| | 0.1% | A088780 001 | Oct 01, 1984 |
| | 0.5% ** | A080745 003 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIAMCINOLONE ACETONIDE

OINTMENT; TOPICAL

| | | | | | |
|-------------------------|---------|--|---------|-----|--------------|
| ARISTOCORT A | | | | | |
| | 0.5% | | A088781 | 001 | Oct 05, 1984 |
| FLUTEX | | | | | |
| IVAX PHARMS | 0.025% | | A087375 | 001 | Nov 01, 1988 |
| | 0.1% | | A087377 | 001 | Nov 01, 1988 |
| | 0.5% | | A087376 | 001 | Nov 01, 1988 |
| KENALOG | | | | | |
| DELCO ASSET CORP | 0.5% ** | | A083944 | 001 | |
| + MYLAN PHARMS INC | 0.025% | | N011600 | 003 | |
| + | 0.1% | | N011600 | 001 | |
| TRIAMCINOLONE ACETONIDE | | | | | |
| ACTAVIS MID ATLANTIC | 0.1% | | A087799 | 001 | Jun 07, 1982 |
| ALPHARMA US PHARMS | 0.5% | | A089913 | 001 | Dec 23, 1988 |
| MORTON GROVE | 0.025% | | A088090 | 001 | Sep 01, 1983 |
| | 0.1% | | A088091 | 001 | Sep 01, 1983 |
| | 0.5% | | A088092 | 001 | Sep 01, 1983 |
| PHARMADERM | 0.025% | | A088692 | 001 | Aug 02, 1984 |
| | 0.1% | | A088690 | 001 | Aug 02, 1984 |
| TARO | 0.025% | | A040040 | 001 | Sep 30, 1994 |
| | 0.025% | | A040374 | 001 | Jun 05, 2001 |
| | 0.1% | | A087902 | 001 | Dec 27, 1982 |
| | 0.5% | | A040386 | 001 | Jun 05, 2001 |
| TRYMEX | | | | | |
| SAVAGE LABS | 0.025% | | A088693 | 001 | Aug 02, 1984 |
| | 0.1% | | A088691 | 001 | Aug 02, 1984 |

PASTE; DENTAL

| | | | | | |
|--------------------|---------|--|---------|-----|--------------|
| KENALOG IN ORABASE | | | | | |
| + DELCO ASSET CORP | 0.1% ** | | N012097 | 001 | |
| ORALONE | | | | | |
| TARO | 0.1% | | A071383 | 001 | Jul 06, 1987 |

SPRAY, METERED; NASAL

| | | | | | |
|-------------------------|---------------|--|---------|-----|--------------|
| ALLERNAZE | | | | | |
| LUPIN ATLANTIS | 0.05MG/SPRAY | | N020120 | 001 | Feb 04, 2000 |
| NASACORT HFA | | | | | |
| SANOFI AVENTIS US | 0.055MG/SPRAY | | N020784 | 001 | Apr 07, 2004 |
| TRIAMCINOLONE ACETONIDE | | | | | |
| PERRIGO ISRAEL | 0.055MG/SPRAY | | A078104 | 001 | Jul 30, 2009 |

TRIAMCINOLONE DIACETATE

INJECTABLE; INJECTION

| | | | | | |
|-------------------------|-----------------|--|---------|-----|--|
| ARISTOCORT | | | | | |
| FOSUN PHARMA | 25MG/ML | | N011685 | 003 | |
| + | 40MG/ML ** | | N012802 | 001 | |
| TRIAMCINOLONE DIACETATE | | | | | |
| AKORN | 25MG/ML | | A085122 | 001 | |
| | 40MG/ML | | A086394 | 001 | |
| WATSON LABS | 40MG/ML | | A084072 | 001 | |
| | 40MG/ML | | A085529 | 001 | |
| SYRUP; ORAL | | | | | |
| ARISTOCORT | | | | | |
| ASTELLAS | 2MG/5ML | | N011960 | 004 | |
| KENACORT | | | | | |
| DELCO ASSET CORP | EQ 4MG BASE/5ML | | N012515 | 001 | |

TRIAZOLAM

TABLET; ORAL

| | | | | | |
|----------------------|---------|--|---------|-----|--------------|
| HALCION | | | | | |
| PHARMACIA AND UPJOHN | 0.5MG | | N017892 | 002 | Nov 15, 1982 |
| TRIAZOLAM | | | | | |
| WATSON LABS | 0.125MG | | A074445 | 001 | Oct 20, 1995 |
| | 0.25MG | | A074445 | 002 | Oct 20, 1995 |

TRICHLORMETHIAZIDE

TABLET; ORAL

| | | | | | |
|-------------------|-----|--|---------|-----|--------------|
| METAHYDRIN | | | | | |
| SANOFI AVENTIS US | 2MG | | N012594 | 001 | Jun 16, 1988 |
| | 4MG | | N012594 | 002 | Jun 16, 1988 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRICHLORMETHIAZIDE

TABLET; ORAL

NAQUA

| | | | |
|----------|-----|---------|-----|
| SCHERING | 2MG | N012265 | 001 |
| | 4MG | N012265 | 002 |

TRICHLOREX

| | | | |
|---------|-----|---------|-----|
| LANNETT | 4MG | A083436 | 001 |
| | 4MG | A085630 | 001 |

TRICHLORMAS

| | | | |
|---------|-----|---------|-----|
| MAST MM | 4MG | A086259 | 001 |
|---------|-----|---------|-----|

TRICHLORMETHIAZIDE

| | | | |
|--------------|-----|---------|-----|
| CHARTWELL RX | 4MG | A085568 | 001 |
| IMPAX LABS | 4MG | A083967 | 001 |
| PAR PHARM | 2MG | A087007 | 001 |
| | 4MG | A087005 | 001 |
| SANDOZ | 4MG | A086171 | 001 |
| WATSON LABS | 2MG | A083847 | 001 |
| | 2MG | A086458 | 001 |
| | 4MG | A083462 | 001 |
| | 4MG | A083855 | 001 |
| | 4MG | A085962 | 001 |

TRICLOFOS SODIUM

SOLUTION; ORAL

TRICLOS

| | | | |
|-------------------|------------|---------|-----|
| SANOFI AVENTIS US | 1.5GM/15ML | N016830 | 001 |
|-------------------|------------|---------|-----|

TABLET; ORAL

TRICLOS

| | | | |
|-------------------|-------|---------|-----|
| SANOFI AVENTIS US | 750MG | N016809 | 002 |
|-------------------|-------|---------|-----|

TRIDIHEXETHYL CHLORIDE

INJECTABLE; INJECTION

PATHILON

| | | | |
|---------|---------|---------|-----|
| LEDERLE | 10MG/ML | N009729 | 001 |
|---------|---------|---------|-----|

TABLET; ORAL

PATHILON

| | | | |
|---------|------|---------|-----|
| LEDERLE | 25MG | N009489 | 005 |
|---------|------|---------|-----|

TRIFLUOPERAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

STELAZINE

| | | | |
|-------------------|--------------------|---------|-----|
| + GLAXOSMITHKLINE | EQ 10MG BASE/ML ** | N011552 | 006 |
|-------------------|--------------------|---------|-----|

TRIFLUOPERAZINE HYDROCHLORIDE

| | | | | |
|--------------|-----------------|---------|-----|--------------|
| FOSUN PHARMA | EQ 10MG BASE/ML | A085787 | 001 | Apr 15, 1982 |
| WOCKHARDT | EQ 10MG BASE/ML | A088143 | 001 | Jul 26, 1983 |

INJECTABLE; INJECTION

STELAZINE

| | | | |
|-------------------|-------------------|---------|-----|
| + GLAXOSMITHKLINE | EQ 2MG BASE/ML ** | N011552 | 005 |
|-------------------|-------------------|---------|-----|

TABLET; ORAL

STELAZINE

| | | | |
|-------------------|-----------------|---------|-----|
| + GLAXOSMITHKLINE | EQ 1MG BASE ** | N011552 | 001 |
| | EQ 2MG BASE ** | N011552 | 002 |
| | EQ 5MG BASE ** | N011552 | 003 |
| | EQ 10MG BASE ** | N011552 | 004 |

TRIFLUOPERAZINE HYDROCHLORIDE

| | | | | |
|---------------------|--------------|---------|-----|--------------|
| DURAMED PHARMS BARR | EQ 1MG BASE | A088967 | 001 | Apr 23, 1985 |
| | EQ 2MG BASE | A088968 | 001 | Apr 23, 1985 |
| | EQ 5MG BASE | A088969 | 001 | Apr 23, 1985 |
| | EQ 10MG BASE | A088970 | 001 | Apr 23, 1985 |
| IVAX PHARMS | EQ 1MG BASE | A087612 | 001 | Nov 19, 1982 |
| | EQ 2MG BASE | A087613 | 001 | Nov 19, 1982 |
| | EQ 5MG BASE | A087328 | 001 | Nov 19, 1982 |
| | EQ 10MG BASE | A087614 | 001 | Nov 19, 1982 |
| SANDOZ | EQ 1MG BASE | A040153 | 001 | Oct 25, 1996 |
| | EQ 2MG BASE | A040153 | 002 | Oct 25, 1996 |
| | EQ 5MG BASE | A040153 | 003 | Oct 25, 1996 |
| | EQ 10MG BASE | A040153 | 004 | Oct 25, 1996 |
| WATSON LABS | EQ 1MG BASE | A085975 | 001 | Jun 23, 1988 |
| | EQ 2MG BASE | A085976 | 001 | Jun 23, 1988 |
| | EQ 5MG BASE | A085973 | 001 | Jun 23, 1988 |
| | EQ 10MG BASE | A088710 | 001 | Jun 23, 1988 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIFLUPROMAZINE

SUSPENSION; ORAL

VESPRIN

APOTHECON EQ 50MG HYDROCHLORIDE/5ML N011491 004

TRIFLUPROMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

VESPRIN

APOTHECON 3MG/ML N011325 005

10MG/ML N011325 004

20MG/ML N011325 001

TABLET; ORAL

VESPRIN

BRISTOL MYERS SQUIBB 10MG N011123 001

25MG N011123 002

50MG N011123 003

TRIHENXYPHENIDYL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

ARTANE

LEDERLE 5MG N006773 010

5MG N012947 001

ELIXIR; ORAL

ARTANE

LEDERLE 2MG/5ML N006773 009

TRIHENXYPHENIDYL HYDROCHLORIDE

PHARM VENTURES 2MG/5ML A089514 001 Apr 07, 1989

TABLET; ORAL

ARTANE

+ LEDERLE 2MG ** N006773 005

+ 5MG ** N006773 003

TREMIM

SCHERING 2MG A080381 001

5MG A080381 003

TRIHENXYPHENIDYL HYDROCHLORIDE

HIKMA PHARMS 2MG A040337 002 Feb 16, 2000

5MG A040337 001 Feb 16, 2000

NYLOS 5MG A085622 001

VANGARD 2MG A088035 001 Jul 30, 1982

WATSON LABS 2MG A040184 001 Feb 06, 1998

2MG A085117 001

5MG A040184 002 Feb 06, 1998

5MG A085105 001

TRILOSTANE

CAPSULE; ORAL

MODRASTANE

BIOENVISION 30MG N018719 002 Dec 31, 1984

60MG N018719 001 Dec 31, 1984

TRIMEPRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

TEMARIL

ALLERGAN HERBERT EQ 5MG BASE N011316 004

SYRUP; ORAL

TEMARIL

ALLERGAN HERBERT EQ 2.5MG BASE/5ML N011316 003

TRIMEPRAZINE TARTRATE

ALPHARMA US PHARMS EQ 2.5MG BASE/5ML A085015 001 Feb 18, 1982

MORTON GROVE EQ 2.5MG BASE/5ML A088285 001 Apr 11, 1985

TABLET; ORAL

TEMARIL

ALLERGAN HERBERT EQ 2.5MG BASE N011316 001

TRIMETHADIONE

CAPSULE; ORAL

TRIDIONE

ABBVIE 300MG N005856 005

SOLUTION; ORAL

TRIDIONE

ABBVIE 200MG/5ML N005856 002

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIMETHAPHAN CAMSYLATE

INJECTABLE; INJECTION

ARFONAD

| | | | |
|-------|---------|---------|-----|
| ROCHE | 50MG/ML | N008983 | 001 |
|-------|---------|---------|-----|

TRIMETHOBENZAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

TRIMETHOBENZAMIDE HYDROCHLORIDE

| | | | | |
|------------------|----------|---------|-----|--------------|
| HOSPIRA | 100MG/ML | A088804 | 001 | Apr 03, 1987 |
| SMITH AND NEPHEW | 100MG/ML | A088960 | 001 | Apr 04, 1986 |
| | 100MG/ML | A089043 | 001 | Apr 04, 1986 |
| SOLOPAK | 100MG/ML | A089094 | 001 | Apr 04, 1986 |
| WATSON LABS | 100MG/ML | A086577 | 001 | Oct 19, 1982 |
| | 100MG/ML | A087939 | 001 | Dec 28, 1982 |

TRIMETHOPRIM

TABLET; ORAL

PROLOPRIM

| | | | | |
|----------------|-------|---------|-----|--------------|
| MONARCH PHARMS | 100MG | N017943 | 001 | |
| | 200MG | N017943 | 003 | Jul 14, 1982 |

TRIMETHOPRIM

| | | | | |
|----------------------|----------|---------|-----|--------------|
| SUN PHARM INDUSTRIES | 100MG | A070494 | 001 | Jan 22, 1986 |
| | 200MG | A070495 | 001 | Sep 24, 1986 |
| TEVA | 200MG ** | A071259 | 001 | Jun 18, 1987 |

TRIMPEX

| | | | | |
|-------|-------|---------|-----|--|
| ROCHE | 100MG | N017952 | 001 | |
|-------|-------|---------|-----|--|

TRIMPEX 200

| | | | | |
|-------|-------|---------|-----|--------------|
| ROCHE | 200MG | N017952 | 002 | Nov 09, 1982 |
|-------|-------|---------|-----|--------------|

TRIMETHOPRIM HYDROCHLORIDE

SOLUTION; ORAL

PRIMSOL

| | | | | |
|---------|------------------|---------|-----|--------------|
| ALLEGIS | EQ 25MG BASE/5ML | N074374 | 001 | Jun 23, 1995 |
|---------|------------------|---------|-----|--------------|

TRIMETREXATE GLUCURONATE

INJECTABLE; INJECTION

NEUTREXIN

| | | | | |
|--------------------|--------------------|---------|-----|--------------|
| MEDIMMUNE ONCOLOGY | EQ 25MG BASE/VIAL | N020326 | 001 | Dec 17, 1993 |
| | EQ 200MG BASE/VIAL | N020326 | 002 | Jul 31, 1998 |

TRIMIPRAMINE MALEATE

CAPSULE; ORAL

TRIMIPRAMINE MALEATE

| | | | | |
|------------|---------------|---------|-----|--------------|
| USL PHARMA | EQ 25MG BASE | A071283 | 001 | Dec 08, 1987 |
| | EQ 50MG BASE | A071284 | 001 | Dec 08, 1987 |
| | EQ 100MG BASE | A071285 | 001 | Dec 08, 1987 |

TRIOXSALEN

TABLET; ORAL

TRISORALEN

| | | | | |
|--------------------|-----|---------|-----|--|
| VALEANT PHARM INTL | 5MG | N012697 | 001 | |
|--------------------|-----|---------|-----|--|

TRIPLENNAMINE CITRATE

ELIXIR; ORAL

PBZ

| | | | | |
|----------|---------------------------|---------|-----|--|
| NOVARTIS | EQ 25MG HYDROCHLORIDE/5ML | N005914 | 004 | |
|----------|---------------------------|---------|-----|--|

TRIPLENNAMINE HYDROCHLORIDE

TABLET; ORAL

PBZ

| | | | | |
|----------|------|---------|-----|--|
| NOVARTIS | 25MG | A083149 | 001 | |
| | 50MG | N005914 | 002 | |

TRIPLENNAMINE HYDROCHLORIDE

| | | | | |
|-------------|------|---------|-----|--|
| ANABOLIC | 50MG | A083037 | 001 | |
| BARR | 50MG | A080744 | 001 | |
| HEATHER | 50MG | A083989 | 001 | |
| IMPAX LABS | 50MG | A080785 | 001 | |
| LANNETT | 50MG | A083557 | 001 | |
| NYLOS | 50MG | A085412 | 001 | |
| PARKE DAVIS | 25MG | A083625 | 001 | |
| | 50MG | A083626 | 001 | |
| WATSON LABS | 50MG | A080713 | 001 | |
| | 50MG | A080790 | 001 | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRIPLENNAMINE HYDROCHLORIDE

TABLET; ORAL

TRIPLENNAMINE HYDROCHLORIDE

50MG

A085188 001

TABLET, EXTENDED RELEASE; ORAL

PBZ-SR

NOVARTIS

50MG

N010533 002

100MG

N010533 001

TRIPLE SULFA (SULFABENZAMIDE; SULFACETAMIDE; SULFATHIAZOLE)

CREAM; VAGINAL

GYNE-SULF

G AND W LABS

3.7%; 2.86%; 3.42%

A088607 001 Jun 09, 1986

SULTRIN

ORTHO MCNEIL PHARM

3.7%; 2.86%; 3.42%

N005794 001

TRIPLE SULFA

ALPHARMA US PHARMS

3.7%; 2.86%; 3.42%

A087864 001 Sep 01, 1982

FOUGERA

3.7%; 2.86%; 3.42%

A086424 001

PERRIGO NEW YORK

3.7%; 2.86%; 3.42%

A087285 001 Nov 15, 1982

TRYSUL

SAVAGE LABS

3.7%; 2.86%; 3.42%

A087887 001 Jul 23, 1982

VAGILIA

G AND W LABS INC

3.7%; 2.86%; 3.42%

A088821 001 Nov 09, 1987

TABLET; VAGINAL

SULTRIN

ORTHO MCNEIL PHARM

184MG; 143.75MG; 172.5MG

N005794 002

TRIPLE SULFA

FOUGERA

184MG; 143.75MG; 172.5MG

A088463 001 Jan 03, 1985

PHARMADERM

184MG; 143.75MG; 172.5MG

A088462 001 Jan 03, 1985

TRIPROLIDINE HYDROCHLORIDE

SYRUP; ORAL

ACTIDIL

GLAXOSMITHKLINE

1.25MG/5ML

N011496 002 Jul 01, 1983

MYIDYL

USL PHARMA

1.25MG/5ML

A087963 001 Jan 18, 1983

TRIPROLIDINE HYDROCHLORIDE

ALPHARMA US PHARMS

1.25MG/5ML

A085940 001

HALSEY

1.25MG/5ML

A088735 001 Jan 17, 1985

PHARM ASSOC

1.25MG/5ML

A087514 001 Feb 10, 1982

TABLET; ORAL

ACTIDIL

GLAXOSMITHKLINE

2.5MG

N011110 002 Jul 01, 1983

TRIPROLIDINE HYDROCHLORIDE

VITARINE

2.5MG

A085610 001

WATSON LABS

2.5MG

A085094 001

TRISULFAPYRIMIDINES (SULFADIAZINE; SULFAMERAZINE; SULFAMETHAZINE)

SUSPENSION; ORAL

LANTRISUL

LANNETT

167MG/5ML; 167MG/5ML; 167MG/5ML

A080123 002

NEOTRIZINE

LILLY

167MG/5ML; 167MG/5ML; 167MG/5ML

N006317 012

SULFALOID

FOREST PHARMS

167MG/5ML; 167MG/5ML; 167MG/5ML

A080100 001

SULFOSE

WYETH AYERST

167MG/5ML; 167MG/5ML; 167MG/5ML

A080013 002

TERFONYL

BRISTOL MYERS SQUIBB

167MG/5ML; 167MG/5ML; 167MG/5ML

N006904 002

TRIPLE SULFA

ALPHARMA US PHARMS

167MG/5ML; 167MG/5ML; 167MG/5ML

A080280 001

TRIPLE SULFAS

LEDERLE

167MG/5ML; 167MG/5ML; 167MG/5ML

N006920 003

TABLET; ORAL

NEOTRIZINE

LILLY

167MG; 167MG; 167MG

N006317 011

SULFA-TRIPLE #2

IMPAX LABS

167MG; 167MG; 167MG

A080079 001

SULFALOID

FOREST PHARMS

167MG; 167MG; 167MG

A080099 001

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TRISULFAPYRIMIDINES (SULFADIAZINE;SULFAMERAZINE;SULFAMETHAZINE)

TABLET;ORAL

SULFOSE

WYETH AYERST 167MG;167MG;167MG A080013 001

TERFONYL

BRISTOL MYERS SQUIBB 167MG;167MG;167MG N006904 001

TRIPLE SULFA

PUREPAC PHARM 167MG;167MG;167MG A080086 001

TRIPLE SULFAS

LEDERLE 167MG;167MG;167MG N006920 002

TRIPLE SULFOID

PAL PAK 167MG;167MG;167MG A080094 001

TROGLITAZONE

TABLET;ORAL

PRELAY

SANKYO 200MG N020719 001 Jan 29, 1997

300MG N020719 003 Aug 04, 1997

400MG N020719 002 Jan 29, 1997

REZULIN

PFIZER PHARMS 200MG N020720 001 Jan 29, 1997

300MG N020720 003 Aug 04, 1997

400MG N020720 002 Jan 29, 1997

TROLAMINE POLYPEPTIDE OLEATE CONDENSATE

SOLUTION/DROPS;OTIC

CERUMENEX

PHARM RES ASSOC 10% N011340 002

TROLEANDOMYCIN

CAPSULE;ORAL

TAO

PFIZER EQ 250MG BASE N050336 002

SUSPENSION;ORAL

TAO

PFIZER EQ 125MG BASE/5ML N050332 001

TROMETHAMINE

INJECTABLE;INJECTION

THAM

+ HOSPIRA 3.6GM/100ML N013025 002

TROPICAMIDE

SOLUTION/DROPS;OPHTHALMIC

MYDRIACYL

ALCON 0.5% ** N012111 002

1% ** N012111 004

MYDRIAFAIR

PHARMAFAIR 0.5% A088274 001 Sep 16, 1983

1% A088230 001 Sep 16, 1983

TROPICAMIDE

AKORN 1% A088447 001 Aug 28, 1985

ALCON PHARMS LTD 1% A089172 001 Dec 28, 1990

MIZA PHARMS USA 0.5% A087636 001 Jul 30, 1982

1% A087637 001 Aug 09, 1982

WATSON LABS 0.5% A089171 001 Dec 28, 1990

TROSPIUM CHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

SANCTURA XR

+ ALLERGAN 60MG ** N022103 001 Aug 03, 2007

TROSPIUM CHLORIDE

UPSHER SMITH LABS 60MG A091635 001 Apr 29, 2015

TABLET;ORAL

SANCTURA

+ ALLERGAN 20MG ** N021595 001 May 28, 2004

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

TROVAFLOXACIN MESYLATE

TABLET; ORAL

TROVAN

PFIZER

EQ 100MG BASE

N020759 001 Dec 18, 1997

EQ 200MG BASE

N020759 002 Dec 18, 1997

TUBOCURARINE CHLORIDE

INJECTABLE; INJECTION

TUBOCURARINE CHLORIDE

BRISTOL MYERS SQUIBB

3MG/ML

N005657 001

HOSPIRA

3MG/ML

N006095 001

LILLY

3MG/ML

N006325 001

TYROPANOATE SODIUM

CAPSULE; ORAL

BILOPAQUE

GE HEALTHCARE

750MG

N013731 001

UNOPROSTONE ISOPROPYL

SOLUTION/DROPS; OPHTHALMIC

RESCULA

+ SUCAMPO PHARMA LLC

0.15% **

N021214 001 Aug 03, 2000

URACIL MUSTARD

CAPSULE; ORAL

URACIL MUSTARD

SHIRE

1MG

N012892 001

UREA

INJECTABLE; INJECTION

STERILE UREA

HOSPIRA

40GM/VIAL

N017698 001

UREAPHIL

HOSPIRA

40GM/VIAL

N012154 001

UREA C-13

FOR SOLUTION; ORAL

BREATHTEK UBT FOR H-PYLORI

OTSUKA AMERICA

EQ 75MG/POUCH

N020586 002 May 10, 2001

HELICOSOL

METABOLIC SOLUTIONS

125MG/VIAL

N021092 001 Dec 17, 1999

MERETEK UBT KIT (W/ PRANACTIN)

OTSUKA AMERICA

125MG/VIAL

N020586 001 Sep 17, 1996

PYLORI-CHEK BREATH TEST

DXS DEVICES

100MG/VIAL

N020900 001 Feb 04, 1999

UROFOLLITROPIN

INJECTABLE; INTRAMUSCULAR

METRODIN

SERONO

75 IU/AMP

N019415 002 Sep 18, 1986

150 IU/AMP

N019415 003 Sep 18, 1986

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

BRAVELLE

+ FERRING

75 IU/VIAL

N021289 001 May 06, 2002

INJECTABLE; SUBCUTANEOUS

FERTINEX

SERONO

75 IU/AMP

N019415 005 Aug 23, 1996

150 IU/AMP

N019415 004 Aug 23, 1996

UROKINASE

INJECTABLE; INJECTION

KINLYTIC

MICROBIX BIOSYSTEMS

5,000 IU/VIAL

N021846 003

9,000 IU/VIAL

N021846 002

250,000 IU/VIAL

N021846 001

URSODIOL

CAPSULE; ORAL

ACTIGALL

ALLERGAN SALES LLC

150MG

N019594 001 Dec 31, 1987

URSODIOL

IMPAX LABS INC

300MG

A077895 001 Jul 27, 2006

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

URSODIOL

TABLET; ORAL

URSODIOL

| | | | |
|-----------------|-------|-------------|--------------|
| TEVA PHARMS USA | 250MG | A079184 001 | May 13, 2009 |
| | 500MG | A079184 002 | May 13, 2009 |

VALACYCLOVIR HYDROCHLORIDE

TABLET; ORAL

VALACYCLOVIR HYDROCHLORIDE

| | | | |
|-------|---------------|-------------|--------------|
| MYLAN | EQ 500MG BASE | A078070 001 | May 24, 2010 |
| | EQ 1GM BASE | A078070 002 | May 24, 2010 |

VALDECOXIB

TABLET; ORAL

BEXTRA

| | | | |
|-----------|------|-------------|--------------|
| GD SEARLE | 10MG | N021341 002 | Nov 16, 2001 |
| | 20MG | N021341 003 | Nov 16, 2001 |

VALPROIC ACID

CAPSULE; ORAL

VALPROIC ACID

| | | | |
|------------|-------|-------------|--------------|
| PAR PHARM | 250MG | A070431 001 | Feb 28, 1986 |
| SCHERER RP | 250MG | A070195 001 | Jul 02, 1987 |
| USL PHARMA | 250MG | A070631 001 | Jun 11, 1987 |

CAPSULE, DELAYED RELEASE; ORAL

STAVZOR

| | | | |
|------------------|----------|-------------|--------------|
| + BIONPHARMA INC | 125MG ** | N022152 001 | Jul 29, 2008 |
| | 250MG ** | N022152 002 | Jul 29, 2008 |
| | 500MG ** | N022152 003 | Jul 29, 2008 |

SYRUP; ORAL

VALPROIC ACID

| | | | |
|------------|-----------|-------------|--------------|
| APOTEX INC | 250MG/5ML | A077105 001 | Jul 29, 2005 |
|------------|-----------|-------------|--------------|

VALSARTAN

CAPSULE; ORAL

DIOVAN

| | | | |
|----------|-------|-------------|--------------|
| NOVARTIS | 80MG | N020665 001 | Dec 23, 1996 |
| | 160MG | N020665 002 | Dec 23, 1996 |

SOLUTION; ORAL

PREXXARTAN

| | | | |
|----------------------|-----------|-------------|--------------|
| + CARMEL BIOSCIENCES | 20MG/5ML | N209139 001 | Dec 19, 2017 |
| | 80MG/20ML | N209139 002 | Dec 19, 2017 |

TABLET; ORAL

VALSARTAN

| | | | |
|-----------------|-------|-------------|--------------|
| WATSON LABS INC | 40MG | A090642 001 | Jan 05, 2015 |
| | 80MG | A090642 002 | Jan 05, 2015 |
| | 160MG | A090642 003 | Jan 05, 2015 |
| | 320MG | A090642 004 | Jan 05, 2015 |

VANCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

VANCOMYCIN HYDROCHLORIDE

| | | | |
|--------------------|---------------|-------------|--------------|
| FRESENIUS KABI USA | EQ 125MG BASE | A065453 001 | Jun 18, 2012 |
| | EQ 250MG BASE | A065453 002 | Jun 18, 2012 |

FOR SOLUTION; ORAL

VANCOCIN HYDROCHLORIDE

| | | | |
|----------------|-------------------|-------------|--------------|
| ANI PHARMS INC | EQ 250MG BASE/5ML | A061667 002 | Jul 13, 1983 |
| | EQ 500MG BASE/6ML | A061667 001 | |

VANCOLED

| | | | |
|---------|-------------------|-------------|--------------|
| LEDERLE | EQ 250MG BASE/5ML | A063321 002 | Oct 15, 1993 |
| | EQ 500MG BASE/6ML | A063321 003 | Oct 15, 1993 |

INJECTABLE; INJECTION

VANCOCIN HYDROCHLORIDE

| | | | |
|----------------|-----------------------|-------------|--------------|
| ANI PHARMS INC | EQ 500MG BASE/VIAL ** | A060180 001 | |
| | EQ 500MG BASE/VIAL | A062476 001 | Mar 15, 1984 |
| | EQ 500MG BASE/VIAL | A062716 001 | Mar 13, 1987 |
| | EQ 500MG BASE/VIAL ** | A062812 001 | Nov 17, 1987 |
| | EQ 1GM BASE/VIAL ** | A060180 002 | Mar 21, 1986 |
| | EQ 1GM BASE/VIAL | A062476 002 | Mar 21, 1986 |
| | EQ 1GM BASE/VIAL | A062716 002 | Mar 13, 1987 |
| | EQ 1GM BASE/VIAL ** | A062812 002 | Nov 17, 1987 |
| | EQ 10GM BASE/VIAL ** | A062812 003 | Nov 17, 1987 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

VANCOLED

| | | | |
|----------------------|-----------------------|-------------|--------------|
| WEST-WARD PHARMS INT | EQ 500MG BASE/VIAL ** | A062682 001 | Jul 22, 1986 |
| | EQ 1GM BASE/VIAL ** | A062682 002 | Mar 30, 1988 |
| | EQ 2GM BASE/VIAL ** | A062682 003 | May 11, 1988 |
| | EQ 5GM BASE/VIAL ** | A062682 004 | May 11, 1988 |
| | EQ 10GM BASE/VIAL ** | A062682 005 | May 11, 1988 |

VANCOMYCIN HYDROCHLORIDE

| | | | |
|----------------------|--------------------|-------------|--------------|
| MYLAN LABS LTD | EQ 10GM BASE/VIAL | A091469 001 | Jul 01, 2011 |
| TEVA PHARMS USA | EQ 500MG BASE/VIAL | A201251 001 | Dec 23, 2015 |
| | EQ 1GM BASE/VIAL | A201251 002 | Dec 23, 2015 |
| | EQ 5GM BASE/VIAL | A201250 001 | Dec 23, 2015 |
| | EQ 10GM BASE/VIAL | A201250 002 | Dec 23, 2015 |
| WEST-WARD PHARMS INT | EQ 500MG BASE/VIAL | A062879 001 | Aug 02, 1988 |
| | EQ 1GM BASE/VIAL | A062879 002 | Aug 02, 1988 |
| XELLIA PHARMS APS | EQ 500MG BASE/VIAL | A091377 001 | Sep 09, 2015 |
| | EQ 1GM BASE/VIAL | A091377 002 | Sep 09, 2015 |
| | EQ 5GM BASE/VIAL | A206243 001 | Dec 23, 2015 |
| | EQ 10GM BASE/VIAL | A206243 002 | Dec 23, 2015 |

VANCOR

| | | | |
|----------------------|--------------------|-------------|--------------|
| PHARMACIA AND UPJOHN | EQ 500MG BASE/VIAL | A062956 001 | Aug 01, 1988 |
| | EQ 1GM BASE/VIAL | A062956 002 | Aug 01, 1988 |

VARDENAFIL HYDROCHLORIDE

TABLET; ORAL

LEVITRA

| | | | |
|------------------|-------|-------------|--------------|
| + BAYER HLTHCARE | 2.5MG | N021400 003 | Aug 19, 2003 |
|------------------|-------|-------------|--------------|

VASOPRESSIN TANNATE

INJECTABLE; INJECTION

PITRESSIN TANNATE

| | | | |
|---------------|----------------------|-------------|--|
| + PARKE DAVIS | 5PRESSOR UNITS/ML ** | N003402 001 | |
|---------------|----------------------|-------------|--|

VECURONIUM BROMIDE

INJECTABLE; INJECTION

NORCURON

| | | | |
|-------------------|--------------|-------------|--------------|
| + ORGANON USA INC | 10MG/VIAL ** | N018776 002 | Apr 30, 1984 |
| | 20MG/VIAL ** | N018776 003 | Jan 03, 1992 |

VECURONIUM BROMIDE

| | | | |
|----------------------|-----------|-------------|--------------|
| HOSPIRA | 4MG/VIAL | A075558 001 | Sep 11, 2001 |
| WATSON LABS | 10MG/VIAL | A074334 001 | Aug 31, 1995 |
| | 20MG/VIAL | A074334 002 | Aug 31, 1995 |
| WEST-WARD PHARMS INT | 10MG/VIAL | A075218 001 | Aug 23, 1999 |
| | 20MG/VIAL | A075218 002 | Aug 23, 1999 |

VELAGLUCERASE ALFA

POWDER; INTRAVENOUS

VPRIV

| | | | |
|---------------------|----------------|-------------|--------------|
| SHIRE HUMAN GENETIC | 200 UNITS/VIAL | N022575 002 | Feb 26, 2010 |
|---------------------|----------------|-------------|--------------|

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

EFFEXOR XR

| | | | |
|--------------|---------------|-------------|--------------|
| WYETH PHARMS | EQ 100MG BASE | N020699 003 | Oct 20, 1997 |
|--------------|---------------|-------------|--------------|

VENLAFAXINE HYDROCHLORIDE

| | | | |
|---------------|----------------|-------------|--------------|
| ANCHEN PHARMS | EQ 37.5MG BASE | A078087 001 | Mar 16, 2012 |
| | EQ 75MG BASE | A078087 002 | Mar 16, 2012 |
| | EQ 150MG BASE | A078087 003 | Mar 16, 2012 |
| MYLAN | EQ 37.5MG BASE | A078789 001 | Jun 01, 2011 |
| | EQ 75MG BASE | A078789 002 | Jun 01, 2011 |
| | EQ 150MG BASE | A078789 003 | Jun 01, 2011 |

TABLET; ORAL

EFFEXOR

| | | | |
|--------------------|-------------------|-------------|--------------|
| + WYETH PHARMS INC | EQ 12.5MG BASE ** | N020151 001 | Dec 28, 1993 |
| | EQ 25MG BASE ** | N020151 002 | Dec 28, 1993 |
| | EQ 37.5MG BASE ** | N020151 006 | Dec 28, 1993 |
| | EQ 50MG BASE ** | N020151 003 | Dec 28, 1993 |
| | EQ 75MG BASE ** | N020151 004 | Dec 28, 1993 |
| | EQ 100MG BASE ** | N020151 005 | Dec 28, 1993 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VENLAFAXINE HYDROCHLORIDE

TABLET; ORAL

VENLAFAXINE HYDROCHLORIDE

| | | | |
|--------------------|----------------|-------------|--------------|
| FOSUN PHARMA | EQ 25MG BASE | A077515 001 | Jun 13, 2008 |
| | EQ 37.5MG BASE | A077515 002 | Jun 13, 2008 |
| | EQ 50MG BASE | A077515 003 | Jun 13, 2008 |
| | EQ 75MG BASE | A077515 004 | Jun 13, 2008 |
| | EQ 100MG BASE | A077515 005 | Jun 13, 2008 |
| PLIVA HRVATSKA DOO | EQ 25MG BASE | A078517 001 | Jun 13, 2008 |
| | EQ 37.5MG BASE | A078517 002 | Jun 13, 2008 |
| | EQ 50MG BASE | A078517 003 | Jun 13, 2008 |
| | EQ 75MG BASE | A078517 004 | Jun 13, 2008 |
| | EQ 100MG BASE | A078517 005 | Jun 13, 2008 |

VERAPAMIL HYDROCHLORIDE

INJECTABLE; INJECTION

CALAN

| | | | |
|-------------------------|-------------|-------------|--------------|
| GD SEARLE LLC | 2.5MG/ML | N019038 001 | Mar 30, 1984 |
| ISOPTIN | | | |
| + MT ADAMS | 2.5MG/ML ** | N018485 001 | |
| VERAPAMIL HYDROCHLORIDE | | | |
| ABRAXIS PHARM | 2.5MG/ML | A070348 001 | May 01, 1986 |
| BEDFORD | 2.5MG/ML | A072888 001 | Jul 28, 1995 |
| HOSPIRA | 2.5MG/ML | A070577 001 | Feb 02, 1987 |
| | 2.5MG/ML | A070739 001 | May 06, 1987 |
| | 2.5MG/ML | A070740 001 | May 06, 1987 |
| INTL MEDICATION | 2.5MG/ML | A070451 001 | Dec 16, 1985 |
| LUITPOLD | 2.5MG/ML | A070225 001 | Nov 12, 1985 |
| | 2.5MG/ML | A070617 001 | Nov 12, 1985 |
| MARSAM PHARMS LLC | 2.5MG/ML | A072233 001 | Feb 26, 1993 |
| | 2.5MG/ML | A073485 001 | Sep 27, 1993 |
| SMITH AND NEPHEW | 2.5MG/ML | A070696 001 | Jul 31, 1987 |
| | 2.5MG/ML | A070697 001 | Jul 31, 1987 |
| SOLOPAK | 2.5MG/ML | A070695 001 | Jul 31, 1987 |

TABLET; ORAL

CALAN

| | | | |
|-------------------------|-------|-------------|--------------|
| GD SEARLE LLC | 40MG | N018817 003 | Feb 23, 1988 |
| | 160MG | N018817 004 | Feb 23, 1988 |
| ISOPTIN | | | |
| + MT ADAMS | 40MG | N018593 003 | Nov 23, 1987 |
| + | 80MG | N018593 001 | Mar 08, 1982 |
| + | 120MG | N018593 002 | Mar 08, 1982 |
| VERAPAMIL HYDROCHLORIDE | | | |
| ACTAVIS ELIZABETH | 80MG | A071019 001 | Sep 24, 1986 |
| | 120MG | A070468 001 | Sep 24, 1986 |
| MUTUAL PHARM | 80MG | A070482 001 | Sep 24, 1986 |
| | 120MG | A070483 001 | Sep 24, 1986 |
| PLIVA | 40MG | A072751 001 | Feb 23, 1996 |
| | 80MG | A072124 001 | Jan 26, 1989 |
| | 120MG | A072125 001 | Jan 26, 1989 |
| SUN PHARM INDUSTRIES | 80MG | A071489 002 | Jan 13, 1988 |
| | 120MG | A071489 001 | Jan 13, 1988 |
| WARNER CHILCOTT | 80MG | A070340 001 | Sep 24, 1986 |
| | 120MG | A070341 001 | Sep 24, 1986 |
| WATSON LABS | 40MG | A072799 001 | Apr 28, 1989 |
| | 40MG | A072923 001 | Jun 29, 1993 |
| | 80MG | A070855 001 | Sep 24, 1986 |
| | 80MG | A071366 001 | Oct 01, 1986 |
| | 120MG | A070856 001 | Sep 24, 1986 |
| | 120MG | A071367 001 | Oct 01, 1986 |
| YAOPHARMA CO LTD | 40MG | A073168 001 | Jul 31, 1992 |
| | 80MG | A071423 001 | May 24, 1988 |
| | 120MG | A071424 001 | May 25, 1988 |

TABLET, EXTENDED RELEASE; ORAL

CALAN SR

| | | | |
|---------------|----------|-------------|--------------|
| + PFIZER | 180MG ** | N019152 002 | Dec 15, 1989 |
| COVERA-HS | | | |
| GD SEARLE LLC | 180MG | N020552 001 | Feb 26, 1996 |
| | 240MG | N020552 002 | Feb 26, 1996 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

VERAPAMIL HYDROCHLORIDE

| | | | |
|-------------|-------|-------------|--------------|
| APOTEX CORP | 120MG | A200878 001 | Apr 20, 2012 |
| | 180MG | A200878 002 | Apr 20, 2012 |
| | 240MG | A200878 003 | Apr 20, 2012 |
| PLIVA | 240MG | A072922 001 | Mar 01, 1996 |

VERATRUM VIRIDE ROOT

TABLET;ORAL

VERTAVIS

| | | | |
|---------------------|-------------|-------------|--|
| MEDPOINTE PHARM HLC | 130CSR UNIT | N005691 002 | |
|---------------------|-------------|-------------|--|

VIDARABINE

INJECTABLE;INJECTION

VIRA-A

| | | | |
|------------|--------------------|-------------|--|
| PARKEDEALE | EQ 187.4MG BASE/ML | N050523 001 | |
|------------|--------------------|-------------|--|

OINTMENT;OPHTHALMIC

VIRA-A

| | | | |
|------------|----|-------------|--|
| PARKEDEALE | 3% | N050486 001 | |
|------------|----|-------------|--|

VINBLASTINE SULFATE

INJECTABLE;INJECTION

VELBAN

+ LILLY

| | | | |
|--|-----------|-------------|--|
| | 10MG/VIAL | N012665 001 | |
|--|-----------|-------------|--|

VINBLASTINE SULFATE

ABRAXIS PHARM

| | | | |
|--|-----------|-------------|--------------|
| | 10MG/VIAL | A089011 001 | Nov 18, 1985 |
|--|-----------|-------------|--------------|

HOSPIRA

| | | | |
|--|-----------|-------------|--------------|
| | 10MG/VIAL | A089565 001 | Aug 18, 1987 |
|--|-----------|-------------|--------------|

VINCRISTINE SULFATE

INJECTABLE;INJECTION

ONCOVIN

LILLY

| | | | |
|--|----------|-------------|--|
| | 1MG/VIAL | N014103 001 | |
|--|----------|-------------|--|

1MG/ML

| | | | |
|--|--|-------------|--------------|
| | | N014103 003 | Mar 07, 1984 |
|--|--|-------------|--------------|

5MG/VIAL

| | | | |
|--|--|-------------|--|
| | | N014103 002 | |
|--|--|-------------|--|

VINCASAR PFS

TEVA PARENTERAL

| | | | |
|--|--------|-------------|--------------|
| | 1MG/ML | A071426 001 | Jul 17, 1987 |
|--|--------|-------------|--------------|

VINCREX

BRISTOL MYERS SQUIBB

| | | | |
|--|----------|-------------|--------------|
| | 5MG/VIAL | A070867 001 | Jul 12, 1988 |
|--|----------|-------------|--------------|

VINCRISTINE SULFATE

ABIC

| | | | |
|--|--------|-------------|--------------|
| | 1MG/ML | A070873 001 | Feb 19, 1987 |
|--|--------|-------------|--------------|

ABRAXIS PHARM

| | | | |
|--|--------|-------------|--------------|
| | 1MG/ML | A070411 001 | Sep 10, 1986 |
|--|--------|-------------|--------------|

FRESENIUS KABI USA

| | | | |
|--|--------|-------------|--------------|
| | 1MG/ML | A076296 001 | Dec 20, 2002 |
|--|--------|-------------|--------------|

HOSPIRA

| | | | |
|--|--------|-------------|--------------|
| | 1MG/ML | A076401 001 | Oct 28, 2003 |
|--|--------|-------------|--------------|

HOSPIRA

| | | | |
|--|----------|-------------|--------------|
| | 1MG/VIAL | A071559 001 | Apr 11, 1988 |
|--|----------|-------------|--------------|

| | | | |
|--|----------|-------------|--------------|
| | 2MG/VIAL | A071560 001 | Apr 11, 1988 |
|--|----------|-------------|--------------|

| | | | |
|--|----------|-------------|--------------|
| | 5MG/VIAL | A071561 001 | Apr 11, 1988 |
|--|----------|-------------|--------------|

VINORELBINE TARTRATE

INJECTABLE;INJECTION

VINORELBINE TARTRATE

EBEWE PHARMA

| | | | |
|--|-----------------|-------------|--------------|
| | EQ 10MG BASE/ML | A078408 001 | Feb 13, 2008 |
|--|-----------------|-------------|--------------|

MYLAN LABS LTD

| | | | |
|--|-----------------|-------------|--------------|
| | EQ 10MG BASE/ML | A200148 001 | Aug 31, 2012 |
|--|-----------------|-------------|--------------|

VIOMYCIN SULFATE

INJECTABLE;INJECTION

VIOCIN SULFATE

PFIZER

| | | | |
|--|------------------|-------------|--|
| | EQ 1GM BASE/VIAL | A061086 001 | |
|--|------------------|-------------|--|

| | | | |
|--|------------------|-------------|--|
| | EQ 5GM BASE/VIAL | A061086 002 | |
|--|------------------|-------------|--|

VITAMIN A

CAPSULE;ORAL

AQUASOL A

ASTRAZENECA

| | | | |
|--|-----------------|-------------|--|
| | 25,000USP UNITS | A083080 002 | |
|--|-----------------|-------------|--|

| | | | |
|--|-----------------|-------------|--|
| | 50,000USP UNITS | A083080 001 | |
|--|-----------------|-------------|--|

VITAMIN A

BANNER PHARMACAPS

| | | | |
|--|-----------------|-------------|--|
| | 50,000USP UNITS | A083973 001 | |
|--|-----------------|-------------|--|

CHASE CHEM

| | | | |
|--|-----------|-------------|--|
| | 50,000 IU | A083351 001 | |
|--|-----------|-------------|--|

EVERYLIFE

| | | | |
|--|-----------|-------------|--|
| | 50,000 IU | A083134 001 | |
|--|-----------|-------------|--|

IMPAX LABS

| | | | |
|--|-----------------|-------------|--|
| | 50,000USP UNITS | A080952 001 | |
|--|-----------------|-------------|--|

WEST WARD

| | | | |
|--|-----------------|-------------|--|
| | 50,000USP UNITS | A080985 001 | |
|--|-----------------|-------------|--|

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

VITAMIN A PALMITATE

CAPSULE; ORAL

AFAXIN

STERLING WINTHROP EQ 50,000 UNITS BASE A083187 001

ALPHALIN

LILLY EQ 50,000 UNITS BASE A080883 001

DEL-VI-A

DEL RAY LABS EQ 50,000 UNITS BASE A080830 001

VI-DOM-A

BAYER PHARMS EQ 50,000 UNITS BASE A080972 001

VITAMIN A

BANNER PHARMACAPS EQ 50,000 UNITS BASE A080702 001

BRISTOL MYERS SQUIBB EQ 50,000 UNITS BASE A080860 001

CHASE CHEM EQ 50,000 UNITS BASE A080746 001

EQ 50,000 UNITS BASE A083207 001

ELKINS SINN EQ 50,000 UNITS BASE A085479 001

EVERYLIFE EQ 50,000 UNITS BASE A080943 001

EQ 50,000 UNITS BASE A083114 001

IMPAX LABS EQ 50,000 UNITS BASE A080953 001

EQ 50,000 UNITS BASE A080955 001

IVAX SUB TEVA PHARMS EQ 50,000 UNITS BASE A083035 001

EQ 50,000 UNITS BASE A083190 001

MK LABS EQ 25,000 UNITS BASE A083457 002

EQ 50,000 UNITS BASE A083457 001

WEST WARD EQ 50,000 UNITS BASE A080967 001

WHARTON LABS EQ 50,000 UNITS BASE A083665 001

VITAMIN A PALMITATE

ARCUM EQ 50,000 UNITS BASE A083311 001

EQ 50,000 UNITS BASE A083321 001

BANNER PHARMACAPS EQ 50,000 UNITS BASE A083948 001

EQ 50,000 UNITS BASE A083981 001

VITAMIN A SOLUBILIZED

TEVA EQ 50,000 UNITS BASE A080921 001

INJECTABLE; INJECTION

VITAMIN A PALMITATE

BEL MAR EQ 50,000 UNITS BASE/ML A080819 001

VORICONAZOLE

FOR SUSPENSION; ORAL

VORICONAZOLE

MYLAN PHARMS INC 200MG/5ML A202361 001 May 28, 2013

VORTIOXETINE HYDROBROMIDE

TABLET; ORAL

TRINTELLIX

+ TAKEDA PHARMS USA EQ 15MG BASE ** N204447 003 Sep 30, 2013

WARFARIN POTASSIUM

TABLET; ORAL

ATHROMBIN-K

PHARM RES ASSOC 2MG N011771 007

5MG N011771 004

10MG N011771 005

25MG N011771 006

WARFARIN SODIUM

INJECTABLE; INJECTION

COUMADIN

BRISTOL MYERS SQUIBB 5MG/VIAL N009218 024 Feb 07, 1995

50MG/VIAL N009218 020

75MG/VIAL N009218 012

TABLET; ORAL

ATHROMBIN

PHARM RES ASSOC 5MG N011771 003

10MG N011771 002

25MG N011771 001

PANWARFIN

ABBOTT 2MG N017020 001

2.5MG N017020 002

5MG N017020 003

7.5MG N017020 004

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

WARFARIN SODIUMTABLET; ORAL
PANWARFIN

| | | | |
|------------------|-------|-------------|--------------|
| | 10MG | N017020 005 | |
| WARFARIN SODIUM | | | |
| MYLAN | 1MG | A040415 001 | Sep 27, 2004 |
| | 2MG | A040415 002 | Sep 27, 2004 |
| | 2.5MG | A040415 003 | Sep 29, 2004 |
| | 3MG | A040415 004 | Sep 27, 2004 |
| | 4MG | A040415 005 | Sep 27, 2004 |
| | 5MG | A040415 006 | Sep 27, 2004 |
| | 6MG | A040415 007 | Sep 27, 2004 |
| | 7.5MG | A040415 008 | Sep 27, 2004 |
| | 10MG | A040415 009 | Sep 27, 2004 |
| USL PHARMA | 2MG | A088719 001 | Jun 27, 1985 |
| | 2.5MG | A088720 001 | Aug 06, 1985 |
| | 5MG | A088721 001 | Jul 02, 1985 |
| WATSON LABS | 2MG | A086123 001 | Aug 17, 1982 |
| | 2.5MG | A086120 001 | Aug 17, 1982 |
| | 5MG | A086119 001 | Aug 17, 1982 |
| | 7.5MG | A086118 001 | Aug 17, 1982 |
| | 10MG | A086122 001 | Aug 17, 1982 |
| YAOPHARMA CO LTD | 1MG | A040196 001 | Sep 30, 1997 |
| | 2MG | A040196 002 | Sep 30, 1997 |
| | 2.5MG | A040196 003 | Sep 30, 1997 |
| | 3MG | A040196 008 | Jul 26, 2000 |
| | 4MG | A040196 004 | Sep 30, 1997 |
| | 5MG | A040196 005 | Sep 30, 1997 |
| | 6MG | A040196 009 | Jul 26, 2000 |
| | 7.5MG | A040196 006 | Sep 30, 1997 |
| | 10MG | A040196 007 | Sep 30, 1997 |

XENON XE-127GAS; INHALATION
XENON XE 127

| | | | |
|--------------|------------|-------------|--------------|
| MALLINCKRODT | 5mCi/VIAL | N018536 001 | Oct 01, 1982 |
| | 10mCi/VIAL | N018536 002 | Oct 01, 1982 |

XENON XE-133GAS; INHALATION
XENON XE 133

| | | | |
|---------------------------------|-------------------|-------------|--|
| GE HEALTHCARE | 1 CI/AMP | N017256 002 | |
| | 10mCi/VIAL | N017687 002 | |
| | 20mCi/VIAL | N017687 003 | |
| GEN ELECTRIC | 5-100 CI/CYLINDER | N017550 001 | |
| | 0.25-5 CI/AMP | N017550 003 | |
| XENON XE 133-V.S.S. | | | |
| GE HEALTHCARE | 10mCi/VIAL | N017687 001 | |
| INJECTABLE; INJECTION | | | |
| XENON XE 133 | | | |
| GE HEALTHCARE | 1.3-1.7 CI/AMP | N017256 001 | |
| LANTHEUS MEDCL | 6.3mCi/ML | N017283 001 | |
| SOLUTION; INHALATION, INJECTION | | | |
| XENEISOL | | | |
| MALLINCKRODT | 18-25mCi/AMP | N017262 002 | |

XYLOSEPOWDER; ORAL
XYLO-PFAN

| | | | |
|-------------|----------|-------------|--------------|
| SAVAGE LABS | 25GM/BOT | N017605 001 | |
| XYLOSE | | | |
| LYNE | 25GM/BOT | N018856 001 | Mar 26, 1987 |

ZALCITABINETABLET; ORAL
HIVID

| | | | |
|-------|---------|-------------|--------------|
| ROCHE | 0.375MG | N020199 001 | Jun 19, 1992 |
| | 0.75MG | N020199 002 | Jun 19, 1992 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ZALEPLON

CAPSULE; ORAL

ZALEPLON

| | | | |
|-------------------|------|-------------|--------------|
| MYLAN | 5MG | A077238 001 | Jun 06, 2008 |
| | 10MG | A077238 002 | Jun 06, 2008 |
| UPSHER SMITH LABS | 5MG | A078095 001 | Jun 06, 2008 |
| | 5MG | A078706 001 | Jun 06, 2008 |
| | 10MG | A078095 002 | Jun 06, 2008 |
| | 10MG | A078706 002 | Jun 06, 2008 |

ZICONOTIDE ACETATE

INJECTABLE; INTRATHECAL

PRIALT

| | | | |
|---------------------|------------------------|-------------|--------------|
| TERSERA THERAPS LLC | 200MCG/2ML (100MCG/ML) | N021060 003 | Dec 28, 2004 |
|---------------------|------------------------|-------------|--------------|

ZIDOVUDINE

INJECTABLE; INJECTION

ZIDOVUDINE

| | | | |
|------------------|---------|-------------|--------------|
| LIAONING CHENGDA | 10MG/ML | A204538 001 | Nov 26, 2013 |
|------------------|---------|-------------|--------------|

TABLET; ORAL

RETROVIR

| | | | |
|----------------|----------|-------------|--------------|
| VIIIV HLTHCARE | 200MG | N020518 001 | Dec 19, 1995 |
| + | 300MG ** | N020518 002 | Oct 04, 1996 |

ZIDOVUDINE

| | | | |
|------------------|-------|-------------|--------------|
| AUROBINDO PHARMA | 60MG | N022294 001 | Jul 23, 2009 |
| HEC PHARM | 300MG | A202058 001 | Oct 07, 2011 |
| MATRIX LABS LTD | 100MG | N200732 001 | Feb 23, 2011 |
| RANBAXY LABS LTD | 300MG | A077327 001 | Sep 19, 2005 |

ZILEUTON

TABLET; ORAL

ZYFLO

| | | | |
|----------------|-------|-------------|--------------|
| CHIESI USA INC | 300MG | N020471 001 | Dec 09, 1996 |
|----------------|-------|-------------|--------------|

ZINC SULFATE

INJECTABLE; INJECTION

ZINC SULFATE

| | | | |
|---------------|----------------|-------------|--------------|
| ABRAXIS PHARM | EQ 1MG ZINC/ML | N019229 002 | May 05, 1987 |
|---------------|----------------|-------------|--------------|

ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL

ZIPRASIDONE HYDROCHLORIDE

| | | | |
|------------------|--------------|-------------|--------------|
| MYLAN PHARMS INC | EQ 20MG BASE | A202395 001 | Oct 10, 2013 |
| | EQ 40MG BASE | A202395 002 | Oct 10, 2013 |
| | EQ 60MG BASE | A202395 003 | Oct 10, 2013 |
| | EQ 80MG BASE | A202395 004 | Oct 10, 2013 |

SUSPENSION; ORAL

GEODON

| | | | |
|------------|-----------------|-------------|--------------|
| PFIZER INC | EQ 10MG BASE/ML | N021483 001 | Mar 29, 2006 |
|------------|-----------------|-------------|--------------|

ZOLEDRONIC ACID

INJECTABLE; IV (INFUSION)

ZOLEDRONIC ACID

| | | | |
|-------------------|------------------|-------------|--------------|
| SUN PHARMA GLOBAL | EQ 4MG BASE/VIAL | A090018 001 | Mar 04, 2013 |
| | EQ 4MG BASE/5ML | A202746 001 | Mar 04, 2013 |

ZOMETA

| | | | | |
|---|----------|---------------------|-------------|--------------|
| + | NOVARTIS | EQ 4MG BASE/VIAL ** | N021223 001 | Aug 20, 2001 |
|---|----------|---------------------|-------------|--------------|

ZOLMITRIPTAN

TABLET; ORAL

ZOLMITRIPTAN

| | | | |
|-------------------|-------|-------------|--------------|
| APOTEX INC | 2.5MG | A202078 001 | May 14, 2013 |
| | 5MG | A202078 002 | May 14, 2013 |
| MYLAN PHARMS INC | 2.5MG | A203186 001 | May 14, 2013 |
| | 5MG | A203186 002 | May 14, 2013 |
| SUN PHARMA GLOBAL | 2.5MG | A203476 001 | Nov 13, 2014 |
| | 5MG | A203476 002 | Nov 13, 2014 |

TABLET, ORALLY DISINTEGRATING; ORAL

ZOLMITRIPTAN

| | | | |
|------------|-------|-------------|--------------|
| APOTEX INC | 2.5MG | A202476 001 | May 14, 2013 |
| | 5MG | A202476 002 | May 14, 2013 |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

DISCONTINUED DRUG PRODUCT LIST

** See List Footnote

ZOLPIDEM TARTRATE

TABLET;ORAL

ZOLPIDEM TARTRATE

| | | | |
|----------------------|------|-------------|--------------|
| DR REDDYS LABS LTD | 5MG | A077985 001 | Apr 23, 2007 |
| | 10MG | A077985 002 | Apr 23, 2007 |
| HIKMA | 5MG | A078129 001 | Apr 30, 2008 |
| | 10MG | A078129 002 | Apr 30, 2008 |
| MYLAN PHARMS INC | 5MG | A078016 001 | Apr 23, 2007 |
| | 10MG | A078016 002 | Apr 23, 2007 |
| STRIDES VIVIMED | 5MG | A076062 001 | Apr 23, 2007 |
| | 10MG | A076062 002 | Apr 23, 2007 |
| SUN PHARM INDS LTD | 5MG | A078055 001 | Apr 23, 2007 |
| | 10MG | A078055 002 | Apr 23, 2007 |
| SUN PHARM INDUSTRIES | 5MG | A077288 001 | Apr 23, 2007 |
| | 10MG | A077288 002 | Apr 23, 2007 |
| SYNTHON PHARMS | 5MG | A077540 001 | Apr 23, 2007 |
| | 10MG | A077540 002 | Apr 23, 2007 |
| WATSON LABS | 5MG | A077773 001 | Apr 23, 2007 |
| | 10MG | A077773 002 | Apr 23, 2007 |

TABLET, EXTENDED RELEASE;ORAL

ZOLPIDEM TARTRATE

| | | | |
|----------------|--------|-------------|--------------|
| SYNTHON PHARMS | 6.25MG | A078483 001 | Apr 12, 2011 |
| | 12.5MG | A078483 002 | Jun 06, 2011 |

TABLET, ORALLY DISINTEGRATING;ORAL

TOVALT ODT

| | | | |
|-------------------|------|-------------|--------------|
| BIOVAIL LABS INTL | 5MG | N021412 001 | Apr 25, 2007 |
| | 10MG | N021412 002 | Apr 25, 2007 |

ZONISAMIDE

CAPSULE;ORAL

ZONEGRAN

| | | | |
|-----------------------|------|-------------|--------------|
| + SUNOVION PHARMS INC | 50MG | N020789 002 | Aug 22, 2003 |
|-----------------------|------|-------------|--------------|

ZONISAMIDE

| | | | |
|----------------------|-------|-------------|--------------|
| ANI PHARMS INC | 25MG | A077639 001 | Dec 22, 2005 |
| | 25MG | A077641 003 | Dec 22, 2005 |
| | 50MG | A077639 002 | Dec 22, 2005 |
| | 50MG | A077641 002 | Dec 22, 2005 |
| | 100MG | A077639 003 | Dec 22, 2005 |
| | 100MG | A077641 001 | Dec 22, 2005 |
| AUROBINDO PHARMA LTD | 25MG | A077645 002 | Sep 29, 2006 |
| | 50MG | A077645 003 | Sep 29, 2006 |
| | 100MG | A077645 001 | Dec 22, 2005 |
| EPIC PHARMA LLC | 25MG | A077876 001 | Feb 21, 2007 |
| | 50MG | A077876 002 | Feb 21, 2007 |
| | 100MG | A077876 003 | Feb 21, 2007 |
| MYLAN PHARMS INC | 25MG | A077647 001 | Dec 22, 2005 |
| | 50MG | A077647 002 | Dec 22, 2005 |
| | 100MG | A077647 003 | Dec 22, 2005 |
| ROXANE | 25MG | A077648 001 | Dec 22, 2005 |
| | 50MG | A077648 002 | Dec 22, 2005 |
| | 100MG | A077648 003 | Dec 22, 2005 |
| SUN PHARM INDUSTRIES | 25MG | A077635 001 | Dec 22, 2005 |
| | 50MG | A077635 002 | Dec 22, 2005 |
| | 100MG | A077635 003 | Dec 22, 2005 |
| UPSHER SMITH LABS | 25MG | A077644 001 | Dec 22, 2005 |
| | 50MG | A077644 002 | Dec 22, 2005 |
| | 100MG | A077644 003 | Dec 22, 2005 |
| WATSON LABS | 25MG | A077650 001 | Apr 20, 2006 |
| | 50MG | A077650 002 | Apr 20, 2006 |
| | 100MG | A077650 003 | Apr 20, 2006 |

ORPHAN PRODUCT DESIGNATIONS AND APPROVALS LIST

The list of Orphan Designations and Approvals is available at:

<http://www.fda.gov/ForIndustry/DevelopingProductsforRareDiseasesConditions/default.htm>

**DRUG PRODUCTS WHICH MUST DEMONSTRATE *IN VIVO* BIOAVAILABILITY
ONLY IF PRODUCT FAILS TO ACHIEVE ADEQUATE DISSOLUTION**

ACETAMINOPHEN;ASPIRIN;BUTALBITAL
CAPSULE OR TABLET; ORAL
160-165MG;160-165MG;50MG
325MG;325MG;50MG

ASPIRIN;CAFFEINE;CARISOPRODOL;
CODEINE PHOSPHATE
TABLET; ORAL
160MG;32MG;200MG;16MG

ACETAMINOPHEN;ASPIRIN;BUTALBITAL;
CAFFEINE
CAPSULE OR TABLET; ORAL
160-165MG;160-165MG;50MG;40MG
325MG;325MG;50MG;40MG

ASPIRIN;CARISOPRODOL
TABLET; ORAL
325MG;200MG

ACETAMINOPHEN;BUTALBITAL
CAPSULE OR TABLET; ORAL
325MG;50MG

ASPIRIN;CARISOPRODOL;
CODEINE PHOSPHATE
TABLET; ORAL
325MG;200MG;16MG

ACETAMINOPHEN;BUTALBITAL;CAFFEINE
CAPSULE OR TABLET; ORAL
325MG;50MG;40MG

ASPIRIN;MEPROBAMATE
TABLET; ORAL
325MG;200MG

AMINOPHYLLINE
TABLET; ORAL
100MG;200MG

ASPIRIN;METHOCARBAMOL
TABLET; ORAL
325MG;400MG

ASPIRIN;BUTALBITAL
CAPSULE OR TABLET; ORAL
325MG;50MG
650MG;50MG

CHLOROTHIAZIDE
TABLET; ORAL
250MG

ASPIRIN;BUTALBITAL;CAFFEINE
CAPSULE OR TABLET; ORAL
325MG;50MG;40MG
650MG;50MG;40MG

HYDROXYZINE HYDROCHLORIDE
TABLET; ORAL
10MG;25MG;
50MG;100MG

ASPIRIN;CAFFEINE;CARISOPRODOL
TABLET; ORAL
160MG;32MG;200MG

PREDNISONE
TABLET; ORAL
1MG;2.5MG;5MG;10MG;
20MG;25MG;50MG

APPENDIX A - PRODUCT NAME INDEX

** A **

A-METHAPRED, METHYLPREDNISOLONE SODIUM SUCCINATE
ABACAVIR SULFATE, ABACAVIR SULFATE
ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE, ABACAVIR SULFATE
ABELCET, AMPHOTERICIN B
ABILIFY, ARIPIPIRAZOLE
ABILIFY MAINTENA KIT, ARIPIPIRAZOLE
ABILIFY MYCITE KIT, ARIPIPIRAZOLE
ABIRATERONE ACETATE, ABIRATERONE ACETATE
ABLYSINOL, ALCOHOL
ABRAXANE, PACLITAXEL
ABREVA, DOCOSANOL (OTC)
ABSORICA, ISOTRETINOIN
ABSTRAL, FENTANYL CITRATE
ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
ACANYA, BENZOYL PEROXIDE
ACARBOSE, ACARBOSE
ACCOLATE, ZAFIRLUKAST
ACCUNEB, ALBUTEROL SULFATE
ACCUPRIL, QUINAPRIL HYDROCHLORIDE
ACCURETIC, HYDROCHLOROTHIAZIDE
ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
ACEPHEN, ACETAMINOPHEN (OTC)
ACETADOTE, ACETYLCYSTEINE
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
ACETAMINOPHEN, ACETAMINOPHEN
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
ACETAMINOPHEN, CAFFEINE AND DIHYDROCODEINE BITARTRATE, ACETAMINOPHEN
ACETASOL HC, ACETIC ACID, GLACIAL
ACETAZOLAMIDE, ACETAZOLAMIDE
ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
ACETIC ACID, ACETIC ACID, GLACIAL
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
ACETYLCYSTEINE, ACETYLCYSTEINE
ACHROMYCIN V, TETRACYCLINE HYDROCHLORIDE
ACIPHEX, RABEPRAZOLE SODIUM
ACIPHEX SPRINKLE, RABEPRAZOLE SODIUM
ACITRETIN, ACITRETIN
ACTHREL, CORTICORELIN OVINE TRIFLUTATE
ACTICLATE, DOXYCYCLINE HYCLATE
ACTIGALL, URSODIOL
ACTIQ, FENTANYL CITRATE
ACTIVELLA, ESTRADIOL
ACTONEL, RISEDRONATE SODIUM
ACTOPLUS MET, METFORMIN HYDROCHLORIDE
ACTOPLUS MET XR, METFORMIN HYDROCHLORIDE
ACTOS, PIOGLITAZONE HYDROCHLORIDE
ACULAR, KETOROLAC TROMETHAMINE
ACULAR LS, KETOROLAC TROMETHAMINE
ACUVAIL, KETOROLAC TROMETHAMINE
ACYCLOVIR, ACYCLOVIR
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
ACZONE, DAPSONE
ADAGEN, PEGADEMASE BOVINE
ADALAT CC, NIFEDIPINE
ADAPALENE, ADAPALENE
ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
ADASUVE, LOXAPINE
ADCIRCA, TADALAFIL
ADDERALL XR 10, AMPHETAMINE ASPARTATE
ADDERALL XR 15, AMPHETAMINE ASPARTATE
ADDERALL XR 20, AMPHETAMINE ASPARTATE
ADDERALL XR 25, AMPHETAMINE ASPARTATE
ADDERALL XR 30, AMPHETAMINE ASPARTATE

APPENDIX A - PRODUCT NAME INDEX

** A **

ADDERALL XR 5, AMPHETAMINE ASPARTATE
ADDYI, FLIBANSERIN
ADEFOVIR DIPIVOXIL, ADEFOVIR DIPIVOXIL
ADEMPAS, RIOCIQUAT
ADENOSINE, ADENOSINE
ADIPEX-P, PHENTERMINE HYDROCHLORIDE
ADLYXIN, LIXISENATIDE
ADMELOG, INSULIN LISPRO
ADMELOG SOLOSTAR, INSULIN LISPRO
ADRENACLICK, EPINEPHRINE
ADRENALIN, EPINEPHRINE
ADREVIEW, IOBENGUANE SULFATE I-123
ADVAIR DISKUS 100/50, FLUTICASONE PROPIONATE
ADVAIR DISKUS 250/50, FLUTICASONE PROPIONATE
ADVAIR DISKUS 500/50, FLUTICASONE PROPIONATE
ADVAIR HFA, FLUTICASONE PROPIONATE
ADVIL, IBUPROFEN (OTC)
ADVIL, IBUPROFEN SODIUM (OTC)
ADVIL ALLERGY AND CONGESTION RELIEF, CHLORPHENIRAMINE MALEATE (OTC)
ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
ADVIL COLD AND SINUS, IBUPROFEN (OTC)
ADVIL CONGESTION RELIEF, IBUPROFEN (OTC)
ADVIL LIQUI-GELS, IBUPROFEN (OTC)
ADVIL MIGRAINE LIQUI-GELS, IBUPROFEN (OTC)
ADVIL MULTI-SYMP TOM COLD & FLU, CHLORPHENIRAMINE MALEATE (OTC)
ADVIL PM, DIPHENHYDRAMINE CITRATE (OTC)
ADVIL PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
ADZENYS ER, AMPHETAMINE
ADZENYS XR-ODT, AMPHETAMINE
AEMCOLO, RIFAMYCIN
AEROSPAN HFA, FLUNISOLIDE
AFINITOR, EVEROLIMUS
AFINITOR DISPERZ, EVEROLIMUS
AFIRMELLE, ETHINYL ESTRADIOL
AFREZZA, INSULIN RECOMBINANT HUMAN
AGGRASTAT, TIROFIBAN HYDROCHLORIDE
AGGRENOLX, ASPIRIN
AGRYLIN, ANAGRELIDE HYDROCHLORIDE
AIRDUO RESPICLICK, FLUTICASONE PROPIONATE
AK-FLUOR 10%, FLUORESCEIN SODIUM
AK-FLUOR 25%, FLUORESCEIN SODIUM
AKBETA, LEVOBUNOLOL HYDROCHLORIDE
AKOAZ, EPHEDRINE SULFATE
AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE
AKTEN, LIDOCAINE HYDROCHLORIDE
AKTIPAK, BENZOYL PEROXIDE
AKTOB, TOBRAMYCIN
AKYNZEO, FOSNETUPITANT CHLORIDE HYDROCHLORIDE
AKYNZEO, NETUPITANT
ALA-CORT, HYDROCORTISONE
ALA-SCALP, HYDROCORTISONE
ALAVERT, LORATADINE (OTC)
ALAWAY, KETOTIFEN FUMARATE (OTC)
ALBENDAZOLE, ALBENDAZOLE
ALBENZA, ALBENDAZOLE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
ALCAINE, PROPARACAINE HYDROCHLORIDE
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
ALDACTAZIDE, HYDROCHLOROTHIAZIDE
ALDACTONE, SPIRONOLACTONE
ALDARA, IMIQUIMOD
ALECENSA, ALECTINIB HYDROCHLORIDE
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ALEVE, NAPROXEN SODIUM (OTC)

APPENDIX A - PRODUCT NAME INDEX

** A **

ALEVE PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
ALEVE-D SINUS & COLD, NAPROXEN SODIUM (OTC)
ALFENTA, ALFENTANIL HYDROCHLORIDE
ALFENTANIL, ALFENTANIL HYDROCHLORIDE
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
ALIMTA, PEMETREXED DISODIUM
ALINIA, NITAZOXANIDE
ALIQOPA, COPANLISIB DIHYDROCHLORIDE
ALKERAN, MELPHALAN
ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
ALLI, ORLISTAT (OTC)
ALLOPURINOL, ALLOPURINOL
ALLOPURINOL SODIUM, ALLOPURINOL SODIUM
ALLZITAL, ACETAMINOPHEN
ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
ALOCRIL, NEDOCROMIL SODIUM
ALOMIDE, LODOXAMIDE TROMETHAMINE
ALOPRIM, ALLOPURINOL SODIUM
ALORA, ESTRADIOL
ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
ALOXI, PALONOSETRON HYDROCHLORIDE
ALPHAGAN P, BRIMONIDINE TARTRATE
ALPRAZOLAM, ALPRAZOLAM
ALPROSTADIL, ALPROSTADIL
ALREX, LOTEPIREDNOL ETABONATE
ALTABAX, RETAPAMULIN
ALTACE, RAMIPRIL
ALTAFLUOR BENOX, BENOXINATE HYDROCHLORIDE
ALTAVERA, ETHINYL ESTRADIOL
ALTOPREV, LOVASTATIN
ALTRENO, TRETINOIN
ALUNBRIG, BRIGATINIB
ALVESCO, CICLESONIDE
ALYACEN 1/35, ETHINYL ESTRADIOL
ALYACEN 7/7/7, ETHINYL ESTRADIOL
AMABELZ, ESTRADIOL
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMARYL, GLIMEPIRIDE
AMBIEN, ZOLPIDEM TARTRATE
AMBIEN CR, ZOLPIDEM TARTRATE
AMBISOME, AMPHOTERICIN B
AMCINONIDE, AMCINONIDE
AMELUZ, AMINOLEVULINIC ACID HYDROCHLORIDE
AMERGE, NARATRIPTAN HYDROCHLORIDE
AMICAR, AMINOCAPROIC ACID
AMIDATE, ETOMIDATE
AMIFOSTINE, AMIFOSTINE
AMIKACIN SULFATE, AMIKACIN SULFATE
AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
AMINO ACIDS, AMINO ACIDS
AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER, GLYCINE
AMINOCAPROIC ACID, AMINOCAPROIC ACID
AMINOCAPROIC ACID IN PLASTIC CONTAINER, AMINOCAPROIC ACID
AMINOPHYLLINE, AMINOPHYLLINE
AMINOSYN 10%, AMINO ACIDS
AMINOSYN 3.5% M, AMINO ACIDS
AMINOSYN 8.5%, AMINO ACIDS
AMINOSYN 8.5% W/ELECTROLYTES, AMINO ACIDS
AMINOSYN II 10% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 15% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN-PF 10%, AMINO ACIDS
AMINOSYN-PF 7%, AMINO ACIDS

APPENDIX A - PRODUCT NAME INDEX

** A **

AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AMITIZA, LUBIPROSTONE
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
AMMONIA N 13, AMMONIA N-13
AMMONIUM CHLORIDE IN PLASTIC CONTAINER, AMMONIUM CHLORIDE
AMMONIUM LACTATE, AMMONIUM LACTATE
AMMONUL, SODIUM BENZOATE
AMNESTEEM, ISOTRETINOIN
AMOXAPINE, AMOXAPINE
AMOXICILLIN, AMOXICILLIN
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN PEDIATRIC, AMOXICILLIN
AMOXIL, AMOXICILLIN
AMPHADASE, HYALURONIDASE
AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
AMPHOTERICIN B, AMPHOTERICIN B
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
AMPICILLIN SODIUM, AMPICILLIN SODIUM
AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE
AMPYRA, DALFAMPRIDINE
AMRINONE LACTATE, INAMRINONE LACTATE
AMRIX, CYCLOBENZAPRINE HYDROCHLORIDE
AMYVID, FLORBETAPIR F-18
AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT
ANADROL-50, OXYMETHOLONE
ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
ANAPROX DS, NAPROXEN SODIUM
ANASTROZOLE, ANASTROZOLE
ANCEF IN PLASTIC CONTAINER, CEFAZOLIN SODIUM
ANCOBON, FLUCYTOSINE
ANDRODERM, TESTOSTERONE
ANDROGEL, TESTOSTERONE
ANDROID 25, METHYLTESTOSTERONE
ANECTINE, SUCCINYLCHOLINE CHLORIDE
ANEXSIA 5/325, ACETAMINOPHEN
ANEXSIA 7.5/325, ACETAMINOPHEN
ANGELIQ, DROSPIRENONE
ANGIOMAX, BIVALIRUDIN
ANNOVERA, ETHINYL ESTRADIOL
ANORO ELLIPTA, UMECLIDINIUM BROMIDE
ANTABUSE, DISULFIRAM
ANTARA (MICRONIZED), FENOFIBRATE
ANTHELIOS 20, AVOBENZONE (OTC)
ANTHELIOS 40, AVOBENZONE (OTC)
ANTHELIOS SX, AVOBENZONE (OTC)
ANTIZOL, FOMEPIZOLE
ANUSOL HC, HYDROCORTISONE
APADAZ, ACETAMINOPHEN
APIDRA, INSULIN GLULISINE RECOMBINANT
APIDRA SOLOSTAR, INSULIN GLULISINE RECOMBINANT
APLENZIN, BUPROPION HYDROBROMIDE
APOKYN, APOMORPHINE HYDROCHLORIDE
APRACLONIDINE HYDROCHLORIDE, APRACLONIDINE HYDROCHLORIDE
APREPITANT, APREPITANT
APRISO, MESALAMINE
APTENSIO XR, METHYLPHENIDATE HYDROCHLORIDE
APTIOM, ESLICARBAZEPINE ACETATE
APTIVUS, TIPRANAVIR

APPENDIX A - PRODUCT NAME INDEX**** A ****

AQUASOL A, VITAMIN A PALMITATE
ARAKODA, TAFENOQUINE SUCCINATE
ARANELLE, ETHINYL ESTRADIOL
ARAVA, LEFLUNOMIDE
ARCAPTA NEOHALER, INDACATEROL MALEATE
ARESTIN, MINOCYCLINE HYDROCHLORIDE
ARGATROBAN, ARGATROBAN
ARGATROBAN IN 0.9% SODIUM CHLORIDE, ARGATROBAN
ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
ARICEPT, DONEPEZIL HYDROCHLORIDE
ARIDOL KIT, MANNITOL
ARIKAYCE KIT, AMIKACIN SULFATE
ARIMIDEX, ANASTROZOLE
ARIPIPIRAZOLE, ARIPIPIRAZOLE
ARISTADA, ARIPIPIRAZOLE LAUROXIL
ARISTADA INITIO KIT, ARIPIPIRAZOLE LAUROXIL
ARISTOSPAN, TRIAMCINOLONE HEXACETONIDE
ARIXTRA, FONDAPARINUX SODIUM
ARMODAFINIL, ARMODAFINIL
ARMONAIR RESPICLICK, FLUTICASONE PROPIONATE
ARNUITY ELLIPTA, FLUTICASONE FUROATE
AROMASIN, EXEMESTANE
ARRANON, NELARABINE
ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
ARTHROTEC, DICLOFENAC SODIUM
ARTICAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, ARTICAINE HYDROCHLORIDE
ASACOL HD, MESALAMINE
ASCLERA, POLIDOCANOL
ASCOR, ASCORBIC ACID
ASENAPINE MALEATE, ASENAPINE MALEATE
ASHLYNA, ETHINYL ESTRADIOL
ASMANEX HFA, MOMETASONE FUROATE
ASMANEX TWISTHALER, MOMETASONE FUROATE
ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
ASTAGRAF XL, TACROLIMUS
ASTELIN, AZELASTINE HYDROCHLORIDE
ASTEPRO, AZELASTINE HYDROCHLORIDE
ASTRAMORPH PF, MORPHINE SULFATE
ATACAND, CANDESARTAN CILEXETIL
ATACAND HCT, CANDESARTAN CILEXETIL
ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
ATELVIA, RISEDRONATE SODIUM
ATENOLOL, ATENOLOL
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
ATHENTIA NEXT, LEVONORGESTREL (OTC)
ATIVAN, LORAZEPAM
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
ATOVAQUONE, ATOVAQUONE
ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRALIN, TRETINOIN
ATRIDOX, DOXYCYCLINE HYCLATE
ATRIPLA, EFAVIRENZ
ATROPEN, ATROPINE SULFATE
ATROPINE SULFATE, ATROPINE SULFATE
ATROPINE SULFATE ANSYR PLASTIC SYRINGE, ATROPINE SULFATE
ATROPINE SULFATE LIFESHIELD ABOBJECT SYRINGE, ATROPINE SULFATE
ATROVENT HFA, IPRATROPIUM BROMIDE
AUBAGIO, TERIFLUNOMIDE
AUGMENTIN '125', AMOXICILLIN
AUGMENTIN '250', AMOXICILLIN
AUGMENTIN '875', AMOXICILLIN
AUGMENTIN XR, AMOXICILLIN

APPENDIX A - PRODUCT NAME INDEX**** A ****

AUROVELA 1.5/30, ETHINYL ESTRADIOL
 AUROVELA 1/20, ETHINYL ESTRADIOL
 AUROVELA 24 FE, ETHINYL ESTRADIOL
 AUROVELA FE 1.5/30, ETHINYL ESTRADIOL
 AUROVELA FE 1/20, ETHINYL ESTRADIOL
 AURYXIA, FERRIC CITRATE
 AUSTEDO, DEUTETRABENAZINE
 AUVI-Q, EPINEPHRINE
 AVAGARD, ALCOHOL (OTC)
 AVAGE, TAZAROTENE
 AVALIDE, HYDROCHLOROTHIAZIDE
 AVANDIA, ROSIGLITAZONE MALEATE
 AVAPRO, IRBESARTAN
 AVC, SULFANILAMIDE
 AVEED, TESTOSTERONE UNDECANOATE
 AVELOX, MOXIFLOXACIN HYDROCHLORIDE
 AVIANE-28, ETHINYL ESTRADIOL
 AVITA, TRETINOIN
 AVODART, DUTASTERIDE
 AVYCAZ, AVIBACTAM SODIUM
 AXERT, ALMOTRIPTAN MALATE
 AXID AR, NIZATIDINE (OTC)
 AXUMIN, FLUCICLOVINE F-18
 AYUNA, ETHINYL ESTRADIOL
 AZACITIDINE, AZACITIDINE
 AZACTAM, AZTREONAM
 AZACTAM IN PLASTIC CONTAINER, AZTREONAM
 AZASAN, AZATHIOPRINE
 AZASITE, AZITHROMYCIN
 AZATHIOPRINE, AZATHIOPRINE
 AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM
 AZEDRA, IOBENGUANE I-131
 AZELAIC ACID, AZELAIC ACID
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZELEX, AZELAIC ACID
 AZILECT, RASAGILINE MESYLATE
 AZITHROMYCIN, AZITHROMYCIN
 AZOPT, BRINZOLAMIDE
 AZOR, AMLODIPINE BESYLATE
 AZTREONAM, AZTREONAM
 AZULFIDINE, SULFASALAZINE
 AZULFIDINE EN-TABS, SULFASALAZINE

**** B ****

BACIIM, BACITRACIN
 BACITRACIN, BACITRACIN
 BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE, BACITRACIN ZINC
 BACLOFEN, BACLOFEN
 BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 BACTOCILL IN PLASTIC CONTAINER, OXACILLIN SODIUM
 BACTRIM, SULFAMETHOXAZOLE
 BACTRIM DS, SULFAMETHOXAZOLE
 BAL, DIMERCAPROL
 BALANCED SALT, CALCIUM CHLORIDE
 BALCOLTRA, ETHINYL ESTRADIOL
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
 BALZIVA-28, ETHINYL ESTRADIOL
 BANZEL, RUFINAMIDE
 BARACLUDE, ENTECAVIR
 BASAGLAR, INSULIN GLARGINE
 BAXDELA, DELAFLOXACIN MEGLUMINE
 BECONASE AQ, BECLOMETHASONE DIPROPIONATE MONOHYDRATE
 BEKYREE, DESOGESTREL

APPENDIX A - PRODUCT NAME INDEX

** B **

BELBUCA, BUPRENORPHINE HYDROCHLORIDE
BELEODAQ, BELINOSTAT
BELRAPZO, BENDAMUSTINE HYDROCHLORIDE
BELSOMRA, SUVOREXANT
BELVIQ, LORCASERIN HYDROCHLORIDE
BELVIQ XR, LORCASERIN HYDROCHLORIDE
BENZAEPRIIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BENZAEPRIIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
BENDEKA, BENDAMUSTINE HYDROCHLORIDE
BENICAR, OLMESARTAN MEDOXOMIL
BENICAR HCT, HYDROCHLOROTHIAZIDE
BENTYL, DICYCLOMINE HYDROCHLORIDE
BENTYL PRESERVATIVE FREE, DICYCLOMINE HYDROCHLORIDE
BENZAACLIN, BENZOYL PEROXIDE
BENZAMYCIN, BENZOYL PEROXIDE
BENZNIDAZOLE, BENZNIDAZOLE
BENZONATATE, BENZONATATE
BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
BEPREVE, BEPOTASTINE BESILATE
BESIVANCE, BESIFLOXACIN HYDROCHLORIDE
BETA-VAL, BETAMETHASONE VALERATE
BETADINE, POVIDONE-IODINE
BETAGAN, LEVOBUNOLOL HYDROCHLORIDE
BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE, BETAMETHASONE ACETATE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
BETAPACE, SOTALOL HYDROCHLORIDE
BETAPACE AF, SOTALOL HYDROCHLORIDE
BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
BETHKIS, TOBRAMYCIN
BETIMOL, TIMOLOL
BETOPTIC, BETAXOLOL HYDROCHLORIDE
BETOPTIC S, BETAXOLOL HYDROCHLORIDE
BEVESPI AEROSPHERE, FORMOTEROL FUMARATE
BEVYXXA, BETRIXABAN
BEXAROTENE, BEXAROTENE
BEYAZ, DROSPIRENONE
BICALUTAMIDE, BICALUTAMIDE
BICILLIN C-R, PENICILLIN G BENZATHINE
BICILLIN C-R 900/300, PENICILLIN G BENZATHINE
BICILLIN L-A, PENICILLIN G BENZATHINE
BICNU, CARMUSTINE
BIDIL, HYDRALAZINE HYDROCHLORIDE
BIJUVA, ESTRADIOL
BIKTARVY, BICTEGRAVIR SODIUM
BILTRICIDE, PRAZIQUANTEL
BIMATOPROST, BIMATOPROST
BINOSTO, ALENDRONATE SODIUM
BIOSCRUB, CHLORHEXIDINE GLUCONATE (OTC)
BISMUTH SUBSALICYLATE, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE, BISMUTH SUBSALICYLATE
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
BIVALIRUDIN, BIVALIRUDIN
BIVALIRUDIN IN 0.9% SODIUM CHLORIDE, BIVALIRUDIN
BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
BLEPH-10, SULFACETAMIDE SODIUM
BLEPHAMIDE, PREDNISOLONE ACETATE
BLEPHAMIDE S.O.P., PREDNISOLONE ACETATE
BLISOVI 24 FE, ETHINYL ESTRADIOL
BLISOVI FE 1.5/30, ETHINYL ESTRADIOL
BLISOVI FE 1/20, ETHINYL ESTRADIOL
BLOXIVERZ, NEOSTIGMINE METHYLSULFATE

APPENDIX A - PRODUCT NAME INDEX

** B **

BONIVA, IBANDRONATE SODIUM
BONJESTA, DOXYLAMINE SUCCINATE
BONTRIL PDM, PHENDIMETRAZINE TARTRATE
BORTEZOMIB, BORTEZOMIB
BOSULIF, BOSUTINIB MONOHYDRATE
BRAFTOVI, ENCORAFENIB
BREGO ELLIPTA, FLUTICASON FUROATE
BRETHINE, TERBUTALINE SULFATE
BRETILUM TOSYLATE, BRETILUM TOSYLATE
BREVIBLOC, ESMOLOL HYDROCHLORIDE
BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
BREVIBLOC IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
BREVICON 28-DAY, ETHINYL ESTRADIOL
BREVITAL SODIUM, METHOHEXITAL SODIUM
BRIAN CARE, CHLORHEXIDINE GLUCONATE (OTC)
BRIDION, SUGAMMADEX SODIUM
BRIELLYN, ETHINYL ESTRADIOL
BRILINTA, TICAGRELOR
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
BRISDELLE, PAROXETINE MESYLATE
BRIVIACT, BRIVARACETAM
BROMFED-DM, BROMPHENIRAMINE MALEATE
BROMFENAC SODIUM, BROMFENAC SODIUM
BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE,
BROMSITE, BROMFENAC SODIUM
BRONCHO SALINE, SODIUM CHLORIDE (OTC)
BROVANA, ARFORMOTEROL TARTRATE
BRYHALI, HALOBETASOL PROPIONATE
BSS, CALCIUM CHLORIDE
BSS PLUS, CALCIUM CHLORIDE
BUDESONIDE, BUDESONIDE (OTC)
BUDESONIDE, BUDESONIDE
BUMETANIDE, BUMETANIDE
BUMEX, BUMETANIDE
BUNAVAIL, BUPRENORPHINE HYDROCHLORIDE
BUPHENYL, SODIUM PHENYLBUTYRATE
BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
BUPIVACAINE HYDROCHLORIDE W/EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
BUPRENEX, BUPRENORPHINE HYDROCHLORIDE
BUPRENORPHINE, BUPRENORPHINE
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
BUSULFAN, BUSULFAN
BUSULFEX, BUSULFAN
BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
BUTAPAP, ACETAMINOPHEN
BUTENAFINE HYDROCHLORIDE, BUTENAFINE HYDROCHLORIDE (OTC)
BUTISOL SODIUM, BUTABARBITAL SODIUM
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
BUTRANS, BUPRENORPHINE
BYDUREON, EXENATIDE SYNTHETIC
BYDUREON BCISE, EXENATIDE
BYDUREON PEN, EXENATIDE SYNTHETIC
BYETTA, EXENATIDE SYNTHETIC
BYSTOLIC, NEBIVOLOL HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** B **

BYVALSON, NEBIVOLOL HYDROCHLORIDE

** C **

CABERGOLINE, CABERGOLINE
 CABOMETYX, CABOZANTINIB S-MALATE
 CADUET, AMLODIPINE BESYLATE
 CAFKIT, CAFFEINE CITRATE
 CAFERGOT, CAFFEINE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CALAN, VERAPAMIL HYDROCHLORIDE
 CALAN SR, VERAPAMIL HYDROCHLORIDE
 CALCIPOTRIENE, CALCIPOTRIENE
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CALCITRIOL, CALCITRIOL
 CALCIUM ACETATE, CALCIUM ACETATE
 CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 CALCIUM DISODIUM VERSENATE, EDTATE CALCIUM DISODIUM
 CALCIUM GLUCONATE, CALCIUM GLUCONATE
 CALCIUM GLUCONATE IN SODIUM CHLORIDE, CALCIUM GLUCONATE
 CALDOLOR, IBUPROFEN
 CALQUENCE, ACALABRUTINIB
 CAMBIA, DICLOFENAC POTASSIUM
 CAMILA, NORETHINDRONE
 CAMPTOSAR, IRINOTECAN HYDROCHLORIDE
 CANASA, MESALAMINE
 CANCIDAS, CASPOFUNGIN ACETATE
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CAPASTAT SULFATE, CAPREOMYCIN SULFATE
 CAPECITABINE, CAPECITABINE
 CAPEX, FLUOCINOLONE ACETONIDE
 CAPITAL SOLEIL 15, AVOBENZONE (OTC)
 CAPRELSA, VANDETANIB
 CAPREOMYCIN SULFATE, CAPREOMYCIN SULFATE
 CAPTOPRIL, CAPTOPRIL
 CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL
 CARAC, FLUOROURACIL
 CARAFATE, SUCRALFATE
 CARBAGLU, CARGLUMIC ACID
 CARBAMAZEPINE, CARBAMAZEPINE
 CARBATROL, CARBAMAZEPINE
 CARBIDOPA, CARBIDOPA
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARBIDOPA, LEVODOPA AND ENTACAPONE, CARBIDOPA
 CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
 CARBOCAINE, MEPIVACAINE HYDROCHLORIDE
 CARBOPLATIN, CARBOPLATIN
 CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
 CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
 CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
 CARDIOGEN-82, RUBIDIUM CHLORIDE RB-82
 CARDIOLITE, TECHNETIUM TC-99M SESTAMIBI KIT
 CARDIOPLEGIC IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 CARDIZEM, DILTIAZEM HYDROCHLORIDE
 CARDIZEM CD, DILTIAZEM HYDROCHLORIDE
 CARDIZEM LA, DILTIAZEM HYDROCHLORIDE
 CARDURA, DOXAZOSIN MESYLATE
 CARDURA XL, DOXAZOSIN MESYLATE
 CARISOPRODOL, CARISOPRODOL
 CARISOPRODOL AND ASPIRIN, ASPIRIN
 CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN
 CARMUSTINE, CARMUSTINE
 CARNITOR, LEVOCARNITINE

APPENDIX A - PRODUCT NAME INDEX

** C **

CARNITOR SF, LEVOCARNITINE
CAROSPIR, SPIRONOLACTONE
CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
CARTIA XT, DILTIAZEM HYDROCHLORIDE
CARVEDILOL, CARVEDILOL
CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE
CASODEX, BICALUTAMIDE
CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
CASPORYN HC, HYDROCORTISONE
CASSIPA, BUPRENORPHINE HYDROCHLORIDE
CATAPRES, CLONIDINE HYDROCHLORIDE
CATAPRES-TTS-1, CLONIDINE
CATAPRES-TTS-2, CLONIDINE
CATAPRES-TTS-3, CLONIDINE
CAVERJECT, ALPROSTADIL
CAVERJECT IMPULSE, ALPROSTADIL
CAYSTON, AZTREONAM
CEFACLOR, CEFACLOR
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CEFAZOLIN AND DEXTROSE, CEFAZOLIN SODIUM
CEFAZOLIN IN PLASTIC CONTAINER, CEFAZOLIN SODIUM
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
CEFDINIR, CEFDINIR
CEFEPIME AND DEXTROSE IN DUPLEX CONTAINER, CEFEPIME HYDROCHLORIDE
CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
CEFEPIME HYDROCHLORIDE IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE
CEFEPIME IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE
CEFIXIME, CEFIXIME
CEFOTAN, CEFOTETAN DISODIUM
CEFOTAXIME, CEFOTAXIME SODIUM
CEFOTAXIME SODIUM, CEFOTAXIME SODIUM
CEFOTETAN, CEFOTETAN DISODIUM
CEFOTETAN AND DEXTROSE IN DUPLEX CONTAINER, CEFOTETAN DISODIUM
CEFOXITIN, CEFOXITIN SODIUM
CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER, CEFOXITIN SODIUM
CEFOXITIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
CEFPROZIL, CEFPROZIL
CEFTAZIDIME, CEFTAZIDIME
CEFTAZIDIME IN DEXTROSE CONTAINER, CEFTAZIDIME
CEFTRIAXONE, CEFTRIAXONE SODIUM
CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER, CEFTRIAXONE SODIUM
CEFTRIAXONE IN PLASTIC CONTAINER, CEFTRIAXONE SODIUM
CEFTRIAXONE SODIUM, CEFTRIAXONE SODIUM
CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER, CEFUROXIME SODIUM
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEFUROXIME SODIUM, CEFUROXIME SODIUM
CELEBREX, CELECOXIB
CELECOXIB, CELECOXIB
CELESTONE SOLUSPAN, BETAMETHASONE ACETATE
CELEXA, CITALOPRAM HYDROBROMIDE
CELLCEPT, MYCOPHENOLATE MOFETIL
CELLCEPT, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
CELONTIN, METHSUXIMIDE
CENTANY, MUPIROCIN
CEPHALEXIN, CEPHALEXIN
CEQUA, CYCLOSPORINE
CERDELGA, ELIGLUSTAT TARTRATE
CEREBYX, FOSPHENYTOIN SODIUM
CERETEC, TECHNETIUM TC-99M EXAMETAZIME KIT
CEREZYME, IMIGLUCERASE
CERINTA, ETHINYL ESTRADIOL
CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE
CERVIDIL, DINOPROSTONE
CESAMET, NABILONE

APPENDIX A - PRODUCT NAME INDEX

** C **

CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CETRAXAL, CIPROFLOXACIN HYDROCHLORIDE
 CETROTIDE, CETRORELIX
 CETYLEV, ACETYLCYSTEINE
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CHANTIX, VARENICLINE TARTRATE
 CHEMET, SUCCIMER
 CHENODIOL, CHENODIOL
 CHG SCRUB, CHLORHEXIDINE GLUCONATE (OTC)
 CHILDREN'S ADVIL, IBUPROFEN (OTC)
 CHILDREN'S ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
 CHILDREN'S ADVIL COLD, IBUPROFEN (OTC)
 CHILDREN'S ADVIL-FLAVORED, IBUPROFEN (OTC)
 CHILDREN'S ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CLARITIN, LORATADINE (OTC)
 CHILDREN'S ELIXSURE, IBUPROFEN (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S IBUPROFEN, IBUPROFEN (OTC)
 CHILDREN'S MOTRIN, IBUPROFEN (OTC)
 CHILDREN'S MOTRIN COLD, IBUPROFEN (OTC)
 CHILDREN'S ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S ZYRTEC HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN
 CHLOR-TRIMETON, CHLORPHENIRAMINE MALEATE (OTC)
 CHLORAMPHENICOL SODIUM SUCCINATE, CHLORAMPHENICOL SODIUM SUCCINATE
 CHLORAPREP ONE-STEP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP ONE-STEP FREPP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP ONE-STEP SEPP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP SINGLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP TRIPLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP WITH TINT, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 CHLOROTHIAZIDE, CHLOROTHIAZIDE
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE (OTC)
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CHLORPROPAMIDE, CHLORPROPAMIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CHLORZOAZONE, CHLORZOAZONE
 CHOLAC, LACTULOSE
 CHOLBAM, CHOLIC ACID
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLETEC, TECHNETIUM TC-99M MEBROFENIN KIT
 CHOLINE C-11, CHOLINE C-11
 CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC
 CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE
 CIALIS, TADALAFIL
 CICLOPIROX, CICLOPIROX
 CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)
 CIDOFOVIR, CIDOFOVIR
 CILOSTAZOL, CILOSTAZOL
 CILOXAN, CIPROFLOXACIN HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** C **

CIMDUO, LAMIVUDINE
 CIMETIDINE, CIMETIDINE (OTC)
 CIMETIDINE, CIMETIDINE
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CINVANTI, APREPITANT
 CIPRO, CIPROFLOXACIN
 CIPRO, CIPROFLOXACIN HYDROCHLORIDE
 CIPRO HC, CIPROFLOXACIN HYDROCHLORIDE
 CIPRODEX, CIPROFLOXACIN
 CIPROFLOXACIN, CIPROFLOXACIN
 CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT
 CIS-PYRO, TECHNETIUM TC-99M PYROPHOSPHATE KIT
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CISPLATIN, CISPLATIN
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CITANEST FORTE DENTAL, EPINEPHRINE BITARTRATE
 CLADRIBINE, CLADRIBINE
 CLARAVIS, ISOTRETINOIN
 CLARINEX, DESLORATADINE
 CLARINEX D 24 HOUR, DESLORATADINE
 CLARINEX-D 12 HOUR, DESLORATADINE
 CLARITHROMYCIN, CLARITHROMYCIN
 CLARITIN, LORATADINE (OTC)
 CLARITIN HIVES RELIEF, LORATADINE (OTC)
 CLARITIN HIVES RELIEF REDITAB, LORATADINE (OTC)
 CLARITIN REDITABS, LORATADINE (OTC)
 CLARITIN-D, LORATADINE (OTC)
 CLARITIN-D 24 HOUR, LORATADINE (OTC)
 CLEMASTINE FUMARATE, CLEMASTINE FUMARATE (OTC)
 CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
 CLENPIQ, CITRIC ACID
 CLEOCIN, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLEOCIN, CLINDAMYCIN PHOSPHATE
 CLEOCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLEOCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 CLEOCIN T, CLINDAMYCIN PHOSPHATE
 CLEVIPREX, CLEVIDIPINE
 CLIMARA, ESTRADIOL
 CLIMARA PRO, ESTRADIOL
 CLINDA-DERM, CLINDAMYCIN PHOSPHATE
 CLINDAGEL, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLINDAMYCIN PHOSPHATE AND TRETINOIN, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PHOSPHATE IN 0.9% SODIUM CHLORIDE, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE
 CLINDESSE, CLINDAMYCIN PHOSPHATE
 CLINDETS, CLINDAMYCIN PHOSPHATE
 CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS

APPENDIX A - PRODUCT NAME INDEX

** C **

CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 2.75/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 2.75/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 2.75/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 4.25/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 4.25/20 SULFITE FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 4.25/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 4.25/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 5/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 5/15 SULFITE FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 5/20 SULFITE FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 5/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,
CLINIMIX E 5/35 SULFITE FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER,
CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
CLOBAZAM, CLOBAZAM
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
CLOBEX, CLOBETASOL PROPIONATE
CLODERM, CLOCORTOLONE PIVALATE
CLOFARABINE, CLOFARABINE
CLOLAR, CLOFARABINE
CLOMIPHENE CITRATE, CLOMIPHENE CITRATE
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CLONAZEPAM, CLONAZEPAM
CLONIDINE, CLONIDINE
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
CLOROTEKAL, CHLOROPROCAINE HYDROCHLORIDE
CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
CLOTRIMAZOLE, CLOTRIMAZOLE
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CLOZAPINE, CLOZAPINE
CLOZARIL, CLOZAPINE
COARTEM, ARTEMETHER
CODEINE SULFATE, CODEINE SULFATE
COGENTIN, BENZTROPINE MESYLATE
COL-PROBENECID, COLCHICINE
COLAZAL, BALSALAZIDE DISODIUM
COLCHICINE, COLCHICINE
COLCRYS, COLCHICINE
COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
COLESTID, COLESTIPOL HYDROCHLORIDE
COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
COLGATE TOTAL, SODIUM FLUORIDE (OTC)
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
COLOCORT, HYDROCORTISONE
COLPREP KIT, MAGNESIUM SULFATE
COLY-MYCIN M, COLISTIMETHATE SODIUM
COLY-MYCIN S, COLISTIN SULFATE
COLYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
COMBIGAN, BRIMONIDINE TARTRATE
COMBIPATCH, ESTRADIOL
COMBIVENT RESPIMAT, ALBUTEROL SULFATE
COMBIVIR, LAMIVUDINE
COMETRIQ, CABOZANTINIB S-MALATE
COMPLERA, EMTRICITABINE
COMPRO, PROCHLORPERAZINE
COMTAN, ENTACAPONE
CONCERTA, METHYLPHENIDATE HYDROCHLORIDE
CONDYLOX, PODOFILOX
CONRAY, IOTHALAMATE MEGLUMINE

APPENDIX A - PRODUCT NAME INDEX

** C **

CONRAY 43, IOTHALAMATE MEGLUMINE
CONSENSI, AMLODIPINE BESYLATE
CONSTILAC, LACTULOSE
CONTRAVE, BUPROPION HYDROCHLORIDE
CONZIP, TRAMADOL HYDROCHLORIDE
COPAXONE, GLATIRAMER ACETATE
COPIKTRA, DUVELISIB
CORDRAN, FLURANDRENOLIDE
CORDRAN SP, FLURANDRENOLIDE
COREG, CARVEDILOL
COREG CR, CARVEDILOL PHOSPHATE
CORGARD, NADOLOL
CORLANOR, IVABRADINE HYDROCHLORIDE
CORLOPAM, FENOLDOPAM MESYLATE
CORMAX, CLOBETASOL PROPIONATE
CORPHEDRA, EPHEDRINE SULFATE
CORTEF, HYDROCORTISONE
CORTENEMA, HYDROCORTISONE
CORTIFOAM, HYDROCORTISONE ACETATE
CORTISONE ACETATE, CORTISONE ACETATE
CORTISPORIN, BACITRACIN ZINC
CORTISPORIN, HYDROCORTISONE ACETATE
CORTROSYN, COSYNTROPIN
CORVERT, IBUTILIDE FUMARATE
CORZIDE, BENDROFLUMETHIAZIDE
COSMEGEN, DACTINOMYCIN
COSOPT, DORZOLAMIDE HYDROCHLORIDE
COSOPT PF, DORZOLAMIDE HYDROCHLORIDE
COSYNTROPIN, COSYNTROPIN
COTELLIC, COBIMETINIB FUMARATE
COTEMPLA XR-ODT, METHYLPHENIDATE
COUMADIN, WARFARIN SODIUM
COZAAR, LOSARTAN POTASSIUM
CREON, PANCRELIPASE (AMYLASE
CRESEMBA, ISAVUCONAZONIUM SULFATE
CRESTOR, ROSUVASTATIN CALCIUM
CRINONE, PROGESTERONE
CRIXIVAN, INDINAVIR SULFATE
CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
CROMOLYN SODIUM, CROMOLYN SODIUM
CROTAN, CROTAMITON
CRYSELLE, ETHINYL ESTRADIOL
CUBICIN, DAPTOMYCIN
CUBICIN RF, DAPTOMYCIN
CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE
CUPRIMINE, PENICILLAMINE
CUROSURF, PORACTANT ALFA
CUTIVATE, FLUTICASONE PROPIONATE
CUVPOSA, GLYCOPYRROLATE
CYANOCOBALAMIN, CYANOCOBALAMIN
CYANOKIT, HYDROXOCOBALAMIN
CYCLAFEM 0.5/35, ETHINYL ESTRADIOL
CYCLAFEM 1/35, ETHINYL ESTRADIOL
CYCLAFEM 7/7/7, ETHINYL ESTRADIOL
CYCLESSA, DESOGESTREL
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
CYCLOGYL, CYCLOPENTOLATE HYDROCHLORIDE
CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE
CYCLOPENTOLATE HYDROCHLORIDE, CYCLOPENTOLATE HYDROCHLORIDE
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
CYCLOSET, BROMOCRIPTINE MESYLATE
CYCLOSPORINE, CYCLOSPORINE
CYKLOKAPRON, TRANEXAMIC ACID
CYMBALTA, DULOXETINE HYDROCHLORIDE
CYONANZ, ETHINYL ESTRADIOL

APPENDIX A - PRODUCT NAME INDEX

** C **

CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 CYSTADANE, BETAINE
 CYSTAGON, CYSTEAMINE BITARTRATE
 CYSTARAN, CYSTEAMINE HYDROCHLORIDE
 CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
 CYSTOGRAFIN, DIATRIZOATE MEGLUMINE
 CYSTOGRAFIN DILUTE, DIATRIZOATE MEGLUMINE
 CYSVIEW KIT, HEXAMINOLEVULINATE HYDROCHLORIDE
 CYTARABINE, CYTARABINE
 CYTOMEL, LIOTHYRONINE SODIUM
 CYTOTEC, MISOPROSTOL
 CYTOVENE, GANCICLOVIR SODIUM

** D **

D.H.E. 45, DIHYDROERGOTAMINE MESYLATE
 DACARBAZINE, DACARBAZINE
 DACOGEN, DECITABINE
 DACTINOMYCIN, DACTINOMYCIN
 DAKLINZA, DACLATASVIR DIHYDROCHLORIDE
 DALFAMPRIDINE, DALFAMPRIDINE
 DALIRESP, ROFLUMILAST
 DALVANCE, DALBAVANCIN HYDROCHLORIDE
 DANAZOL, DANAZOL
 DANTRIUM, DANTROLENE SODIUM
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DAPSONE, DAPSONE
 DAPTOMYCIN, DAPTOMYCIN
 DARAPRIM, PYRIMETHAMINE
 DARIFENACIN, DARIFENACIN HYDROBROMIDE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DARUNAVIR ETHANOLATE, DARUNAVIR ETHANOLATE
 DASETTA 1/35, ETHINYL ESTRADIOL
 DASETTA 7/7/7, ETHINYL ESTRADIOL
 DATSCAN, IOFLUPANE I-123
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DAURISMO, GLASDEGIB
 DAYPRO, OXAPROZIN
 DAYSEE, ETHINYL ESTRADIOL
 DAYTRANA, METHYLPHENIDATE
 DDAVP, DESMOPRESSIN ACETATE
 DDAVP (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 DECITABINE, DECITABINE
 DEFERASIROX, DEFERASIROX
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEFINITY, PERFLUTREN
 DEFITELIO, DEFIBROTIDE SODIUM
 DELESTROGEN, ESTRADIOL VALERATE
 DELFLEX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELSTRIGO, DORAVIRINE
 DELSYM, DEXTROMETHORPHAN POLISTIREX (OTC)
 DELZICOL, MESALAMINE
 DEMADAX, TORSEMIDE
 DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 DEMEROL, MEPERIDINE HYDROCHLORIDE
 DEMSER, METYROSINE
 DENAVIR, PENCICLOVIR
 DEPACON, VALPROATE SODIUM

APPENDIX A - PRODUCT NAME INDEX

** D **

DEPAKENE, VALPROIC ACID
 DEPAKOTE, DIVALPROEX SODIUM
 DEPAKOTE ER, DIVALPROEX SODIUM
 DEPEN, PENICILLAMINE
 DEPO-ESTRADIOL, ESTRADIOL CYPIONATE
 DEPO-MEDROL, METHYLPREDNISOLONE ACETATE
 DEPO-PROVERA, MEDROXYPROGESTERONE ACETATE
 DEPO-SUBQ PROVERA 104, MEDROXYPROGESTERONE ACETATE
 DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE
 DERMA-SMOOTH/FS, FLUOCINOLONE ACETONIDE
 DERMABET, BETAMETHASONE VALERATE
 DERMATOP, PREDNICARBATE
 DERMATOP E EMOLLIENT, PREDNICARBATE
 DERMOTIC, FLUOCINOLONE ACETONIDE
 DESCOVY, EMTRICITABINE
 DESFERAL, DEFEROXAMINE MESYLATE
 DESFLURANE, DESFLURANE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DESLORATADINE, DESLORATADINE
 DESLORATADINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR, DESLORATADINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 DESOGEN, DESOGESTREL
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DESONATE, DESONIDE
 DESONIDE, DESONIDE
 DESOWEN, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DESOXYN, METHAMPHETAMINE HYDROCHLORIDE
 DESVENLAFAXINE, DESVENLAFAXINE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DETROL, TOLTERODINE TARTRATE
 DETROL LA, TOLTERODINE TARTRATE
 DEXAMETHASONE, DEXAMETHASONE
 DEXAMETHASONE INTENSOL, DEXAMETHASONE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXASPORIN, DEXAMETHASONE
 DEXCHLORPHENIRAMINE MALEATE, DEXCHLORPHENIRAMINE MALEATE
 DEXEDRINE, DEXTROAMPHETAMINE SULFATE
 DEXFERRUM, IRON DEXTRAN
 DEXILANT, DEXLANSOPRAZOLE
 DEXLANSOPRAZOLE, DEXLANSOPRAZOLE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 DEXTENZA, DEXAMETHASONE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE, DEXTROMETHORPHAN HYDROBROMIDE
 DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)
 DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DEXTROSE 20% IN PLASTIC CONTAINER, DEXTROSE

APPENDIX A - PRODUCT NAME INDEX

** D **

DEXTROSE 25%, DEXTROSE
 DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 30% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 40% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND ELECTROLYTE NO. 48 IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER,
 DEXTROSE 50%, DEXTROSE
 DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
 DEXYCU KIT, DEXAMETHASONE
 DIABETA, GLYBURIDE
 DIACOMIT, STIRIPENTOL
 DIAMOX, ACETAZOLAMIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIASTAT, DIAZEPAM
 DIASTAT ACUDIAL, DIAZEPAM
 DIAZEPAM, DIAZEPAM
 DIAZEPAM INTENSOL, DIAZEPAM
 DIBENZYLINE, PHENOXYBENZAMINE HYDROCHLORIDE
 DICLEGIS, DOXYLAMINE SUCCINATE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE), DICYCLOMINE HYDROCHLORIDE
 DIDANOSINE, DIDANOSINE

APPENDIX A - PRODUCT NAME INDEX

** D **

DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
DIFFERIN, ADAPALENE (OTC)
DIFFERIN, ADAPALENE
DIFICID, FIDAXOMICIN
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
DIFLUCAN, FLUCONAZOLE
DIFLUNISAL, DIFLUNISAL
DIGOXIN, DIGOXIN
DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
DILANTIN, PHENYTOIN
DILANTIN, PHENYTOIN SODIUM
DILANTIN-125, PHENYTOIN
DILATRATE-SR, ISOSORBIDE DINITRATE
DILAUDID, HYDROMORPHONE HYDROCHLORIDE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DILTZAC, DILTIAZEM HYDROCHLORIDE
DIMENHYDRINATE, DIMENHYDRINATE
DIMETHYL SULFOXIDE, DIMETHYL SULFOXIDE
DIOVAN, VALSARTAN
DIOVAN HCT, HYDROCHLOROTHIAZIDE
DIPENTUM, OLSALAZINE SODIUM
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
DIPRIVAN, PROPOFOL
DIPROLENE, BETAMETHASONE DIPROPIONATE
DIPROLENE AF, BETAMETHASONE DIPROPIONATE
DIPYRIDAMOLE, DIPYRIDAMOLE
DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
DISULFIRAM, DISULFIRAM
DITROPAN XL, OXYBUTYNIN CHLORIDE
DIURIL, CHLOROTHIAZIDE
DIURIL, CHLOROTHIAZIDE SODIUM
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DIVIGEL, ESTRADIOL
DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
DOCETAXEL, DOCETAXEL
DOCETAXEL, DOCETAXEL
DOCOSANOL, DOCOSANOL (OTC)
DOFETILIDE, DOFETILIDE
DOLOPHINE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
DOPRAM, DOXAPRAM HYDROCHLORIDE
DOPTelet, AVATROMBOPAG MALEATE
DORAL, QUAZEPAM
DORYX, DOXYCYCLINE HYCLATE
DORYX MPC, DOXYCYCLINE HYCLATE
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
DOTAREM, GADOTERATE MEGLUMINE
DOVONEX, CALCIPOTRIENE
DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
DOXERCALCIFEROL, DOXERCALCIFEROL
DOXIL (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
DOXY 100, DOXYCYCLINE HYCLATE
DOXY 200, DOXYCYCLINE HYCLATE

APPENDIX A - PRODUCT NAME INDEX

** D **

DOXYCYCLINE, DOXYCYCLINE
 DOXYCYCLINE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
 DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
 DRAX EXAMETAZIME, TECHNETIUM TC-99M EXAMETAZIME KIT
 DRAXIMAGE MDP-25, TECHNETIUM TC-99M MEDRONATE
 DRISDOL, ERGOCALCIFEROL
 DRONABINOL, DRONABINOL
 DROPERIDOL, DROPERIDOL
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONE
 DROXIA, HYDROXYUREA
 DSUVIA, SUFENTANIL CITRATE
 DTPA, TECHNETIUM TC-99M PENTETATE KIT
 DUAC, BENZOYL PEROXIDE
 DUAVEE, BAZEDOXIFENE ACETATE
 DUETACT, GLIMEPIRIDE
 DUEXIS, FAMOTIDINE
 DULERA, FORMOTEROL FUMARATE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUODOTE, ATROPINE
 DUOPA, CARBIDOPA
 DURACLON, CLONIDINE HYDROCHLORIDE
 DURAGESIC-100, FENTANYL
 DURAGESIC-12, FENTANYL
 DURAGESIC-25, FENTANYL
 DURAGESIC-37, FENTANYL
 DURAGESIC-50, FENTANYL
 DURAGESIC-75, FENTANYL
 DURAMORPH PF, MORPHINE SULFATE
 DURAPREP, IODINE POVACRYLEX (OTC)
 DUREZOL, DIFLUPREDNATE
 DURLAZA, ASPIRIN
 DUTASTERIDE, DUTASTERIDE
 DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
 DUTOPROL, HYDROCHLOROTHIAZIDE
 DUVOID, BETHANECHOL CHLORIDE
 DUZALLO, ALLOPURINOL
 DYANAVEL XR, AMPHETAMINE
 DYAZIDE, HYDROCHLOROTHIAZIDE
 DYCLOPRO, DYCLONINE HYDROCHLORIDE
 DYMISTA, AZELASTINE HYDROCHLORIDE
 DYNACIN, MINOCYCLINE HYDROCHLORIDE
 DYRENIUM, TRIAMTERENE

** E **

E-Z SCRUB 201, POVIDONE-IODINE (OTC)
 E-Z SCRUB 241, POVIDONE-IODINE (OTC)
 E-Z-HD, BARIUM SULFATE
 E-Z-PAQUE, BARIUM SULFATE
 E.E.S., ERYTHROMYCIN ETHYLSUCCINATE
 E.E.S. 400, ERYTHROMYCIN ETHYLSUCCINATE
 EC-NAPROSYN, NAPROXEN
 ECONAZOLE NITRATE, ECONAZOLE NITRATE
 ECOZA, ECONAZOLE NITRATE
 EDARBI, AZILSARTAN KAMEDOXOMIL
 EDARBYCLOR, AZILSARTAN KAMEDOXOMIL
 EDECRIN, ETHACRYNATE SODIUM
 EDECRIN, ETHACRYNIC ACID
 EDEX, ALPROSTADIL
 EDLUAR, ZOLPIDEM TARTRATE
 EDURANT, RILPIVIRINE HYDROCHLORIDE
 EFAVIRENZ, EFAVIRENZ
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ

APPENDIX A - PRODUCT NAME INDEX

** E **

EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE
EFFIENT, PRASUGREL HYDROCHLORIDE
EFUDEX, FLUOROURACIL
EGRIFTA, TESAMORELIN ACETATE
ELELYSO, TALIGLUCERASE ALFA
ELEPSIA XR, LEVETIRACETAM
ELESTAT, EPINASTINE HYDROCHLORIDE
ELESTRIN, ESTRADIOL
ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
ELIDEL, PIMECROLIMUS
ELIFEMME, ETHINYL ESTRADIOL
ELIGARD, LEUPROLIDE ACETATE
ELIMITE, PERMETHRIN
ELINEST, ETHINYL ESTRADIOL
ELIQUIS, APIXABAN
ELIXOPHYLLIN, THEOPHYLLINE
ELLA, ULIPRISTAL ACETATE
ELLENCEN, EPIRUBICIN HYDROCHLORIDE
ELLIOTTS B SOLUTION, CALCIUM CHLORIDE
ELMIRON, PENTOSAN POLYSULFATE SODIUM
ELOCON, MOMETASONE FUROATE
ELOXATIN, OXALIPLATIN
EMADINE, EMEDASTINE DIFUMARATE
EMBEDA, MORPHINE SULFATE
EMBELINE, CLOBETASOL PROPIONATE
EMBELINE E, CLOBETASOL PROPIONATE
EMCYT, ESTRAMUSTINE PHOSPHATE SODIUM
EMEND, APREPITANT
EMEND, FOSAPREPITANT DIMEGLUMINE
EMFLAZA, DEFLAZACORT
EMLA, LIDOCAINE
EMOQUETTE, DESOGESTREL
EMSAM, SELEGILINE
EMTRICITABINE, EMTRICITABINE
EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
EMTRIVA, EMTRICITABINE
EMVERM, MEBENDAZOLE
ENABLEX, DARIFENACIN HYDROBROMIDE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
ENALAPRILAT, ENALAPRILAT
ENDARI, L-GLUTAMINE
ENDOMETRIN, PROGESTERONE
ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
ENPRESSE-28, ETHINYL ESTRADIOL
ENSKYCE, DESOGESTREL
ENSTILAR, BETAMETHASONE DIPROPIONATE
ENTACAPONE, ENTACAPONE
ENTECAVIR, ENTECAVIR
ENTEREG, ALVIMOPAN
ENTOCORT EC, BUDESONIDE
ENTRESTO, SACUBITRIL
ENULOSE, LACTULOSE
ENVARUS XR, TACROLIMUS
EOVIST, GADOXETATE DISODIUM
EPANED, ENALAPRIL MALEATE
EPANED KIT, ENALAPRIL MALEATE
EPCLUSA, SOFOSBUVIR
EPHEDRINE SULFATE, EPHEDRINE SULFATE
EPIDIOLEX, CANNABIDIOL
EPIDUO, ADAPALENE
EPIDUO FORTE, ADAPALENE
EPIFOAM, HYDROCORTISONE ACETATE
EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** E **

EPINEPHRINE, EPINEPHRINE
EPINEPHRINE (AUTOINJECTOR), EPINEPHRINE
EPIPEN, EPINEPHRINE
EPIPEN JR., EPINEPHRINE
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
EPITOL, CARBAMAZEPINE
EPIVIR, LAMIVUDINE
EPIVIR-HBV, LAMIVUDINE
EPLERENONE, EPLERENONE
EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM
EPROSARTAN MESYLATE, EPROSARTAN MESYLATE
EPTIFIBATIDE, EPTIFIBATIDE
EPZICOM, ABACAVIR SULFATE
EQUETRO, CARBAMAZEPINE
ERAXIS, ANIDULAFUNGIN
ERGOCALCIFEROL, ERGOCALCIFEROL
ERGOLOID MESYLATES, ERGOLOID MESYLATES
ERGOMAR, ERGOTAMINE TARTRATE
ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
ERIVEDGE, VISMODEGIB
ERLEADA, APALUTAMIDE
ERRIN, NORETHINDRONE
ERTACZO, SERTACONAZOLE NITRATE
ERTAPENEM SODIUM, ERTAPENEM SODIUM
ERY-TAB, ERYTHROMYCIN
ERYC, ERYTHROMYCIN
ERYGEL, ERYTHROMYCIN
ERYPED, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROCIN, ERYTHROMYCIN LACTOBIONATE
ERYTHROCIN STEARATE, ERYTHROMYCIN STEARATE
ERYTHROMYCIN, ERYTHROMYCIN
ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
ESBRIET, PIRFENIDONE
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
ESKATA, HYDROGEN PEROXIDE
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
ESOMEPRAZOLE STRONTIUM, ESOMEPRAZOLE STRONTIUM
ESTARYLLA, ETHINYL ESTRADIOL
ESTAZOLAM, ESTAZOLAM
ESTRACE, ESTRADIOL
ESTRADIOL, ESTRADIOL
ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
ESTRADIOL AND NORGESTIMATE, ESTRADIOL
ESTRADIOL VALERATE, ESTRADIOL VALERATE
ESTRING, ESTRADIOL
ESTROGEL, ESTRADIOL
ESTROPIPATE, ESTROPIPATE
ESTROSTEP FE, ETHINYL ESTRADIOL
ESZOPICLONE, ESZOPICLONE
ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
ETHACRYNIC ACID, ETHACRYNIC ACID
ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
ETHAMOLIN, ETHANOLAMINE OLEATE
ETHOSUXIMIDE, ETHOSUXIMIDE
ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
ETHYOL, AMIFOSTINE
ETIDRONATE DISODIUM, ETIDRONATE DISODIUM
ETODOLAC, ETODOLAC
ETOMIDATE, ETOMIDATE

APPENDIX A - PRODUCT NAME INDEX

** E **

ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE
 ETOPOSIDE, ETOPOSIDE
 EUCRISA, CRISABOROLE
 EURAX, CROTAMITON
 EUTHYROX, LEVOTHYROXINE SODIUM **
 EVAMIST, ESTRADIOL
 EVEKEO, AMPHETAMINE SULFATE
 EVEROLIMUS, EVEROLIMUS
 EVISTA, RALOXIFENE HYDROCHLORIDE
 EVOCLIN, CLINDAMYCIN PHOSPHATE
 EVOMELA, MELPHALAN HYDROCHLORIDE
 EVOTAZ, ATAZANAVIR SULFATE
 EVOXAC, CEVIMELINE HYDROCHLORIDE
 EVZIO, NALOXONE HYDROCHLORIDE
 EXALGO, HYDROMORPHONE HYDROCHLORIDE
 EXCEDRIN (MIGRAINE), ACETAMINOPHEN (OTC)
 EXELDERM, SULCONAZOLE NITRATE
 EXELON, RIVASTIGMINE
 EXELON, RIVASTIGMINE TARTRATE
 EXEMESTANE, EXEMESTANE
 EXFORGE, AMLODIPINE BESYLATE
 EXFORGE HCT, AMLODIPINE BESYLATE
 EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)
 EXJADE, DEFERASIROX
 EXONDYS 51, ETEPLIRSEN
 EXPAREL, BUPIVACAINE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 EXTINA, KETOCONAZOLE
 EXTRANEAL, ICODextrin
 EZALLOR, ROSUVASTATIN CALCIUM
 EZETIMIBE, EZETIMIBE
 EZETIMIBE AND ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE

** F **

FABIOR, TAZAROTENE
 FACTIVE, GEMIFLOXACIN MESYLATE
 FALLBACK SOLO, LEVONORGESTREL (OTC)
 FALMINA, ETHINYL ESTRADIOL
 FAMCICLOVIR, FAMCICLOVIR
 FAMOTIDINE, FAMOTIDINE (OTC)
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER, FAMOTIDINE
 FAMOTIDINE, CALCIUM CARBONATE, AND MAGNESIUM HYDROXIDE, CALCIUM CARBONATE (OTC)
 FANAPT, ILOPERIDONE
 FARESTON, TOREMIFENE CITRATE
 FARXIGA, DAPAGLIFLOZIN
 FARYDAK, PANOBINOSTAT LACTATE
 FASLODEX, FULVESTRANT
 FAYOSIM, ETHINYL ESTRADIOL
 FAZACLO ODT, CLOZAPINE
 FELBAMATE, FELBAMATE
 FELBATOL, FELBAMATE
 FELDENE, PIROXICAM
 FELODIPINE, FELODIPINE
 FEMARA, LETROZOLE
 FEMCON FE, ETHINYL ESTRADIOL
 FEMHRT, ETHINYL ESTRADIOL
 FEMRING, ESTRADIOL ACETATE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FENOGLIDE, FENOFIBRATE
 FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE

APPENDIX A - PRODUCT NAME INDEX

** F **

FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
 FENTANYL CITRATE, FENTANYL CITRATE
 FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
 FENTANYL-100, FENTANYL
 FENTANYL-12, FENTANYL
 FENTANYL-25, FENTANYL
 FENTANYL-37, FENTANYL
 FENTANYL-50, FENTANYL
 FENTANYL-62, FENTANYL
 FENTANYL-75, FENTANYL
 FENTANYL-87, FENTANYL
 FENTORA, FENTANYL CITRATE
 FERAHEME, FERUMOXYTOL
 FERRIPROX, DEFERIPRONE
 FERRLECIT, SODIUM FERRIC GLUCONATE COMPLEX
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FETZIMA, LEVOMILNACIPRAN HYDROCHLORIDE
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FIASP, INSULIN ASPART
 FIASP FLEXTOUCH, INSULIN ASPART
 FIBRICOR, FENOFIBRIC ACID
 FINACEA, AZELAIC ACID
 FINASTERIDE, FINASTERIDE
 FIORICET W/ CODEINE, ACETAMINOPHEN
 FIORINAL, ASPIRIN
 FIORINAL W/CODEINE, ASPIRIN
 FIRAZYR, ICATIBANT ACETATE
 FIRDAPSE, AMIFAMPRIDINE PHOSPHATE
 FIRMAGON, DEGARELIX ACETATE
 FIRVANQ KIT, VANCOMYCIN HYDROCHLORIDE
 FLAC, FLUOCINOLONE ACETONIDE
 FLAGYL, METRONIDAZOLE
 FLAGYL I.V. RTU IN PLASTIC CONTAINER, METRONIDAZOLE
 FLAREX, FLUOROMETHOLONE ACETATE
 FLAVORED COLESTID, COLESTIPOL HYDROCHLORIDE
 FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLECTOR, DICLOFENAC EPOLAMINE
 FLOLAN, EPOPROSTENOL SODIUM
 FLOLIPID, SIMVASTATIN
 FLOMAX, TAMSULOSIN HYDROCHLORIDE
 FLONASE ALLERGY RELIEF, FLUTICASONE PROPIONATE (OTC)
 FLONASE SENSIMIST ALLERGY RELIEF, FLUTICASONE FUROATE (OTC)
 FLOVENT DISKUS 100, FLUTICASONE PROPIONATE
 FLOVENT DISKUS 250, FLUTICASONE PROPIONATE
 FLOVENT DISKUS 50, FLUTICASONE PROPIONATE
 FLOVENT HFA, FLUTICASONE PROPIONATE
 FLOXURIDINE, FLOXURIDINE
 FLUCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUCONAZOLE, FLUCONAZOLE
 FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 FLUCYTOSINE, FLUCYTOSINE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 FLUMADINE, RIMANTADINE HYDROCHLORIDE
 FLUMAZENIL, FLUMAZENIL
 FLUNISOLIDE, FLUNISOLIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE

APPENDIX A - PRODUCT NAME INDEX

** F **

FLUOCINONIDE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUORESCITE, FLUORESCEIN SODIUM
 FLUOROPLEX, FLUOROURACIL
 FLUOROURACIL, FLUOROURACIL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUOXYMESTERONE, FLUOXYMESTERONE
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FLURANDRENOLIDE, FLURANDRENOLIDE
 FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 FLURBIPROFEN, FLURBIPROFEN
 FLURBIPROFEN SODIUM, FLURBIPROFEN SODIUM
 FLUTAMIDE, FLUTAMIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 FML, FLUOROMETHOLONE
 FML FORTE, FLUOROMETHOLONE
 FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 FOLLISTIM AQ, FOLLITROPIN ALFA/BETA
 FOLOTYN, PRALATREXATE
 FOMEPIZOLE, FOMEPIZOLE
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 FORANE, ISOFLURANE
 FORFIVO XL, BUPROPION HYDROCHLORIDE
 FORTAMET, METFORMIN HYDROCHLORIDE
 FORTAZ, CEFTAZIDIME
 FORTEO, TERIPARATIDE RECOMBINANT HUMAN
 FORTESTA, TESTOSTERONE
 FOSAMAX, ALENDRONATE SODIUM
 FOSAMAX PLUS D, ALENDRONATE SODIUM
 FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FOSCAVIR, FOSCARNET SODIUM
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FOSRENOL, LANTHANUM CARBONATE
 FRAGMIN, DALTEPARIN SODIUM
 FREAMINE HBC 6.9%, AMINO ACIDS
 FREAMINE III 10%, AMINO ACIDS
 FREAMINE III 3% W/ ELECTROLYTES, AMINO ACIDS
 FREAMINE III 8.5%, AMINO ACIDS
 FREAMINE III 8.5% W/ ELECTROLYTES, AMINO ACIDS
 FROVA, FROVATRIPTAN SUCCINATE
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 FURADANTIN, NITROFURANTOIN
 FUROSEMIDE, FUROSEMIDE
 FUSILEV, LEVOLEUCOVORIN CALCIUM
 FUZEON, ENFUVIRTIDE
 FYAVOLV, ETHINYL ESTRADIOL
 FYCOMPA, PERAMPANEL

** G **

GABAPENTIN, GABAPENTIN
 GABITRIL, TIAGABINE HYDROCHLORIDE
 GABLOFEN, BACLOFEN
 GADAVIST, GADOBUTROL
 GALAFOLD, MIGALASTAT HYDROCHLORIDE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67

APPENDIX A - PRODUCT NAME INDEX

** G **

GALZIN, ZINC ACETATE
GANCICLOVIR, GANCICLOVIR
GANCICLOVIR, GANCICLOVIR SODIUM
GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
GANIRELIX ACETATE, GANIRELIX ACETATE
GASTROCROM, CROMOLYN SODIUM
GASTROGRAFIN, DIATRIZOATE MEGLUMINE
GATIFLOXACIN, GATIFLOXACIN
GATTEX KIT, TEDUGLUTIDE RECOMBINANT
GAVISCON, ALUMINUM HYDROXIDE (OTC)
GELNIQUE, OXYBUTYNIN CHLORIDE
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
GEMFIBROZIL, GEMFIBROZIL
GEMIFLOXACIN MESYLATE, GEMIFLOXACIN MESYLATE
GEMZAR, GEMCITABINE HYDROCHLORIDE
GEN-XENE, CLORAZEPATE DIPOTASSIUM
GENERLAC, LACTULOSE
GENGRAF, CYCLOSPORINE
GENOPTIC, GENTAMICIN SULFATE
GENOTROPIN, SOMATROPIN RECOMBINANT
GENOTROPIN PRESERVATIVE FREE, SOMATROPIN RECOMBINANT
GENTAK, GENTAMICIN SULFATE
GENTAMICIN SULFATE, GENTAMICIN SULFATE
GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
GENVOYA, COBICISTAT
GEODON, ZIPRASIDONE HYDROCHLORIDE
GEODON, ZIPRASIDONE MESYLATE
GIAPREZA, ANGIOTENSIN II ACETATE
GIAZO, BALSALAZIDE DISODIUM
GILDAGIA, ETHINYL ESTRADIOL
GILDESS 1.5/30, ETHINYL ESTRADIOL
GILDESS 1/20, ETHINYL ESTRADIOL
GILDESS 24 FE, ETHINYL ESTRADIOL
GILDESS FE 1.5/30, ETHINYL ESTRADIOL
GILDESS FE 1/20, ETHINYL ESTRADIOL
GILENYA, FINGOLIMOD HYDROCHLORIDE
GILOTRIF, AFATINIB DIMALEATE
GLATIRAMER ACETATE, GLATIRAMER ACETATE
GLATOPA, GLATIRAMER ACETATE
GLEEVEC, IMATINIB MESYLATE
GLEOLAN, AMINOLEVULINIC ACID HYDROCHLORIDE
GLEOSTINE, LOMUSTINE
GLIADEL, CARMUSTINE
GLIMEPIRIDE, GLIMEPIRIDE
GLIPIZIDE, GLIPIZIDE
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
GLOFIL-125, IOTHALAMATE SODIUM I-125
GLUCAGEN, GLUCAGON HYDROCHLORIDE
GLUCAGON, GLUCAGON
GLUCAGON, GLUCAGON HYDROCHLORIDE
GLUCOPHAGE, METFORMIN HYDROCHLORIDE
GLUCOPHAGE XR, METFORMIN HYDROCHLORIDE
GLUCOTROL, GLIPIZIDE
GLUCOTROL XL, GLIPIZIDE
GLUMETZA, METFORMIN HYDROCHLORIDE
GLYBURIDE, GLYBURIDE
GLYBURIDE (MICRONIZED), GLYBURIDE
GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
GLYCOLAX, POLYETHYLENE GLYCOL 3350 (OTC)
GLYCOLAX, POLYETHYLENE GLYCOL 3350
GLYCOPYRROLATE, GLYCOPYRROLATE
GLYDO, LIDOCAINE HYDROCHLORIDE
GLYNASE, GLYBURIDE
GLYRX-PF, GLYCOPYRROLATE

APPENDIX A - PRODUCT NAME INDEX

** G **

GLYSET, MIGLITOL
 GLYXAMBI, EMPAGLIFLOZIN
 GOCOVRI, AMANTADINE HYDROCHLORIDE
 GOLYTELY, POLYETHYLENE GLYCOL 3350
 GONAL-F, FOLLITROPIN ALFA/BETA
 GONAL-F RFF, FOLLITROPIN ALFA/BETA
 GONAL-F RFF REDI-JECT, FOLLITROPIN ALFA/BETA
 GONITRO, NITROGLYCERIN
 GOPRELTO, COCAINE HYDROCHLORIDE
 GRALISE, GABAPENTIN
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE, GRANISETRON HYDROCHLORIDE
 GRIS-PEG, GRISEOFULVIN, ULTRAMICROSIZE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
 GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
 GUAIFENESIN, GUAIFENESIN (OTC)
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 GUANABENZ ACETATE, GUANABENZ ACETATE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 GUANIDINE HYDROCHLORIDE, GUANIDINE HYDROCHLORIDE
 GYNAZOLE-1, BUTOCONAZOLE NITRATE

** H **

H.P. ACTHAR GEL, CORTICOTROPIN
 HABITROL, NICOTINE (OTC)
 HAILEY 1.5/30, ETHINYL ESTRADIOL
 HAILEY FE 1.5/30, ETHINYL ESTRADIOL
 HAILEY FE 1/20, ETHINYL ESTRADIOL
 HALAVEN, ERIBULIN MESYLATE
 HALCION, TRIAZOLAM
 HALDOL, HALOPERIDOL DECANOATE
 HALDOL, HALOPERIDOL LACTATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HALOG, HALCINONIDE
 HALOPERIDOL, HALOPERIDOL
 HALOPERIDOL, HALOPERIDOL LACTATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HARVONI, LEDIPASVIR
 HEATHER, NORETHINDRONE
 HECTOROL, DOXERCALCIFEROL
 HEMABATE, CARBOPROST TROMETHAMINE
 HEMANGEOL, PROPRANOLOL HYDROCHLORIDE
 HEPARIN SODIUM, HEPARIN SODIUM
 HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPATAMINE 8%, AMINO ACIDS
 HEPATOLITE, TECHNETIUM TC-99M DISOFENIN KIT
 HEPSERA, ADEFOVIR DIPIVOXIL
 HER STYLE, LEVONORGESTREL (OTC)
 HETLIOZ, TASIMELTEON
 HEXALEN, ALTRETAMINE
 HIBICLENS, CHLORHEXIDINE GLUCONATE (OTC)
 HIBISTAT, CHLORHEXIDINE GLUCONATE (OTC)

APPENDIX A - PRODUCT NAME INDEX

** H **

HICON, SODIUM IODIDE I-131
 HIPREX, METHENAMINE HIPPURATE
 HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE
 HORIZANT, GABAPENTIN ENACARBIL
 HUMALOG, INSULIN LISPRO RECOMBINANT
 HUMALOG KWIKPEN, INSULIN LISPRO RECOMBINANT
 HUMALOG MIX 50/50, INSULIN LISPRO PROTAMINE RECOMBINANT
 HUMALOG MIX 50/50 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT
 HUMALOG MIX 75/25, INSULIN LISPRO PROTAMINE RECOMBINANT
 HUMALOG MIX 75/25 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT
 HUMATROPE, SOMATROPIN RECOMBINANT
 HUMULIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
 HUMULIN 70/30 PEN, INSULIN RECOMBINANT HUMAN (OTC)
 HUMULIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)
 HUMULIN R, INSULIN HUMAN
 HUMULIN R, INSULIN RECOMBINANT HUMAN (OTC)
 HUMULIN R KWIKPEN, INSULIN HUMAN
 HUMULIN R PEN, INSULIN RECOMBINANT HUMAN (OTC)
 Hycamtin, Topotecan Hydrochloride
 HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDREA, HYDROXYUREA
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 HYDROCODONE BITARTRATE AND PSEUDOEPHEDRINE HYDROCHLORIDE, HYDROCODONE BITARTRATE
 HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE,
 HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX, CHLORPHENIRAMINE POLISTIREX
 HYDROCODONE POLISTIREX AND CHLORPHENIRAMNE POLISTIREX, CHLORPHENIRAMINE POLISTIREX
 HYDROCORTISONE, HYDROCORTISONE
 HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 HYDROCORTISONE IN ABSORBASE, HYDROCORTISONE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 HYDROXOCOBALAMIN, HYDROXOCOBALAMIN
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE
 HYDROXYUREA, HYDROXYUREA
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 HYLENEX RECOMBINANT, HYALURONIDASE RECOMBINANT HUMAN
 HYSINGLA, HYDROCODONE BITARTRATE
 HYZAAR, HYDROCHLOROTHIAZIDE

** I **

IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IBRANCE, PALBOCICLIB
 IBU-TAB, IBUPROFEN
 IBU-TAB 200, IBUPROFEN (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
 IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN LYSINE, IBUPROFEN LYSINE
 IBUPROFEN SODIUM, IBUPROFEN SODIUM (OTC)
 IBUPROHM, IBUPROFEN (OTC)
 IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)
 IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
 IC-GREEN, INDOCYANINE GREEN
 ICLEVIA, ETHINYL ESTRADIOL

APPENDIX A - PRODUCT NAME INDEX

** I **

ICLUSIG, PONATINIB HYDROCHLORIDE
IDAMYCIN PFS, IDARUBICIN HYDROCHLORIDE
IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
IDARUBICIN HYDROCHLORIDE PFS, IDARUBICIN HYDROCHLORIDE
IDHIFA, ENASIDENIB MESYLATE
IDKIT:HP, CITRIC ACID
IFEX, IFOSFAMIDE
IFOSFAMIDE, IFOSFAMIDE
ILEVRO, NEPAFENAC
ILOPERIDONE, ILOPERIDONE
ILUVIEN, FLUOCINOLONE ACETONIDE
IMATINIB MESYLATE, IMATINIB MESYLATE
IMBRUVICA, IBRUTINIB
IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
IMIQUIMOD, IMIQUIMOD
IMITREX, SUMATRIPTAN
IMITREX, SUMATRIPTAN SUCCINATE
IMITREX STATDOSE, SUMATRIPTAN SUCCINATE
IMODIUM A-D, LOPERAMIDE HYDROCHLORIDE (OTC)
IMODIUM MULTI-SYMPOM RELIEF, LOPERAMIDE HYDROCHLORIDE (OTC)
IMPAVIDO, MILTEFOSINE
IMPOYZ, CLOBETASOL PROPIONATE
IMURAN, AZATHIOPRINE
IMVEXXY, ESTRADIOL
INAPSINE, DROPERIDOL
INBRIJA, LEVODOPA
INCASSIA, NORETHINDRONE
INCRELEX, MECASERMIN RECOMBINANT
INCRUSE ELLIPTA, UMECLIDINIUM BROMIDE
INDAPAMIDE, INDAPAMIDE
INDERAL LA, PROPRANOLOL HYDROCHLORIDE
INDICLOR, INDIUM IN-111 CHLORIDE
INDIUM IN 111 CHLORIDE, INDIUM IN-111 CHLORIDE
INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE
INDOCIN, INDOMETHACIN
INDOCIN, INDOMETHACIN SODIUM
INDOCYANINE GREEN, INDOCYANINE GREEN
INDOMETHACIN, INDOMETHACIN
INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
INFANTS' FEVERALL, ACETAMINOPHEN (OTC)
INFASURF PRESERVATIVE FREE, CALFACTANT
INFED, IRON DEXTRAN
INFUGEM, GEMCITABINE HYDROCHLORIDE
INFUMORPH, MORPHINE SULFATE
INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE
INFUVITE PEDIATRIC, ASCORBIC ACID
INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID
INGENOL MEBUTATE, INGENOL MEBUTATE
INGREZZA, VALBENZAZINE TOSYLATE
INJECTAFER, FERRIC CARBOXYMALTOSE
INLYTA, AXITINIB
INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE
INOMAX, NITRIC OXIDE
INSPIRA, EPLERENONE
INTEGRILIN, EPTIFIBATIDE
INTELENCE, ETRAVIRINE
INTERMEZZO, ZOLPIDEM TARTRATE
INTRALIPID 10%, SOYBEAN OIL
INTRALIPID 20%, SOYBEAN OIL
INTRALIPID 30%, SOYBEAN OIL
INTRAROSA, PRASTERONE
INTROVALE, ETHINYL ESTRADIOL
INTUNIV, GUANFACINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** I **

INVANZ, ERTAPENEM SODIUM
 INVEGA, PALIPERIDONE
 INVEGA SUSTENNA, PALIPERIDONE PALMITATE
 INVEGA TRINZA, PALIPERIDONE PALMITATE
 INVELTYS, LOTEPREDNOL ETABONATE
 INVIRASE, SAQUINAVIR MESYLATE
 INVOKAMET, CANAGLIFLOZIN
 INVOKAMET XR, CANAGLIFLOZIN
 INVOKANA, CANAGLIFLOZIN
 IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 IOPIDINE, APRACLONIDINE HYDROCHLORIDE
 IOSAT, POTASSIUM IODIDE (OTC)
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 IRBESARTAN, IRBESARTAN
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRESSA, GEFITINIB
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 ISENTRESS, RALTEGRAVIR POTASSIUM
 ISENTRESS HD, RALTEGRAVIR POTASSIUM
 ISIBLOOM, DESOGESTREL
 ISOFLURANE, ISOFLURANE
 ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 ISOLYTE S IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 ISOLYTE S PH 7.4 IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 ISONIAZID, ISONIAZID
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 ISOPTO ATROPINE, ATROPINE SULFATE
 ISOPTO CARPINE, Pilocarpine Hydrochloride
 ISORDIL, ISOSORBIDE DINITRATE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 ISOSULFAN BLUE, ISOSULFAN BLUE
 ISOTRETINOIN, ISOTRETINOIN
 ISOVUE-200, IOPAMIDOL
 ISOVUE-250, IOPAMIDOL
 ISOVUE-300, IOPAMIDOL
 ISOVUE-370, IOPAMIDOL
 ISOVUE-M 200, IOPAMIDOL
 ISOVUE-M 300, IOPAMIDOL
 ISRADIPINE, ISRADIPINE
 ISTALOL, TIMOLOL MALEATE
 ISTODAX, ROMIDEPSIN
 ISUPREL, ISOPROTERENOL HYDROCHLORIDE
 ITRACONAZOLE, ITRACONAZOLE
 IVERMECTIN, IVERMECTIN
 IVY BLOCK, BENTOQUATAM (OTC)
 IXEMPRA KIT, IXABEPILONE

** J **

JADENU, DEFERASIROX
 JADENU SPRINKLE, DEFERASIROX
 JAIMIESS, ETHINYL ESTRADIOL
 JAKAFI, RUXOLITINIB PHOSPHATE
 JALYN, DUTASTERIDE
 JANTOVEN, WARFARIN SODIUM
 JANUMET, METFORMIN HYDROCHLORIDE
 JANUMET XR, METFORMIN HYDROCHLORIDE
 JANUVIA, SITAGLIPTIN PHOSPHATE
 JARDIANCE, EMPAGLIFLOZIN
 JEANATOPE, ALBUMIN IODINATED I-125 SERUM
 JENCYCLA, NORETHINDRONE
 JENTADUETO, LINAGLIPTIN
 JENTADUETO XR, LINAGLIPTIN
 JEVTANA KIT, CABAZITAXEL
 JORNAY PM, METHYLPHENIDATE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** J **

JUBLIA, EFINACONAZOLE
 JULUCA, DOLUTEGRAVIR SODIUM
 JUNEL 1.5/30, ETHINYL ESTRADIOL
 JUNEL 1/20, ETHINYL ESTRADIOL
 JUNEL FE 1.5/30, ETHINYL ESTRADIOL
 JUNEL FE 1/20, ETHINYL ESTRADIOL
 JUNIOR STRENGTH ADVIL, IBUPROFEN (OTC)
 JUNIOR STRENGTH IBUPROFEN, IBUPROFEN (OTC)
 JUNIOR STRENGTH MOTRIN, IBUPROFEN (OTC)
 JUXTAPID, LOMITAPIDE MESYLATE
 JYNARQUE, TOLVAPTAN

** K **

K-TAB, POTASSIUM CHLORIDE
 KABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
 KADIAN, MORPHINE SULFATE
 KAITLIB FE, ETHINYL ESTRADIOL
 KALETRA, LOPINAVIR
 KALEXATE, SODIUM POLYSTYRENE SULFONATE
 KALLIGA, DESOGESTREL
 KALYDECO, IVACAFTOR
 KAPSPARGO SPRINKLE, METOPROLOL SUCCINATE
 KAPVAY, CLONIDINE HYDROCHLORIDE
 KARBINAL ER, CARBINOXAMINE MALEATE
 KARIVA, DESOGESTREL
 KAZANO, ALOGLIPTIN BENZOATE
 KEFLEX, CEPHALEXIN
 KELNOR, ETHINYL ESTRADIOL
 KENALOG, TRIAMCINOLONE ACETONIDE
 KENALOG-10, TRIAMCINOLONE ACETONIDE
 KENALOG-40, TRIAMCINOLONE ACETONIDE
 KENGREAL, CANGRELOR
 KEPPRA, LEVETIRACETAM
 KEPPRA XR, LEVETIRACETAM
 KERYDIN, TAVABOROLE
 KETALAR, KETAMINE HYDROCHLORIDE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 KETOCONAZOLE, KETOCONAZOLE
 KETOPROFEN, KETOPROFEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 KETOZOLE, KETOCONAZOLE
 KEVEYIS, DICHLORPHENAMIDE
 KHAPZORY, LEVOLEUCOVORIN
 KHEDEZLA, DESVENLAFAXINE
 KIMIDESS, DESOGESTREL
 KINEVAC, SINCALIDE
 KIONEX, SODIUM POLYSTYRENE SULFONATE
 KISQALI, RIBOCICLIB SUCCINATE
 KISQALI FEMARA CO-PACK (COPACKAGED), LETROZOLE
 KITABIS PAK, TOBRAMYCIN
 KLARON, SULFACETAMIDE SODIUM
 KLONOPIN, CLONAZEPAM
 KLOR-CON, POTASSIUM CHLORIDE
 KLOR-CON M10, POTASSIUM CHLORIDE
 KLOR-CON M15, POTASSIUM CHLORIDE
 KLOR-CON M20, POTASSIUM CHLORIDE
 KOMBIGLYZE XR, METFORMIN HYDROCHLORIDE
 KORLYM, MIFEPRISTONE
 KOVANAZE, OXYMETAZOLINE HYDROCHLORIDE
 KRINTAFEL, TAFENOQUINE SUCCINATE
 KURVELO, ETHINYL ESTRADIOL
 KUVAN, SAPROPTERIN DIHYDROCHLORIDE
 KYBELLA, DEOXYCHOLIC ACID
 KYLEENA, LEVONORGESTREL

APPENDIX A - PRODUCT NAME INDEX

** K **

KYPROLIS, CARFILZOMIB

** L **

LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LACRISERT, HYDROXYPROPYL CELLULOSE
 LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LACTULOSE, LACTULOSE
 LAMICTAL, LAMOTRIGINE
 LAMICTAL CD, LAMOTRIGINE
 LAMICTAL ODT, LAMOTRIGINE
 LAMICTAL XR, LAMOTRIGINE
 LAMISIL, TERBINAFINE HYDROCHLORIDE (OTC)
 LAMISIL, TERBINAFINE HYDROCHLORIDE
 LAMISIL AT, TERBINAFINE (OTC)
 LAMISIL AT, TERBINAFINE HYDROCHLORIDE (OTC)
 LAMIVUDINE, LAMIVUDINE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMOTRIGINE, LAMOTRIGINE
 LANIAZID, ISONIAZID
 LANORINAL, ASPIRIN
 LANOXIN, DIGOXIN
 LANOXIN PEDIATRIC, DIGOXIN
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LANSOPRAZOLE, LANSOPRAZOLE
 LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN, AMOXICILLIN
 LANTHANUM CARBONATE, LANTHANUM CARBONATE
 LANTUS, INSULIN GLARGINE RECOMBINANT
 LANTUS SOLOSTAR, INSULIN GLARGINE RECOMBINANT
 LARIN 1.5/30, ETHINYL ESTRADIOL
 LARIN 1/20, ETHINYL ESTRADIOL
 LARIN 24 FE, ETHINYL ESTRADIOL
 LARIN FE 1.5/30, ETHINYL ESTRADIOL
 LARIN FE 1/20, ETHINYL ESTRADIOL
 LAROTID, AMOXICILLIN
 LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE
 LASIX, FUROSEMIDE
 LASTACAPT, ALCAFTADINE
 LATANOPROST, LATANOPROST
 LATISSE, BIMATOPROST
 LATUDA, LURASIDONE HYDROCHLORIDE
 LAX-LYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
 LAZANDA, FENTANYL CITRATE
 LEFLUNOMIDE, LEFLUNOMIDE
 LENVIMA, LENVATINIB MESYLATE
 LERIBANE, ETHINYL ESTRADIOL
 LESCOL XL, FLUVASTATIN SODIUM
 LESSINA-28, ETHINYL ESTRADIOL
 LETAIRIS, AMBRISENTAN
 LETROZOLE, LETROZOLE
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEUKERAN, CHLORAMBUCIL
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 LEVEMIR, INSULIN DETEMIR RECOMBINANT
 LEVEMIR FLEXTOUCH, INSULIN DETEMIR RECOMBINANT
 LEVETIRACETAM, LEVETIRACETAM
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LEVITRA, VARDENAFIL HYDROCHLORIDE
 LEVO-T, LEVOTHYROXINE SODIUM **
 LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
 LEVOCARNITINE, LEVOCARNITINE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** L **

LEVOFLOXACIN, LEVOFLOXACIN
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 LEVONEST, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LEVONORGESTREL, LEVONORGESTREL
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVOPHED, NOREPINEPHRINE BITARTRATE
 LEVORA 0.15/30-28, ETHINYL ESTRADIOL
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LEVOXYL, LEVOTHYROXINE SODIUM **
 LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE
 LEXAPRO, ESCITALOPRAM OXALATE
 LEXISCAN, REGADENOSON
 LEXIVA, FOSAMPRENAVIR CALCIUM
 LIALDA, MESALAMINE
 LIBRAX, CHLORDIAZEPOXIDE HYDROCHLORIDE
 LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE
 LICART, DICLOFENAC EPOLAMINE
 LIDEX, FLUOCINONIDE
 LIDEX-E, FLUOCINONIDE
 LIDOCAINE, LIDOCAINE
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE
 LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE VISCOUS, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE
 LIDODERM, LIDOCAINE
 LIGNOSPAN FORTE, EPINEPHRINE BITARTRATE
 LIGNOSPAN STANDARD, EPINEPHRINE BITARTRATE
 LILETTA, LEVONORGESTREL
 LINCOCIN, LINCOMYCIN HYDROCHLORIDE
 LINCOMYCIN, LINCOMYCIN HYDROCHLORIDE
 LINDANE, LINDANE
 LINEZOLID, LINEZOLID
 LINEZOLID IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, LINEZOLID
 LINZESS, LINALOTIDE
 LIORESAL, BACLOFEN
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 LIPIODOL, ETHIODIZED OIL
 LIPITOR, ATORVASTATIN CALCIUM
 LIPOFEN, FENOFIBRATE
 LIQUID E-Z-PAQUE, BARIUM SULFATE
 LISINOPRIL, LISINOPRIL
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 LITHIUM CITRATE, LITHIUM CITRATE
 LITHOBID, LITHIUM CARBONATE
 LITHOSTAT, ACETOHYDROXAMIC ACID
 LIVALO, PITAVASTATIN CALCIUM
 LO LOESTRIN FE, ETHINYL ESTRADIOL
 LO SIMPESE, ETHINYL ESTRADIOL
 LO-ZUMANDIMINE, DROSPIRENONE
 LOCID, HYDROCORTISONE BUTYRATE
 LOCID LIPOCREAM, HYDROCORTISONE BUTYRATE

APPENDIX A - PRODUCT NAME INDEX

** L **

LODOSYN, CARBIDOPA
LOESTRIN 21 1.5/30, ETHINYL ESTRADIOL
LOESTRIN 21 1/20, ETHINYL ESTRADIOL
LOESTRIN 24 FE, ETHINYL ESTRADIOL
LOESTRIN FE 1.5/30, ETHINYL ESTRADIOL
LOESTRIN FE 1/20, ETHINYL ESTRADIOL
LOGILIA, ULIPRISTAL ACETATE
LOKELMA, SODIUM ZIRCONIUM CYCLOSILICATE
LOMAIRA, PHENTERMINE HYDROCHLORIDE
LOMOTIL, ATROPINE SULFATE
LONHALA MAGNAIR KIT, GLYCOPYRROLATE
LONSURF, TIPIRACIL HYDROCHLORIDE
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
LOPID, GEMFIBROZIL
LOPINA VIR AND RITONAVIR, LOPINA VIR
LOPRESSOR, METOPROLOL TARTRATE
LOPRESSOR HCT, HYDROCHLOROTHIAZIDE
LOPROX, CICLOPIROX
LOPURIN, ALLOPURINOL
LORATADINE, LORATADINE (OTC)
LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
LORATADINE REDIDOSE, LORATADINE (OTC)
LORAZEPAM, LORAZEPAM
LORAZEPAM INTENSOL, LORAZEPAM
LORBRENA, LORLATINIB
LORYNA, DROSPIRENONE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LOSEASONIQUE, ETHINYL ESTRADIOL
LOTEMAX, LOTE PREDNOL ETABONATE
LOTENSIN, BENAZEPRIL HYDROCHLORIDE
LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
LOTREL, AMLODIPINE BESYLATE
LOTRIMIN ULTRA, BUTENAFINE HYDROCHLORIDE (OTC)
LOTRISONE, BETAMETHASONE DIPROPIONATE
LOTRONEX, ALOSETRON HYDROCHLORIDE
LOVASTATIN, LOVASTATIN
LOVAZA, OMEGA-3-ACID ETHYL ESTERS
LOVENOX, ENOXAPARIN SODIUM
LOVENOX (PRESERVATIVE FREE), ENOXAPARIN SODIUM
LOW-OGESTREL-28, ETHINYL ESTRADIOL
LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
LUCEMYRA, LOFEXIDINE HYDROCHLORIDE
LUMASON, SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES
LUMIFY, BRIMONIDINE TARTRATE (OTC)
LUMIGAN, BIMATOPROST
LUNESTA, ESZOPICLONE
LUPANETA PACK, LEUPROLIDE ACETATE
LUPRON DEPOT, LEUPROLIDE ACETATE
LUPRON DEPOT-PED, LEUPROLIDE ACETATE
LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
LUTATHERA, LUTETIUM DOTATATE LU-177
LUTRATE DEPOT KIT, LEUPROLIDE ACETATE
LUVOX, FLUVOXAMINE MALEATE
LUXIQ, BETAMETHASONE VALERATE
LUZU, LULICONAZOLE
LYMPHOSEEK KIT, TECHNETIUM TC-99M TILMANOCEPT
LYNPARZA, OLAPARIB
LYRICA, PREGABALIN
LYRICA CR, PREGABALIN
LYSODREN, MITOTANE
LYSTEDA, TRANEXAMIC ACID

APPENDIX A - PRODUCT NAME INDEX

** M **

M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
M.V.I. ADULT, ASCORBIC ACID
M.V.I. ADULT (PHARMACY BULK PACKAGE), ASCORBIC ACID
M.V.I. PEDIATRIC, ASCORBIC ACID
MACRILEN, MACIMORELIN ACETATE
MACROBID, NITROFURANTOIN
MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE
MACUGEN, PEGAPTANIB SODIUM
MAFENIDE ACETATE, MAFENIDE ACETATE
MAGNESIUM SULFATE, MAGNESIUM SULFATE
MAGNESIUM SULFATE, MAGNESIUM SULFATE
MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
MAGNEVIST, GADOPENTETATE DIMEGLUMINE
MAKENA, HYDROXYPROGESTERONE CAPROATE
MAKENA (AUTOINJECTOR), HYDROXYPROGESTERONE CAPROATE
MAKENA PRESERVATIVE FREE, HYDROXYPROGESTERONE CAPROATE
MALARONE, ATOVAQUONE
MALARONE PEDIATRIC, ATOVAQUONE
MALATHION, MALATHION
MALMOREDE, ETHINYL ESTRADIOL
MANGANESE CHLORIDE IN PLASTIC CONTAINER, MANGANESE CHLORIDE
MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 25%, MANNITOL
MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL
MAPROTILINE HYDROCHLORIDE, MAPROTILINE HYDROCHLORIDE
MARCAINE, BUPIVACAINE HYDROCHLORIDE
MARCAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
MARCAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
MARCAINE HYDROCHLORIDE W/ EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
MARINOL, DRONABINOL
MARLISSA, ETHINYL ESTRADIOL
MARPLAN, ISOCARBOXAZID
MARQIBO KIT, VINCRISTINE SULFATE
MATULANE, PROCARBAZINE HYDROCHLORIDE
MAVYRET, GLECAPREVIR
MAXALT, RIZATRIPTAN BENZOATE
MAXALT-MLT, RIZATRIPTAN BENZOATE
MAXIDEX, DEXAMETHASONE
MAXIPIME, CEFEPIME HYDROCHLORIDE
MAXITROL, DEXAMETHASONE
MAXZIDE, HYDROCHLOROTHIAZIDE
MAXZIDE-25, HYDROCHLOROTHIAZIDE
MD-GASTROVIEW, DIATRIZOATE MEGLUMINE
MECAMYLAMINE HYDROCHLORIDE, MECAMYLAMINE HYDROCHLORIDE
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM
MEDROL, METHYLPREDNISOLONE
MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
MEFENAMIC ACID, MEFENAMIC ACID
MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
MEFOXIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM
MEGACE ES, MEGESTROL ACETATE
MEGATOPE, ALBUMIN IODINATED I-131 SERUM
MEGESTROL ACETATE, MEGESTROL ACETATE
MEKINIST, TRAMETINIB DIMETHYL SULFOXIDE
MEKTOVI, BINIMETINIB
MELAMISA, DROSPIRENONE
MELOXICAM, MELOXICAM
MELPHALAN, MELPHALAN
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** M **

MEMBRANEBLUE, TRYPAN BLUE
MEN'S ROGAINE, MINOXIDIL (OTC)
MENEST, ESTROGENS, ESTERIFIED
MENOPUR, MENOTROPINS (FSH)
MENOSTAR, ESTRADIOL
MENTAX, BUTENAFINE HYDROCHLORIDE
MEPERIDINE HYDROCHLORIDE, MEPEPERIDINE HYDROCHLORIDE
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPEPERIDINE HYDROCHLORIDE
MEPHYTON, PHYTONADIONE
MEPROBAMATE, MEPROBAMATE
MEPRON, ATOVAQUONE
MERCAPTOPYRINE, MERCAPTOPYRINE
MEROPENEM, MEROPENEM
MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER, MEROPENEM
MERREM, MEROPENEM
MESALAMINE, MESALAMINE
MESNA, MESNA
MESNEX, MESNA
MESTINON, PYRIDOSTIGMINE BROMIDE
METADATE CD, METHYLPHENIDATE HYDROCHLORIDE
METADATE ER, METHYLPHENIDATE HYDROCHLORIDE
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
METASTRON, STRONTIUM CHLORIDE SR-89
METAXALONE, METAXALONE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METHADONE HYDROCHLORIDE INTENSOL, METHADONE HYDROCHLORIDE
METHADOSE, METHADONE HYDROCHLORIDE
METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
METHAZOLAMIDE, METHAZOLAMIDE
METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
METHERGINE, METHYLERGONOVINE MALEATE
METHIMAZOLE, METHIMAZOLE
METHOCARBAMOL, METHOCARBAMOL
METHOCARBAMOL AND ASPIRIN, ASPIRIN
METHOTREXATE PRESERVATIVE FREE, METHOTREXATE SODIUM
METHOTREXATE SODIUM, METHOTREXATE SODIUM
METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
METHOXSALEN, METHOXSALEN
METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
METHYLCLOTHIAZIDE, METHYLCLOTHIAZIDE
METHYLDOPA, METHYLDOPA
METHYLDOPA AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
METHYLDOPATE HYDROCHLORIDE, METHYLDOPATE HYDROCHLORIDE
METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
METHYLIN, METHYLPHENIDATE HYDROCHLORIDE
METHYLIN ER, METHYLPHENIDATE HYDROCHLORIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
METHYLPREDNISOLONE, METHYLPREDNISOLONE
METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
METHYLTESTOSTERONE, METHYLTESTOSTERONE
METIPRANOLOL, METIPRANOLOL HYDROCHLORIDE
METOCLOPRAMIDE, METOCLOPRAMIDE HYDROCHLORIDE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
METOLAZONE, METOLAZONE
METOPIRONE, METYRAPONE
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
METRO I.V. IN PLASTIC CONTAINER, METRONIDAZOLE
METROCREAM, METRONIDAZOLE
METROGEL, METRONIDAZOLE
METROGEL-VAGINAL, METRONIDAZOLE
METROLOTION, METRONIDAZOLE

APPENDIX A - PRODUCT NAME INDEX

** M **

METRONIDAZOLE, METRONIDAZOLE
METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
MIACALCIN, CALCITONIN SALMON
MIBELAS 24 FE, ETHINYL ESTRADIOL
MICARDIS, TELMISARTAN
MICARDIS HCT, HYDROCHLOROTHIAZIDE
MICONAZOLE 3, MICONAZOLE NITRATE (OTC)
MICONAZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MICONAZOLE 7, MICONAZOLE NITRATE (OTC)
MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
MICONAZOLE NITRATE, MICONAZOLE NITRATE
MICONAZOLE NITRATE COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MICORT-HC, HYDROCORTISONE ACETATE
MICRO-K, POTASSIUM CHLORIDE
MICRO-K 10, POTASSIUM CHLORIDE
MICROGESTIN 1.5/30, ETHINYL ESTRADIOL
MICROGESTIN 1/20, ETHINYL ESTRADIOL
MICROGESTIN FE 1.5/30, ETHINYL ESTRADIOL
MICROGESTIN FE 1/20, ETHINYL ESTRADIOL
MICRONOR, NORETHINDRONE
MICROZIDE, HYDROCHLOROTHIAZIDE
MIDAMOR, AMILORIDE HYDROCHLORIDE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
MIDOL LIQUID GELS, IBUPROFEN (OTC)
MIDOZALAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MIFEPRX, MIFEPRISTONE
MIGERGOT, CAFFEINE
MIGLITOL, MIGLITOL
MIGLUSTAT, MIGLUSTAT
MIGRANAL, DIHYDROERGOTAMINE MESYLATE
MILI, ETHINYL ESTRADIOL
MILRINONE LACTATE, MILRINONE LACTATE
MILRINONE LACTATE IN DEXTROSE 5%, MILRINONE LACTATE
MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
MILRINONE LACTATE IN PLASTIC CONTAINER, MILRINONE LACTATE
MINASTRIN 24 FE, ETHINYL ESTRADIOL
MINIPRESS, PRAZOSIN HYDROCHLORIDE
MINIRIN, DESMOPRESSIN ACETATE
MINITRAN, NITROGLYCERIN
MINIVELLE, ESTRADIOL
MINOCIN, MINOCYCLINE HYDROCHLORIDE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MINOLIRA, MINOCYCLINE HYDROCHLORIDE
MINOXIDIL, MINOXIDIL (OTC)
MINOXIDIL, MINOXIDIL
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
MIOCHOL-E, ACETYLCHOLINE CHLORIDE
MIOSTAT, CARBACHOL
MIRALAX, POLYETHYLENE GLYCOL 3350 (OTC)
MIRAPEX, PRAMIPEXOLE DIHYDROCHLORIDE
MIRAPEX ER, PRAMIPEXOLE DIHYDROCHLORIDE
MIRENA, LEVONORGESTREL
MIRTAZAPINE, MIRTAZAPINE
MIRVASO, BRIMONIDINE TARTRATE
MISOPROSTOL, MISOPROSTOL
MITIGARE, COLCHICINE
MITIGO, MORPHINE SULFATE
MITOMYCIN, MITOMYCIN
MITOSOL, MITOMYCIN
MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** M **

MOBIC, MELOXICAM
 MODAFINIL, MODAFINIL
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 MOLINDONE HYDROCHLORIDE, MOLINDONE HYDROCHLORIDE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MONISTAT 1 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MONISTAT 3, MICONAZOLE NITRATE (OTC)
 MONISTAT 3, MICONAZOLE NITRATE
 MONISTAT 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MONISTAT 3 COMBINATION PACK (PREFILLED), MICONAZOLE NITRATE (OTC)
 MONISTAT 7, MICONAZOLE NITRATE (OTC)
 MONISTAT 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MONO-LINYAH, ETHINYL ESTRADIOL
 MONODOX, DOXYCYCLINE
 MONOKET, ISOSORBIDE MONONITRATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MONUROL, FOSFOMYCIN TROMETHAMINE
 MORPHABOND ER, MORPHINE SULFATE
 MORPHINE SULFATE, MORPHINE SULFATE
 MOTEGRITY, PRUCALOPRIDE SUCCINATE
 MOTOFEN, ATROPINE SULFATE
 MOTRIN IB, IBUPROFEN (OTC)
 MOVANTIK, NALOXEGOL OXALATE
 MOVIPREP, ASCORBIC ACID
 MOXEZA, MOXIFLOXACIN HYDROCHLORIDE
 MOXIDECTIN, MOXIDECTIN
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER, MOXIFLOXACIN
 MOZOBIL, PLERIXAFOR
 MPI INDIUM DTPA IN 111, INDIUM IN-111 PENTETATE DISODIUM
 MS CONTIN, MORPHINE SULFATE
 MUCINEX, GUAIFENESIN (OTC)
 MUCINEX D, GUAIFENESIN (OTC)
 MUCINEX DM, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 MULPLETA, LUSUTROMBOPAG
 MULTAQ, DRONEDARONE HYDROCHLORIDE
 MULTIHANCE, GADOBENATE DIMEGLUMINE
 MULTIHANCE MULTIPACK, GADOBENATE DIMEGLUMINE
 MUPIROCIN, MUPIROCIN
 MUPIROCIN, MUPIROCIN CALCIUM
 MUSE, ALPROSTADIL
 MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE
 MYCAMINE, MICAFUNGIN SODIUM
 MYCOBUTIN, RIFABUTIN
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
 MYDAYIS, AMPHETAMINE ASPARTATE
 MYDRIACYL, TROPICAMIDE
 MYFORTIC, MYCOPHENOLIC ACID
 MYKACET, NYSTATIN
 MYLERAN, BUSULFAN
 MYORISAN, ISOTRETINOIN
 MYOVIEW 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT
 MYRBETRIQ, MIRABEGRON
 MYSOLINE, PRIMIDONE
 MYTESI, CROFELEMER
 MYZILRA, ETHINYL ESTRADIOL

** N **

NABUMETONE, NABUMETONE
 NADOLOL, NADOLOL
 NADOLOL AND BENDROFLUMETHIAZIDE, BENDROFLUMETHIAZIDE
 NAFACILLIN SODIUM, NAFACILLIN SODIUM

APPENDIX A - PRODUCT NAME INDEX

** N **

NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 NAFTIN, NAFTIFINE HYDROCHLORIDE
 NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE
 NALFON, FENOPROFEN CALCIUM
 NALLPEN IN PLASTIC CONTAINER, NAFICILLIN SODIUM
 NALOXONE, NALOXONE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 NAMENDA, MEMANTINE HYDROCHLORIDE
 NAMENDA XR, MEMANTINE HYDROCHLORIDE
 NAMZARIC, DONEPEZIL HYDROCHLORIDE
 NANDROLONE DECANOATE, NANDROLONE DECANOATE
 NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 NAPHCN-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
 NAPRELAN, NAPROXEN SODIUM
 NAPROSYN, NAPROXEN
 NAPROXEN, NAPROXEN
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NARCAN, NALOXONE HYDROCHLORIDE
 NARDIL, PHENELZINE SULFATE
 NAROPIN, ROPIVACAINE HYDROCHLORIDE
 NASACORT ALLERGY 24 HOUR, TRIAMCINOLONE ACETONIDE (OTC)
 NASCOBAL, CYANOCOBALAMIN
 NASONEX, MOMETASONE FUROATE
 NATACYN, NATAMYCIN
 NATAZIA, DIENOGEST
 NATEGLINIDE, NATEGLINIDE
 NATESTO, TESTOSTERONE
 NATRECOR, NESIRITIDE RECOMBINANT
 NATROBA, SPINOSAD
 NAVALBINE, VINORELBINE TARTRATE
 NEBUPENT, PENTAMIDINE ISETHIONATE
 NEDOCROMIL SODIUM, NEDOCROMIL SODIUM
 NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
 NEMBUTAL SODIUM, PENTOBARBITAL SODIUM
 NEO-SYNALAR, FLUOCINOLONE ACETONIDE
 NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
 NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
 NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
 NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE, BACITRACIN ZINC
 NEOMYCIN SULFATE, NEOMYCIN SULFATE
 NEOPAP, ACETAMINOPHEN (OTC)
 NEOPROFEN, IBUPROFEN LYSINE
 NEORAL, CYCLOSPORINE
 NEOSPORIN, BACITRACIN ZINC
 NEOSPORIN, GRAMICIDIN
 NEOSPORIN G.U. IRRIGANT, NEOMYCIN SULFATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NEPHRAMINE 5.4%, AMINO ACIDS
 NERLYNX, NERATINIB MALEATE
 NESACAINE, CHLOROPROCAINE HYDROCHLORIDE
 NESACAINE-MPF, CHLOROPROCAINE HYDROCHLORIDE
 NESINA, ALOGLIPTIN BENZOATE
 NETSPOT, GALLIUM DOTATATE GA-68
 NEUPRO, ROTIGOTINE
 NEURACEQ, FLORBETABEN F-18
 NEUROLITE, TECHNETIUM TC-99M BICISATE KIT
 NEURONTIN, GABAPENTIN

APPENDIX A - PRODUCT NAME INDEX

** N **

NEVANAC, NEPAFENAC
NEVIRAPINE, NEVIRAPINE
NEXAVAR, SORAFENIB TOSYLATE
NEXESTA FE, ETHINYL ESTRADIOL
NEXIUM, ESOMEPRAZOLE MAGNESIUM
NEXIUM 24HR, ESOMEPRAZOLE MAGNESIUM (OTC)
NEXIUM IV, ESOMEPRAZOLE SODIUM
NEXPLANON, ETNOGESTREL
NEXTERONE, AMIODARONE HYDROCHLORIDE
NIACIN, NIACIN
NIACOR, NIACIN
NIASPAN, NIACIN
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
NICODERM CQ, NICOTINE (OTC)
NICORETTE, NICOTINE POLACRILEX (OTC)
NICORETTE (MINT), NICOTINE POLACRILEX (OTC)
NICOTINE, NICOTINE (OTC)
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
NICOTROL, NICOTINE
NIFEDIPINE, NIFEDIPINE
NIKKI, DROSPIRENONE
NILANDRON, NILUTAMIDE
NILUTAMIDE, NILUTAMIDE
NIMBEX, CISATRACURIUM BESYLATE
NIMBEX PRESERVATIVE FREE, CISATRACURIUM BESYLATE
NIMODIPINE, NIMODIPINE
NINLARO, IXAZOMIB CITRATE
NIPENT, PENTOSTATIN
NIPRIDE RTU IN SODIUM CHLORIDE 0.9%, SODIUM NITROPRUSSIDE
NISOLDIPINE, NISOLDIPINE
NITHIODOTE, SODIUM NITRITE
NITRO-DUR, NITROGLYCERIN
NITROFURANTOIN, NITROFURANTOIN
NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
NITROGLYCERIN, NITROGLYCERIN
NITROGLYCERIN IN DEXTROSE 5%, NITROGLYCERIN
NITROLINGUAL PUMPSPRAY, NITROGLYCERIN
NITROMIST, NITROGLYCERIN
NITROPRESS, SODIUM NITROPRUSSIDE
NITROSTAT, NITROGLYCERIN
NITYR, NITISINONE
NIX, PERMETHRIN (OTC)
NIZATIDINE, NIZATIDINE
NIZORAL, KETOCONAZOLE
NIZORAL A-D, KETOCONAZOLE (OTC)
NOCDURNA, DESMOPRESSIN ACETATE
NOCTIVA, DESMOPRESSIN ACETATE
NOR-QD, NORETHINDRONE
NORCO, ACETAMINOPHEN
NORDITROPIN FLEXPOR, SOMATROPIN RECOMBINANT
NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
NORETHINDRONE, NORETHINDRONE
NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
NORETHINDRONE AND ETHINYL ESTRADIOL (10/11), ETHINYL ESTRADIOL
NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
NORINYL 1+35 21-DAY, ETHINYL ESTRADIOL
NORINYL 1+35 28-DAY, ETHINYL ESTRADIOL
NORINYL 1+50 28-DAY, MESTRANOL
NORITATE, METRONIDAZOLE
NORMOCARB HF 25, MAGNESIUM CHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** N **

NORMOCARB HF 35, MAGNESIUM CHLORIDE
 NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 NORMOSOL-R IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 NORPACE, DISOPYRAMIDE PHOSPHATE
 NORPACE CR, DISOPYRAMIDE PHOSPHATE
 NORPRAMIN, DESIPRAMINE HYDROCHLORIDE
 NORTHERA, DROXIDOPA
 NORTREL 0.5/35-28, ETHINYL ESTRADIOL
 NORTREL 1/35-21, ETHINYL ESTRADIOL
 NORTREL 1/35-28, ETHINYL ESTRADIOL
 NORTREL 7/7/7, ETHINYL ESTRADIOL
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NORVASC, AMLODIPINE BESYLATE
 NORVIR, RITONAVIR
 NOVOLIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
 NOVOLIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)
 NOVOLIN R, INSULIN RECOMBINANT HUMAN (OTC)
 NOVOLOG, INSULIN ASPART RECOMBINANT
 NOVOLOG FLEXPEN, INSULIN ASPART RECOMBINANT
 NOVOLOG MIX 70/30, INSULIN ASPART PROTAMINE RECOMBINANT
 NOVOLOG MIX 70/30 FLEXPEN, INSULIN ASPART PROTAMINE RECOMBINANT
 NOVOLOG PENFILL, INSULIN ASPART RECOMBINANT
 NOXAFIL, POSACONAZOLE
 NOXIVENT, NITRIC OXIDE
 NUCYNTA, TAPENTADOL HYDROCHLORIDE
 NUCYNTA ER, TAPENTADOL HYDROCHLORIDE
 NUEDEXTA, DEXTROMETHORPHAN HYDROBROMIDE
 NULYTELY, POLYETHYLENE GLYCOL 3350
 NULYTELY-FLAVORED, POLYETHYLENE GLYCOL 3350
 NUPLAZID, PIMAVANSERIN TARTRATE
 NUTRESTORE, L-GLUTAMINE
 NUTRILIPID 10%, SOYBEAN OIL
 NUTRILIPID 20%, SOYBEAN OIL
 NUTROPIN AQ NUSPIN, SOMATROPIN RECOMBINANT
 NUVARING, ETHINYL ESTRADIOL
 NUVESSA, METRONIDAZOLE
 NUVIGIL, ARMODAFINIL
 NUZYRA, OMADACYCLINE TOSYLATE
 NYLIA 1/35, ETHINYL ESTRADIOL
 NYLIA 7/7/7, ETHINYL ESTRADIOL
 NYMALIZE, NIMODIPINE
 NYSTATIN, NYSTATIN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTOP, NYSTATIN

** O **

OBREDON, GUAIFENESIN
 OCALIVA, OBETICHOLIC ACID
 OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCUFEN, FLURBIPROFEN SODIUM
 OCUFLOX, OFLOXACIN
 ODEFSEY, EMTRICITABINE
 ODOMZO, SONIDEGIB PHOSPHATE
 OFEV, NINTEDANIB ESYLATE
 OFIRMEV, ACETAMINOPHEN
 OFLOXACIN, OFLOXACIN
 OGEN 5, ESTROPIPATE
 OGESTREL 0.5/50-28, ETHINYL ESTRADIOL
 OLANZAPINE, OLANZAPINE
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

APPENDIX A - PRODUCT NAME INDEX

** O **

OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
OLUMIANT, BARICITINIB
OLUX, CLOBETASOL PROPIONATE
OLUX E, CLOBETASOL PROPIONATE
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
OMEGAVEN, FISH OIL TRIGLYCERIDES
OMEPRAZOLE, OMEPRAZOLE (OTC)
OMEPRAZOLE, OMEPRAZOLE
OMEPRAZOLE AND CLARITHROMYCIN AND AMOXICILLIN, AMOXICILLIN
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
OMIDRIA, KETOROLAC TROMETHAMINE
OMNARIS, CICLESONIDE
OMNIPAQUE 12, IOHEXOL
OMNIPAQUE 140, IOHEXOL
OMNIPAQUE 180, IOHEXOL
OMNIPAQUE 240, IOHEXOL
OMNIPAQUE 300, IOHEXOL
OMNIPAQUE 350, IOHEXOL
OMNIPAQUE 9, IOHEXOL
OMNIPRED, PREDNISOLONE ACETATE
OMNISCAN, GADODIAMIDE
OMNITROPE, SOMATROPIN RECOMBINANT
ONDANSETRON, ONDANSETRON
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
ONEXTON, BENZOYL PEROXIDE
ONFI, CLOBAZAM
ONGLYZA, SAXAGLIPTIN HYDROCHLORIDE
ONIVYDE, IRINOTECAN HYDROCHLORIDE
ONMEL, ITRACONAZOLE
ONPATTRO, PATISIRAN SODIUM
ONZETRA XSAIL, SUMATRIPTAN SUCCINATE
OPANA, OXYMORPHONE HYDROCHLORIDE
OPCICON ONE-STEP, LEVONORGESTREL (OTC)
OPCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
OPSUMIT, MACITENTAN
OPTIPRANOLOL, METIPRANOLOL HYDROCHLORIDE
OPTIRAY 240, IOVERSOL
OPTIRAY 300, IOVERSOL
OPTIRAY 320, IOVERSOL
OPTIRAY 350, IOVERSOL
OPTISON, ALBUMIN HUMAN
ORABLOC, ARTICAININE HYDROCHLORIDE
ORACEA, DOXYCYCLINE
ORALTAG, IOHEXOL
ORAP, PIMOZIDE
ORAPRED ODT, PREDNISOLONE SODIUM PHOSPHATE
ORAQIX, LIDOCAINE
ORAVERSER, PHENTOLAMINE MESYLATE
ORAVIG, MICONAZOLE
ORBACTIV, ORITAVANCIN DIPHOSPHATE
ORENITRAM, TREPROSTINIL DIOLAMINE
ORFADIN, NITISINONE
ORILISSA, ELAGOLIX SODIUM
ORKAMBI, IVACAFTOR
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE, ASPIRIN
ORSYTHIA, ETHINYL ESTRADIOL
ORTHO CYCLEN-28, ETHINYL ESTRADIOL
ORTHO TRI-CYCLEN, ETHINYL ESTRADIOL
ORTHO TRI-CYCLEN LO, ETHINYL ESTRADIOL
ORTHO-NOVUM 1/35-28, ETHINYL ESTRADIOL

APPENDIX A - PRODUCT NAME INDEX

** O **

ORTHO-NOVUM 7/7/7-28, ETHINYL ESTRADIOL
 ORVATEN, MIDODRINE HYDROCHLORIDE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OSENI, ALOGLIPTIN BENZOATE
 OSMITROL 10% IN WATER, MANNITOL
 OSMITROL 10% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 15% IN WATER, MANNITOL
 OSMITROL 15% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 20% IN WATER, MANNITOL
 OSMITROL 20% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 5% IN WATER, MANNITOL
 OSMITROL 5% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMOLEX ER, AMANTADINE HYDROCHLORIDE
 OSMOPREP, SODIUM PHOSPHATE, DIBASIC, ANHYDROUS
 OSPHENA, OSPEMIFENE
 OTEZLA, APREMILAST
 OTICAIR, HYDROCORTISONE
 OTIPRIO, CIPROFLOXACIN
 OTOVEL, CIPROFLOXACIN HYDROCHLORIDE
 OTREXUP, METHOTREXATE
 OVIDE, MALATHION
 OVIDREL, CHORIOGONADOTROPIN ALFA
 OXACILLIN SODIUM, OXACILLIN SODIUM
 OXALIPLATIN, OXALIPLATIN
 OXANDROLONE, OXANDROLONE
 OXAPROZIN, OXAPROZIN
 OXAYDO, OXYCODONE HYDROCHLORIDE
 OXAZEPAM, OXAZEPAM
 OXCARBAZEPINE, OXCARBAZEPINE
 OXICONAZOLE NITRATE, OXICONAZOLE NITRATE
 OXISTAT, OXICONAZOLE NITRATE
 OXSORALEN-ULTRA, METHOXSALLEN
 OXTELLAR XR, OXCARBAZEPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCET, ACETAMINOPHEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE AND ASPIRIN, ASPIRIN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYCODONE HYDROCHLORIDE AND IBUPROFEN, IBUPROFEN
 OXYCONTIN, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 OXYTOCIN, OXYTOCIN
 OXYTROL, OXYBUTYNIN
 OXYTROL FOR WOMEN, OXYBUTYNIN (OTC)
 OZEMPIC, SEMAGLUTIDE
 OZURDEX, DEXAMETHASONE

** P **

PACERONE, AMIODARONE HYDROCHLORIDE
 PACITAXEL, PACLITAXEL
 PACLITAXEL, PACLITAXEL
 PALIPERIDONE, PALIPERIDONE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMELOR, NORTRIPTYLINE HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PANCREAZE, PANCRELIPASE (AMYLASE)
 PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
 PANDEL, HYDROCORTISONE PROBUTATE
 PANRETIN, ALITRETINOIN
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARAGARD T 380A, COPPER
 PAREMYD, HYDROXYAMPHETAMINE HYDROBROMIDE
 PARICALCITOL, PARICALCITOL
 PARLODEL, BROMOCRIPTINE MESYLATE
 PARNATE, TRANYLCPROMINE SULFATE

APPENDIX A - PRODUCT NAME INDEX

** P **

PAROEX, CHLORHEXIDINE GLUCONATE
PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
PAROXETINE, PAROXETINE HYDROCHLORIDE
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
PAROXETINE MESYLATE, PAROXETINE MESYLATE
PARSABIV, ETELICALCETIDE
PASER, AMINOSALICYLIC ACID
PATADAY, OLOPATADINE HYDROCHLORIDE
PATANASE, OLOPATADINE HYDROCHLORIDE
PATANOL, OLOPATADINE HYDROCHLORIDE
PAXIL, PAROXETINE HYDROCHLORIDE
PAXIL CR, PAROXETINE HYDROCHLORIDE
PAZEO, OLOPATADINE HYDROCHLORIDE
PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE
PEDIATRIC ADVIL, IBUPROFEN (OTC)
PEG 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL
PEGANONE, ETHOTOIN
PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
PENICILLIN G POTASSIUM IN PLASTIC CONTAINER, PENICILLIN G POTASSIUM
PENICILLIN G PROCAINE, PENICILLIN G PROCAINE
PENICILLIN G SODIUM, PENICILLIN G SODIUM
PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
PENICILLIN-VK, PENICILLIN V POTASSIUM
PENLAC, CICLOPIROX
PENNSAID, DICLOFENAC SODIUM
PENTAM, PENTAMIDINE ISETHIONATE
PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE
PENTASA, MESALAMINE
PENTETATE CALCIUM TRISODIUM, PENTETATE CALCIUM TRISODIUM
PENTETATE ZINC TRISODIUM, PENTETATE ZINC TRISODIUM
PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
PENTOLAIR, CYCLOPENTOLATE HYDROCHLORIDE
PENTOSTATIN, PENTOSTATIN
PENTOXIFYLLINE, PENTOXIFYLLINE
PENTOXIL, PENTOXIFYLLINE
PEPCID, FAMOTIDINE
PEPCID AC, FAMOTIDINE (OTC)
PEPCID COMPLETE, CALCIUM CARBONATE (OTC)
PERCOCET, ACETAMINOPHEN
PERCODAN, ASPIRIN
PERFOROMIST, FORMOTEROL FUMARATE
PERIDEX, CHLORHEXIDINE GLUCONATE
PERIKABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
PERIOCHIP, CHLORHEXIDINE GLUCONATE
PERIOGARD, CHLORHEXIDINE GLUCONATE
PERMETHRIN, PERMETHRIN (OTC)
PERMETHRIN, PERMETHRIN
PERPHENAZINE, PERPHENAZINE
PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
PERSANTINE, DIPYRIDAMOLE
PERSERIS KIT, RISPERIDONE
PERTZYE, PANCRELIPASE (AMYLASE)
PEXEVA, PAROXETINE MESYLATE
PFIZERPEN, PENICILLIN G POTASSIUM
PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
PHENELZINE SULFATE, PHENELZINE SULFATE
PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE
PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
PHENYTEK, PHENYTOIN SODIUM
PHENYTOIN, PHENYTOIN

APPENDIX A - PRODUCT NAME INDEX

** P **

PHENYTOIN SODIUM, PHENYTOIN SODIUM
PHILITH, ETHINYL ESTRADIOL
PHOSLO GELCAPS, CALCIUM ACETATE
PHOSLYRA, CALCIUM ACETATE
PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE
PHOTOFRIN, PORFIMER SODIUM
PHOTREXA, RIBOFLAVIN 5'-PHOSPHATE SODIUM
PHOTREXA VISCOUS IN DEXTRAN 20%, RIBOFLAVIN 5'-PHOSPHATE SODIUM
PHOXILLUM B22K 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PHYSIOLYTE IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PHYSIOSOL IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PHYTONADIONE, PHYTONADIONE
PICATO, INGENOL MEBUTATE
PIFELTRO, DORAVIRINE
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
PIMECROLIMUS, PIMECROLIMUS
PIMOZIDE, PIMOZIDE
PIMTREA, DESOGESTREL
PINDOLOL, PINDOLOL
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
PIOGLITAZONE HYDROCHLORIDE AND GLIMEPIRIDE, GLIMEPIRIDE
PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
PIPERACILLIN, PIPERACILLIN SODIUM
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
PIRMELLA 1/35, ETHINYL ESTRADIOL
PIRMELLA 7/7/7, ETHINYL ESTRADIOL
PIROXICAM, PIROXICAM
PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
PITOCIN, OXYTOCIN
PLAN B ONE-STEP, LEVONORGESTREL (OTC)
PLAQUENIL, HYDROXYCHLOROQUINE SULFATE
PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PLASMA-LYTE A IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PLAVIX, CLOPIDOGREL BISULFATE
PLEGISOL IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PLENVU, ASCORBIC ACID
PLIAGLIS, LIDOCAINE
PODOFILOX, PODOFILOX
POLMON, DEXCHLORPHENIRAMINE MALEATE
POLOCAINE, MEPIVACAINE HYDROCHLORIDE
POLOCAINE-MPF, MEPIVACAINE HYDROCHLORIDE
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350
POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
POLYTRIM, POLYMYXIN B SULFATE
POMALYST, POMALIDOMIDE
PONSTEL, MEFENAMIC ACID
PORTIA-28, ETHINYL ESTRADIOL
POTASSIUM ACETATE, POTASSIUM ACETATE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,

APPENDIX A - PRODUCT NAME INDEX

** P **

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CITRATE, POTASSIUM CITRATE
 POTASSIUM IODIDE, POTASSIUM IODIDE (OTC)
 POVIDONE IODINE, POVIDONE-IODINE (OTC)
 PRADAXA, DABIGATRAN ETEXILATE MESYLATE
 PRALIDOXIME CHLORIDE, PRALIDOXIME CHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRAMOSONE, HYDROCORTISONE ACETATE
 PRANDIN, REPAGLINIDE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRAVACHOL, PRAVASTATIN SODIUM
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PRAZIQUANTEL, PRAZIQUANTEL
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PRE-OP, HEXACHLOROPHENE
 PRE-OP II, HEXACHLOROPHENE
 PRE-PEN, BENZYL PENICILLOYL POLYLYSINE
 PRECEDEX, DEXMEDETOMIDINE HYDROCHLORIDE
 PRECOSE, ACARBOSE
 PRED FORTE, PREDNISOLONE ACETATE
 PRED MILD, PREDNISOLONE ACETATE
 PRED-G, GENTAMICIN SULFATE
 PREDNICARBATE, PREDNICARBATE
 PREDNISOLONE, PREDNISOLONE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PREDNISON, PREDNISON
 PREDNISON INTENSOL, PREDNISON
 PREGNYL, GONADOTROPIN, CHORIONIC
 PRELONE, PREDNISOLONE
 PREMARIN, ESTROGENS, CONJUGATED
 PREMASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
 PREMASOL 6% IN PLASTIC CONTAINER, AMINO ACIDS
 PREMPhase 14/14, ESTROGENS, CONJUGATED
 PREMPRO, ESTROGENS, CONJUGATED
 PREPIDIL, DINOPROSTONE
 PREPOPIK, CITRIC ACID
 PRESTALIA, AMLODIPINE BESYLATE
 PREVACID, LANSOPRAZOLE
 PREVACID 24 HR, LANSOPRAZOLE (OTC)
 PREVALITE, CHOLESTYRAMINE
 PREVANTICS MAXI SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 PREVANTICS SWAB, CHLORHEXIDINE GLUCONATE (OTC)
 PREVANTICS SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 PREVIFEM, ETHINYL ESTRADIOL
 PREVMIS, LETERMOVIR
 PREZCOBIX, COBICISTAT

APPENDIX A - PRODUCT NAME INDEX

** P **

PREZISTA, DARUNAVIR ETHANOLATE
PRIALT, ZICONOTIDE ACETATE
PRIFTIN, RIFAPENTINE
PRILOCAINE HYDROCHLORIDE, PRILOCAINE HYDROCHLORIDE
PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, EPINEPHRINE BITARTRATE
PRILOSEC, OMEPRAZOLE MAGNESIUM
PRILOSEC OTC, OMEPRAZOLE MAGNESIUM (OTC)
PRIMAQUINE, PRIMAQUINE PHOSPHATE
PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE
PRIMATENE MIST, EPINEPHRINE (OTC)
PRIMAXIN, CILASTATIN SODIUM
PRIMIDONE, PRIMIDONE
PRIMSOL, TRIMETHOPRIM HYDROCHLORIDE
PRINIVIL, LISINAPRIL
PRISMASOL B22GK 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 0/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 2/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 2/3.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 4/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BK 0/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISTIQ, DESVENLAFAXINE SUCCINATE
PROAIR HFA, ALBUTEROL SULFATE
PROAIR RESPICLICK, ALBUTEROL SULFATE
PROBALAN, PROBENECID
PROBENECID, PROBENECID
PROBENECID AND COLCHICINE, COLCHICINE
PROBUPHINE, BUPRENORPHINE HYDROCHLORIDE
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
PROCALAMINE, AMINO ACIDS
PROCARDIA, NIFEDIPINE
PROCARDIA XL, NIFEDIPINE
PROCHLORPERAZINE, PROCHLORPERAZINE
PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
PROCOMP, PROCHLORPERAZINE MALEATE
PROCTOFOAM HC, HYDROCORTISONE ACETATE
PROCYSBI, CYSTEAMINE BITARTRATE
PROFERDEX, IRON DEXTRAN
PROGESTERONE, PROGESTERONE
PROGLYCEM, DIAZOXIDE
PROGRAF, TACROLIMUS
PROHANCE, GADOTERIDOL
PROHANCE MULTIPACK, GADOTERIDOL
PROLENSA, BROMFENAC SODIUM
PROMACTA, ELTROMBOPAG OLAMINE
PROMACTA KIT, ELTROMBOPAG OLAMINE
PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE PHOSPHATE
PROMETH VC W/ CODEINE, CODEINE PHOSPHATE
PROMETHAZINE DM, DEXTROMETHORPHAN HYDROBROMIDE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN
PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
PROMETHAZINE PLAIN, PROMETHAZINE HYDROCHLORIDE
PROMETHAZINE W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE
PROMETHAZINE WITH CODEINE, CODEINE PHOSPHATE
PROMETHEGAN, PROMETHAZINE HYDROCHLORIDE
PROMETRIUM, PROGESTERONE
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
PROPANTHELINE BROMIDE, PROPANTHELINE BROMIDE
PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
PROPECIA, FINASTERIDE

APPENDIX A - PRODUCT NAME INDEX

** P **

PROPOFOL, PROPOFOL
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 PROSCAR, FINASTERIDE
 PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
 PROSTIN E2, DINOPROSTONE
 PROSTIN VR PEDIATRIC, ALPROSTADIL
 PROTAMINE SULFATE, PROTAMINE SULFATE
 PROTONIX, PANTOPRAZOLE SODIUM
 PROTONIX IV, PANTOPRAZOLE SODIUM
 PROTOPAM CHLORIDE, PRALIDOXIME CHLORIDE
 PROTOPIC, TACROLIMUS
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 PROVAYBLUE, METHYLENE BLUE
 PROVENTIL-HFA, ALBUTEROL SULFATE
 PROVERA, MEDROXYPROGESTERONE ACETATE
 PROVIGIL, MODAFINIL
 PROVOCHOLINE, METHACHOLINE CHLORIDE
 PROZAC, FLUOXETINE HYDROCHLORIDE
 PROZAC WEEKLY, FLUOXETINE HYDROCHLORIDE
 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 PULMICORT FLEXHALER, BUDESONIDE
 PULMICORT RESPULES, BUDESONIDE
 PUR-WASH, PURIFIED WATER (OTC)
 PURINETHOL, MERCAPTOPURINE
 PURIXAN, MERCAPTOPURINE
 PYLERA, BISMUTH SUBCITRATE POTASSIUM
 PYRAZINAMIDE, PYRAZINAMIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
 PYTEST, UREA, C-14
 PYTEST KIT, UREA, C-14

** Q **

QBRELIS, LISINAPRIL
 QBREXZA, GLYCOPYRRONIUM TOSYLATE
 QMIIZ ODT, MELOXICAM
 QNASL, BECLOMETHASONE DIPROPIONATE
 QOLIANA, BRIMONIDINE TARTRATE
 QSYMIA, PHENTERMINE HYDROCHLORIDE
 QTERN, DAPAGLIFLOZIN
 QUADRAMET, SAMARIUM SM-153 LEXIDRONAM PENTASODIUM
 QUALAQUIN, QUININE SULFATE
 QUARTETTE, ETHINYL ESTRADIOL
 QUASENSE, ETHINYL ESTRADIOL
 QUDEXY XR, TOPIRAMATE
 QUELICIN, SUCCINYLCHOLINE CHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUILICHEW ER, METHYLPHENIDATE HYDROCHLORIDE
 QUILLIVANT XR, METHYLPHENIDATE HYDROCHLORIDE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 QUINARETIC, HYDROCHLOROTHIAZIDE
 QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
 QUINIDINE SULFATE, QUINIDINE SULFATE
 QUININE SULFATE, QUININE SULFATE
 QUTENZA, CAPSAICIN
 QVAR REDHALER, BECLOMETHASONE DIPROPIONATE

** R **

R-GENE 10, ARGININE HYDROCHLORIDE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RADICAVA, EDARAVONE
 RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE(II)

APPENDIX A - PRODUCT NAME INDEX

** R **

RADIOGENIX SYSTEM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
RAMELTEON, RAMELTEON
RAMIPRIL, RAMIPRIL
RANEXA, RANOLAZINE
RANITIDINE, RANITIDINE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RAPAFLO, SILODOSIN
RAPAMUNE, SIROLIMUS
RAPIVAB, PERAMIVIR
RASAGILINE MESYLATE, RASAGILINE MESYLATE
RASUVO, METHOTREXATE
RAVICTI, GLYCEROL PHENYL BUTYRATE
RAYALDEE, CALCIFEDIOL
RAYOS, PREDNISONE
RAZADYNE, GALANTAMINE HYDROBROMIDE
RAZADYNE ER, GALANTAMINE HYDROBROMIDE
READI-CAT 2, BARIUM SULFATE
READI-CAT 2 SMOOTHIES, BARIUM SULFATE
READYPREP CHG, CHLORHEXIDINE GLUCONATE (OTC)
REBETOL, RIBAVIRIN
RECLAST, ZOLEDRONIC ACID
RECTIV, NITROGLYCERIN
REGLAN, METOCLOPRAMIDE HYDROCHLORIDE
REGONOL, PYRIDOSTIGMINE BROMIDE
RELENZA, ZANAMIVIR
RELISTOR, METHYLNALTREXONE BROMIDE
RELPAX, ELETRIPTAN HYDROBROMIDE
REMERON, MIRTAZAPINE
REMERON SOLTAB, MIRTAZAPINE
REMIFENTANIL HYDROCHLORIDE, REMIFENTANIL HYDROCHLORIDE
REMODULIN, TREPROSTINIL
RENACIDIN, CITRIC ACID
RENAGEL, SEVELAMER HYDROCHLORIDE
RENOVA, TRETINOIN
RENEVELA, SEVELAMER CARBONATE
REPAGLINIDE, REPAGLINIDE
REPREXAIN, HYDROCODONE BITARTRATE
REQUIP, ROPINIROLE HYDROCHLORIDE
REQUIP XL, ROPINIROLE HYDROCHLORIDE
RESCRIPTOR, DELAVIRDINE MESYLATE
RESECTISOL IN PLASTIC CONTAINER, MANNITOL
RESTASIS, CYCLOSPORINE
RESTASIS MULTIDOSE, CYCLOSPORINE
RESTORIL, TEMAZEPAM
RETIN-A, TRETINOIN
RETIN-A MICRO, TRETINOIN
RETIN-A-MICRO, TRETINOIN
RETISERT, FLUOCINOLONE ACETONIDE
RETROVIR, ZIDOVUDINE
REVATIO, SILDENAFIL CITRATE
REVLIMID, LENALIDOMIDE
REVONTO, DANTROLENE SODIUM
REXULTI, BREXPIRAZOLE
REYATAZ, ATAZANAVIR SULFATE
REZIRA, HYDROCODONE BITARTRATE
RHINOCORT ALLERGY, BUDESONIDE (OTC)
RHOFADE, OXYMETAZOLINE HYDROCHLORIDE
RHOPRESSA, NETARSUDIL DIMESYLATE
RIBASPHERE, RIBAVIRIN
RIBAVIRIN, RIBAVIRIN
RIDAURA, AURANOFIN
RIFABUTIN, RIFABUTIN
RIFADIN, RIFAMPIN

APPENDIX A - PRODUCT NAME INDEX

** R **

RIFAMATE, ISONIAZID
 RIFAMPIN, RIFAMPIN
 RIFATER, ISONIAZID
 RILUTEK, RILUZOLE
 RILUZOLE, RILUZOLE
 RIMACTANE, RIFAMPIN
 RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
 RIMSO-50, DIMETHYL SULFOXIDE
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 RIOMET, METFORMIN HYDROCHLORIDE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RISPERDAL, RISPERIDONE
 RISPERDAL CONSTA, RISPERIDONE
 RISPERIDONE, RISPERIDONE
 RITALIN, METHYLPHENIDATE HYDROCHLORIDE
 RITALIN LA, METHYLPHENIDATE HYDROCHLORIDE
 RITONAVIR, RITONAVIR
 RIVASTIGMINE, RIVASTIGMINE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROBAXIN, METHOCARBAMOL
 ROBAXIN-750, METHOCARBAMOL
 ROCALTROL, CALCITRIOL
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROFLUMILAST, ROFLUMILAST
 ROGAINE (FOR MEN), MINOXIDIL (OTC)
 ROGAINE (FOR WOMEN), MINOXIDIL (OTC)
 ROGAINE EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 ROWASA, MESALAMINE
 ROWEEPRA, LEVETIRACETAM
 ROXICET, ACETAMINOPHEN
 ROXICODONE, OXYCODONE HYDROCHLORIDE
 ROZEREM, RAMELTEON
 RUBRACA, RUCAPARIB CAMSYLATE
 RUBY-FILL, RUBIDIUM CHLORIDE RB-82
 RUFINAMIDE, RUFINAMIDE
 RYANODEX, DANTROLENE SODIUM
 RYDAPT, MIDOSTAURIN
 RYTARY, CARBIDOPA
 RYTHMOL SR, PROPAFENONE HYDROCHLORIDE
 RYZODEG 70/30, INSULIN ASPART

** S **

SABRIL, VIGABATRIN
 SAFYRAL, DROSPIRENONE
 SAIZEN, SOMATROPIN RECOMBINANT
 SALAGEN, PILOCARPINE HYDROCHLORIDE
 SALONPAS, MENTHOL (OTC)
 SAMSCA, TOLVAPTAN
 SANCUSO, GRANISETRON
 SANDIMMUNE, CYCLOSPORINE
 SANDOSTATIN, OCTREOTIDE ACETATE
 SANDOSTATIN LAR, OCTREOTIDE ACETATE
 SAPHRIS, ASENAPINE MALEATE
 SARAFEM, FLUOXETINE HYDROCHLORIDE
 SAVAYSA, EDOXABAN TOSYLATE
 SAVELLA, MILNACIPRAN HYDROCHLORIDE
 SAXENDA, LIRAGLUTIDE RECOMBINANT
 SCANDONEST L, LEVONORDEFRIN
 SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
 SCANLUX-300, IOPAMIDOL
 SCANLUX-370, IOPAMIDOL

APPENDIX A - PRODUCT NAME INDEX

** S **

SCLEROSOL, TALC
SCOPOLAMINE, SCOPOLAMINE
SEASONALE, ETHINYL ESTRADIOL
SEASONIQUE, ETHINYL ESTRADIOL
SECONAL SODIUM, SECOBARBITAL SODIUM
SEEBRI, GLYCOPYRROLATE
SEGLUROMET, ERTUGLIFLOZIN
SEIZALAM, MIDAZOLAM HYDROCHLORIDE
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
SELENIUM SULFIDE, SELENIUM SULFIDE
SELFEMRA, FLUOXETINE HYDROCHLORIDE
SELZENTRY, MARAVIROC
SEMPREX-D, ACRIVASTINE
SENSIPAR, CINACALCET HYDROCHLORIDE
SENSORCAINE, BUPIVACAINE HYDROCHLORIDE
SEPTOCAINE, ARTICAINE HYDROCHLORIDE
SEPTRA, SULFAMETHOXAZOLE
SEPTRA DS, SULFAMETHOXAZOLE
SEREVENT, SALMETEROL XINAFOATE
SERNIVO, BETAMETHASONE DIPROPIONATE
SEROMYCIN, CYCLOSERINE
SEROQUEL, QUETIAPINE FUMARATE
SEROQUEL XR, QUETIAPINE FUMARATE
SEROSTIM, SOMATROPIN RECOMBINANT
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SETLAKIN, ETHINYL ESTRADIOL
SEVELAMER CARBONATE, SEVELAMER CARBONATE
SEVOFLURANE, SEVOFLURANE
SEYSARA, SARECYCLINE HYDROCHLORIDE
SFROWASA, MESALAMINE
SIGNIFOR, PASIREOTIDE DIASPARTATE
SIGNIFOR LAR KIT, PASIREOTIDE PAMOATE
SIKLOS, HYDROXYUREA
SILDENAFIL CITRATE, SILDENAFIL CITRATE
SILENOR, DOXEPIN HYDROCHLORIDE
SILODOSIN, SILODOSIN
SILVADENE, SILVER SULFADIAZINE
SIMBRINZA, BRIMONIDINE TARTRATE
SIMPESSE, ETHINYL ESTRADIOL
SIMVASTATIN, SIMVASTATIN
SINE-AID IB, IBUPROFEN (OTC)
SINEMET, CARBIDOPA
SINEMET CR, CARBIDOPA
SINGULAIR, MONTELUKAST SODIUM
SINUVA, MOMETASONE FUROATE
SIROLIMUS, SIROLIMUS
SIRTURO, BEDAQUILINE FUMARATE
SITAVIG, ACYCLOVIR
SIVEXTRO, TEDIZOLID PHOSPHATE
SKELAXIN, METAXALONE
SKLICE, IVERMECTIN
SKYLA, LEVONORGESTREL
SMOFLIPID 20%, FISH OIL
SODIUM ACETATE, SODIUM ACETATE
SODIUM BICARBONATE, SODIUM BICARBONATE
SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE, SODIUM FERRIC GLUCONATE COMPLEX
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18
SODIUM IODIDE I 123, SODIUM IODIDE I-123

APPENDIX A - PRODUCT NAME INDEX

** S **

SODIUM IODIDE I 131, SODIUM IODIDE I-131
SODIUM LACTATE IN PLASTIC CONTAINER, SODIUM LACTATE
SODIUM NITRITE, SODIUM NITRITE
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
SODIUM OXYBATE, SODIUM OXYBATE
SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE
SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE
SODIUM PHOSPHATES IN PLASTIC CONTAINER, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE
SODIUM THIOSULFATE, SODIUM THIOSULFATE
SOJOURN, SEVOFLURANE
SOLARAZE, DICLOFENAC SODIUM
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
SOLIQUA 100/33, INSULIN GLARGINE
SOLODYN, MINOCYCLINE HYDROCHLORIDE
SOLOSEC, SECNIDAZOLE
SOLTAMOX, TAMOXIFEN CITRATE
SOLU-CORTEF, HYDROCORTISONE SODIUM SUCCINATE
SOLU-MEDROL, METHYLPREDNISOLONE SODIUM SUCCINATE
SOLUPREP, CHLORHEXIDINE GLUCONATE (OTC)
SOMA, CARISOPRODOL
SOMATULINE DEPOT, LANREOTIDE ACETATE
SOMAVERT, PEGVISOMANT
SONATA, ZALEPLON
SOOLANTRA, IVERMECTIN
SORBITOL 3% IN PLASTIC CONTAINER, SORBITOL
SORBITOL 3.3% IN PLASTIC CONTAINER, SORBITOL
SORBITOL-MANNITOL IN PLASTIC CONTAINER, MANNITOL
SORIATANE, ACITRETIN
SORILUX, CALCIPOTRIENE
SORINE, SOTALOL HYDROCHLORIDE
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
SOTRADECOL, SODIUM TETRADECYL SULFATE
SOTYLIZE, SOTALOL HYDROCHLORIDE
SOVALDI, SOFOSBUVIR
SPECTAZOLE, ECONAZOLE NITRATE
SPINRAZA, NUSINERSEN SODIUM
SPIRIVA, TIOTROPIUM BROMIDE
SPIRIVA RESPIMAT, TIOTROPIUM BROMIDE
SPIRONOLACTONE, SPIRONOLACTONE
SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
SPORANOX, ITRACONAZOLE
SPRINTEC, ETHINYL ESTRADIOL
SPRITAM, LEVETIRACETAM
SPRIX, KETOROLAC TROMETHAMINE
SPRYCEL, DASATINIB
SPS, SODIUM POLYSTYRENE SULFONATE
SPY AGENT GREEN KIT, INDOCYANINE GREEN
SSD, SILVER SULFADIAZINE
STALEVO 100, CARBIDOPA
STALEVO 125, CARBIDOPA
STALEVO 150, CARBIDOPA
STALEVO 200, CARBIDOPA
STALEVO 50, CARBIDOPA
STALEVO 75, CARBIDOPA
STARLIX, NATEGLINIDE
STAVUDINE, STAVUDINE
STAXYN, VARDENAFIL HYDROCHLORIDE
STEGLATRO, ERTUGLIFLOZIN
STEGLUJAN, ERTUGLIFLOZIN
STENDRA, AVANAFIL
STERILE WATER, STERILE WATER FOR IRRIGATION
STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION

APPENDIX A - PRODUCT NAME INDEX

** S **

STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION
 STERITALC, TALC
 STIE-CORT, HYDROCORTISONE
 STIMATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 STIOLTO RESPIMAT, OLODATEROL HYDROCHLORIDE
 STIVARGA, REGORAFENIB
 STRATTERA, ATOMOXETINE HYDROCHLORIDE
 STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
 STRIANT, TESTOSTERONE
 STRIBILD, COBICISTAT
 STRIVERDI RESPIMAT, OLODATEROL HYDROCHLORIDE
 STROMEKTOL, IVERMECTIN
 STRONTIUM CHLORIDE SR-89, STRONTIUM CHLORIDE SR-89
 SUBLIMAZE PRESERVATIVE FREE, FENTANYL CITRATE
 SUBLOCADE, BUPRENORPHINE
 SUBOXONE, BUPRENORPHINE HYDROCHLORIDE
 SUBSYS, FENTANYL
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 SUCRAID, SACROSIDASE
 SUCRALFATE, SUCRALFATE
 SUDAFED 12 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 SUDAFED 24 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 SUFENTA PRESERVATIVE FREE, SUFENTANIL CITRATE
 SUFENTANIL CITRATE, SUFENTANIL CITRATE
 SULAR, NISOLDIPINE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 SULFADIAZINE, SULFADIAZINE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH, SULFAMETHOXAZOLE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH, SULFAMETHOXAZOLE
 SULFAMYLON, MAFENIDE ACETATE
 SULFASALAZINE, SULFASALAZINE
 SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE
 SULINDAC, SULINDAC
 SUMATRIPTAN, SUMATRIPTAN
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 SUPPRELIN LA, HISTRELIN ACETATE
 SUPRANE, DESFLURANE
 SUPRAX, CEFIXIME
 SUPREP BOWEL PREP KIT, MAGNESIUM SULFATE
 SURMONTIL, TRIMIPRAMINE MALEATE
 SURVANTA, BERACTANT
 SUSTIVA, EFAVIRENZ
 SUSTOL, GRANISETRON
 SUTENT, SUNITINIB MALATE
 SYEDA, DROSPIRENONE
 SYMBICORT, BUDESONIDE
 SYMBYAX, FLUOXETINE HYDROCHLORIDE
 SYMDEKO (COPACKAGED), IVACAFTOR
 SYMFI, EFAVIRENZ
 SYMFI LO, EFAVIRENZ
 SYMJEPI, EPINEPHRINE
 SYMLIN, PRAMLINTIDE ACETATE
 SYMPAZAN, CLOBAZAM
 SYMPROIC, NALDEMEDINE TOSYLATE
 SYMTUZA, COBICISTAT
 SYNALAR, FLUOCINOLONE ACETONIDE
 SYNAREL, NAFARELIN ACETATE
 SYNDROS, DRONABINOL
 SYNERA, LIDOCAINE
 SYNERCID, DALFOPRISTIN
 SYNJARDY, EMPAGLIFLOZIN
 SYNJARDY XR, EMPAGLIFLOZIN

APPENDIX A - PRODUCT NAME INDEX

** S **

SYNRIBO, OMACETAXINE MEPESUCCINATE
 SYNTHROID, LEVOTHYROXINE SODIUM **
 SYPRINE, TRIENTINE HYDROCHLORIDE

** T **

TAB-PROFEN, IBUPROFEN (OTC)
 TACLONEX, BETAMETHASONE DIPROPIONATE
 TACROLIMUS, TACROLIMUS
 TADALAFIL, TADALAFIL
 TAFINLAR, DABRAFENIB MESYLATE
 TAGAMET HB, CIMETIDINE (OTC)
 TAGITOL V, BARIUM SULFATE
 TAGRISSO, OSIMERTINIB MESYLATE
 TALC, TALC
 TALZENNA, TALAZOPARIB TOSYLATE
 TAMBOCOR, FLECAINIDE ACETATE
 TAMIFLU, OSELTAMIVIR PHOSPHATE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TAPAZOLE, METHIMAZOLE
 TARCEVA, ERLOTINIB HYDROCHLORIDE
 TARGRETIN, BEXAROTENE
 TARKA, TRANDOLAPRIL
 TASIGNA, NILOTINIB HYDROCHLORIDE
 TASMAR, TOLCAPONE
 TAVALISSE, FOSTAMATINIB DISODIUM
 TAXOL, PACLITAXEL
 TAXOTERE, DOCETAXEL
 TAYTULLA, ETHINYL ESTRADIOL
 TAZAROTENE, TAZAROTENE
 TAZICEF, CEFTAZIDIME
 TAZORAC, TAZAROTENE
 TAZTIA XT, DILTIAZEM HYDROCHLORIDE
 TECFIDERA, DIMETHYL FUMARATE
 TECHNELITE, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 TECHNESCAN, TECHNETIUM TC-99M OXIDRONATE KIT
 TECHNESCAN MAG3, TECHNETIUM TC-99M MERTIATIDE KIT
 TECHNESCAN PYP KIT, TECHNETIUM TC-99M PYROPHOSPHATE KIT
 TECHNETIUM TC 99M ALBUMIN AGGREGATED KIT, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT
 TECHNETIUM TC 99M GENERATOR, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
 TECHNETIUM TC-99M MEBROFENIN, TECHNETIUM TC-99M MEBROFENIN KIT
 TECHNIVIE, OMBITASVIR
 TEFLARO, CEFTAROLINE FOSAMIL
 TEGRETOL, CARBAMAZEPINE
 TEGRETOL-XR, CARBAMAZEPINE
 TEGSEDI, INOTERSEN SODIUM
 TEKTURNA, ALISKIREN HEMIFUMARATE
 TEKTURNA HCT, ALISKIREN HEMIFUMARATE
 TELMISARTAN, TELMISARTAN
 TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TEMAZEPAM, TEMAZEPAM
 TEMIXYS, LAMIVUDINE
 TEMODAR, TEMOZOLOMIDE
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TEMSIROLIMUS, TEMSIROLIMUS
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TENORETIC 100, ATENOLOL
 TENORETIC 50, ATENOLOL
 TENORMIN, ATENOLOL
 TENUATE, DIETHYLPROPION HYDROCHLORIDE
 TENUATE DOSPAN, DIETHYLPROPION HYDROCHLORIDE
 TEPADINA, THIOTEPA
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** T **

TERBINAFFINE HYDROCHLORIDE, TERBINAFFINE HYDROCHLORIDE (OTC)
 TERBINAFFINE HYDROCHLORIDE, TERBINAFFINE HYDROCHLORIDE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 TERCONAZOLE, TERCONAZOLE
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TERIL, CARBAMAZEPINE
 TESSALON, BENZONATATE
 TESTIM, TESTOSTERONE
 TESTOPEL, TESTOSTERONE
 TESTOSTERONE, TESTOSTERONE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
 TESTRED, METHYLTESTOSTERONE
 TETRABENAZINE, TETRABENAZINE
 TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 TEXACORT, HYDROCORTISONE
 THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
 THALOMID, THALIDOMIDE
 THEO-24, THEOPHYLLINE
 THEOCHRON, THEOPHYLLINE
 THEOPHYLLINE, THEOPHYLLINE
 THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THERMAZENE, SILVER SULFADIAZINE
 THEROXIDIL, MINOXIDIL (OTC)
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 THIOGUANINE, THIOGUANINE
 THIOLA, TIOPRONIN
 THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
 THIOTEPA, THIOTEPA
 THIOTHIXENE, THIOTHIXENE
 THYROGEN, THYROTROPIN ALFA
 THYROSAFE, POTASSIUM IODIDE (OTC)
 THYROSHIELD, POTASSIUM IODIDE (OTC)
 TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE
 TIAZAC, DILTIAZEM HYDROCHLORIDE
 TIBSOVO, IVOSIDENIB
 TICAGRELOR, TICAGRELOR
 TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
 TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE
 TIGECYCLINE, TIGECYCLINE
 TIGLUTIK KIT, RILUZOLE
 TIKOSYN, DOFETILIDE
 TIMOLOL, TIMOLOL
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIMOPTIC, TIMOLOL MALEATE
 TIMOPTIC IN OCUDOSE, TIMOLOL MALEATE
 TIMOPTIC-XE, TIMOLOL MALEATE
 TINDAMAX, TINIDAZOLE
 TINIDAZOLE, TINIDAZOLE
 TIOCONAZOLE, TIOCONAZOLE (OTC)
 TIROSINT, LEVOTHYROXINE SODIUM
 TIS-U-SOL, MAGNESIUM SULFATE
 TIS-U-SOL IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 TIVICAY, DOLUTEGRAVIR SODIUM
 TIVORBEX, INDOMETHACIN
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOBI, TOBRAMYCIN
 TOBI PODHALER, TOBRAMYCIN
 TOBRADEX, DEXAMETHASONE
 TOBRADEX ST, DEXAMETHASONE
 TOBRAMYCIN, TOBRAMYCIN

APPENDIX A - PRODUCT NAME INDEX

** T **

TOBRAMYCIN AND DEXAMETHASONE, DEXAMETHASONE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
TOBRAMYCIN SULFATE (PHARMACY BULK), TOBRAMYCIN SULFATE
TOBRAMYCIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, TOBRAMYCIN SULFATE
TOBREX, TOBRAMYCIN
TODAY, NONOXYNOL-9 (OTC)
TOFRANIL, IMIPRAMINE HYDROCHLORIDE
TOLAK, FLUOROURACIL
TOLAZAMIDE, TOLAZAMIDE
TOLBUTAMIDE, TOLBUTAMIDE
TOLCAPONE, TOLCAPONE
TOLMETIN SODIUM, TOLMETIN SODIUM
TOLSURA, ITRACONAZOLE
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
TOPAMAX, TOPIRAMATE
TOPICORT, DESOXIMETASONE
TOPIRAMATE, TOPIRAMATE
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
TOPROL-XL, METOPROLOL SUCCINATE
TOREMIFENE CITRATE, TOREMIFENE CITRATE
TORISEL, TEMSIROLIMUS
TORSEMIDE, TORSEMIDE
TOTECT, DEXRAZOXANE HYDROCHLORIDE
TOUJEO MAX SOLOSTAR, INSULIN GLARGINE RECOMBINANT
TOUJEO SOLOSTAR, INSULIN GLARGINE RECOMBINANT
TOVIAZ, FESOTERODINE FUMARATE
TPN ELECTROLYTES IN PLASTIC CONTAINER, CALCIUM CHLORIDE
TPOXX, TECOVIRIMAT
TRACLEER, BOSENTAN
TRADJENTA, LINAGLIPTIN
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
TRANDATE, LABETALOL HYDROCHLORIDE
TRANDOLAPRIL, TRANDOLAPRIL
TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE, TRANDOLAPRIL
TRANEXAMIC ACID, TRANEXAMIC ACID
TRANSDERM SCOP, SCOPOLAMINE
TRANXENE, CLORAZEPATE DIPOTASSIUM
TRANLYCYPROMINE SULFATE, TRANLYCYPROMINE SULFATE
TRAVASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 5.5% IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 8.5% IN PLASTIC CONTAINER, AMINO ACIDS
TRAVATAN Z, TRAVOPROST
TRAVOPROST, TRAVOPROST
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
TREANDA, BENDAMUSTINE HYDROCHLORIDE
TRECATOR, ETHIONAMIDE
TRELEGY ELLIPTA, FLUTICASONE FUROATE
TRELSTAR, TRIPTORELIN PAMOATE
TREPASTINIL, TREPASTINIL
TRESIBA, INSULIN DEGLUDEC
TRETINOIN, TRETINOIN
Trexall, METHOTREXATE SODIUM
Treximet, NAPROXEN SODIUM
Trezix, ACETAMINOPHEN
TRI LO SPRINTec, ETHINYL ESTRADIOL
TRI-ESTARYLLA, ETHINYL ESTRADIOL
TRI-LEGEST 21, ETHINYL ESTRADIOL
TRI-LEGEST FE, ETHINYL ESTRADIOL
TRI-LINYAH, ETHINYL ESTRADIOL
TRI-LO-ESTARYLLA, ETHINYL ESTRADIOL
TRI-LO-MILI, ETHINYL ESTRADIOL
TRI-LUMA, FLUOCINOLONE ACETONIDE
TRI-MILI, ETHINYL ESTRADIOL
TRI-NORINYL 28-DAY, ETHINYL ESTRADIOL

APPENDIX A - PRODUCT NAME INDEX

** T **

TRI-PREVIFEM, ETHINYL ESTRADIOL
 TRI-SPRINTEC, ETHINYL ESTRADIOL
 TRIACIN-C, CODEINE PHOSPHATE
 TRIAMCINOLONE ACETATE, TRIAMCINOLONE ACETONIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE (OTC)
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIAMCINOLONE ACETONIDE IN ABSORBASE, TRIAMCINOLONE ACETONIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIAZOLAM, TRIAZOLAM
 TRIBENZOR, AMLODIPINE BESYLATE
 TRICOR, FENOFIBRATE
 TRIDERM, TRIAMCINOLONE ACETONIDE
 TRIDIONE, TRIMETHADIONE
 TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE
 TRISENCE, TRIAMCINOLONE ACETONIDE
 TRIFERIC, FERRIC PYROPHOSPHATE CITRATE
 TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 TRIFLURIDINE, TRIFLURIDINE
 TRIGLIDE, FENOFIBRATE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 TRILEPTAL, OXCARBAZEPINE
 TRILIPIX, CHOLINE FENOFIBRATE
 TRILYTE, POLYETHYLENE GLYCOL 3350
 TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
 TRIMETHOBENZAMIDE HYDROCHLORIDE PRESERVATIVE FREE, TRIMETHOBENZAMIDE HYDROCHLORIDE
 TRIMETHOPRIM, TRIMETHOPRIM
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE
 TRINTELLIX, VORTIOXETINE HYDROBROMIDE
 TRIOSTAT, LIOTHYRONINE SODIUM
 TRIPTODUR KIT, TRIPTORELIN PAMOATE
 TRISENOX, ARSENIC TRIOXIDE
 TRIUMEQ, ABACAVIR SULFATE
 TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)
 TRIVORA-28, ETHINYL ESTRADIOL
 TRIZIVIR, ABACAVIR SULFATE
 TROKENDI XR, TOPIRAMATE
 TROPHAMINE, AMINO ACIDS
 TROPHAMINE 10%, AMINO ACIDS
 TROPICACYL, TROPICAMIDE
 TROPICAMIDE, TROPICAMIDE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
 TRULANCE, PLECANATIDE
 TRUSOPT, DORZOLAMIDE HYDROCHLORIDE
 TRUVADA, EMTRICITABINE
 TUDORZA PRESSAIR, ACLIDINIUM BROMIDE
 TUSSICAPS, CHLORPHENIRAMINE POLISTIREX
 TUSSIGON, HOMATROPINE METHYLBROMIDE
 TUXARIN ER, CHLORPHENIRAMINE MALEATE
 TUZISTRA XR, CHLORPHENIRAMINE POLISTIREX
 TWYNSTA, AMLODIPINE BESYLATE
 TYBOST, COBICISTAT
 TYDEMY, DROSPIRENONE
 TYGACIL, TIGECYCLINE
 TYKERB, LAPATINIB DITOSYLATE
 TYLENOL, ACETAMINOPHEN (OTC)
 TYLENOL W/ CODEINE NO. 3, ACETAMINOPHEN
 TYLENOL W/ CODEINE NO. 4, ACETAMINOPHEN
 TYMLOS, ABALOPARATIDE
 TYVASO, TREPROSTINIL
 TYZINE, TETRAHYDROZOLINE HYDROCHLORIDE

** U **

U-CORT, HYDROCORTISONE ACETATE
 UCERIS, BUDESONIDE

APPENDIX A - PRODUCT NAME INDEX

** U **

ULESFIA, BENZYL ALCOHOL
 ULORIC, FEBUXOSTAT
 ULTACAN, ARTICAINA HYDROCHLORIDE
 ULTACAN FORTE, ARTICAINA HYDROCHLORIDE
 ULTANE, SEVOFLURANE
 ULTIVA, REMIFENTANIL HYDROCHLORIDE
 ULTRA-TECHNEKOW FM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 ULTRACET, ACETAMINOPHEN
 ULTRAM, TRAMADOL HYDROCHLORIDE
 ULTRATAG, TECHNETIUM TC-99M RED BLOOD CELL KIT
 ULTRAVATE, HALOBETASOL PROPIONATE
 ULTRAVIST (PHARMACY BULK), IOPROMIDE
 ULTRAVIST 240, IOPROMIDE
 ULTRAVIST 300, IOPROMIDE
 ULTRAVIST 370, IOPROMIDE
 UNASYN, AMPICILLIN SODIUM
 UNISOM, DOXYLAMINE SUCCINATE (OTC)
 UNITHROID, LEVOTHYROXINE SODIUM **
 UPTRAVI, SELEXIPAG
 URECHOLINE, BETHANECHOL CHLORIDE
 UREX, METHENAMINE HIPPURATE
 UROCIT-K, POTASSIUM CITRATE
 UROXATRAL, ALFUZOSIN HYDROCHLORIDE
 URSO 250, URSODIOL
 URSO FORTE, URSODIOL
 URSODIOL, URSODIOL
 UTIBRON, GLYCOPYRROLATE
 UVADEX, METHOXSALEN

** V **

VABOMERE, MEROPENEM
 VAGIFEM, ESTRADIOL
 VAGISTAT-1, TIOCONAZOLE (OTC)
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALCHLOR, MECHLORETHAMINE HYDROCHLORIDE
 VALCYTE, VALGANCICLOVIR HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VALIUM, DIAZEPAM
 VALNAC, BETAMETHASONE VALERATE
 VALPROATE SODIUM, VALPROATE SODIUM
 VALPROIC ACID, VALPROIC ACID
 VALSARTAN, VALSARTAN
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSTAR PRESERVATIVE FREE, VALRUBICIN
 VALTREX, VALACYCLOVIR HYDROCHLORIDE
 VANCOCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VANCOCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
 VANDAZOLE, METRONIDAZOLE
 VANIQA, EFLORNITHINE HYDROCHLORIDE
 VANOS, FLUOCINONIDE
 VANTAS, HISTRELIN ACETATE
 VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER, CONIVAPTAN HYDROCHLORIDE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VARIBAR HONEY, BARIUM SULFATE
 VARIBAR NECTAR, BARIUM SULFATE
 VARIBAR PUDDING, BARIUM SULFATE
 VARIBAR THIN HONEY, BARIUM SULFATE
 VARITHENA, POLIDOCANOL
 VARUBI, ROLAPITANT HYDROCHLORIDE
 VASCEPA, ICOSAPENT ETHYL
 VASERETIC, ENALAPRIL MALEATE
 VASOSTRICT, VASOPRESSIN
 VASOTEC, ENALAPRIL MALEATE

APPENDIX A - PRODUCT NAME INDEX

** v **

VAZALORE, ASPIRIN (OTC)
 VAZCULEP, PHENYLEPHRINE HYDROCHLORIDE
 VECTICAL, CALCITRIOL
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VELCADE, BORTEZOMIB
 VELETRI, EPOPROSTENOL SODIUM
 VELIVET, DESOGESTREL
 VELPHORO, SUCROFERRIC OXYHYDROXIDE
 VELTASSA, PATIROMER SORBITE X CALCIUM
 VELTIN, CLINDAMYCIN PHOSPHATE
 VEMLIDY, TENOFOVIR ALAFENAMIDE FUMARATE
 VENCLEXTA, VENETOCLAX
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VENOFER, IRON SUCROSE
 VENTAVIS, ILOPROST
 VENTOLIN HFA, ALBUTEROL SULFATE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 VERDESO, DESONIDE
 VEREGEN, SINECATECHINS
 VERELAN, VERAPAMIL HYDROCHLORIDE
 VERELAN PM, VERAPAMIL HYDROCHLORIDE
 VERSACLOZ, CLOZAPINE
 VERZENIO, ABEMACICLIB
 VESICARE, SOLIFENACIN SUCCINATE
 VFEND, VORICONAZOLE
 VIAGRA, SILDENAFIL CITRATE
 VIBATIV, TELAVANCIN HYDROCHLORIDE
 VIBERZI, ELUXADOLINE
 VIBISONE, CYANOCOBALAMIN
 VIBRAMYCIN, DOXYCYCLINE
 VIBRAMYCIN, DOXYCYCLINE CALCIUM
 VIBRAMYCIN, DOXYCYCLINE HYCLATE
 VICTOZA, LIRAGLUTIDE RECOMBINANT
 VIDAZA, AZACITIDINE
 VIDEX, DIDANOSINE
 VIDEX EC, DIDANOSINE
 VIEKIRA PAK (COPACKAGED), DASABUVIR SODIUM
 VIEKIRA XR, DASABUVIR SODIUM
 VIENVA, ETHINYL ESTRADIOL
 VIGABATRIN, VIGABATRIN
 VIGADRONE, VIGABATRIN
 VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE
 VIIBRYD, VILAZODONE HYDROCHLORIDE
 VIMOVO, ESOMEPRAZOLE MAGNESIUM
 VIMPAT, LACOSAMIDE
 VINBLASTINE SULFATE, VINBLASTINE SULFATE
 VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE
 VINORELBINE TARTRATE, VINORELBINE TARTRATE
 VIOKACE, PANCRELIPASE (AMYLASE)
 VIORELE, DESOGESTREL
 VIRACEPT, NELFINAVIR MESYLATE
 VIRAMUNE, NEVIRAPINE
 VIRAMUNE XR, NEVIRAPINE
 VIRAZOLE, RIBAVIRIN
 VIREAD, TENOFOVIR DISOPROXIL FUMARATE
 VIROPTIC, TRIFLURIDINE
 VISINE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 VISINE L.R., OXYMETAZOLINE HYDROCHLORIDE (OTC)
 VISIONBLUE, TRYPAN BLUE
 VISIPAQUE 270, IODIXANOL
 VISIPAQUE 320, IODIXANOL
 VISTARIL, HYDROXYZINE PAMOATE
 VISTOGARD, URIDINE TRIACETATE
 VISUDYNE, VERTEPORFIN
 VITAMIN D, ERGOCALCIFEROL

APPENDIX A - PRODUCT NAME INDEX

** V **

VITAMIN K1, PHYTONADIONE
 VITRAKVI, LAROTRECTINIB
 VITRASE, HYALURONIDASE
 VITUZ, CHLORPHENIRAMINE MALEATE
 VIVACTIL, PROTRIPTYLINE HYDROCHLORIDE
 VIVELLE-DOT, ESTRADIOL
 VIVITROL, NALTREXONE
 VIVLODEX, MELOXICAM
 VIZAMYL, FLUTEMETAMOL F-18
 VIZIMPRO, DACOMITINIB
 VOGELXO, TESTOSTERONE
 VOLNEA, DESOGESTREL
 VOLTAREN, DICLOFENAC SODIUM
 VORICONAZOLE, VORICONAZOLE
 VOSEVI, SOFOSBUVIR
 VOSOL, ACETIC ACID, GLACIAL
 VOSOL HC, ACETIC ACID, GLACIAL
 VOSPIRE ER, ALBUTEROL SULFATE
 VOTRIENT, PAZOPANIB HYDROCHLORIDE
 VPRIV, VELAGLUCERASE ALFA
 VRAYLAR, CARIPRAZINE HYDROCHLORIDE
 VUSION, MICONAZOLE NITRATE
 VYFEMLA, ETHINYL ESTRADIOL
 VYTORIN, EZETIMIBE
 VYVANSE, LISDEXAMFETAMINE DIMESYLATE
 VYXEOS, CYTARABINE
 VYZULTA, LATANOPROSTENE BUNOD

** W **

WARFARIN SODIUM, WARFARIN SODIUM
 WELCHOL, COLESEVELAM HYDROCHLORIDE
 WELLBUTRIN SR, BUPROPION HYDROCHLORIDE
 WELLBUTRIN XL, BUPROPION HYDROCHLORIDE
 WERA, ETHINYL ESTRADIOL
 WOMEN'S ROGAINE, MINOXIDIL (OTC)

** X **

XADAGO, SAFINAMIDE MESYLATE
 XALATAN, LATANOPROST
 XALKORI, CRIZOTINIB
 XANAX, ALPRAZOLAM
 XANAX XR, ALPRAZOLAM
 XARELTO, RIVAROXABAN
 XATMEP, METHOTREXATE SODIUM
 XELJANZ, TOFACITINIB CITRATE
 XELJANZ XR, TOFACITINIB CITRATE
 XELODA, CAPECITABINE
 XELPROS, LATANOPROST
 XENAZINE, TETRABENAZINE
 XENICAL, ORLISTAT
 XENON XE 133, XENON XE-133
 XEPI, OZENOXACIN
 XERAVA, ERAVACYCLINE DIHYDROCHLORIDE
 XERESE, ACYCLOVIR
 XERMELLO, TELOTTRISTAT ETIPRATE
 XHANCE, FLUTICASONE PROPIONATE
 XIFAXAN, RIFAXIMIN
 XIGDUO XR, DAPAGLIFLOZIN
 XIIDRA, LIFITEGRAST
 XIMINO, MINOCYCLINE HYDROCHLORIDE
 XOFIGO, RADIUM RA-223 DICHLORIDE
 XOFLUZA, BALOXAVIR MARBOXIL
 XOLEGEL, KETOCONAZOLE
 XOPENEX, LEVALBUTEROL HYDROCHLORIDE
 XOPENEX HFA, LEVALBUTEROL TARTRATE

APPENDIX A - PRODUCT NAME INDEX

** X **

XOSPATA, GILTERITINIB FUMARATE
 XTAMPZA ER, OXYCODONE
 XTANDI, ENZALUTAMIDE
 XTORO, FINAFLOXACIN
 XTRELUS, GUAIFENESIN
 XULANE, ETHINYL ESTRADIOL
 XULTOPHY 100/3.6, INSULIN DEGLUDEC
 XURIDEN, URIDINE TRIACETATE
 XYLOCAINE, LIDOCAINE HYDROCHLORIDE
 XYLOCAINE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
 XYOSTED (AUTOINJECTOR), TESTOSTERONE ENANTHATE
 XYREM, SODIUM OXYBATE
 XYZAL, LEVOCETIRIZINE DIHYDROCHLORIDE
 XYZAL ALLERGY 24HR, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)

** Y **

YAELA, DROSPIRENONE
 YASMIN, DROSPIRENONE
 YAZ, DROSPIRENONE
 YONDELIS, TRABECTEDIN
 YONSA, ABIRATERONE ACETATE
 YOSPRALA, ASPIRIN
 YUPELRI, REVEFENACIN
 YUTIQ, FLUOCINOLONE ACETONIDE

** Z **

ZAFIRLUKAST, ZAFIRLUKAST
 ZALEPLON, ZALEPLON
 ZANAFLEX, TIZANIDINE HYDROCHLORIDE
 ZANOSAR, STREPTOZOCIN
 ZANTAC, RANITIDINE HYDROCHLORIDE
 ZANTAC 150, RANITIDINE HYDROCHLORIDE (OTC)
 ZANTAC 75, RANITIDINE HYDROCHLORIDE (OTC)
 ZARONTIN, ETHOSUXIMIDE
 ZAROXOLYN, METOLAZONE
 ZAVESCA, MIGLUSTAT
 ZEGERID, OMEPRAZOLE
 ZEGERID OTC, OMEPRAZOLE (OTC)
 ZEJULA, NIRAPARIB TOSYLATE
 ZELAPAR, SELEGILINE HYDROCHLORIDE
 ZELBORAF, VEMURAFENIB
 ZEMBRACE SYMTOUCH, SUMATRIPTAN SUCCINATE
 ZEMDRI, PLAZOMICIN SULFATE
 ZEMPLAR, PARICALCITOL
 ZENATANE, ISOTRETINOIN
 ZENPEP, PANCRELIPASE (AMYLASE)
 ZEPATIER, ELBASVIR
 ZERBAXA, CEFTOLOZANE SULFATE
 ZERIT, STAVUDINE
 ZERVIAE, CETIRIZINE HYDROCHLORIDE
 ZESTORETIC, HYDROCHLOROTHIAZIDE
 ZESTRIL, LISINOPRIL
 ZETIA, EZETIMIBE
 ZETONNA, CICLESONIDE
 ZIAC, BISOPROLOL FUMARATE
 ZIAGEN, ABACAVIR SULFATE
 ZIANA, CLINDAMYCIN PHOSPHATE
 ZIDOVUDINE, ZIDOVUDINE
 ZILEUTON, ZILEUTON
 ZILRETTA, TRIAMCINOLONE ACETONIDE
 ZINACEF, CEFUROXIME SODIUM
 ZINC CHLORIDE IN PLASTIC CONTAINER, ZINC CHLORIDE
 ZINECARD, DEXRAZOXANE HYDROCHLORIDE
 ZINGO, LIDOCAINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

** Z **

ZIOPTAN, TAFLUPROST
ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
ZIPSOR, DICLOFENAC POTASSIUM
ZIRGAN, GANCICLOVIR
ZITHROMAX, AZITHROMYCIN
ZOCOR, SIMVASTATIN
ZOFRAN, ONDANSETRON HYDROCHLORIDE
ZOFRAN ODT, ONDANSETRON
ZOHYDRO ER, HYDROCODONE BITARTRATE
ZOLADEX, GOSERELIN ACETATE
ZOLEDRONIC, ZOLEDRONIC ACID
ZOLEDRONIC ACID, ZOLEDRONIC ACID
ZOLINZA, VORINOSTAT
ZOLMITRIPTAN, ZOLMITRIPTAN
ZOLOFT, SERTRALINE HYDROCHLORIDE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
ZOLPIMIST, ZOLPIDEM TARTRATE
ZOMACTON, SOMATROPIN
ZOMETA, ZOLEDRONIC ACID
ZOMIG, ZOLMITRIPTAN
ZOMIG-ZMT, ZOLMITRIPTAN
ZONALON, DOXEPIN HYDROCHLORIDE
ZONEGRAN, ZONISAMIDE
ZONISAMIDE, ZONISAMIDE
ZONTIVITY, VORAPAXAR SULFATE
ZORBTIVE, SOMATROPIN RECOMBINANT
ZORTRESS, EVEROLIMUS
ZORVOLEX, DICLOFENAC
ZOSYN, PIPERACILLIN SODIUM
ZOSYN IN PLASTIC CONTAINER, PIPERACILLIN SODIUM
ZOVIA 1/35E-28, ETHINYL ESTRADIOL
ZOVIA 1/50E-28, ETHINYL ESTRADIOL
ZOVIRAX, ACYCLOVIR
ZTLIDO, LIDOCAINE
ZUBSOLV, BUPRENORPHINE HYDROCHLORIDE
ZUMANDIMINE, DROSPIRENONE
ZUPLENZ, ONDANSETRON
ZURAMPIC, LESINURAD
ZUTRIPRO, CHLORPHENIRAMINE MALEATE
ZYBAN, BUPROPION HYDROCHLORIDE
ZYCLARA, IMIQUIMOD
ZYDELIG, IDELALISIB
ZYFLO, ZILEUTON
ZYFLO CR, ZILEUTON
ZYKADIA, CERITINIB
ZYLET, LOTEPREDNOL ETABONATE
ZYLOPRIM, ALLOPURINOL
ZYMAR, GATIFLOXACIN
ZYMAXID, GATIFLOXACIN
ZYPITAMAG, PITAVASTATIN MAGNESIUM
ZYPREXA, OLANZAPINE
ZYPREXA RELPREVV, OLANZAPINE PAMOATE
ZYPREXA ZYDIS, OLANZAPINE
ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
ZYRTEC-D 12 HOUR, CETIRIZINE HYDROCHLORIDE (OTC)
ZYTIGA, ABIRATERONE ACETATE
ZYVOX, LINEZOLID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** 3 ******3D IMAGING DRUG**

- * 3D IMAGING DRUG DESIGN AND DEVELOPMENT LLC
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

3M

- * 3M CO
PERIDEX, CHLORHEXIDINE GLUCONATE
- * 3M HEALTH CARE INC
AVAGARD, ALCOHOL (OTC)
DURAPREP, IODINE POVACRYLEX (OTC)

3M DRUG DELIVERY

- * 3M DRUG DELIVERY SYSTEMS
FENTANYL-100, FENTANYL
FENTANYL-12, FENTANYL
FENTANYL-25, FENTANYL
FENTANYL-50, FENTANYL
FENTANYL-75, FENTANYL
PROVENTIL-HFA, ALBUTEROL SULFATE

3M HEALTH CARE

- * 3M HEALTH CARE INFECTION PREVENTION DIV
SOLUPREP, CHLORHEXIDINE GLUCONATE (OTC)

**** 6 ******60 DEGREES PHARMS**

- * 60 DEGREES PHARMACEUTICALS LLC
ARAKODA, TAFENOQUINE SUCCINATE

**** A ******AAA USA INC**

- * ADVANCED ACCELERATOR APPLICATIONS USA INC
LUTATHERA, LUTETIUM DOTATATE LU-177
NETSPOT, GALLIUM DOTATATE GA-68

AAIPHARMA LLC

- * AAIPHARMA LLC
AZASAN, AZATHIOPRINE

ABBVIE

- * ABBVIE INC
ANDROGEL, TESTOSTERONE
CREON, PANCRELIPASE (AMYLASE)
CYCLOSPORINE, CYCLOSPORINE
DEPACON, VALPROATE SODIUM
DEPAKENE, VALPROIC ACID
DEPAKOTE ER, DIVALPROEX SODIUM
DEPAKOTE, DIVALPROEX SODIUM
GENGRAF, CYCLOSPORINE
K-TAB, POTASSIUM CHLORIDE
KALETRA, LOPINAVIR
MARINOL, DRONABINOL
NIASPAN, NIACIN
NIMBEX PRESERVATIVE FREE, CISATRACURIUM BESYLATE
NIMBEX, CISATRACURIUM BESYLATE
NORVIR, RITONAVIR
SURVANTA, BERACTANT
SYNTHROID, LEVOTHYROXINE SODIUM **
TARKA, TRANDOLAPRIL
TRICOR, FENOFIBRATE
TRIDIONE, TRIMETHADIONE
TRILIPIX, CHOLINE FENOFIBRATE
ULTANE, SEVOFLURANE
ZEMPLAR, PARICALCITOL

ABBVIE ENDOCRINE

- * ABBVIE ENDOCRINE INC
LUPANETA PACK, LEUPROLIDE ACETATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******ABBVIE ENDOCRINE INC**

* ABBVIE ENDOCRINE INC
LUPRON DEPOT, LEUPROLIDE ACETATE
LUPRON DEPOT-PED, LEUPROLIDE ACETATE

ABBVIE INC

* ABBVIE INC
DUOPA, CARBIDOPA
MAVYRET, GLECAPREVIR
NORVIR, RITONAVIR
ORILISSA, ELAGOLIX SODIUM
TECHNIVIE, OMBITASVIR
VENCLEXTA, VENETOCLAX
VIEKIRA PAK (COPACKAGED), DASABUVIR SODIUM
VIEKIRA XR, DASABUVIR SODIUM

ABHAI INC

* ABHAI INC
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

ABHAI LLC

* ABHAI LLC
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
URSODIOL, URSODIOL

ABON PHARMS LLC

* ABON PHARMACEUTICALS LLC
CLOFARABINE, CLOFARABINE

ABRAXIS BIOSCIENCE

* ABRAXIS BIOSCIENCE LLC
ABRAXANE, PACLITAXEL

ABRAXIS PHARM

* ABRAXIS PHARMACEUTICAL PRODUCTS
CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE

ACADEMIC PHARMS INC

* ACADEMIC PHARMACEUTICALS INC
BRETILIUM TOSYLATE, BRETILIUM TOSYLATE

ACADIA PHARMS INC

* ACADIA PHARMACEUTICALS INC
NUPLAZID, PIMAVANSERIN TARTRATE

ACCELRX LABS

* ACCELRX LABS LLC
CARISOPRODOL, CARISOPRODOL

ACCORD HLTHCARE

* ACCORD HEALTHCARE INC
ACETAZOLAMIDE, ACETAZOLAMIDE
ALLOPURINOL, ALLOPURINOL
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
ANASTROZOLE, ANASTROZOLE
ARIPIPIRAZOLE, ARIPIPIRAZOLE
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
AZACITIDINE, AZACITIDINE
BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
BICALUTAMIDE, BICALUTAMIDE
BIVALIRUDIN, BIVALIRUDIN
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CAPECITABINE, CAPECITABINE
CARBIDOPA AND LEVODOPA, CARBIDOPA
CARBOPLATIN, CARBOPLATIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ACCORD HEALTHCARE INC
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CISPLATIN, CISPLATIN
 CLONAZEPAM, CLONAZEPAM
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CLOZAPINE, CLOZAPINE
 DALFAMPRIDINE, DALFAMPRIDINE
 DECITABINE, DECITABINE
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOCETAXEL, DOCETAXEL
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 ENTECAVIR, ENTECAVIR
 EPLERENONE, EPLERENONE
 EPTIFIBATIDE, EPTIFIBATIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 ETOPOSIDE, ETOPOSIDE
 EZETIMIBE, EZETIMIBE
 FINASTERIDE, FINASTERIDE
 FLUOROURACIL, FLUOROURACIL
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLIMEPIRIDE, GLIMEPIRIDE
 GLIPIZIDE, GLIPIZIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 ITRACONAZOLE, ITRACONAZOLE
 LETROZOLE, LETROZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LISINAPRIL, LISINAPRIL
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHYLDOPA, METHYLDOPA
 MITOMYCIN, MITOMYCIN
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE, NORETHINDRONE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PACLITAXEL, PACLITAXEL
 PARICALCITOL, PARICALCITOL
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RAMIPRIL, RAMIPRIL
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SIMVASTATIN, SIMVASTATIN
 SPIRONOLACTONE, SPIRONOLACTONE
 TACROLIMUS, TACROLIMUS
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TEMSIROLIMUS, TEMSIROLIMUS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ACCORD HEALTHCARE INC**

TERIFLUNOMIDE, TERIFLUNOMIDE
 TOPIRAMATE, TOPIRAMATE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

ACCORD HLTHCARE INC*** ACCORD HEALTHCARE INC USA**

AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 TIGECYCLINE, TIGECYCLINE

ACELLA PHARMS LLC*** ACELLA PHARMACEUTICALS LLC**

BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 GABAPENTIN, GABAPENTIN
 HYDROCODONE BITARTRATE AND CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE
 PHENYTOIN SODIUM, PHENYTOIN SODIUM

ACELRX PHARMS*** ACELRX PHARMACEUTICALS INC**

DSUVIA, SUFENTANIL CITRATE

ACHAOGEN INC*** ACHAOGEN INC**

ZEMDRI, PLAZOMICIN SULFATE

ACI HEALTHCARE LTD*** ACI HEALTHCARE LTD**

DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 LEVETIRACETAM, LEVETIRACETAM
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

ACIC PHARMS*** ACIC PHARMACEUTICALS INC**

LEVETIRACETAM, LEVETIRACETAM
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID

ACLARIS*** ACLARIS THERAPEUTICS INC**

ESKATA, HYDROGEN PEROXIDE
 RHOFAD, OXYMETAZOLINE HYDROCHLORIDE

ACORDA*** ACORDA THERAPEUTICS INC**

AMPYRA, DALFAMPRIDINE
 INBRIJA, LEVODOPA

ACS DOBFAR*** ACS DOBFAR SPA**

AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
 CEFOXITIN, CEFOXITIN SODIUM
 CEFTAZIDIME, CEFTAZIDIME
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
 MEROPENEM, MEROPENEM

ACS DOBFAR SPA*** ACS DOBFAR SPA**

AMPICILLIN SODIUM, AMPICILLIN SODIUM
 CEFUROXIME SODIUM, CEFUROXIME SODIUM
 ERTAPENEM SODIUM, ERTAPENEM SODIUM
 MEROPENEM, MEROPENEM
 PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM

ACTAVIS ELIZABETH*** ACTAVIS ELIZABETH LLC**

ALPRAZOLAM, ALPRAZOLAM
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CARBIDOPA AND LEVODOPA, CARBIDOPA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ACTAVIS ELIZABETH LLC
 CLONAZEPAM, CLONAZEPAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DEFERASIROX, DEFERASIROX
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE, DEXTROMETHORPHAN HYDROBROMIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 GABAPENTIN, GABAPENTIN
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 INDAPAMIDE, INDAPAMIDE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LOVASTATIN, LOVASTATIN
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NIFEDIPINE, NIFEDIPINE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 OXAZEPAM, OXAZEPAM
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE AND IBUPROFEN, IBUPROFEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 TEMAZEPAM, TEMAZEPAM
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

* ACTAVIS ELIZABETH LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 ALPRAZOLAM, ALPRAZOLAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 LAMOTRIGINE, LAMOTRIGINE
 MORPHINE SULFATE, MORPHINE SULFATE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

ACTAVIS INC

* ACTAVIS INC
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 METHOXSALEN, METHOXSALEN
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

ACTAVIS LABS

* ACTAVIS LABORATORIES INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 PERMETHRIN, PERMETHRIN

ACTAVIS LABS FL

* ACTAVIS LABORATORIES FL INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 GUAIFENESIN, GUAIFENESIN (OTC)
 MESALAMINE, MESALAMINE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

ACTAVIS LABS FL INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ACTAVIS LABORATORIES FL INC
 BUDESONIDE, BUDESONIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CABERGOLINE, CABERGOLINE
 CARTIA XT, DILTIAZEM HYDROCHLORIDE
 CLARITHROMYCIN, CLARITHROMYCIN
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DALFAMPRIDINE, DALFAMPRIDINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
 DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
 DUTASTERIDE, DUTASTERIDE
 FENTANYL CITRATE, FENTANYL CITRATE
 FLUTAMIDE, FLUTAMIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 LEVETIRACETAM, LEVETIRACETAM
 LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
 LORATADINE, LORATADINE (OTC)
 METAXALONE, METAXALONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 NITROGLYCERIN, NITROGLYCERIN
 OMEPRAZOLE, OMEPRAZOLE
 OXYCODONE AND ASPIRIN, ASPIRIN
 PALIPERIDONE, PALIPERIDONE
 PAROXETINE MESYLATE, PAROXETINE MESYLATE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREDNISONE, PREDNISONE
 RAMELTEON, RAMELTEON
 RISPERIDONE, RISPERIDONE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TAZTIA XT, DILTIAZEM HYDROCHLORIDE
 TETRABENAZINE, TETRABENAZINE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

ACTAVIS LABS NY INC

* ACTAVIS LABORATORIES NY INC
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)

ACTAVIS LABS UT INC

* ACTAVIS LABORATORIES UT INC
 AZELAIC ACID, AZELAIC ACID
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLONIDINE, CLONIDINE
 DOCOSANOL, DOCOSANOL (OTC)
 EMLA, LIDOCAINE
 FLORICET W/ CODEINE, ACETAMINOPHEN
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 LIDOCAINE, LIDOCAINE
 NORINYL 1+50 28-DAY, MESTRANOL
 PROGESTERONE, PROGESTERONE
 TESTOSTERONE, TESTOSTERONE

* ACTAVIS LABORATORIES UT INC INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 PIMECROLIMUS, PIMECROLIMUS
 TESTOSTERONE, TESTOSTERONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******ACTAVIS LLC**

- * ACTAVIS LLC
 - AZACITIDINE, AZACITIDINE
 - DAPSONE, DAPSONE
 - FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 - LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 - MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
- * ACTAVIS LLC AN INDIRECT WHOLLY-OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - BUSULFAN, BUSULFAN
 - DOCETAXEL, DOCETAXEL
 - HYDROXOCOBALAMIN, HYDROXOCOBALAMIN
 - OXALIPLATIN, OXALIPLATIN

ACTAVIS MID ATLANTIC

- * ACTAVIS MID ATLANTIC LLC
 - ACETASOL HC, ACETIC ACID, GLACIAL
 - ACYCLOVIR, ACYCLOVIR
 - ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 - ADAPALENE, ADAPALENE
 - BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 - CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 - CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 - CICLOPIROX, CICLOPIROX
 - CLINDAMYCIN PHOSPHATE AND TRETINOIN, CLINDAMYCIN PHOSPHATE
 - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 - CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
 - DESOXIMETASONE, DESOXIMETASONE
 - DICLOFENAC SODIUM, DICLOFENAC SODIUM
 - ENULOSE, LACTULOSE
 - GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 - HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 - HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 - HYDROCORTISONE, HYDROCORTISONE
 - IBUPROFEN, IBUPROFEN (OTC)
 - LEVETIRACETAM, LEVETIRACETAM
 - MICONAZOLE 7, MICONAZOLE NITRATE (OTC)
 - MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 - MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 - NITROFURANTOIN, NITROFURANTOIN
 - NYSTATIN, NYSTATIN
 - PROMETH VC W/ CODEINE, CODEINE PHOSPHATE
 - PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 - VALNAC, BETAMETHASONE VALERATE
- * ACTAVIS MID ATLANTIC LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - IBUPROFEN, IBUPROFEN
 - M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 - PERMETHRIN, PERMETHRIN (OTC)

ACTAVIS PHARMA

- * ACTAVIS PHARMA INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - MICONAZOLE NITRATE, MICONAZOLE NITRATE
 - MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)

ACTAVIS TOTOWA

- * ACTAVIS TOTOWA LLC
 - DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 - EPIDUBICIN HYDROCHLORIDE, EPIDUBICIN HYDROCHLORIDE
 - FINASTERIDE, FINASTERIDE
 - FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 - OXALIPLATIN, OXALIPLATIN
 - PACLITAXEL, PACLITAXEL
 - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 - REPAGLINIDE, REPAGLINIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * ACTAVIS TOTOWA LLC
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
VINORELBINE TARTRATE, VINORELBINE TARTRATE
- ACTAVIS TOTOWA TEVA**
- * ACTAVIS TOTOWA LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
FINASTERIDE, FINASTERIDE
- ACTELION PHARMS**
- * ACTELION PHARMACEUTICALS LTD
TRACLEER, BOSENTAN
- ACTELION PHARMS LTD**
- * ACTELION PHARMACEUTICALS LTD
OPSUMIT, MACITENTAN
TRACLEER, BOSENTAN
UPTRAVI, SELEXIPAG
VELETRI, EPOPROSTENOL SODIUM
VENTAVIS, ILOPROST
ZAVESCA, MIGLUSTAT
- ACTIENT PHARMS**
- * ACTIENT PHARMACEUTICALS LLC
THEO-24, THEOPHYLLINE
- ADAMAS PHARMA**
- * ADAMAS PHARMA LLC
GOCOVRI, AMANTADINE HYDROCHLORIDE
- ADAMIS PHARMS CORP**
- * ADAMIS PHARMACEUTICALS CORP
SYMJEPI, EPINEPHRINE
- ADAPT**
- * ADAPT PHARMA OPERATIONS LTD
NARCAN, NALOXONE HYDROCHLORIDE
- ADARE PHARMS INC**
- * ADARE PHARMACEUTICALS INC
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
- ADDMEDICA SAS**
- * ADDMEDICA SAS
SIKLOS, HYDROXYUREA
- ADIENNE SA**
- * ADIENNE SA
TEPADINA, THIOTEPA
- AEGERION**
- * AEGERION PHARMACEUTICALS INC
JUXTAPID, LOMITAPIDE MESYLATE
- AERIE PHARMS INC**
- * AERIE PHARMACEUTICALS INC
RHOPRESSA, NETARSUDIL DIMESYLATE
- AGIOS PHARMS INC**
- * AGIOS PHARMACEUTICALS INC
TIBSOVO, IVOSIDENIB
- AGOURON PHARMS**
- * AGOURON PHARMACEUTICALS LLC
VIRACEPT, NELFINAVIR MESYLATE
- AILEX PHARMS LLC**
- * AILEX PHARMACEUTICALS LLC
BISMUTH SUBSALICYLATE, METRONIDAZOLE AND TETRACYCLINE HYDROCHLORIDE, BISMUTH
CROMOLYN SODIUM, CROMOLYN SODIUM
SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE
- AIPING PHARM INC**
- * AIPING PHARMACEUTICAL INC
BENZONATATE, BENZONATATE
- AJANTA PHARMA LTD**
- * AJANTA PHARMA LTD
ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
ARIPIPIRAZOLE, ARIPIPIRAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AJANTA PHARMA LTD**

CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 ENTACAPONE, ENTACAPONE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRATE, FENOFIBRATE
 LANSOPRAZOLE, LANSOPRAZOLE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 VORICONAZOLE, VORICONAZOLE
 ZOLMITRIPTAN, ZOLMITRIPTAN

AKARX INC*** AKARX INC**

DOPTELET, AVATROMBOPAG MALEATE

AKCEA THERAPS*** AKCEA THERAPEUTICS INC**

TEGSEDI, INOTERSEN SODIUM

AKORN*** AKORN INC**

ADENOSINE, ADENOSINE
 AK-FLUOR 10%, FLUORESCEIN SODIUM
 AK-FLUOR 25%, FLUORESCEIN SODIUM
 AKBETA, LEVOBUNOLOL HYDROCHLORIDE
 AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE
 AKTEN, LIDOCAINE HYDROCHLORIDE
 AKTOB, TOBRAMYCIN
 ALFENTA, ALFENTANIL HYDROCHLORIDE
 ATROPINE SULFATE, ATROPINE SULFATE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 BACITRACIN, BACITRACIN
 BAL, DIMERCAPROL
 BALANCED SALT, CALCIUM CHLORIDE
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 CALCITRIOL, CALCITRIOL
 CAPASTAT SULFATE, CAPREOMYCIN SULFATE
 CARBOPLATIN, CARBOPLATIN
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 DESOXIMETASONE, DESOXIMETASONE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 EPTIFIBATIDE, EPTIFIBATIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 GENTAK, GENTAMICIN SULFATE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 IC-GREEN, INDOCYANINE GREEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AKORN INC**

IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 LATANOPROST, LATANOPROST
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LORAZEPAM, LORAZEPAM
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NEDOCROMIL SODIUM, NEDOCROMIL SODIUM
 NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
 NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE, BACITRACIN ZINC
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OFLOXACIN, OFLOXACIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 PAREMYD, HYDROXYAMPHETAMINE HYDROBROMIDE
 PARICALCITOL, PARICALCITOL
 PYRAZINAMIDE, PYRAZINAMIDE
 RIFAMPIN, RIFAMPIN
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUBLIMAZE PRESERVATIVE FREE, FENTANYL CITRATE
 SUFENTA PRESERVATIVE FREE, SUFENTANIL CITRATE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIMOLOL, TIMOLOL
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TROPICACYL, TROPICAMIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

AKORN INC*** AKORN INC**

ACETYLCYSTEINE, ACETYLCYSTEINE
 APRACLONIDINE HYDROCHLORIDE, APRACLONIDINE HYDROCHLORIDE
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 CYCLOPENTOLATE HYDROCHLORIDE, CYCLOPENTOLATE HYDROCHLORIDE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DRONABINOL, DRONABINOL
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 INAPSINE, DROPERIDOL
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AKORN INC**

ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 TOBRAMYCIN, TOBRAMYCIN
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

ALCON*** ALCON LABORATORIES INC**

BSS PLUS, CALCIUM CHLORIDE
 BSS, CALCIUM CHLORIDE
 MIOSTAT, CARBACHOL
 NAPHCN-A, NAPHAZOLINE HYDROCHLORIDE (OTC)

ALCON LABS*** ALCON LABORATORIES LTD**

TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE

ALCON LABS INC*** ALCON LABORATORIES INC**

ALCAINE, PROPARACAINE HYDROCHLORIDE
 CYCLOGYL, CYCLOPENTOLATE HYDROCHLORIDE
 CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE
 FLUORESCITE, FLUORESCHEIN SODIUM
 ISOPTO ATROPINE, ATROPINE SULFATE

ALCON PHARMS LTD*** ALCON PHARMACEUTICALS LTD**

BETADINE, POVIDONE-IODINE
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)

ALEMBIC LTD*** ALEMBIC LTD**

LITHIUM CARBONATE, LITHIUM CARBONATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE

ALEMBIC PHARMS LTD*** ALEMBIC PHARMACEUTICALS LTD**

ACYCLOVIR, ACYCLOVIR
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CELECOXIB, CELECOXIB
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DESVENLAFAXINE, DESVENLAFAXINE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 FAMOTIDINE, FAMOTIDINE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 ITRACONAZOLE, ITRACONAZOLE
 LAMOTRIGINE, LAMOTRIGINE
 LEFLUNOMIDE, LEFLUNOMIDE
 LINEZOLID, LINEZOLID
 LITHIUM CARBONATE, LITHIUM CARBONATE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MEPROBAMATE, MEPROBAMATE
 METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE, METRONIDAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ALEMBIC PHARMACEUTICALS LTD**

MODAFINIL, MODAFINIL
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMAZEPAM, TEMAZEPAM
 THEOPHYLLINE, THEOPHYLLINE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN

ALEOR DERMACEUTICALS*** ALEOR DERMACEUTICALS LTD**

LIDOCAINE, LIDOCAINE

ALIGNSCIENCE PHARMA*** ALIGNSCIENCE PHARMA INC**

METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE

ALIMERA SCIENCES INC*** ALIMERA SCIENCES INC**

ILUVIEN, FLUOCINOLONE ACETONIDE

ALKALOIDA ZRT*** ALKALOIDA CHEMICAL CO ZRT**

HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE

ALKEM*** ALKEM LABORATORIES LTD**

AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 GABAPENTIN, GABAPENTIN
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

ALKEM LABS LTD*** ALKEM LABORATORIES LTD**

AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 CAPECITABINE, CAPECITABINE
 CEFIXIME, CEFIXIME
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CEPHALEXIN, CEPHALEXIN
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DALFAMPRIDINE, DALFAMPRIDINE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 FINASTERIDE, FINASTERIDE
 GABAPENTIN, GABAPENTIN
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 ITRACONAZOLE, ITRACONAZOLE
 LAMOTRIGINE, LAMOTRIGINE
 LIDOCAINE, LIDOCAINE
 LINEZOLID, LINEZOLID
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALKEM LABORATORIES LTD
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 RILUZOLE, RILUZOLE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

ALKERMES

* ALKERMES INC
 VIVITROL, NALTREXONE

ALKERMES INC

* ALKERMES INC
 ARISTADA INITIO KIT, ARIPIPRAZOLE LAUROXIL
 ARISTADA, ARIPIPRAZOLE LAUROXIL

ALLEGIANCE HLTHCARE

* ALLEGIANCE HEALTHCARE CORP
 POVIDONE IODINE, POVIDONE-IODINE (OTC)

ALLEGIS

* ALLEGIS HOLDINGS LLC
 PRIMISOL, TRIMETHOPRIM HYDROCHLORIDE

ALLERGAN

* ALLERGAN
 ACULAR LS, KETOROLAC TROMETHAMINE
 ALPHAGAN P, BRIMONIDINE TARTRATE
 BLEPH-10, SULFACETAMIDE SODIUM
 GENOPTIC, GENTAMICIN SULFATE
 ZYMAXID, GATIFLOXACIN

* ALLERGAN INC
 ACULAR, KETOROLAC TROMETHAMINE
 ACUVAIL, KETOROLAC TROMETHAMINE
 ACZONE, DAPSONE
 ALOCRIIL, NEDOCROMIL SODIUM
 ALPHAGAN P, BRIMONIDINE TARTRATE
 AVAGE, TAZAROTENE
 COMBIGAN, BRIMONIDINE TARTRATE
 ELESTAT, EPINASTINE HYDROCHLORIDE
 LASTACAFT, ALCAFTADINE
 LATISSE, BIMATOPROST
 LUMIGAN, BIMATOPROST
 OCUFLOX, OFLOXACIN
 OZURDEX, DEXAMETHASONE
 POLYTRIM, POLYMYXIN B SULFATE
 RESTASIS MULTIDOSE, CYCLOSPORINE
 RESTASIS, CYCLOSPORINE
 TAZORAC, TAZAROTENE
 ZYMAR, GATIFLOXACIN

* ALLERGAN PHARMACEUTICAL
 BETAGAN, LEVOBUNOLOL HYDROCHLORIDE
 BLEPHAMIDE S.O.P., PREDNISOLONE ACETATE
 BLEPHAMIDE, PREDNISOLONE ACETATE
 FML FORTE, FLUOROMETHOLONE
 FML, FLUOROMETHOLONE
 OCUFEN, FLURBIPROFEN SODIUM
 PRED FORTE, PREDNISOLONE ACETATE
 PRED MILD, PREDNISOLONE ACETATE
 PRED-G, GENTAMICIN SULFATE

ALLERGAN HOLDINGS

* ALLERGAN HOLDINGS UNLTD CO
 VIBERZI, ELUXADOLINE

ALLERGAN SALES LLC

* ALLERGAN SALES LLC
 ACTIGALL, URSODIOL
 ALORA, ESTRADIOL
 ANDRODERM, TESTOSTERONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALLERGAN SALES LLC
 AVYCAZ, AVIBACTAM SODIUM
 BENTYL PRESERVATIVE FREE, DICYCLOMINE HYDROCHLORIDE
 BENTYL, DICYCLOMINE HYDROCHLORIDE
 BREVICON 28-DAY, ETHINYL ESTRADIOL
 BYSTOLIC, NEBIVOLOL HYDROCHLORIDE
 BYVALSON, NEBIVOLOL HYDROCHLORIDE
 CANASA, MESALAMINE
 CARAFATE, SUCRALFATE
 CELEXA, CITALOPRAM HYDROBROMIDE
 CONDYLOX, PODOFILOX
 CRINONE, PROGESTERONE
 DALVANCE, DALBAVANCIN HYDROCHLORIDE
 ESTRACE, ESTRADIOL
 FETZIMA, LEVOMILNACIPRAN HYDROCHLORIDE
 FIORINAL W/CODEINE, ASPIRIN
 FIORINAL, ASPIRIN
 GELNIQUE, OXYBUTYNIN CHLORIDE
 INFED, IRON DEXTRAN
 KADIAN, MORPHINE SULFATE
 LEXAPRO, ESCITALOPRAM OXALATE
 LINZESS, LINACLOTIDE
 MICROZIDE, HYDROCHLOROTHIAZIDE
 NAMENDA, MEMANTINE HYDROCHLORIDE
 NAMZARIC, DONEPEZIL HYDROCHLORIDE
 NORINYL 1+35 21-DAY, ETHINYL ESTRADIOL
 NORINYL 1+35 28-DAY, ETHINYL ESTRADIOL
 OXYTROL FOR WOMEN, OXYBUTYNIN (OTC)
 OXYTROL, OXYBUTYNIN
 PYLERA, BISMUTH SUBCITRATE POTASSIUM
 RAPAFLO, SILODOSIN
 RECTIV, NITROGLYCERIN
 SAVELLA, MILNACIPRAN HYDROCHLORIDE
 TEFLARO, CEFTAROLINE FOSAMIL
 TRELSTAR, TRIPTORELIN PAMOATE
 URSO 250, URSODIOL
 URSO FORTE, URSODIOL
 VIIBRYD, VILAZODONE HYDROCHLORIDE
 VRAYLAR, CARIPRAZINE HYDROCHLORIDE

ALLERQUEST

* ALLERQUEST LLC
 PRE-PEN, BENZYLPENICILLOYL POLYLYSINE

ALLIED

* ALLIED PHARMA INC
 CLARITHROMYCIN, CLARITHROMYCIN
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

ALLIED PHARMA INC

* ALLIED PHARMA INC
 CLARITHROMYCIN, CLARITHROMYCIN

ALLOS

* ALLOS THERAPEUTICS INC
 FOLOTYN, PRALATREXATE

ALMIRALL

* ALMIRALL LLC
 SEYSARA, SARECYCLINE HYDROCHLORIDE

ALNYLAM PHARMS INC

* ALNYLAM PHARMACEUTICALS INC
 ONPATRO, PATISIRAN SODIUM

ALPHARMA PHARMS

* ALPHARMA PHARMACEUTICALS LLC
 EMBEDA, MORPHINE SULFATE

ALRA

* ALRA LABORATORIES INC
 CHOLAC, LACTULOSE
 CONSTILAC, LACTULOSE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALRA LABORATORIES INC
 GEN-XENE, CLORAZEPATE DIPOTASSIUM
 IBU-TAB 200, IBUPROFEN (OTC)
 IBU-TAB, IBUPROFEN

ALTAIRE PHARMS INC

* ALTAIRE PHARMACEUTICALS INC
 ALTAFLUOR BENOX, BENOXINATE HYDROCHLORIDE
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 OFLOXACIN, OFLOXACIN

ALTATHERA PHARMS LLC

* ALTATHERA PHARMACEUTICALS LLC
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

ALVOGEN

* ALVOGEN GROUP HOLDINGS 2 LLC
 DAPSONE, DAPSONE
 ETHACRYNIC ACID, ETHACRYNIC ACID
 OFLOXACIN, OFLOXACIN

* ALVOGEN GROUP HOLDINGS 3 LLC
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FORFIVO XL, BUPROPION HYDROCHLORIDE

* ALVOGEN GROUP HOLDINGS LLC
 ADALAT CC, NIFEDIPINE

* ALVOGEN INC
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

ALVOGEN INC

* ALVOGEN INC
 ACETYLCYSTEINE, ACETYLCYSTEINE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE
 VORICONAZOLE, VORICONAZOLE

ALVOGEN MALTA

* ALVOGEN MALTA OPERATIONS LTD
 ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 ATENOLOL, ATENOLOL
 BUDESONIDE, BUDESONIDE
 CARBIDOPA, CARBIDOPA
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DISULFIRAM, DISULFIRAM
 EXEMESTANE, EXEMESTANE
 FELBAMATE, FELBAMATE
 MACROBID, NITROFURANTOIN
 MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE
 MELPHALAN, MELPHALAN
 NAPRELAN, NAPROXEN SODIUM
 NATEGLINIDE, NATEGLINIDE
 NEVIRAPINE, NEVIRAPINE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RIVASTIGMINE, RIVASTIGMINE
 SODIUM PHENYL BUTYRATE, SODIUM PHENYL BUTYRATE
 SPECTAZOLE, ECONAZOLE NITRATE
 TENORETIC 100, ATENOLOL
 TENORETIC 50, ATENOLOL
 TENORMIN, ATENOLOL
 ZESTORETIC, HYDROCHLOROTHIAZIDE
 ZESTRIL, LISINAPRIL

ALVOGEN PINE BROOK

* ALVOGEN PINE BROOK LLC
 ESTRADIOL, ESTRADIOL
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******AM ANTIBIOTICS**

* AMERICAN ANTIBIOTICS INC
 AMOXICILLIN, AMOXICILLIN

AMAG PHARMA USA

* AMAG PHARMA USA INC
 MAKENA (AUTOINJECTOR), HYDROXYPROGESTERONE CAPROATE
 MAKENA PRESERVATIVE FREE, HYDROXYPROGESTERONE CAPROATE
 MAKENA, HYDROXYPROGESTERONE CAPROATE

AMAG PHARMS INC

* AMAG PHARMACEUTICALS INC
 FERAHME, FERUMOXYTOL
 INTRAROSA, PRASTERONE

AMARIN PHARMS

* AMARIN PHARMACEUTICALS IRELAND LTD
 VASCEPA, ICOSAPENT ETHYL

AMERIGEN PHARMS LTD

* AMERIGEN PHARMACEUTICALS LTD
 BEXAROTENE, BEXAROTENE
 CARBIDOPA, CARBIDOPA
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 INDAPAMIDE, INDAPAMIDE
 MIGLUSTAT, MIGLUSTAT
 TEMOZOLOMIDE, TEMOZOLOMIDE

AMGEN

* AMGEN INC
 SENSIPAR, CINACALCET HYDROCHLORIDE

AMGEN INC

* AMGEN INC
 CORLANOR, IVABRADINE HYDROCHLORIDE

AMICUS THERAPS US

* AMICUS THERAPEUTICS US INC
 GALAFOLD, MIGALASTAT HYDROCHLORIDE

AMNEAL PHARM

* AMNEAL PHARMACEUTICAL
 ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
 BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FOLIC ACID, FOLIC ACID
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 PRIMIDONE, PRIMIDONE

AMNEAL PHARMS

* AMNEAL PHARMACEUTICALS
 ACYCLOVIR, ACYCLOVIR
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ARIPIRAZOLE, ARIPIRAZOLE
 ATOVAQUONE, ATOVAQUONE
 CALCITRIOL, CALCITRIOL
 CALCIUM ACETATE, CALCIUM ACETATE
 CAPECITABINE, CAPECITABINE
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 ENTECAVIR, ENTECAVIR
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESTRADIOL, ESTRADIOL
 FELBAMATE, FELBAMATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AMNEAL PHARMACEUTICALS**

GABAPENTIN, GABAPENTIN
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 INDOMETHACIN, INDOMETHACIN
 ITRACONAZOLE, ITRACONAZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LIDOCAINE, LIDOCAINE
 LINEZOLID, LINEZOLID
 LORAZEPAM, LORAZEPAM
 MEROPENEM, MEROPENEM
 METAXALONE, METAXALONE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NIACIN, NIACIN
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 NITROFURANTOIN, NITROFURANTOIN
 NIZATIDINE, NIZATIDINE
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXCARBAZEPINE, OXCARBAZEPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
 QUININE SULFATE, QUININE SULFATE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TELMISARTAN, TELMISARTAN
 TEMAZEPAM, TEMAZEPAM
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 WARFARIN SODIUM, WARFARIN SODIUM

*** AMNEAL PHARMACEUTICALS HOLDINGS GMBH**

DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE

*** AMNEAL PHARMACEUTICALS OF NEW YORK LLC**

ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 AMPHETAMINE SULFATE, AMPHETAMINE SULFATE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BUDESONIDE, BUDESONIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CELECOXIB, CELECOXIB
 DUTASTERIDE, DUTASTERIDE
 EPTIFIBATIDE, EPTIFIBATIDE
 EXEMESTANE, EXEMESTANE
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 GUAIFENESIN, GUAIFENESIN (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 IRBESARTAN, IRBESARTAN
 LAMOTRIGINE, LAMOTRIGINE
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AMNEAL PHARMACEUTICALS OF NEW YORK LLC**

METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE, NORETHINDRONE
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PARICALCITOL, PARICALCITOL
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RIVASTIGMINE, RIVASTIGMINE
 SPIRONOLACTONE, SPIRONOLACTONE
 TOBRAMYCIN, TOBRAMYCIN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VALSARTAN, VALSARTAN
 VIGABATRIN, VIGABATRIN

AMNEAL PHARMS CO*** AMNEAL PHARMACEUTICALS CO GMBH**

ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ARGATROBAN, ARGATROBAN
 BUMETANIDE, BUMETANIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 BUSULFAN, BUSULFAN
 CARMUSTINE, CARMUSTINE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 CLOFARABINE, CLOFARABINE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DOCETAXEL, DOCETAXEL
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ERYTHROMYCIN, ERYTHROMYCIN
 ETHACRYNIC ACID, ETHACRYNIC ACID
 ETODOLAC, ETODOLAC
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 FUROSEMIDE, FUROSEMIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 NADOLOL, NADOLOL
 NAPROXEN SODIUM AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OXAPROZIN, OXAPROZIN
 PARICALCITOL, PARICALCITOL
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PHYTONADIONE, PHYTONADIONE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMNEAL PHARMACEUTICALS CO GMBH
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SILODOSIN, SILODOSIN
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 URSODIOL, URSODIOL
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

AMNEAL PHARMS LLC

* AMNEAL PHARMACEUTICALS LLC
 ACTIVELLA, ESTRADIOL
 AZATHIOPRINE, AZATHIOPRINE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)
 FENOFIBRATE, FENOFIBRATE
 FLUOCINONIDE, FLUOCINONIDE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 RITONAVIR, RITONAVIR
 TIGECYCLINE, TIGECYCLINE

AMNEAL PHARMS NY

* AMNEAL PHARMACEUTICALS NY LLC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ALPRAZOLAM, ALPRAZOLAM
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 GABAPENTIN, GABAPENTIN
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NAPROXEN, NAPROXEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 REPRESXAIN, HYDROCODONE BITARTRATE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

* AMNEAL PHARMACEUTICALS OF NY LLC
 BEXAROTENE, BEXAROTENE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 ISOTRETINOIN, ISOTRETINOIN
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PROGESTERONE, PROGESTERONE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

AMPHASTAR PHARM

* AMPHASTAR PHARMACEUTICAL INC
 AMPHADASE, HYALURONIDASE
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE

AMPHASTAR PHARMS INC

* AMPHASTAR PHARMACEUTICALS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AMPHASTAR PHARMACEUTICALS INC
 CORTROSYN, COSYNTROPIN
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE

AMRING PHARMS

* AMRING PHARMACEUTICALS INC
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 LATANOPROST, LATANOPROST
 NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE

ANACOR PHARMS INC

* ANACOR PHARMACEUTICALS INC
 EUCRISA, CRISABOROLE
 KERYDIN, TAVABOROLE

ANBEX

* ANBEX INC
 IOSAT, POTASSIUM IODIDE (OTC)

ANBISON LAB

* ANBISON LABORATORY CO LTD
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 MONTELUKAST SODIUM, MONTELUKAST SODIUM

ANCHEN PHARMS

* ANCHEN PHARMACEUTICALS INC
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRETINOIN, TRETINOIN
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

* ANCHEN PHARMACEUTICALS TAIWAN INC
 DIVALPROEX SODIUM, DIVALPROEX SODIUM

* ANCHEN PHARMACEUTICALS, INC
 ALPRAZOLAM, ALPRAZOLAM
 CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN

ANDA REPOSITORY

* ANDA REPOSITORY LLC
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 IMIQUIMOD, IMIQUIMOD
 LEVOCARNITINE, LEVOCARNITINE
 LORAZEPAM, LORAZEPAM
 MOMETASONE FUROATE, MOMETASONE FUROATE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 PRIMIDONE, PRIMIDONE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

ANDRX LABS LLC

* ANDRX LABS LLC
 FORTAMET, METFORMIN HYDROCHLORIDE

ANI PHARMS

* ANI PHARMACEUTICALS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ANI PHARMACEUTICALS INC
 CORTENEMA, HYDROCORTISONE
 LACTULOSE, LACTULOSE
 LUVOX, FLUVOXAMINE MALEATE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 REGLAN, METOCLOPRAMIDE HYDROCHLORIDE

ANI PHARMS INC

* ANI PHARMACEUTICALS INC
 ALPRAZOLAM, ALPRAZOLAM
 ARIMIDEX, ANASTROZOLE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 ATACAND HCT, CANDESARTAN CILEXETIL
 ATACAND, CANDESARTAN CILEXETIL
 BRETHINE, TERBUTALINE SULFATE
 CASODEX, BICALUTAMIDE
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
 ETODOLAC, ETODOLAC
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 FELBAMATE, FELBAMATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 GLIPIZIDE, GLIPIZIDE
 GUANABENZ ACETATE, GUANABENZ ACETATE
 INDAPAMIDE, INDAPAMIDE
 INDERAL LA, PROPRANOLOL HYDROCHLORIDE
 INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 LITHOBID, LITHIUM CARBONATE
 LORAZEPAM, LORAZEPAM
 METHAZOLAMIDE, METHAZOLAMIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NILUTAMIDE, NILUTAMIDE
 NIZATIDINE, NIZATIDINE
 OXCARBAZEPINE, OXCARBAZEPINE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
 PINDOLOL, PINDOLOL
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALPROIC ACID, VALPROIC ACID
 VANCOCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

ANTARES PHARMA INC

* ANTARES PHARMA INC
 OTREXUP, METHOTREXATE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 XYOSTED (AUTOINJECTOR), TESTOSTERONE ENANTHATE

ANTIBIOTICE

* ANTIBIOTICE SA
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 NAFCILLIN SODIUM, NAFCILLIN SODIUM

ANTRIM PHARMS LLC

* ANTRIM PHARMACEUTICALS LLC
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE

APC PHARMS LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* APC PHARMACEUTICALS LLC

DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

APEX PHARMS INC

* APEX PHARMACEUTICALS INC

CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE

APGDI

* ASTELLAS PHARMA GLOBAL DEVELOPMENT INC

MYRBETRIQ, MIRABEGRON

APICORE US

* APICORE US LLC

TETRABENAZINE, TETRABENAZINE

APIL

* ALLERGAN PHARMACEUTICALS INTERNATIONAL LTD

ACTONEL, RISEDRONATE SODIUM

ASACOL HD, MESALAMINE

ATELVIA, RISEDRONATE SODIUM

DELZICOL, MESALAMINE

ENABLEX, DARIFENACIN HYDROBROMIDE

ESTROSTEP FE, ETHINYL ESTRADIOL

FEMCON FE, ETHINYL ESTRADIOL

FEMHRT, ETHINYL ESTRADIOL

LO LOESTRIN FE, ETHINYL ESTRADIOL

LOESTRIN 21 1.5/30, ETHINYL ESTRADIOL

LOESTRIN 21 1/20, ETHINYL ESTRADIOL

LOESTRIN 24 FE, ETHINYL ESTRADIOL

LOESTRIN FE 1.5/30, ETHINYL ESTRADIOL

LOESTRIN FE 1/20, ETHINYL ESTRADIOL

MINASTRIN 24 FE, ETHINYL ESTRADIOL

NOR-QD, NORETHINDRONE

NORCO, ACETAMINOPHEN

NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL

SARAFEM, FLUOXETINE HYDROCHLORIDE

TAYTULLA, ETHINYL ESTRADIOL

APNAR PHARMA LP

* APNAR PHARMA LP

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)

CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE

POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)

PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE

SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE

APOLLO PHARMS INC

* APOLLO PHARMACEUTICALS INC

PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM

APOPHARMA INC

* APOPHARMA INC

FERRIPROX, DEFERIPRONE

APOTEX

* APOTEX INC

ALENDRONATE SODIUM, ALENDRONATE SODIUM

AMLODIPINE BESYLATE, AMLODIPINE BESYLATE

CARBIDOPA AND LEVODOPA, CARBIDOPA

CEFUROXIME AXETIL, CEFUROXIME AXETIL

CIMETIDINE, CIMETIDINE

CIMETIDINE, CIMETIDINE (OTC)

CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE

CYCLOSPORINE, CYCLOSPORINE

DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM

DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE

DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE

ENALAPRIL MALEATE, ENALAPRIL MALEATE

EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE

EPLERENONE, EPLERENONE

ETODOLAC, ETODOLAC

FAMCICLOVIR, FAMCICLOVIR

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* APOTEX INC**

FAMOTIDINE, FAMOTIDINE
 FLUCONAZOLE, FLUCONAZOLE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 GEMFIBROZIL, GEMFIBROZIL
 GLIPIZIDE, GLIPIZIDE
 LAMIVUDINE, LAMIVUDINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 OMEPRAZOLE, OMEPRAZOLE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PENTOXIFYLLINE, PENTOXIFYLLINE
 RAMIPRIL, RAMIPRIL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
 TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE

APOTEX CORP*** APOTEX CORP**

CLARITHROMYCIN, CLARITHROMYCIN
 QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 RILUZOLE, RILUZOLE

APOTEX INC*** APOTEX INC**

ABACAVIR SULFATE, ABACAVIR SULFATE
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACYCLOVIR, ACYCLOVIR
 ADEFOVIR DIPIVOXIL, ADEFOVIR DIPIVOXIL
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 ALKERAN, MELPHALAN
 ALPRAZOLAM, ALPRAZOLAM
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 ANASTROZOLE, ANASTROZOLE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 ATOVAQUONE, ATOVAQUONE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BENZONATATE, BENZONATATE
 BICALUTAMIDE, BICALUTAMIDE
 BIMATOPROST, BIMATOPROST
 BIVALIRUDIN, BIVALIRUDIN
 BUDESONIDE, BUDESONIDE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CARBAMAZEPINE, CARBAMAZEPINE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CEFPROZIL, CEFPROZIL
 CELECOXIB, CELECOXIB
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CICLOPIROX, CICLOPIROX
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIVALPROEX SODIUM, DIVALPROEX SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* APOTEX INC
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
DUTASTERIDE, DUTASTERIDE
ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
EZETIMIBE, EZETIMIBE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
IBANDRONATE SODIUM, IBANDRONATE SODIUM
IMATINIB MESYLATE, IMATINIB MESYLATE
IMIQUIMOD, IMIQUIMOD
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LAMIVUDINE, LAMIVUDINE
LAMOTRIGINE, LAMOTRIGINE
LETROZOLE, LETROZOLE
LEVETIRACETAM, LEVETIRACETAM
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
LEVOFLOXACIN, LEVOFLOXACIN
LEVONORGESTREL, LEVONORGESTREL (OTC)
LOVASTATIN, LOVASTATIN
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
MODAFINIL, MODAFINIL
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
MOMETASONE FUROATE, MOMETASONE FUROATE
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
NABUMETONE, NABUMETONE
NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
OFLOXACIN, OFLOXACIN
OLANZAPINE, OLANZAPINE
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
OXCARBAZEPINE, OXCARBAZEPINE
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
RISEDRONATE SODIUM, RISEDRONATE SODIUM
RISPERIDONE, RISPERIDONE
RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
TEMOZOLOMIDE, TEMOZOLOMIDE
TENOFIVIR DISOPROXIL FUMARATE, TENOFIVIR DISOPROXIL FUMARATE
TERIFLUNOMIDE, TERIFLUNOMIDE
TIGECYCLINE, TIGECYCLINE
TIMOLOL MALEATE, TIMOLOL MALEATE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
TRANEXAMIC ACID, TRANEXAMIC ACID
TRAVOPROST, TRAVOPROST
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
ZOLEDRONIC ACID, ZOLEDRONIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * APOTEX INC
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
- * APOTEX INC ETOBICOKE SITE
ACYCLOVIR, ACYCLOVIR
ALLOPURINOL, ALLOPURINOL
BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CARBAMAZEPINE, CARBAMAZEPINE
CARVEDILOL, CARVEDILOL
CILOSTAZOL, CILOSTAZOL
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DILTZAC, DILTIAZEM HYDROCHLORIDE
ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
ETODOLAC, ETODOLAC
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
GABAPENTIN, GABAPENTIN
LEFLUNOMIDE, LEFLUNOMIDE
LISINOPRIL, LISINOPRIL
LORATADINE, LORATADINE (OTC)
MELOXICAM, MELOXICAM
MIRTAZAPINE, MIRTAZAPINE
OXAPROZIN, OXAPROZIN
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
TOPIRAMATE, TOPIRAMATE
TORSEMIDE, TORSEMIDE
ZONISAMIDE, ZONISAMIDE
- * APOTEX INC RICHMOND HILL
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
BUDESONIDE, BUDESONIDE (OTC)
DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
RISPERIDONE, RISPERIDONE
- * APOTEX INC.
DILTZAC, DILTIAZEM HYDROCHLORIDE
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
- APOTEX TECHNOLOGIES**
- * APOTEX TECHNOLOGIES INC
PAXIL CR, PAROXETINE HYDROCHLORIDE
PAXIL, PAROXETINE HYDROCHLORIDE
- APOTHECON**
- * APOTHECON INC DIV BRISTOL MYERS SQUIBB
KENALOG-10, TRIAMCINOLONE ACETONIDE
KENALOG-40, TRIAMCINOLONE ACETONIDE
- APP PHARMS**
- * APP PHARMACEUTICALS LLC
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
- APPCO PHARMA LLC**
- * APPCO PHARMA LLC
CHLORTHALIDONE, CHLORTHALIDONE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
- APRECIA PHARMS**
- * APRECIA PHARMACEUTICALS LLC
SPRITAM, LEVETIRACETAM
- APTAPHARMA INC**
- * APTAPHARMA INC
IBUPROFEN, IBUPROFEN (OTC)
- AQUA PHARMS**
- * AQUA PHARMACEUTICALS
CORDRAN SP, FLURANDRENOLIDE
CORDRAN, FLURANDRENOLIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AQUA PHARMACEUTICALS**MONODOX, DOXYCYCLINE
VERDESO, DESONIDE*** AQUA PHARMACEUTICALS LLC**FLUOROPLEX, FLUOROURACIL
XOLEGEL, KETOCONAZOLE**AQUA PHARMS LLC***** AQUA PHARMACEUTICALS LLC**ACTICLATE, DOXYCYCLINE HYCLATE
ACZONE, DAPSONE
ALTABAX, RETAPAMULIN
AZELEX, AZELAIC ACID
CORDRAN, FLURANDRENOLIDE
VELTIN, CLINDAMYCIN PHOSPHATE**AQUESTIVE THERAP***** AQUESTIVE THERAPEUTICS**

SYMPAZAN, CLOBAZAM

ARALEZ PHARMS*** ARALEZ PHARMACEUTICALS TRADING DAC**TOPROL-XL, METOPROLOL SUCCINATE
ZONTIVITY, VORAPAXAR SULFATE**ARALEZ PHARMS INC***** ARALEZ PHARMACEUTICALS INC**

FIBRICOR, FENOFIBRIC ACID

ARBOR PHARMS LLC*** ARBOR PHARMACEUTICALS LLC**BIDIL, HYDRALAZINE HYDROCHLORIDE
CETYLEV, ACETYLCYSTEINE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
E.E.S. 400, ERYTHROMYCIN ETHYLSUCCINATE
E.E.S., ERYTHROMYCIN ETHYLSUCCINATE
EDARBI, AZILSARTAN KAMEDOXOMIL
EDARBYCLOR, AZILSARTAN KAMEDOXOMIL
ERY-TAB, ERYTHROMYCIN
ERYPED, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROCIN STEARATE, ERYTHROMYCIN STEARATE
ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROMYCIN, ERYTHROMYCIN
EVEKEO, AMPHETAMINE SULFATE
GLIADEL, CARMUSTINE
HORIZANT, GABAPENTIN ENACARBIL
NYMALIZE, NIMODIPINE
SKLICE, IVERMECTIN
SOTYLIZE, SOTALOL HYDROCHLORIDE
TRIPTODUR KIT, TRIPTORELIN PAMOATE**ARCO PHARMS LLC***** ARCO PHARMACEUTICALS LLC**

THYROSHIELD, POTASSIUM IODIDE (OTC)

AREVA PHARMS*** AREVA PHARMACEUTICALS INC**FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM**ARIAD***** ARIAD PHARMACEUTICALS INC**ALUNBRIG, BRIGATINIB
ICLUSIG, PONATINIB HYDROCHLORIDE**ARISE PHARMS***** ARISE PHARMACEUTICALS LLC**IBUPROFEN, IBUPROFEN (OTC)
LAMIVUDINE, LAMIVUDINE**ARMSTRONG PHARMS***** ARMSTRONG PHARMACEUTICALS INC**

PRIMATENE MIST, EPINEPHRINE (OTC)

ARRAY BIOPHARMA INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ARRAY BIOPHARMA INC

BRAFTOVI, ENCORAFENIB
MEKTOVI, BINIMETINIB**ASCEND THERAPS US**

* ASCEND THERAPEUTICS US LLC

ESTROGEL, ESTRADIOL

ASCENT PHARMS INC

* ASCENT PHARMACEUTICALS INC

DUTASTERIDE, DUTASTERIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
IBUPROFEN, IBUPROFEN (OTC)
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE**ASPEN GLOBAL**

* ASPEN GLOBAL INC

MYLERAN, BUSULFAN

ASPEN GLOBAL INC

* ASPEN GLOBAL INC

BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
CYCLESSA, DESOGESTREL
HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE
LEUKERAN, CHLORAMBUCIL
MYLERAN, BUSULFAN
THIOGUANINE, THIOGUANINE**ASSERTIO**

* ASSERTIO THERAPEUTICS INC

CAMBIA, DICLOFENAC POTASSIUM
GRALISE, GABAPENTIN
ZIPSOR, DICLOFENAC POTASSIUM**ASTELLAS**

* ASTELLAS PHARMA US INC

AMBISOME, AMPHOTERICIN B
ASTAGRAF XL, TACROLIMUS
CRESEMBA, ISAVUCONAZONIUM SULFATE
LEXISCAN, REGADENOSON
MYCAMINE, MICAFUNGIN SODIUM
PROGRAF, TACROLIMUS
VESICARE, SOLIFENACIN SUCCINATE
XOSPATA, GILTERITINIB FUMARATE
XTANDI, ENZALUTAMIDE**ASTRAL**

* ASTRAL STERITECH PVT LTD

AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
CEFTRIAZONE SODIUM, CEFTRIAZONE SODIUM
CEFTRIAZONE, CEFTRIAZONE SODIUM**ASTRAZENECA**

* ASTRAZENECA LP

PULMICORT FLEXHALER, BUDESONIDE
SYMBICORT, BUDESONIDE

* ASTRAZENECA PHARMACEUTICALS LP

FASLODEX, FULVESTRANT
ZOMIG, ZOLMITRIPTAN
ZOMIG-ZMT, ZOLMITRIPTAN

* ASTRAZENECA UK LTD

CALQUENCE, ACALABRUTINIB
SEROQUEL XR, QUETIAPINE FUMARATE**ASTRAZENECA AB**

* ASTRAZENECA AB

BYDUREON BCISE, EXENATIDE
BYDUREON PEN, EXENATIDE SYNTHETIC
BYDUREON, EXENATIDE SYNTHETIC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* ASTRAZENECA AB**

BYETTA, EXENATIDE SYNTHETIC
 FARXIGA, DAPAGLIFLOZIN
 KOMBIGLYZE XR, METFORMIN HYDROCHLORIDE
 ONGLYZA, SAXAGLIPTIN HYDROCHLORIDE
 QTERN, DAPAGLIFLOZIN
 SYMLIN, PRAMLINTIDE ACETATE
 XIGDUO XR, DAPAGLIFLOZIN

ASTRAZENECA LP

* ASTRAZENECA LP
 NEXIUM 24HR, ESOMEPRAZOLE MAGNESIUM (OTC)

ASTRAZENECA PHARMS

* ASTRAZENECA PHARMACEUTICALS LP
 ALVESCO, CICLESONIDE
 BEVESPI AEROSPHERE, FORMOTEROL FUMARATE
 BRILINTA, TICAGRELOR
 DALIRESP, ROFLUMILAST
 IRESSA, GEFITINIB
 LOKELMA, SODIUM ZIRCONIUM CYCLOSILICATE
 LYNPARZA, OLAPARIB
 MOVANTIK, NALOXEGOL OXALATE
 NEXIUM IV, ESOMEPRAZOLE SODIUM
 NEXIUM, ESOMEPRAZOLE MAGNESIUM
 OMNARIS, CICLESONIDE
 PRILOSEC OTC, OMEPRAZOLE MAGNESIUM (OTC)
 PULMICORT RESPULES, BUDESONIDE
 RHINOCORT ALLERGY, BUDESONIDE (OTC)
 SEROQUEL, QUETIAPINE FUMARATE
 TAGRISSO, OSIMERTINIB MESYLATE
 TUDORZA PRESSAIR, ACLIDINIUM BROMIDE
 ZETONNA, CICLESONIDE

ATHENEX INC

* ATHENEX INC
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE
 ENALAPRILAT, ENALAPRILAT
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 VALPROATE SODIUM, VALPROATE SODIUM

ATLAS PHARMS LLC

* ATLAS PHARMACEUTICALS LLC
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

ATNAHS PHARMA US

* ATNAHS PHARMA US LTD
 ANAPROX DS, NAPROXEN SODIUM
 EC-NAPROSYN, NAPROXEN
 NAPROSYN, NAPROXEN

ATON

* ATON PHARMA INC
 CUPRIMINE, PENICILLAMINE
 EDECRIN, ETHACRYNATE SODIUM
 EDECRIN, ETHACRYNIC ACID
 LACRISERT, HYDROXYPROPYL CELLULOSE
 LODOSYN, CARBIDOPA
 SYPRINE, TRIENTINE HYDROCHLORIDE
 TIMOPTIC IN OCUDOSE, TIMOLOL MALEATE
 TIMOPTIC, TIMOLOL MALEATE

ATON PHARMA VPNA

* ATON PHARMA DIV VALEANT PHARMACEUTICALS NORTH AMERICA LLC
 DEMSER, METYROSINE

AUCTA PHARMS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUCTA PHARMACEUTICALS LLC
VIGADRONE, VIGABATRIN

AUROBINDO

* AUROBINDO PHARMA LTD
AMOXICILLIN, AMOXICILLIN
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLARITHROMYCIN, CLARITHROMYCIN
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LISINOPRIL, LISINOPRIL
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
NEVIRAPINE, NEVIRAPINE
ZIDOVUDINE, ZIDOVUDINE

AUROBINDO PHARMA

* AUROBINDO PHARMA
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM

* AUROBINDO PHARMA LTD
ALENDRONATE SODIUM, ALENDRONATE SODIUM
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
AMPICILLIN SODIUM, AMPICILLIN SODIUM
ATENOLOL, ATENOLOL
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
CARISOPRODOL, CARISOPRODOL
CARVEDILOL, CARVEDILOL
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CEFDINIR, CEFDINIR
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
CEFPROZIL, CEFPROZIL
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
DIDANOSINE, DIDANOSINE
FINASTERIDE, FINASTERIDE
FLUCONAZOLE, FLUCONAZOLE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
GLYBURIDE, GLYBURIDE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LAMOTRIGINE, LAMOTRIGINE
LEVETIRACETAM, LEVETIRACETAM
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
MELOXICAM, MELOXICAM
METOPROLOL TARTRATE, METOPROLOL TARTRATE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON, ONDANSETRON
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
RIBAVIRIN, RIBAVIRIN
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SIMVASTATIN, SIMVASTATIN
STAVUDINE, STAVUDINE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
TOPIRAMATE, TOPIRAMATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * AUROBINDO PHARMA LTD
 TORSEMIDE, TORSEMIDE
 TRANDOLAPRIL, TRANDOLAPRIL
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZALEPLON, ZALEPLON
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
- AUROBINDO PHARMA LTD**
- * AUROBINDO PHARMA LIMITED
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
- * AUROBINDO PHARMA LTD
 ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
 ABACAVIR SULFATE, ABACAVIR SULFATE
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 ADENOSINE, ADENOSINE
 AFIRMELLE, ETHINYL ESTRADIOL
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 ALPRAZOLAM, ALPRAZOLAM
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ARMODAFINIL, ARMODAFINIL
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 ATHENTIA NEXT, LEVONORGESTREL (OTC)
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 AUROVELA 1.5/30, ETHINYL ESTRADIOL
 AUROVELA 1/20, ETHINYL ESTRADIOL
 AUROVELA 24 FE, ETHINYL ESTRADIOL
 AUROVELA FE 1.5/30, ETHINYL ESTRADIOL
 AUROVELA FE 1/20, ETHINYL ESTRADIOL
 AYUNA, ETHINYL ESTRADIOL
 AZITHROMYCIN, AZITHROMYCIN
 BIVALIRUDIN, BIVALIRUDIN
 BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CEFIXIME, CEFIXIME
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CEFPROZIL, CEFPROZIL
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CELECOXIB, CELECOXIB
 CEPHALEXIN, CEPHALEXIN
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CLOZAPINE, CLOZAPINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LTD
 CYONANZ, ETHINYL ESTRADIOL
 DALFAMPRIDINE, DALFAMPRIDINE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 EFAVIRENZ, EFAVIRENZ
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 ENTACAPONE, ENTACAPONE
 ENTECAVIR, ENTECAVIR
 EPTIFIBATIDE, EPTIFIBATIDE
 ERTAPENEM SODIUM, ERTAPENEM SODIUM
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ESZOPICLONE, ESZOPICLONE
 ETOMIDATE, ETOMIDATE
 EZETIMIBE, EZETIMIBE
 FAMCICLOVIR, FAMCICLOVIR
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FELODIPINE, FELODIPINE
 FENOFIBRATE, FENOFIBRATE
 FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
 FINASTERIDE, FINASTERIDE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUCONAZOLE, FLUCONAZOLE
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 GABAPENTIN, GABAPENTIN
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GEMFIBROZIL, GEMFIBROZIL
 GLIMEPIRIDE, GLIMEPIRIDE
 GLIPIZIDE, GLIPIZIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 ICLEVIA, ETHINYL ESTRADIOL
 INCASSIA, NORETHINDRONE
 INDOMETHACIN, INDOMETHACIN
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 ISOSULFAN BLUE, ISOSULFAN BLUE
 KALLIGA, DESOGESTREL
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUROBINDO PHARMA LTD
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LINEZOLID, LINEZOLID
 LO SIMPESE, ETHINYL ESTRADIOL
 LO-ZUMANDIMINE, DROSPIRENONE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE, LORATADINE (OTC)
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MEROPENEM, MEROPENEM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 METHOCARBAMOL, METHOCARBAMOL
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 METRONIDAZOLE, METRONIDAZOLE
 MILI, ETHINYL ESTRADIOL
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 MODAFINIL, MODAFINIL
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NADOLOL, NADOLOL
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NAPROXEN, NAPROXEN
 NEVIRAPINE, NEVIRAPINE
 NEXESTA FE, ETHINYL ESTRADIOL
 NIACIN, NIACIN
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 NYLIA 1/35, ETHINYL ESTRADIOL
 NYLIA 7/7/7, ETHINYL ESTRADIOL
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 OMEPRAZOLE, OMEPRAZOLE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXACILLIN SODIUM, OXACILLIN SODIUM
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARICALCITOL, PARICALCITOL
 PHENYTOIN SODIUM, PHENYTOIN SODIUM
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RAMIPRIL, RAMIPRIL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******* AUROBINDO PHARMA LTD**

REPAGLINIDE, REPAGLINIDE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RISPERIDONE, RISPERIDONE
 RITONAVIR, RITONAVIR
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 SIMPESE, ETHINYL ESTRADIOL
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRI-LO-MILI, ETHINYL ESTRADIOL
 TRI-MILI, ETHINYL ESTRADIOL
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VORICONAZOLE, VORICONAZOLE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID
 ZOLMITRIPTAN, ZOLMITRIPTAN
 ZUMANDIMINE, DROSPIRENONE

*** AUROBINDO PHARMA LTD INC**

ZIDOVUDINE, ZIDOVUDINE

AUROLIFE PHARMA LLC*** AUROLIFE PHARMA LLC**

ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DUTASTERIDE, DUTASTERIDE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 LORAZEPAM, LORAZEPAM
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

AUSTARPHARMA LLC*** AUSTARPHARMA LLC**

METHOCARBAMOL, METHOCARBAMOL
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE

AUXILIUM PHARMS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* AUXILIUM PHARMACEUTICALS INC
TESTOPEL, TESTOSTERONE
THEO-24, THEOPHYLLINE

AUXILIUM PHARMS LLC

* AUXILIUM PHARMACEUTICALS LLC
DILATRATE-SR, ISOSORBIDE DINITRATE
EDEX, ALPROSTADIL
ROBAXIN, METHOCARBAMOL
ROBAXIN-750, METHOCARBAMOL
SEMPREX-D, ACRIVASTINE
STRIANT, TESTOSTERONE
TESTIM, TESTOSTERONE
THEO-24, THEOPHYLLINE

AVACOR PRODS

* AVACOR PRODUCTS LLC
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)

AVADEL LEGACY

* AVADEL LEGACY PHARMACEUTICALS LLC
BLOXIVERZ, NEOSTIGMINE METHYLSULFATE
VAZCULEP, PHENYLEPHRINE HYDROCHLORIDE

AVADEL SPECILT

* AVADEL SPECIALTY PHARMACEUTICALS LLC
NOCTIVA, DESMOPRESSIN ACETATE

AVANIR PHARMS

* AVANIR PHARMACEUTICALS
ONZETRA XSAIL, SUMATRIPTAN SUCCINATE
* AVANIR PHARMACEUTICALS INC
NUEDEXTA, DEXTROMETHORPHAN HYDROBROMIDE

AVANTHI INC

* AVANTHI INC
CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE (OTC)
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE
INDOMETHACIN, INDOMETHACIN
LOMAIRA, PHENTERMINE HYDROCHLORIDE
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE

AVEDRO INC

* AVEDRO INC
PHOTREXA VISCOUS IN DEXTRAN 20%, RIBOFLAVIN 5'-PHOSPHATE SODIUM
PHOTREXA, RIBOFLAVIN 5'-PHOSPHATE SODIUM

AVEMA PHARMA

* AVEMA PHARMA SOLUTIONS
IBUPROFEN, IBUPROFEN (OTC)

AVENT

* AVENT INC
PYTEST KIT, UREA, C-14
PYTEST, UREA, C-14

AVERITAS

* AVERITAS PHARMA INC
QUTENZA, CAPSAICIN

AVEVA

* AVEVA DRUG DELIVERY SYSTEMS INC
CLONIDINE, CLONIDINE
FENTANYL-100, FENTANYL
FENTANYL-12, FENTANYL
FENTANYL-25, FENTANYL
FENTANYL-37, FENTANYL
FENTANYL-50, FENTANYL
FENTANYL-62, FENTANYL
FENTANYL-75, FENTANYL
FENTANYL-87, FENTANYL
NICOTINE, NICOTINE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******AVID RADIOPHARMS INC**

* AVID RADIOPHARMACEUTICALS INC
AMYVID, FLORBETAPIR F-18

AVION PHARMS

* AVION PHARMACEUTICALS LLC
BALCOLTRA, ETHINYL ESTRADIOL

AVONDALE PHARMS

* AVONDALE PHARMACEUTICALS LLC
NIACOR, NIACIN

AYTU

* AYTU BIOSCIENCE INC
NATESTO, TESTOSTERONE
TUZISTRA XR, CHLORPHENIRAMINE POLISTIREX

**** B ******B BRAUN**

* B BRAUN MEDICAL INC
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
AMINO ACIDS, AMINO ACIDS
BALANCED SALT, CALCIUM CHLORIDE
CEFAZOLIN AND DEXTROSE, CEFAZOLIN SODIUM
CEFEPIME AND DEXTROSE IN DUPLEX CONTAINER, CEFEPIME HYDROCHLORIDE
CEFOTETAN AND DEXTROSE IN DUPLEX CONTAINER, CEFOTETAN DISODIUM
CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER, CEFOXITIN SODIUM
CEFTAZIDIME IN DEXTROSE CONTAINER, CEFTAZIDIME
CEFTRIAOXONE AND DEXTROSE IN DUPLEX CONTAINER, CEFTRIAOXONE SODIUM
CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER, CEFUROXIME SODIUM
DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM
DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%, DOPAMINE HYDROCHLORIDE
FREAMINE HBC 6.9%, AMINO ACIDS
FREAMINE III 10%, AMINO ACIDS
FREAMINE III 3% W/ ELECTROLYTES, AMINO ACIDS
FREAMINE III 8.5% W/ ELECTROLYTES, AMINO ACIDS
FREAMINE III 8.5%, AMINO ACIDS
GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPATAMINE 8%, AMINO ACIDS
ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
ISOLYTE S IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
ISOLYTE S PH 7.4 IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* B BRAUN MEDICAL INC

LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL
METRO I.V. IN PLASTIC CONTAINER, METRONIDAZOLE
NEPHRAMINE 5.4%, AMINO ACIDS
NUTRILIPID 10%, SOYBEAN OIL
NUTRILIPID 20%, SOYBEAN OIL
PHYSIOLYTE IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******* B BRAUN MEDICAL INC**

POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROCALAMINE, AMINO ACIDS
 RESECTISOL IN PLASTIC CONTAINER, MANNITOL
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
 SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SORBITOL 3.3% IN PLASTIC CONTAINER, SORBITOL
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION
 THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 TROPHAMINE 10%, AMINO ACIDS
 TROPHAMINE, AMINO ACIDS

B BRAUN MEDICAL INC*** B BRAUN MEDICAL INC**

CLOROTEKAL, CHLOROPROCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER, MEROPENEM

BARR*** BARR LABORATORIES INC**

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 ARANELLE, ETHINYL ESTRADIOL
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 BALZIVA-28, ETHINYL ESTRADIOL
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 DANAZOL, DANAZOL
 DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIAZEPAM, DIAZEPAM
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 DUTASTERIDE, DUTASTERIDE
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 ESTRADIOL AND NORGESTIMATE, ESTRADIOL
 ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 HYDROXYUREA, HYDROXYUREA
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 ISONIAZID, ISONIAZID
 JUNEL 1.5/30, ETHINYL ESTRADIOL
 JUNEL 1/20, ETHINYL ESTRADIOL
 JUNEL FE 1.5/30, ETHINYL ESTRADIOL
 JUNEL FE 1/20, ETHINYL ESTRADIOL
 KARIVA, DESOGESTREL
 KELNOR, ETHINYL ESTRADIOL
 LEFLUNOMIDE, LEFLUNOMIDE
 LESSINA-28, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BARR LABORATORIES INC
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 NIACIN, NIACIN
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORTREL 0.5/35-28, ETHINYL ESTRADIOL
 NORTREL 1/35-21, ETHINYL ESTRADIOL
 NORTREL 1/35-28, ETHINYL ESTRADIOL
 NORTREL 7/7/7, ETHINYL ESTRADIOL
 ONDANSETRON, ONDANSETRON
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PORTIA-28, ETHINYL ESTRADIOL
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 SPRINTEC, ETHINYL ESTRADIOL
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TREXALL, METHOTREXATE SODIUM
 TRI-LEGEST 21, ETHINYL ESTRADIOL
 TRI-LEGEST FE, ETHINYL ESTRADIOL
 TRI-SPRINTEC, ETHINYL ESTRADIOL
 WARFARIN SODIUM, WARFARIN SODIUM
- * BARR PHARMACEUTICALS
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
- BARR LABS DIV TEVA**
- * BARR LABORATORIES INC SUB TEVA PHARMACEUTICALS USA
 ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
 BUDESONIDE, BUDESONIDE
- BARR LABS INC**
- * BARR LABORATORIES INC
 ACITRETIN, ACITRETIN
 CLOZAPINE, CLOZAPINE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 ESTRADIOL, ESTRADIOL
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 OLANZAPINE, OLANZAPINE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 OXYCODONE HYDROCHLORIDE AND IBUPROFEN, IBUPROFEN
 TRETINOIN, TRETINOIN
 TRI LO SPRINTEC, ETHINYL ESTRADIOL
- BAUSCH AND LOMB**
- * BAUSCH AND LOMB INC
 ALAWAY, KETOTIFEN FUMARATE (OTC)
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALREX, LOTEPIREDNOL ETABONATE
 BESIVANCE, BESIFLOXACIN HYDROCHLORIDE
 CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 FLURBIPROFEN SODIUM, FLURBIPROFEN SODIUM
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 ISTALOL, TIMOLOL MALEATE
 LATANOPROST, LATANOPROST
 LOTEMAX, LOTEPIREDNOL ETABONATE
 MIOCHOL-E, ACETYLCHOLINE CHLORIDE
 OFLOXACIN, OFLOXACIN
 OPCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
 PROLENSA, BROMFENAC SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BAUSCH AND LOMB INC
 RETISERT, FLUOCINOLONE ACETONIDE
 SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TROPICAMIDE, TROPICAMIDE
 VITRASE, HYALURONIDASE
 VYZULTA, LATANOPROSTENE BUNOD
 ZIRGAN, GANCICLOVIR
 ZYLET, LOTEPREDNOL ETABONATE
- * BAUSCH AND LOMB PHARMACEUTICALS INC
 BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXASPORIN, DEXAMETHASONE
 ERYTHROMYCIN, ERYTHROMYCIN
 FLUNISOLIDE, FLUNISOLIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
 NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
 NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
 NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE, BACITRACIN ZINC
 OFLOXACIN, OFLOXACIN
 OPTIPRANOLOL, METIPRANOLOL HYDROCHLORIDE
 OTICAIR, HYDROCORTISONE
 PENTOLAIR, CYCLOPENTOLATE HYDROCHLORIDE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TOBRAMYCIN AND DEXAMETHASONE, DEXAMETHASONE
 TOBRAMYCIN, TOBRAMYCIN
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 TROPICAMIDE, TROPICAMIDE

BAUSCH AND LOMB INC

- * BAUSCH AND LOMB INC
 BEPREVE, BEPOTASTINE BESILATE
 LOTE MAX, LOTEPREDNOL ETABONATE
 LUMIFY, BRIMONIDINE TARTRATE (OTC)

BAXTER HLTHCARE

- * BAXTER HEALTHCARE CORP
 ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
 AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER, GLYCINE
 ANCEF IN PLASTIC CONTAINER, CEFAZOLIN SODIUM
 BACTOCILL IN PLASTIC CONTAINER, OXACILLIN SODIUM
 BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 BREVIBLOC IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 BREVIBLOC, ESMOLOL HYDROCHLORIDE
 CARDIOPLEGIC IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 CEFEPIME IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE
 CEFTRIAOXONE IN PLASTIC CONTAINER, CEFTRIAOXONE SODIUM
 CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* BAXTER HEALTHCARE CORP

CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 2.75/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC
 CLINIMIX E 2.75/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC
 CLINIMIX E 2.75/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 4.25/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC
 CLINIMIX E 4.25/20 SULFITE FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC
 CLINIMIX E 4.25/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC
 CLINIMIX E 4.25/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/15 SULFITE FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/20 SULFITE FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINIMIX E 5/35 SULFITE FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER,
 CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND ELECTROLYTE NO. 48 IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER,
 DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC
 DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
 DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
 DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
 EXTRANEAL, ICODEXTRIN
 FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER, FAMOTIDINE
 FLAGYL I.V. RTU IN PLASTIC CONTAINER, METRONIDAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 FORANE, ISOFLURANE
 GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
 HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 IFEX, IFOSFAMIDE
 LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* BAXTER HEALTHCARE CORP

LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 MESNEX, MESNA
 MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 NALLPEN IN PLASTIC CONTAINER, NAFCILLIN SODIUM
 NEXTERONE, AMIODARONE HYDROCHLORIDE
 NITROGLYCERIN IN DEXTROSE 5%, NITROGLYCERIN
 OSMITROL 10% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 10% IN WATER, MANNITOL
 OSMITROL 15% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 15% IN WATER, MANNITOL
 OSMITROL 20% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 20% IN WATER, MANNITOL
 OSMITROL 5% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 5% IN WATER, MANNITOL
 PENICILLIN G POTASSIUM IN PLASTIC CONTAINER, PENICILLIN G POTASSIUM
 PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 PLASMA-LYTE A IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, POTASSIUM
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM
 POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 PREMASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
 PREMASOL 6% IN PLASTIC CONTAINER, AMINO ACIDS
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 SEVOFLURANE, SEVOFLURANE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SORBITOL 3% IN PLASTIC CONTAINER, SORBITOL
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION
 STERILE WATER, STERILE WATER FOR IRRIGATION
 SUPRANE, DESFLURANE
 TIS-U-SOL IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 TIS-U-SOL, MAGNESIUM SULFATE
 TRAVASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
 TRAVASOL 5.5% IN PLASTIC CONTAINER, AMINO ACIDS
 TRAVASOL 8.5% IN PLASTIC CONTAINER, AMINO ACIDS
 VANCOCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE

* BAXTER HEALTHCARE INTERNATIONAL SPECIALTY THERAPIES DIV

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BAXTER HEALTHCARE INTERNATIONAL SPECIALTY THERAPIES DIV
PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS

BAXTER HLTHCARE CORP

- * BAXTER HEALTHCARE CORP
BIVALIRUDIN IN 0.9% SODIUM CHLORIDE, BIVALIRUDIN
BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
CEFAZOLIN IN PLASTIC CONTAINER, CEFAZOLIN SODIUM
CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
CIPROFLOXACIN, CIPROFLOXACIN
CLINDAMYCIN PHOSPHATE IN 0.9% SODIUM CHLORIDE, CLINDAMYCIN PHOSPHATE
CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
FUROSEMIDE, FUROSEMIDE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
LEVOFLOXACIN, LEVOFLOXACIN
METOPROLOL TARTRATE, METOPROLOL TARTRATE
METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
PHOXILLUM B22K 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL B22GK 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 0/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 2/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 2/3.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 4/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PRISMASOL BK 0/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
- * BAXTER HEALTHCARE CORP ANESTHESIA AND CRITICAL CARE
PROTOPAM CHLORIDE, PRALIDOXIME CHLORIDE

BAYER

- * BAYER HEALTHCARE LLC
ALEVE, NAPROXEN SODIUM (OTC)
ALEVE-D SINUS & COLD, NAPROXEN SODIUM (OTC)

BAYER HEALTHCARE

- * BAYER HEALTHCARE PHARMACEUTICALS INC
ALIQOPA, COPANLISIB DIHYDROCHLORIDE

BAYER HEALTHCARE LLC

- * BAYER HEALTHCARE LLC
CHILDREN'S CLARITIN, LORATADINE (OTC)
CHLOR-TRIMETON, CHLORPHENIRAMINE MALEATE (OTC)
CLARITIN HIVES RELIEF REDITAB, LORATADINE (OTC)
CLARITIN HIVES RELIEF, LORATADINE (OTC)
CLARITIN REDITABS, LORATADINE (OTC)
CLARITIN, LORATADINE (OTC)
CLARITIN-D 24 HOUR, LORATADINE (OTC)
CLARITIN-D, LORATADINE (OTC)
LOTRIMIN ULTRA, BUTENAFINE HYDROCHLORIDE (OTC)
MIRALAX, POLYETHYLENE GLYCOL 3350 (OTC)
ZEGERID OTC, OMEPRAZOLE (OTC)

BAYER HLTHCARE

- * BAYER HEALTHCARE CONSUMER CARE
ALEVE PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
- * BAYER HEALTHCARE PHARMACEUTICALS INC
ADEMPAS, RIOCIQUAT
ANGELIQ, DROSPIRENONE
AVELOX, MOXIFLOXACIN HYDROCHLORIDE
BEYAZ, DROSPIRENONE
BILTRICIDE, PRAZIQUANTEL
CIPRO, CIPROFLOXACIN
CIPRO, CIPROFLOXACIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******* BAYER HEALTHCARE PHARMACEUTICALS INC**

CLIMARA PRO, ESTRADIOL
 CLIMARA, ESTRADIOL
 EOVIIST, GADOXETATE DISODIUM
 GADAVIST, GADOBUTROL
 KYLEENA, LEVONORGESTREL
 LEVITRA, VARDENAFIL HYDROCHLORIDE
 MAGNEVIST, GADOPENTETATE DIMEGLUMINE
 MENOSTAR, ESTRADIOL
 MIRENA, LEVONORGESTREL
 NATAZIA, DIENOGEST
 NEXAVAR, SORAFENIB TOSYLATE
 PRECOSE, ACARBOSE
 SAFYRAL, DROSPIRENONE
 SKYLA, LEVONORGESTREL
 STAXYN, VARDENAFIL HYDROCHLORIDE
 STIVARGA, REGORAFENIB
 ULTRAVIST (PHARMACY BULK), IOPROMIDE
 ULTRAVIST 240, IOPROMIDE
 ULTRAVIST 300, IOPROMIDE
 ULTRAVIST 370, IOPROMIDE
 XOFIGO, RADIUM RA-223 DICHLORIDE
 YASMIN, DROSPIRENONE
 YAZ, DROSPIRENONE

BAYSHORE PHARMS LLC*** BAYSHORE PHARMACEUTICALS LLC**

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
 PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE

BDSI*** BIODELIVERY SCIENCES INTERNATIONAL INC**

BELBUCA, BUPRENORPHINE HYDROCHLORIDE
 BUNAVAIL, BUPRENORPHINE HYDROCHLORIDE

BECTON DICKINSON*** BECTON DICKINSON AND CO**

CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)
 E-Z SCRUB 201, POVIDONE-IODINE (OTC)
 E-Z SCRUB 241, POVIDONE-IODINE (OTC)

BECTON DICKINSON CO*** BECTON DICKINSON AND CO**

CHLORAPREP ONE-STEP FREPP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP ONE-STEP SEPP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP ONE-STEP, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP SINGLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP TRIPLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 CHLORAPREP WITH TINT, CHLORHEXIDINE GLUCONATE (OTC)

BEIJING YILING*** BEIJING YILING BIO-ENGINEERING AND TECHNOLOGY CO LTD**

ANASTROZOLE, ANASTROZOLE
 LETROZOLE, LETROZOLE

BELCHER PHARMS*** BELCHER PHARMACEUTICALS LLC**

CEPHALEXIN, CEPHALEXIN
 DESLORATADINE, DESLORATADINE

BELCHER PHARMS LLC*** BELCHER PHARMACEUTICALS LLC**

ABLYSINOL, ALCOHOL
 CEFIXIME, CEFIXIME
 EPINEPHRINE, EPINEPHRINE
 MEFENAMIC ACID, MEFENAMIC ACID
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 TACROLIMUS, TACROLIMUS

BELOTECA INC*** BELOTECA INC**

ISOSULFAN BLUE, ISOSULFAN BLUE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******BEXIMCO PHARMS USA**

* BEXIMCO PHARMACEUTICALS USA INC
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHOCARBAMOL, METHOCARBAMOL
 NADOLOL, NADOLOL
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

BEXIMCO USA

* BEXIMCO PHARMACEUTICALS USA INC
 CARVEDILOL, CARVEDILOL

BI-COASTAL PHARMA

* BI-COASTAL PHARMA INTERNATIONAL LLC
 DUVOID, BETHANECHOL CHLORIDE

BIO NUCLEONICS

* BIO NUCLEONICS INC
 STRONTIUM CHLORIDE SR-89, STRONTIUM CHLORIDE SR-89

BIO PHARM INC

* BIO PHARM INC
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE

BIO-PHARM INC

* BIO-PHARM INC
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 LACTULOSE, LACTULOSE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE

BIOCODEX SA

* BIOCODEX SA
 DIACOMIT, STIRIPENTOL

BIOCON LIMITED

* BIOCON LIMITED
 SIMVASTATIN, SIMVASTATIN

BIOCON LTD

* BIOCON LTD
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

BIOCRYST

* BIOCRYST PHARMACEUTICALS INC
 RAPIVAB, PERAMIVIR

BIOFRONTERA

* BIOFRONTERA BIOSCIENCE GMBH
 AMELUZ, AMINOLEVULINIC ACID HYDROCHLORIDE

BIOGEN IDEC

* BIOGEN IDEC INC
 SPINRAZA, NUSINERSEN SODIUM

BIOGEN IDEC INC

* BIOGEN IDEC INC
 TECFIDERA, DIMETHYL FUMARATE

BIOMARIN PHARM

* BIOMARIN PHARMACEUTICAL INC
 KUVAN, SAPROPTERIN DIHYDROCHLORIDE

BIOMEDCL RES FDN

* BIOMEDICAL RESEARCH FOUNDATION NORTHWEST LOUISIANA
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

BIONPHARMA INC

* BIONPHARMA INC
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AZITHROMYCIN, AZITHROMYCIN
 BENZONATATE, BENZONATATE
 BEXAROTENE, BEXAROTENE
 CALCITRIOL, CALCITRIOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******* BIONPHARMA INC**

CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CLOBAZAM, CLOBAZAM
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DOFETILIDE, DOFETILIDE
 DUTASTERIDE, DUTASTERIDE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE, LORATADINE (OTC)
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MIDOL LIQUID GELS, IBUPROFEN (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NIMODIPINE, NIMODIPINE
 PARICALCITOL, PARICALCITOL
 PROGESTERONE, PROGESTERONE
 TETRABENAZINE, TETRABENAZINE
 VALPROIC ACID, VALPROIC ACID
 VITAMIN D, ERGOCALCIFEROL

BLAIREX*** BLAIREX LABORATORIES INC**

BRONCHO SALINE, SODIUM CHLORIDE (OTC)

BLUE EARTH*** BLUE EARTH DIAGNOSTICS LTD**

AXUMIN, FLUCICLOVINE F-18

BLUEPHARMA*** BLUEPHARMA US INC**

GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE, GRANISETRON HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 ZONISAMIDE, ZONISAMIDE

BOEHRINGER INGELHEIM*** BOEHRINGER INGELHEIM**

CATAPRES, CLONIDINE HYDROCHLORIDE
 CATAPRES-TTS-1, CLONIDINE
 CATAPRES-TTS-2, CLONIDINE
 CATAPRES-TTS-3, CLONIDINE
 GILOTRIF, AFATINIB DIMALEATE
 GLYXAMBI, EMPAGLIFLOZIN
 MICARDIS HCT, HYDROCHLOROTHIAZIDE
 MICARDIS, TELMISARTAN
 MIRAPEX, PRAMIPEXOLE DIHYDROCHLORIDE

*** BOEHRINGER INGELHEIM PHARMACEUTICALS INC**

AGGRENOX, ASPIRIN
 APTIVUS, TIPRANAVIR
 ATROVENT HFA, IPRATROPIUM BROMIDE
 COMBIVENT RESPIMAT, ALBUTEROL SULFATE
 JARDIANCE, EMPAGLIFLOZIN
 JENTADUETO XR, LINAGLIPTIN
 JENTADUETO, LINAGLIPTIN
 MIRAPEX ER, PRAMIPEXOLE DIHYDROCHLORIDE
 MOBIC, MELOXICAM
 OFEV, NINTEDANIB ESYLATE
 PERSANTINE, DIPYRIDAMOLE
 PRADAXA, DABIGATRAN ETEXILATE MESYLATE
 SPIRIVA RESPIMAT, TIOTROPIUM BROMIDE
 SPIRIVA, TIOTROPIUM BROMIDE
 STIOLTO RESPIMAT, OLODATEROL HYDROCHLORIDE
 STRIVERDI RESPIMAT, OLODATEROL HYDROCHLORIDE
 SYNJARDY XR, EMPAGLIFLOZIN
 SYNJARDY, EMPAGLIFLOZIN
 TRADJENTA, LINAGLIPTIN
 TWYNSTA, AMLODIPINE BESYLATE
 VIRAMUNE XR, NEVIRAPINE
 VIRAMUNE, NEVIRAPINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******BOSCOGEN**

* BOSCOGEN INC
 ACYCLOVIR, ACYCLOVIR
 ANASTROZOLE, ANASTROZOLE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 BICALUTAMIDE, BICALUTAMIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 REPAGLINIDE, REPAGLINIDE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE

BPI LABS LLC

* BPI LABS LLC
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

BRACCO

* BRACCO DIAGNOSTICS INC
 CARDIOGEN-82, RUBIDIUM CHLORIDE RB-82
 CHOLETEC, TECHNETIUM TC-99M MEBROFENIN KIT
 CYSTOGRAFIN DILUTE, DIATRIZOATE MEGLUMINE
 CYSTOGRAFIN, DIATRIZOATE MEGLUMINE
 E-Z-HD, BARIUM SULFATE
 E-Z-PAQUE, BARIUM SULFATE
 GASTROGRAFIN, DIATRIZOATE MEGLUMINE
 ISOVUE-200, IOPAMIDOL
 ISOVUE-250, IOPAMIDOL
 ISOVUE-300, IOPAMIDOL
 ISOVUE-370, IOPAMIDOL
 ISOVUE-M 200, IOPAMIDOL
 ISOVUE-M 300, IOPAMIDOL
 KINEVAC, SINCALIDE
 LIQUID E-Z-PAQUE, BARIUM SULFATE
 LUMASON, SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES
 MULTIHANCE MULTIPACK, GADOBENATE DIMEGLUMINE
 MULTIHANCE, GADOBENATE DIMEGLUMINE
 PROHANCE MULTIPACK, GADOTERIDOL
 PROHANCE, GADOTERIDOL
 READI-CAT 2 SMOOTHIES, BARIUM SULFATE
 READI-CAT 2, BARIUM SULFATE
 TAGITOL V, BARIUM SULFATE
 VARIBAR HONEY, BARIUM SULFATE
 VARIBAR NECTAR, BARIUM SULFATE
 VARIBAR PUDDING, BARIUM SULFATE
 VARIBAR THIN HONEY, BARIUM SULFATE

BRAINTREE

* BRAINTREE LABORATORIES INC
 GOLYTELY, POLYETHYLENE GLYCOL 3350
 NULYTELY, POLYETHYLENE GLYCOL 3350
 NULYTELY-FLAVORED, POLYETHYLENE GLYCOL 3350

BRAINTREE LABS

* BRAINTREE LABORATORIES INC
 SUPREP BOWEL PREP KIT, MAGNESIUM SULFATE

BRECKENRIDGE PHARM

* BRECKENRIDGE PHARMACEUTICAL INC
 ALPRAZOLAM, ALPRAZOLAM
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CILOSTAZOL, CILOSTAZOL
 CLOBAZAM, CLOBAZAM
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUTASTERIDE, DUTASTERIDE
 ENTECAVIR, ENTECAVIR
 EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 EPLERENONE, EPLERENONE
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 LANSOPRAZOLE, LANSOPRAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BRECKENRIDGE PHARMACEUTICAL INC
 LEVETIRACETAM, LEVETIRACETAM
 MEFENAMIC ACID, MEFENAMIC ACID
 MEGESTROL ACETATE, MEGESTROL ACETATE
 MELOXICAM, MELOXICAM
 METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 NEOMYCIN SULFATE, NEOMYCIN SULFATE
 OMEPRAZOLE, OMEPRAZOLE
 OXCARBAZEPINE, OXCARBAZEPINE
 PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PIROXICAM, PIROXICAM
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 ROFLUMILAST, ROFLUMILAST
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

BRIGHAM WOMENS

* BRIGHAM AND WOMENS HOSP
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

BRIGHAM WOMENS HOSP

* BRIGHAM AND WOMENS HOSP INC
 AMMONIA N 13, AMMONIA N-13

BRISTOL MYERS SQUIBB

* BRISTOL MYERS SQUIBB
 AZACTAM, AZTREONAM
 BARACLUDE, ENTECAVIR
 PRAVACHOL, PRAVASTATIN SODIUM

* BRISTOL MYERS SQUIBB CO
 AZACTAM IN PLASTIC CONTAINER, AZTREONAM
 DROXIA, HYDROXYUREA
 GLUCOPHAGE XR, METFORMIN HYDROCHLORIDE
 HYDREA, HYDROXYUREA
 REYATAZ, ATAZANAVIR SULFATE
 SPRYCEL, DASATINIB
 SUSTIVA, EFAVIRENZ
 VIDEX EC, DIDANOSINE

* BRISTOL MYERS SQUIBB CO PHARMACEUTICAL RESEARCH INSTITUTE
 ELIQUIS, APIXABAN
 ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE
 GLUCOPHAGE, METFORMIN HYDROCHLORIDE
 ZERIT, STAVUDINE

* BRISTOL MYERS SQUIBB PHARMA CO
 COUMADIN, WARFARIN SODIUM

BRISTOL-MYERS SQUIBB

* BRISTOL-MYERS SQUIBB CO
 DAKLINZA, DACLATASVIR DIHYDROCHLORIDE
 EVOTAZ, ATAZANAVIR SULFATE
 VIDEX, DIDANOSINE
 ZERIT, STAVUDINE

**** C ******CADILA PHARMS LTD**

* CADILA PHARMACEUTICALS LTD
 ACYCLOVIR, ACYCLOVIR
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 GEMFIBROZIL, GEMFIBROZIL
 GLYBURIDE, GLYBURIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

- * CADILA PHARMACEUTICALS LTD
 OFLOXACIN, OFLOXACIN
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 TELMISARTAN, TELMISARTAN
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
- CADISTA PHARMS**
- * CADISTA PHARMACEUTICALS INC
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
- CALL INC**
- * CALL INC DBA ROCHESTER PHARMACEUTICALS
 ADAPALENE, ADAPALENE
- CAPELLON PHARMS LLC**
- * CAPELLON PHARMACEUTICALS LLC
 POLMON, DEXCHLORPHENIRAMINE MALEATE
- CARDINAL HEALTH 414**
- * CARDINAL HEALTH 414 LLC CARDINAL HEALTH NUCLEAR PHARMACY SERVICES
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 LYMPHOSEEK KIT, TECHNETIUM TC-99M TILMANOCEPT
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18
 TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
- CARDINAL HEALTH 418**
- * CARDINAL HEALTH 418 INC
 SODIUM IODIDE I 123, SODIUM IODIDE I-123
- CARIBE HOLDINGS**
- * CARIBE HOLDINGS CAYMAN CO LTD DBA PURACAP CARIBE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 GEMFIBROZIL, GEMFIBROZIL
- CARLSBAD**
- * CARLSBAD TECHNOLOGY INC
 ACYCLOVIR, ACYCLOVIR
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 FAMOTIDINE, FAMOTIDINE
 GLIMEPIRIDE, GLIMEPIRIDE
 LOVASTATIN, LOVASTATIN
- CARLSBAD TECHNOLOGY**
- * CARLSBAD TECHNOLOGY INC
 ACYCLOVIR, ACYCLOVIR
- CASI PHARMS INC**
- * CASI PHARMACEUTICALS INC
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CEFPROZIL, CEFPROZIL
 CILOSTAZOL, CILOSTAZOL
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 ECONAZOLE NITRATE, ECONAZOLE NITRATE
 ENTECAVIR, ENTECAVIR
 EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 HEPARIN SODIUM, HEPARIN SODIUM
 LISINOPRIL, LISINOPRIL
 METHIMAZOLE, METHIMAZOLE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 NABUMETONE, NABUMETONE
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 REPAGLINIDE, REPAGLINIDE
 RIBAVIRIN, RIBAVIRIN
 SPIRONOLACTONE, SPIRONOLACTONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CASI PHARMACEUTICALS INC
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

CASPER PHARMA LLC

* CASPER PHARMA LLC
 AQUASOL A, VITAMIN A PALMITATE
 CASPORYN HC, HYDROCORTISONE
 FURADANTIN, NITROFURANTOIN
 NEOSPORIN, BACITRACIN ZINC
 ZYLOPRIM, ALLOPURINOL

CATALENT

* CATALENT PHARMA SOLUTIONS LLC
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 VALPROIC ACID, VALPROIC ACID

CATALYST PHARMS

* CATALYST PHARMACEUTICALS INC
 FIRDAPSE, AMIFAMPRIDINE PHOSPHATE

CE DIPROF INC

* CE DIPROF INC
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 LEVO-T, LEVOTHYROXINE SODIUM **

CELATOR PHARMS

* CELATOR PHARMACEUTICALS INC
 VYXEOS, CYTARABINE

CELERITY PHARMS

* CELERITY PHARMACEUTICALS LLC
 EPTIFIBATIDE, EPTIFIBATIDE

CELGENE

* CELGENE CORP
 ISTODAX, ROMIDEPSIN
 POMALYST, POMALIDOMIDE
 REVLIMID, LENALIDOMIDE
 THALOMID, THALIDOMIDE
 VIDAZA, AZACITIDINE

CELGENE CORP

* CELGENE CORP
 IDHIFA, ENASIDENIB MESYLATE
 OTEZLA, APREMILAST

CELLTRION

* CELLTRION INC
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 TEMIXYS, LAMIVUDINE

CEPHALON

* CEPHALON INC
 ACTIQ, FENTANYL CITRATE
 FENTORA, FENTANYL CITRATE
 GABITRIL, TIAGABINE HYDROCHLORIDE
 NUVIGIL, ARMODAFINIL
 PROVIGIL, MODAFINIL
 TREANDA, BENDAMUSTINE HYDROCHLORIDE
 TRISENOX, ARSENIC TRIOXIDE

CERECOR INC

* CERECOR INC
 ACIPHEX SPRINKLE, RABEPRAZOLE SODIUM

CEYONE

* CEYONE PHARMA LLC
 RISPERIDONE, RISPERIDONE

CHANGZHOU PHARM

* CHANGZHOU PHARMACEUTICAL FACTORY
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

CHARTWELL LIFE SCI

* CHARTWELL LIFE SCIENCE LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CHARTWELL LIFE SCIENCE LLC
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DOXYCYCLINE, DOXYCYCLINE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

CHARTWELL MOLECULAR

* CHARTWELL MOLECULAR HOLDINGS LLC
CARVEDILOL, CARVEDILOL
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
FOLIC ACID, FOLIC ACID
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
IRBESARTAN, IRBESARTAN
RAMIPRIL, RAMIPRIL
RISPERIDONE, RISPERIDONE

CHARTWELL MOLECULES

* CHARTWELL MOLECULES LLC
DISULFIRAM, DISULFIRAM
GEMFIBROZIL, GEMFIBROZIL
NABUMETONE, NABUMETONE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

CHARTWELL RX

* CHARTWELL RX SCIENCES LLC
CALCIUM ACETATE, CALCIUM ACETATE
CILOSTAZOL, CILOSTAZOL
GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
INDOMETHACIN, INDOMETHACIN
LEVETIRACETAM, LEVETIRACETAM
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE

CHARTWELL TETRA

* CHARTWELL TETRA LLC
TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

CHATTEM

* CHATTEM INC
UNISOM, DOXYLAMINE SUCCINATE (OTC)

CHEMI SPA

* CHEMI SPA
DECITABINE, DECITABINE
TEMOZOLOMIDE, TEMOZOLOMIDE

CHEMISCH FBRK KRSSLR

* CHEMISCHE FABRIK KREUSSLER & CO. GMBH
ASCLERA, POLIDOCANOL

CHEMO RESEARCH SL

* CHEMO RESEARCH SL
BENZNIDAZOLE, BENZNIDAZOLE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
NUVESSA, METRONIDAZOLE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN

CHEPLAPHARM

* CHEPLAPHARM ARZNEIMITTEL GMBH
XENICAL, ORLISTAT

CHIESI USA INC

* CHIESI USA INC
BETHKIS, TOBRAMYCIN
CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE
CLEVIPREX, CLEVIDIPINE
CUROSURF, PORACTANT ALFA
KENGREAL, CANGRELOR
ZYFLO CR, ZILEUTON
ZYFLO, ZILEUTON

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******CHILDRENS HOSP MI**

* CHILDRENS HOSP MICHIGAN
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

CHINA RESOURCES

* CHINA RESOURCES SAIKE PHARMACEUTICAL CO LTD
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE

CHIRHOCLIN

* CHIRHOCLIN INC
CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN

CINTEX SVCS

* CINTEX SERVICES LLC
FLURANDRENOLIDE, FLURANDRENOLIDE

CIPHER PHARMS INC

* CIPHER PHARMACEUTICALS INC
CONZIP, TRAMADOL HYDROCHLORIDE
LIPOFEN, FENOFIBRATE

CIPLA

* CIPLA LTD
ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
ABACAVIR SULFATE, ABACAVIR SULFATE
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
ANASTROZOLE, ANASTROZOLE
ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
AZACITIDINE, AZACITIDINE
BIVALIRUDIN, BIVALIRUDIN
BUDESONIDE, BUDESONIDE
CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
CELECOXIB, CELECOXIB
CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
DECITABINE, DECITABINE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DOCETAXEL, DOCETAXEL
EFAVIRENZ, EFAVIRENZ
EMTRICITABINE, EMTRICITABINE
ENTECAVIR, ENTECAVIR
EXEMESTANE, EXEMESTANE
FAMCICLOVIR, FAMCICLOVIR
FENOFIBRATE, FENOFIBRATE
FINASTERIDE, FINASTERIDE
FLUTAMIDE, FLUTAMIDE
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
LAMIVUDINE, LAMIVUDINE
LAMOTRIGINE, LAMOTRIGINE
LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
MELOXICAM, MELOXICAM
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
MONTELUKAST SODIUM, MONTELUKAST SODIUM
NEVIRAPINE, NEVIRAPINE
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
OXALIPLATIN, OXALIPLATIN
PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
TENOFVIR DISOPROXIL FUMARATE, TENOFVIR DISOPROXIL FUMARATE
TERBINAFFINE HYDROCHLORIDE, TERBINAFFINE HYDROCHLORIDE
TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******* CIPLA LTD**

TESTOSTERONE, TESTOSTERONE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 ZIDOVUDINE, ZIDOVUDINE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

CIPLA LTD*** CIPLA LTD**

ALBENDAZOLE, ALBENDAZOLE
 CARBOPLATIN, CARBOPLATIN
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 TOPIRAMATE, TOPIRAMATE
 ZALEPLON, ZALEPLON
 ZIDOVUDINE, ZIDOVUDINE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

CLINIGEN HLTHCARE*** CLINIGEN HEALTHCARE LTD**

ETHYOL, AMIFOSTINE
 FOSCAVIR, FOSCARNET SODIUM
 TOTECT, DEXRAZOXANE HYDROCHLORIDE

CLOVER PHARMS*** CLOVER PHARMACEUTICALS CORP**

AMICAR, AMINOCAPROIC ACID

CLOVIS ONCOLOGY INC*** CLOVIS ONCOLOGY INC**

RUBRACA, RUCAPARIB CAMSYLATE

CMP DEV LLC*** CMP DEVELOPMENT LLC**

CAROSPIR, SPIRONOLACTONE

CMP PHARMA INC*** CMP PHARMA INC**

AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 HYDROCORTISONE IN ABSORBASE, HYDROCORTISONE
 ISONIAZID, ISONIAZID
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 SPS, SODIUM POLYSTYRENE SULFONATE
 TRIAMCINOLONE ACETONIDE IN ABSORBASE, TRIAMCINOLONE ACETONIDE

CNTY LINE PHARMS*** COUNTY LINE PHARMACEUTICALS LLC**

BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CICLOPIROX, CICLOPIROX
 DYNACIN, MINOCYCLINE HYDROCHLORIDE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRATE, FENOFIBRATE
 FLUOCINONIDE, FLUOCINONIDE
 LIDEX, FLUOCINONIDE
 LIDEX-E, FLUOCINONIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 TAMBOCOR, FLECAINIDE ACETATE
 TRANDATE, LABETALOL HYDROCHLORIDE
 TRANLYCYPROMINE SULFATE, TRANLYCYPROMINE SULFATE
 UREX, METHENAMINE HIPPURATE

COLGATE PALMOLIVE*** COLGATE PALMOLIVE**

COLGATE TOTAL, SODIUM FLUORIDE (OTC)

COLGATE PALMOLIVE CO*** COLGATE PALMOLIVE CO**

PERIOGARD, CHLORHEXIDINE GLUCONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******COLGATE-PALMOLIVE CO**

* COLGATE-PALMOLIVE CO
PERIOGARD, CHLORHEXIDINE GLUCONATE

COLLEGIUM PHARM INC

* COLLEGIUM PHARMACEUTICAL INC
XTAMPZA ER, OXYCODONE

COMBE

* COMBE INC
VAGISTAT-1, TIOCONAZOLE (OTC)

CONCORD BIOTECH LTD

* CONCORD BIOTECH LTD
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL

CONCORDIA LABS INC

* CONCORDIA LABORATORIES INC
PHOTOFRIN, PORFIMER SODIUM

CONCORDIA PHARMS INC

* CONCORDIA PHARMACEUTICALS INC
DIBENZYLINE, PHENOXYBENZAMINE HYDROCHLORIDE
DUTOPROL, HYDROCHLOROTHIAZIDE
DYRENIUM, TRIAMTERENE
KAPVAY, CLONIDINE HYDROCHLORIDE
LANOXIN, DIGOXIN
NILANDRON, NILUTAMIDE
ORAPRED ODT, PREDNISOLONE SODIUM PHOSPHATE
PARNATE, TRANLYCYPROMINE SULFATE
PLAQUENIL, HYDROXYCHLOROQUINE SULFATE
UROXATRAL, ALFUZOSIN HYDROCHLORIDE

CONTRACT PHARMACAL

* CONTRACT PHARMACAL CORP
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

COOPERSURGICAL

* COOPERSURGICAL INC
PARAGARD T 380A, COPPER

CORCEPT THERAP

* CORCEPT THERAPEUTICS INC
KORLYM, MIFEPRISTONE

CORDEN PHARMA

* CORDEN PHARMA LATINA SPA
GLEOSTINE, LOMUSTINE

COSMO TECHNOLOGIES

* COSMO TECHNOLOGIES LTD
AEMCOLO, RIFAMYCIN

COVIS PHARMA BV

* COVIS PHARMA BV
ALTOPREV, LOVASTATIN
BETAPACE AF, SOTALOL HYDROCHLORIDE
BETAPACE, SOTALOL HYDROCHLORIDE
LANOXIN PEDIATRIC, DIGOXIN
LANOXIN, DIGOXIN
PRILOSEC, OMEPRAZOLE MAGNESIUM
RILUTEK, RILUZOLE
SULAR, NISOLDIPINE
ZANAFLEX, TIZANIDINE HYDROCHLORIDE

CPDC

* CENTRE FOR PROBE DEVELOPMENT AND COMMERCIALIZATION
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

CPPI CV

* CP PHARMACEUTICALS INTERNATIONAL CV
SUTENT, SUNITINIB MALATE

CRANE PHARMS LLC

* CRANE PHARMACEUTICALS LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CRANE PHARMACEUTICALS LLC
DAPTOMYCIN, DAPTOMYCIN

CROSSMEDIKA SA

* CROSSMEDIKA SA
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE
VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE

CROWN LABS

* CROWN LABORATORIES INC
ALA-CORT, HYDROCORTISONE
ALA-SCALP, HYDROCORTISONE
TRIDERM, TRIAMCINOLONE ACETONIDE

CROWN LABS INC

* CROWN LABORATORIES INC
NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
NYSTATIN, NYSTATIN

CSPC NBP PHARM CO

* CSPC NBP PHARMACEUTICAL CO LTD
BENZONATATE, BENZONATATE

CSPC OUYI PHARM CO

* CSPC OUYI PHARMACEUTICAL CO LTD
AZITHROMYCIN, AZITHROMYCIN
CELECOXIB, CELECOXIB
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
GABAPENTIN, GABAPENTIN
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
MONTELUKAST SODIUM, MONTELUKAST SODIUM
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

CUBIST PHARMS

* CUBIST PHARMACEUTICALS INC
ENTEREG, ALVIMOPAN

CUBIST PHARMS LLC

* CUBIST PHARMACEUTICALS LLC
CUBICIN RF, DAPTOMYCIN
CUBICIN, DAPTOMYCIN
DIFICID, FIDAXOMICIN
SIVEXTRO, TEDIZOLID PHOSPHATE
ZERBAXA, CEFTOLOZANE SULFATE

CUMBERLAND PHARMS

* CUMBERLAND PHARMACEUTICALS INC
ACETADOTE, ACETYLCYSTEINE
CALDOLOR, IBUPROFEN
LACTULOSE, LACTULOSE
OMEPRAZOLE AND CLARITHROMYCIN AND AMOXICILLIN, AMOXICILLIN
VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER, CONIVAPTAN HYDROCHLORIDE
VIBATIV, TELAVANCIN HYDROCHLORIDE

CUSTOPHARM INC

* CUSTOPHARM INC
ACETAMINOPHEN, ACETAMINOPHEN
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM

CUTANEA

* CUTANEA LIFE SCIENCES INC
AKTIPAK, BENZOYL PEROXIDE

CYCLE PHARMS LTD

* CYCLE PHARMACEUTICALS LTD
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
NITYR, NITISINONE

CYPRESS PHARM

* CYPRESS PHARMACEUTICAL INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

* CYPRESS PHARMACEUTICAL INC
 REZIRA, HYDROCODONE BITARTRATE
 VITUZ, CHLORPHENIRAMINE MALEATE
 ZUTRIPRO, CHLORPHENIRAMINE MALEATE

**** D ******DAEWOONG PHARM CO**

* DAEWOONG PHARMACEUTICAL CO LTD
 MEROPENEM, MEROPENEM

DAIICHI SANKYO

* DAIICHI SANKYO INC
 AZOR, AMLODIPINE BESYLATE
 BENICAR HCT, HYDROCHLOROTHIAZIDE
 BENICAR, OLMESARTAN MEDOXOMIL
 TRIBENZOR, AMLODIPINE BESYLATE
 WELCHOL, COLESEVELAM HYDROCHLORIDE

DAIICHI SANKYO INC

* DAIICHI SANKYO INC
 EVOXAC, CEVIMELINE HYDROCHLORIDE
 MORPHABOND ER, MORPHINE SULFATE
 SAVAYSA, EDOXABAN TOSYLATE

DAITO PHARMS CO LTD

* DAITO PHARMACEUTICALS CO LTD
 RILUZOLE, RILUZOLE

DANCO LABS LLC

* DANCO LABORATORIES LLC
 MIFEPREX, MIFEPRISTONE

DAVA INTL INC

* DAVA INTERNATIONAL INC
 ALPRAZOLAM, ALPRAZOLAM

DAVA PHARMS INC

* DAVA PHARMACEUTICALS INC
 ACYCLOVIR, ACYCLOVIR
 AMOXICILLIN, AMOXICILLIN
 AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE
 ATENOLOL, ATENOLOL
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 GLYBURIDE (MICRONIZED), GLYBURIDE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 MORPHINE SULFATE, MORPHINE SULFATE
 PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 PROPYLTHIOURACIL, PROPYLTHIOURACIL
 PYRAZINAMIDE, PYRAZINAMIDE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 VOSPIRE ER, ALBUTEROL SULFATE

DAVIS AND GECK

* DAVIS AND GECK DIV AMERICAN CYANAMID CO
 PRE-OP II, HEXACHLOROPHENE
 PRE-OP, HEXACHLOROPHENE

DBL PHARMS

* DBL PHARMACEUTICALS INC
 METHOCARBAMOL, METHOCARBAMOL

DENTSPLY PHARM

* DENTSPLY PHARMACEUTICAL INC
 CITANEST FORTE DENTAL, EPINEPHRINE BITARTRATE
 ORAQIX, LIDOCAINE

DEPO NF

* DEPO NF SUB LLC A SUB OF ASSERTIO THERAPEUTICS INC
 NUCYN TA ER, TAPENTADOL HYDROCHLORIDE
 NUCYN TA, TAPENTADOL HYDROCHLORIDE

DEPROCO

* DEPROCO INC
 LIGNOSPAN FORTE, EPINEPHRINE BITARTRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ******* DEPROCO INC**

LIGNOSPAN STANDARD, EPINEPHRINE BITARTRATE
 SCANDONEST L, LEVONORDEFIN
 SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
 SEPTOCAINE, ARTICAINE HYDROCHLORIDE

DERMIRA INC

* DERMIRA INC
 QBREXZA, GLYCOPYRRONIUM TOSYLATE

DEVA HOLDING AS

* DEVA HOLDING AS
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 TEMOZOLOMIDE, TEMOZOLOMIDE

DEXCEL LTD

* DEXCEL LTD
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE

DEXCEL PHARMA

* DEXCEL PHARMA TECHNOLOGIES LTD
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LEVETIRACETAM, LEVETIRACETAM
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 PERIOCHIP, CHLORHEXIDINE GLUCONATE

DFB ONCOLOGY LTD

* DFB ONCOLOGY LTD
 DOCETAXEL, DOCETAXEL

DIAGNOSTIC GREEN

* DIAGNOSTIC GREEN GMBH
 INDOCYANINE GREEN, INDOCYANINE GREEN

DIALYSIS SUPS

* DIALYSIS SUPPLIES INC
 NORMOCARB HF 25, MAGNESIUM CHLORIDE
 NORMOCARB HF 35, MAGNESIUM CHLORIDE

DIGESTIVE CARE INC

* DIGESTIVE CARE INC
 PERTZYE, PANCRELIPASE (AMYLASE)

DORC

* DORC INTERNATIONAL BV
 MEMBRANEBLUE, TRYPAN BLUE
 VISIONBLUE, TRYPAN BLUE

DOUGLAS PHARMS

* DOUGLAS PHARMACEUTICALS AMERICA LTD
 MYORISAN, ISOTRETINOIN

DOW PHARM

* DOW PHARMACEUTICAL SCIENCES
 ACANYA, BENZOYL PEROXIDE
 ALTRENO, TRETINOIN
 ATRALIN, TRETINOIN
 BRYHALI, HALOBETASOL PROPIONATE
 ONEXTON, BENZOYL PEROXIDE
 OXSORALEN-ULTRA, METHOXSALEN

DR REDDYS LA

* DR REDDYS LABORATORIES LOUISIANA LLC
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 LOPURIN, ALLOPURINOL
 SSD, SILVER SULFADIAZINE

DR REDDYS LABS INC

* DR REDDYS LABORATORIES INC
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 AUGMENTIN '875', AMOXICILLIN
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 FINASTERIDE, FINASTERIDE
 FLUCONAZOLE, FLUCONAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

- * DR REDDYS LABORATORIES INC
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 LEVOFLOXACIN, LEVOFLOXACIN
 MELOXICAM, MELOXICAM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NITROGLYCERIN, NITROGLYCERIN
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PROGESTERONE, PROGESTERONE
 PROPOFOL, PROPOFOL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 SIMVASTATIN, SIMVASTATIN
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
- DR REDDYS LABS LTD**
- * DR REDDYS LABORATORIES LIMITED
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
- * DR REDDYS LABORATORIES LTD
 AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZACITIDINE, AZACITIDINE
 BIVALIRUDIN, BIVALIRUDIN
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CARVEDILOL, CARVEDILOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLOFARABINE, CLOFARABINE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DECITABINE, DECITABINE
 DESLORATADINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR, DESLORATADINE
 DESLORATADINE, DESLORATADINE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOCETAXEL, DOCETAXEL
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
 ESZOPICLONE, ESZOPICLONE
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FINASTERIDE, FINASTERIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLIMEPIRIDE, GLIMEPIRIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ******* DR REDDYS LABORATORIES LTD**

GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LAMOTRIGINE, LAMOTRIGINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LATANOPROST, LATANOPROST
 LETROZOLE, LETROZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NATEGLINIDE, NATEGLINIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NIZATIDINE, NIZATIDINE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OFLOXACIN, OFLOXACIN
 OLANZAPINE, OLANZAPINE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 OMEPRAZOLE, OMEPRAZOLE
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXAPROZIN, OXAPROZIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARICALCITOL, PARICALCITOL
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RAMIPRIL, RAMIPRIL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 RISPERIDONE, RISPERIDONE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 SIROLIMUS, SIROLIMUS
 TACROLIMUS, TACROLIMUS
 TETRABENAZINE, TETRABENAZINE
 THIOTEPA, THIOTEPA
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VIGABATRIN, VIGABATRIN
 VINOURELBINE TARTRATE, VINOURELBINE TARTRATE
 ZAFIRLUKAST, ZAFIRLUKAST
 ZEMBRACE SYMTOUCH, SUMATRIPTAN SUCCINATE
 ZENATANE, ISOTRETINOIN
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

DR REDDYS LABS SA*** DR REDDYS LABORATORIES SA**

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

* DR REDDYS LABORATORIES SA
 HABITROL, NICOTINE (OTC)
 MERCAPTOPYRINE, MERCAPTOPYRINE
 RAMELTEON, RAMELTEON
 TOBRAMYCIN, TOBRAMYCIN

DRAximAGE

* DRAximAGE INC
 TECHNETIUM TC 99M ALBUMIN AGGREGATED KIT, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

DUCHESNAY

* DUCHESNAY INC
 BONJESTA, DOXYLAMINE SUCCINATE
 DICLEGIS, DOXYLAMINE SUCCINATE
 OSPHENA, OSPHEMIFENE

DURAMED PHARMS BARR

* DURAMED PHARMACEUTICALS INC SUB BARR LABORATORIES INC
 AVIANE-28, ETHINYL ESTRADIOL
 CRYSELLE, ETHINYL ESTRADIOL
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 ENPRESSE-28, ETHINYL ESTRADIOL
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VELIVET, DESOGESTREL

DUSA

* DUSA PHARMACEUTICALS INC
 LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE

REDDYS

* DOCTOR REDDYS LABORATORIES LTD
 DESLORATADINE, DESLORATADINE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE

**** E ******EAGLE PHARMS**

* EAGLE PHARMACEUTICALS INC
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 BELRAPZO, BENDAMUSTINE HYDROCHLORIDE
 BENDEKA, BENDAMUSTINE HYDROCHLORIDE
 DOCETAXEL, DOCETAXEL
 RYANODEX, DANTROLENE SODIUM

ECI PHARMS LLC

* ECI PHARMACEUTICALS LLC
 BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 LAMIVUDINE, LAMIVUDINE
 LEVETIRACETAM, LEVETIRACETAM
 METHIMAZOLE, METHIMAZOLE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 VALPROIC ACID, VALPROIC ACID
 XTRELUS, GUAIFENESIN

ECOLAB

* ECOLAB INC
 CHG SCRUB, CHLORHEXIDINE GLUCONATE (OTC)
 CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)

ECR

* ECR PHARMACEUTICALS
 DEXAMETHASONE, DEXAMETHASONE

ECR PHARMA

* ECR PHARMA
 TUSSICAPS, CHLORPHENIRAMINE POLISTIREX

EDENBRIDGE PHARMS

* EDENBRIDGE PHARMACEUTICALS LLC
 CARBIDOPA, CARBIDOPA
 ETHACRYNIC ACID, ETHACRYNIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EDENBRIDGE PHARMACEUTICALS LLC
 ETODOLAC, ETODOLAC
 IVERMECTIN, IVERMECTIN
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 TINIDAZOLE, TINIDAZOLE

EDISON THERAPS LLC

* EDISON THERAPEUTICS LLC
 METHERGINE, METHYLERGONOVINE MALEATE

EGALET US INC

* EGALET US INC
 OXAYDO, OXYCODONE HYDROCHLORIDE
 SPRIX, KETOROLAC TROMETHAMINE

EI INC

* EI INC
 THEROXIDIL, MINOXIDIL (OTC)

EISAI INC

* EISAI INC
 ACIPHEX, RABEPRAZOLE SODIUM
 ARICEPT, DONEPEZIL HYDROCHLORIDE
 BANZEL, RUFINAMIDE
 BELVIQ XR, LORCASERIN HYDROCHLORIDE
 BELVIQ, LORCASERIN HYDROCHLORIDE
 FYCOMPA, PERAMPANEL
 HALAVEN, ERIBULIN MESYLATE
 HEXALEN, ALTRETAMINE
 LENVIMA, LENVATINIB MESYLATE
 PANRETIN, ALITRETINOIN
 SALAGEN, PILOCARPINE HYDROCHLORIDE

ELEFSEE PHARMS INTL

* ELEFSEE PHARMACEUTICALS INTERNATIONAL LTD
 LAZANDA, FENTANYL CITRATE

ELI LILLY AND CO

* ELI LILLY AND CO
 BASAGLAR, INSULIN GLARGINE
 EFFIENT, PRASUGREL HYDROCHLORIDE
 HUMALOG KWIKPEN, INSULIN LISPRO RECOMBINANT
 OLUMIANT, BARICITINIB
 PROZAC, FLUOXETINE HYDROCHLORIDE
 VERZENIO, ABEMACICLIB

ELI LILLY CO

* ELI LILLY CO
 ADCIRCA, TADALAFIL
 ZYPREXA RELPREVV, OLANZAPINE PAMOATE

ELITE LABS

* ELITE LABORATORIES INC
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

ELITE LABS INC

* ELITE LABORATORIES INC
 DANTROLENE SODIUM, DANTROLENE SODIUM
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 ISRADIPINE, ISRADIPINE
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE

EMCURE PHARMS

* EMCURE PHARMACEUTICALS LTD
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE

EMCURE PHARMS LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ******* EMCURE PHARMACEUTICALS LTD**

ACARBOSE, ACARBOSE
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 ADENOSINE, ADENOSINE
 AMIKACIN SULFATE, AMIKACIN SULFATE
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BICNU, CARMUSTINE
 CIDOFOVIR, CIDOFOVIR
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ETOMIDATE, ETOMIDATE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FUROSEMIDE, FUROSEMIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 METOCLOPRAMIDE, METOCLOPRAMIDE HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 RIFAMPIN, RIFAMPIN
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

EMD SERONO*** EMD SERONO INC**

GONAL-F RFF REDI-JECT, FOLLITROPIN ALFA/BETA
 GONAL-F RFF, FOLLITROPIN ALFA/BETA
 GONAL-F, FOLLITROPIN ALFA/BETA
 OVIDREL, CHORIOGONADOTROPIN ALFA
 SAIZEN, SOMATROPIN RECOMBINANT
 SEROSTIM, SOMATROPIN RECOMBINANT
 ZORBTIVE, SOMATROPIN RECOMBINANT

EMD SERONO INC*** EMD SERONO INC**

CETROTIDE, CETRORELIX

EMERALD INTL LTD*** EMERALD INTERNATIONAL LTD**

BACLOFEN, BACLOFEN

EMMAUS MEDCL*** EMMAUS MEDICAL INC**

ENDARI, L-GLUTAMINE
 NUTRESTORE, L-GLUTAMINE

ENCORE DERMAT*** ENCORE DERMATOLOGY INC**

IMPOYZ, CLOBETASOL PROPIONATE

ENCUBE ETHICALS*** ENCUBE ETHICALS PVT LTD**

FLUOCINONIDE, FLUOCINONIDE

ENDO PHARM*** ENDO PHARMACEUTICAL SOLUTIONS INC**

SUPPRELIN LA, HISTRELIN ACETATE
 VALSTAR PRESERVATIVE FREE, VALRUBICIN
 VANTAS, HISTRELIN ACETATE

ENDO PHARMS*** ENDO PHARMACEUTICALS INC**

FORTESTA, TESTOSTERONE
 FROVA, FROVATRIPTAN SUCCINATE
 OPANA, OXYMORPHONE HYDROCHLORIDE
 PERCODAN, ASPIRIN

ENDO PHARMS INC*** ENDO PHARMACEUTICALS INC**

AVEED, TESTOSTERONE UNDECANOATE
 COLY-MYCIN S, COLISTIN SULFATE
 MEGACE ES, MEGESTROL ACETATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* ENDO PHARMACEUTICALS INC
 NASCOBAL, CYANOCOBALAMIN
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE

EPI HLTH

* EPI HEALTH LLC
 CLODERM, CLOCORTOLONE PIVALATE
 MINOLIRA, MINOCYCLINE HYDROCHLORIDE
 SITAVIG, ACYCLOVIR

EPIC PHARMA

* EPIC PHARMA INC
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE

* EPIC PHARMA LLC
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
 SULINDAC, SULINDAC
 TRANDOLAPRIL, TRANDOLAPRIL
 URSODIOL, URSODIOL

EPIC PHARMA INC

* EPIC PHARMA INC
 ESTRADIOL, ESTRADIOL
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

EPIC PHARMA LLC

* EPIC PHARMA LLC
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AZITHROMYCIN, AZITHROMYCIN
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 DEMECLOXYCLINE HYDROCHLORIDE, DEMECLOXYCLINE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 GABAPENTIN, GABAPENTIN
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 GLYBURIDE, GLYBURIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 MOLINDONE HYDROCHLORIDE, MOLINDONE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PHENYTOIN, PHENYTOIN
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE

ESPERO

* ESPERO BIOPHARMA INC
 DURLAZA, ASPIRIN

ESSENTIAL ISOTOPES

* ESSENTIAL ISOTOPES LLC
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

ETHYPHARM

* ETHYPHARM
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE

ETHYPHARM USA CORP

* ETHYPHARM USA CORP
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE

EUGIA PHARMA

* EUGIA PHARMA SPECIALITIES LTD
 CAPECITABINE, CAPECITABINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* EUGIA PHARMA SPECIALITIES LTD
 CARBOPLATIN, CARBOPLATIN
 LETROZOLE, LETROZOLE
 OXALIPLATIN, OXALIPLATIN
 PROGESTERONE, PROGESTERONE

EUROHLTH INTL SARL

* EUROHEALTH INTERNATIONAL SARL
 DROPERIDOL, DROPERIDOL
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE

EXALENZ BIOSCIENCE

* EXALENZ BIOSCIENCE LTD
 IDKIT:HP, CITRIC ACID

EXELA HOLDINGS

* EXELA HOLDINGS INC
 DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM

EXELA PHARMA SCIENCE

* EXELA PHARMA SCIENCES
 CAFFEINE CITRATE, CAFFEINE CITRATE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE

EXELA PHARMA SCS LLC

* EXELA PHARMA SCIENCES LLC
 CAFFEINE CITRATE, CAFFEINE CITRATE
 GANCICLOVIR, GANCICLOVIR
 GLYRX-PF, GLYCOPYRROLATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 NIPRIDE RTU IN SODIUM CHLORIDE 0.9%, SODIUM NITROPRUSSIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 POTASSIUM ACETATE, POTASSIUM ACETATE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

EXELIXIS

* EXELIXIS INC
 COMETRIQ, CABOZANTINIB S-MALATE

EXELIXIS INC

* EXELIXIS INC
 CABOMETYX, CABOZANTINIB S-MALATE

EYEPOINT PHARMS

* EYEPOINT PHARMACEUTICALS INC
 DEXYCU KIT, DEXAMETHASONE
 YUTIQ, FLUOCINOLONE ACETONIDE

EYEVANCE PHARMS

* EYEVANCE PHARMACEUTICALS LLC
 ZERVIATE, CETIRIZINE HYDROCHLORIDE

LILLY

* ELI LILLY AND CO
 ALIMTA, PEMETREXED DISODIUM
 CIALIS, TADALAFIL
 CYMBALTA, DULOXETINE HYDROCHLORIDE
 EVISTA, RALOXIFENE HYDROCHLORIDE
 FORTEO, TERIPARATIDE RECOMBINANT HUMAN
 GEMZAR, GEMCITABINE HYDROCHLORIDE
 GLUCAGON, GLUCAGON
 HUMALOG KWIKPEN, INSULIN LISPRO RECOMBINANT
 HUMALOG MIX 50/50 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT
 HUMALOG MIX 50/50, INSULIN LISPRO PROTAMINE RECOMBINANT
 HUMALOG MIX 75/25 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT
 HUMALOG MIX 75/25, INSULIN LISPRO PROTAMINE RECOMBINANT
 HUMALOG, INSULIN LISPRO RECOMBINANT
 HUMATROPE, SOMATROPIN RECOMBINANT
 HUMULIN 70/30 PEN, INSULIN RECOMBINANT HUMAN (OTC)
 HUMULIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
 HUMULIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

* ELI LILLY AND CO
 HUMULIN R KWIKPEN, INSULIN HUMAN
 HUMULIN R PEN, INSULIN RECOMBINANT HUMAN (OTC)
 HUMULIN R, INSULIN HUMAN
 HUMULIN R, INSULIN RECOMBINANT HUMAN (OTC)
 PROZAC WEEKLY, FLUOXETINE HYDROCHLORIDE
 STRATTERA, ATOMOXETINE HYDROCHLORIDE
 SYMBYAX, FLUOXETINE HYDROCHLORIDE
 ZYPREXA ZYDIS, OLANZAPINE
 ZYPREXA, OLANZAPINE

**** F ******FDC LTD**

* FDC LTD
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 LATANOPROST, LATANOPROST
 OFLOXACIN, OFLOXACIN
 TIMOLOL MALEATE, TIMOLOL MALEATE

FDN CONSUMER

* FOUNDATION CONSUMER HEALTHCARE LLC
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 PLAN B ONE-STEP, LEVONORGESTREL (OTC)

FEINSTEIN

* FEINSTEIN INSTITUTE MEDICAL RESEARCH
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

FERA PHARMS

* FERA PHARMACEUTICALS LLC
 TOBRAMYCIN, TOBRAMYCIN

FERA PHARMS LLC

* FERA PHARMACEUTICALS LLC
 DEXAMETHASONE, DEXAMETHASONE

FERRER INTERNACIONAL

* FERRER INTERNACIONAL SA
 XEPI, OZENOXACIN

FERRING

* FERRING PHARMACEUTICALS INC
 ACTHREL, CORTICORELIN OVINE TRIFLUTATE
 CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC
 ENDOMETRIN, PROGESTERONE
 FIRMAGON, DEGARELIX ACETATE
 MENOPUR, MENOTROPINS (FSH)
 MINIRIN, DESMOPRESSIN ACETATE
 ZOMACTON, SOMATROPIN

FERRING PHARMS INC

* FERRING PHARMACEUTICALS INC
 CERVIDIL, DINOPROSTONE
 CLENPIQ, CITRIC ACID
 DDAVP (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 DDAVP, DESMOPRESSIN ACETATE
 LYSTEDA, TRANEXAMIC ACID
 NOCDURNA, DESMOPRESSIN ACETATE
 PREPOPIK, CITRIC ACID
 STIMATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE

FLAMEL IRELAND LTD

* FLAMEL IRELAND LIMITED
 AKOVAZ, EPHEDRINE SULFATE

FLAMINGO PHARMS

* FLAMINGO PHARMACEUTICALS LTD
 METRONIDAZOLE, METRONIDAZOLE
 PIROXICAM, PIROXICAM

FLEXION THERAPS INC

* FLEXION THERAPEUTICS INC
 ZILRETTA, TRIAMCINOLONE ACETONIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ******FOREST LABS INC**

* FOREST LABORATORIES INC
 VIOKACE, PANCRELIPASE (AMYLASE)
 ZENPEP, PANCRELIPASE (AMYLASE)

FOREST LABS LLC

* FOREST LABORATORIES LLC
 NAMENDA XR, MEMANTINE HYDROCHLORIDE
 SAPHRIS, ASENAPINE MALEATE

FOUGERA PHARMS

* FOUGERA PHARMACEUTICALS INC
 ADAPALENE, ADAPALENE
 ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
 AMCINONIDE, AMCINONIDE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE, CALCIPOTRIENE
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 CUTIVATE, FLUTICASONE PROPIONATE
 DESONIDE, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 ERYTHROMYCIN, ERYTHROMYCIN
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HYDROCORTISONE, HYDROCORTISONE
 IMIQUIMOD, IMIQUIMOD
 KETOCONAZOLE, KETOCONAZOLE
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 METRONIDAZOLE, METRONIDAZOLE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MUPIROXIN, MUPIROXIN
 NYSTATIN, NYSTATIN
 OXISTAT, OXICONAZOLE NITRATE
 PANDEL, HYDROCORTISONE PROBUTATE
 PREDNICARBATE, PREDNICARBATE
 SOLARAZE, DICLOFENAC SODIUM
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TERCONAZOLE, TERCONAZOLE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TYZINE, TETRAHYDROZOLINE HYDROCHLORIDE

FOUGERA PHARMS INC

* FOUGERA PHARMACEUTICALS INC
 ACYCLOVIR, ACYCLOVIR
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HYDROCORTISONE, HYDROCORTISONE
 LIDOCAINE, LIDOCAINE
 NITROGLYCERIN, NITROGLYCERIN
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 TACROLIMUS, TACROLIMUS
 VEREGEN, SINECATECHINS

FRESENIUS

* FRESENIUS KABI DEUTSCHLAND GMBH

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

- * FRESENIUS KABI DEUTSCHLAND GMBH
 - INTRALIPID 10%, SOYBEAN OIL
 - INTRALIPID 20%, SOYBEAN OIL
 - INTRALIPID 30%, SOYBEAN OIL

FRESENIUS KABI

- * FRESENIUS KABI ANTI INFECTIVES SRL
 - PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
- * FRESENIUS KABI AUSTRIA GMBH
 - LACTULOSE, LACTULOSE

FRESENIUS KABI ONCOL

- * FRESENIUS KABI ONCOLOGY PLC
 - ANASTROZOLE, ANASTROZOLE
 - BICALUTAMIDE, BICALUTAMIDE
 - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 - LETROZOLE, LETROZOLE

FRESENIUS KABI USA

- * FRESENIUS KABI USA LLC
 - ACETAMINOPHEN, ACETAMINOPHEN
 - ACETYLCYSTEINE, ACETYLCYSTEINE
 - ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 - ADENOSINE, ADENOSINE
 - AMIKACIN SULFATE, AMIKACIN SULFATE
 - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 - ARGATROBAN, ARGATROBAN
 - ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 - ASTRAMORPH PF, MORPHINE SULFATE
 - ATROPINE SULFATE, ATROPINE SULFATE
 - AZITHROMYCIN, AZITHROMYCIN
 - AZTREONAM, AZTREONAM
 - BACITRACIN, BACITRACIN
 - BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 - BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 - BIVALIRUDIN, BIVALIRUDIN
 - BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 - BORTEZOMIB, BORTEZOMIB
 - CAFFEINE CITRATE, CAFFEINE CITRATE
 - CALCIUM GLUCONATE, CALCIUM GLUCONATE
 - CARBOPLATIN, CARBOPLATIN
 - CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 - CEFOTETAN, CEFOTETAN DISODIUM
 - CHLORAMPHENICOL SODIUM SUCCINATE, CHLORAMPHENICOL SODIUM SUCCINATE
 - CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 - CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC
 - CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 - CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 - CISPLATIN, CISPLATIN
 - CLADRIBINE, CLADRIBINE
 - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 - COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 - CYTARABINE, CYTARABINE
 - DACARBAZINE, DACARBAZINE
 - DAPTOMYCIN, DAPTOMYCIN
 - DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 - DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 - DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
 - DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 - DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
 - DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 - DILAUDID, HYDROMORPHONE HYDROCHLORIDE
 - DIMENHYDRINATE, DIMENHYDRINATE
 - DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
 - DIPRIVAN, PROPOFOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FRESENIUS KABI USA LLC
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DOXY 100, DOXYCYCLINE HYCLATE
 DOXY 200, DOXYCYCLINE HYCLATE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ETOPOSIDE, ETOPOSIDE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FLOXURIDINE, FLOXURIDINE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUMAZENIL, FLUMAZENIL
 FLUOROURACIL, FLUOROURACIL
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FUROSEMIDE, FUROSEMIDE
 GANCICLOVIR, GANCICLOVIR SODIUM
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GLUCAGON, GLUCAGON HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE, GRANISETRON HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
 IFOSFAMIDE, IFOSFAMIDE
 INDOMETHACIN, INDOMETHACIN
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 KABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LINEZOLID, LINEZOLID
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MANNITOL 25%, MANNITOL
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MESNA, MESNA
 METHOCARBAMOL, METHOCARBAMOL
 METHOTREXATE PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

* FRESENIUS KABI USA LLC
MORPHINE SULFATE, MORPHINE SULFATE
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
NAROPIN, ROPIVACAINE HYDROCHLORIDE
NEBUPENT, PENTAMIDINE ISETHIONATE
NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
NESACAINE, CHLOROPROCAINE HYDROCHLORIDE
NESACAINE-MPF, CHLOROPROCAINE HYDROCHLORIDE
OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
OMEGAVEN, FISH OIL TRIGLYCERIDES
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
OXALIPLATIN, OXALIPLATIN
OXYTOCIN, OXYTOCIN
PACLITAXEL, PACLITAXEL
PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
PENTAM, PENTAMIDINE ISETHIONATE
PERIKABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
POLOCAINE, MEPIVACAINE HYDROCHLORIDE
POLOCAINE-MPF, MEPIVACAINE HYDROCHLORIDE
POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
POTASSIUM CHLORIDE IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PROGESTERONE, PROGESTERONE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
PROTAMINE SULFATE, PROTAMINE SULFATE
PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
REMIFENTANIL HYDROCHLORIDE, REMIFENTANIL HYDROCHLORIDE
RIFAMPIN, RIFAMPIN
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
SENSORCAINE, BUPIVACAINE HYDROCHLORIDE
SENSORCAINE, BUPIVACAINE HYDROCHLORIDE
SMOFLIPID 20%, FISH OIL
SODIUM ACETATE, SODIUM ACETATE
SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
TERBUTALINE SULFATE, TERBUTALINE SULFATE
THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
TIGECYCLINE, TIGECYCLINE
TOBRAMYCIN SULFATE (PHARMACY BULK), TOBRAMYCIN SULFATE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
TRANEXAMIC ACID, TRANEXAMIC ACID
VALPROATE SODIUM, VALPROATE SODIUM
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
VIBISONE, CYANOCOBALAMIN
VINBLASTINE SULFATE, VINBLASTINE SULFATE
VINORELBINE TARTRATE, VINORELBINE TARTRATE
XYLOCAINE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
XYLOCAINE, LIDOCAINE HYDROCHLORIDE
ZOLEDRONIC ACID, ZOLEDRONIC ACID

FRESENIUS MEDCL

* FRESENIUS MEDICAL CARE NORTH AMERICA
DELFLEX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM
DELFLEX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

- * FRESENIUS MEDICAL CARE NORTH AMERICA
 DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM
 DELFLEX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM
 PHOSLO GELCAPS, CALCIUM ACETATE
 PHOSLYRA, CALCIUM ACETATE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

FRONTIDA BIOPHARM

- * FRONTIDA BIOPHARM INC
 CILOSTAZOL, CILOSTAZOL
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE

**** G ******G AND W LABS**

- * G AND W LABORATORIES INC
 ACEPHEN, ACETAMINOPHEN (OTC)
 CICLOPIROX, CICLOPIROX
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 INDOMETHACIN, INDOMETHACIN
 METRONIDAZOLE, METRONIDAZOLE
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 MOMETASONE FUROATE, MOMETASONE FUROATE
 PROCHLORPERAZINE, PROCHLORPERAZINE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROMETHEGAN, PROMETHAZINE HYDROCHLORIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

G AND W LABS INC

- * G AND W LABORATORIES INC
 ACYCLOVIR, ACYCLOVIR
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 BETA-VAL, BETAMETHASONE VALERATE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CALCIPOTRIENE, CALCIPOTRIENE
 CICLOPIROX, CICLOPIROX
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DESONIDE, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DOXYCYCLINE, DOXYCYCLINE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUOCINONIDE, FLUOCINONIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 LIDOCAINE, LIDOCAINE
 METRONIDAZOLE, METRONIDAZOLE
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 MYKACET, NYSTATIN
 NYSTATIN, NYSTATIN
 TAZAROTENE, TAZAROTENE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

GALDERMA LABS

- * GALDERMA LABORATORIES INC
 CLOBEX, CLOBETASOL PROPIONATE
 EPIDUO FORTE, ADAPALENE

GALDERMA LABS LP

- * GALDERMA LABORATORIES L P
 CLOBEX, CLOBETASOL PROPIONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GALDERMA LABORATORIES LP
 CAPEX, FLUOCINOLONE ACETONIDE
 CLOBEX, CLOBETASOL PROPIONATE
 DESOWEN, DESONIDE
 DIFFERIN, ADAPALENE
 DIFFERIN, ADAPALENE (OTC)
 EPIDUO, ADAPALENE
 METROCREAM, METRONIDAZOLE
 METROGEL, METRONIDAZOLE
 METROLOTION, METRONIDAZOLE
 MIRVASO, BRIMONIDINE TARTRATE
 ORACEA, DOXYCYCLINE
 SOOLANTRA, IVERMECTIN
 TRI-LUMA, FLUOCINOLONE ACETONIDE
 VECTICAL, CALCITRIOL

GALEN SPECIALTY

* GALEN SPECIALTY PHARMA US LLC
 SYNERA, LIDOCAINE

GALEN UK

* GALEN LTD
 ADASUVE, LOXAPINE

GALT PHARMS

* GALT PHARMACEUTICALS LLC
 DORAL, QUAZEPAM

GATE PHARMS

* GATE PHARMACEUTICALS
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 LINEZOLID, LINEZOLID

GATOR PHARMS

* GATOR PHARMACEUTICALS INC
 COLPREP KIT, MAGNESIUM SULFATE

GAVIS PHARMS

* GAVIS PHARMACEUTICALS LLC
 NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 QUINARETIC, HYDROCHLOROTHIAZIDE
 TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE

GAVIS PHARMS LLC

* GAVIS PHARMACEUTICALS LLC
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE

GD SEARLE

* GD SEARLE LLC
 CELEBREX, CELECOXIB
 DAYPRO, OXAPROZIN

GD SEARLE LLC

* GD SEARLE LLC
 ALDACTAZIDE, HYDROCHLOROTHIAZIDE
 ALDACTONE, SPIRONOLACTONE
 ARTHROTEC, DICLOFENAC SODIUM
 CALAN, VERAPAMIL HYDROCHLORIDE
 CYTOTEC, MISOPROSTOL
 FLAGYL, METRONIDAZOLE
 INSPRA, EPLERENONE
 LOMOTIL, ATROPINE SULFATE
 NORPACE CR, DISOPYRAMIDE PHOSPHATE
 NORPACE, DISOPYRAMIDE PHOSPHATE
 SYNAREL, NAFARELIN ACETATE

GE HEALTHCARE

* GE HEALTHCARE
 ADREVIEW, IOBENGUANE SULFATE I-123
 CERETEC, TECHNETIUM TC-99M EXAMETAZIME KIT
 INDICLOR, INDIUM IN-111 CHLORIDE
 INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ******* GE HEALTHCARE**

METASTRON, STRONTIUM CHLORIDE SR-89
 MPI INDIUM DTPA IN 111, INDIUM IN-111 PENTETATE DISODIUM
 MYOVUE 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT
 OMNIPAQUE 12, IOHEXOL
 OMNIPAQUE 140, IOHEXOL
 OMNIPAQUE 180, IOHEXOL
 OMNIPAQUE 240, IOHEXOL
 OMNIPAQUE 300, IOHEXOL
 OMNIPAQUE 350, IOHEXOL
 OMNIPAQUE 9, IOHEXOL
 OMNISCAN, GADODIAMIDE
 OPTISON, ALBUMIN HUMAN
 TECHNETIUM TC 99M GENERATOR, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
 VISIPAQUE 270, IODIXANOL
 VISIPAQUE 320, IODIXANOL
 VIZAMYL, FLUTEMETAMOL F-18

GE HLTHCARE INC

* GE HEALTHCARE INC
 DATSCAN, IOFLUPANE I-123

GEMINI LABS LLC

* GEMINI LABORATORIES LLC
 PRANDIN, REPAGLINIDE

GENENTECH

* GENENTECH INC
 ERIVEDGE, VISMODEGIB
 NUTROPIN AQ NUSPIN, SOMATROPIN RECOMBINANT

GENENTECH INC

* GENENTECH INC
 COCELLIC, COBIMETINIB FUMARATE
 ESBRIET, PIRFENIDONE
 XOFLUZA, BALOXAVIR MARBOXIL

GENERIC

* GENERICS INTERNATIONAL VENTURES ENTERPRISES LLC
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

GENEYORK PHARMS

* GENEYORK PHARMACEUTICALS GROUP LLC
 PREDNISONE, PREDNISONE

GENUS LIFESCIENCES

* GENUS LIFE SCIENCES INC
 GOPRELTO, COCAINE HYDROCHLORIDE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 YOSPRALA, ASPIRIN
 * GENUS LIFESCIENCES INC
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN

GENZYME

* GENZYME CORP
 CEREZYME, IMIGLUCERASE
 CLOLAR, CLOFARABINE
 MOZOBIL, PLERIXAFOR
 RENAGEL, SEVELAMER HYDROCHLORIDE
 RENVELA, SEVELAMER CARBONATE
 THYROGEN, THYROTROPIN ALFA

GENZYME CORP

* GENZYME CORP
 CAPRELSA, VANDETANIB
 CERDELGA, ELIGLUSTAT TARTRATE

GILEAD

* GILEAD SCIENCES INC
 CAYSTON, AZTREONAM
 EMTRIVA, EMTRICITABINE
 HEPSERA, ADEFOVIR DIPIVOXIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GILEAD SCIENCES INC
 LETAIRIS, AMBRISENTAN
 RANEXA, RANOLAZINE
 TRUVADA, EMTRICITABINE

GILEAD SCIENCES

* GILEAD SCIENCES LLC
 ATRIPLA, EFAVIRENZ

GILEAD SCIENCES INC

* GILEAD SCIENCES INC
 BIKTARVY, BICTEGRAVIR SODIUM
 COMPLERA, EMTRICITABINE
 DESCOVY, EMTRICITABINE
 EPCLUSA, SOFOSBUVIR
 GENVOYA, COBICISTAT
 HARVONI, LEDIPASVIR
 ODEFSEY, EMTRICITABINE
 SOVALDI, SOFOSBUVIR
 STRIBILD, COBICISTAT
 TYBOST, COBICISTAT
 VEMLIDY, TENOFOVIR ALAFENAMIDE FUMARATE
 VIREAD, TENOFOVIR DISOPROXIL FUMARATE
 VOSEVI, SOFOSBUVIR
 ZYDELIG, IDELALISIB

GLAND PHARMA LTD

* GLAND PHARMA LTD
 ADENOSINE, ADENOSINE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 AZITHROMYCIN, AZITHROMYCIN
 CARBOPLATIN, CARBOPLATIN
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 CISPLATIN, CISPLATIN
 CLOFARABINE, CLOFARABINE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 ETOMIDATE, ETOMIDATE
 FLUOROURACIL, FLUOROURACIL
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM, HEPARIN SODIUM
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 MEROPENEM, MEROPENEM
 MESNA, MESNA
 METHOCARBAMOL, METHOCARBAMOL
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PACITAXEL, PACLITAXEL
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID
 ZOLEDRONIC, ZOLEDRONIC ACID

GLASSHOUSE PHARMS

* GLASSHOUSE PHARMACEUTICALS LTD CANADA
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GLASSHOUSE PHARMACEUTICALS LTD CANADA
FLUOCINONIDE, FLUOCINONIDE

GLAXO GRP ENGLAND

* GLAXO GROUP LTD ENGLAND DBA GLAXOSMITHKLINE
INCRUSE ELLIPTA, UMECLIDINIUM BROMIDE

GLAXO GRP LTD

* GLAXO GROUP LTD DBA GLAXOSMITHKLINE
FLOVENT HFA, FLUTICASONE PROPIONATE

* GLAXO GROUP LTD ENGLAND DBA GLAXOSMITHKLINE
ADVAIR DISKUS 100/50, FLUTICASONE PROPIONATE
ADVAIR DISKUS 250/50, FLUTICASONE PROPIONATE
ADVAIR DISKUS 500/50, FLUTICASONE PROPIONATE
ADVAIR HFA, FLUTICASONE PROPIONATE
BREQ ELLIPTA, FLUTICASONE FUROATE
FLOVENT DISKUS 100, FLUTICASONE PROPIONATE
FLOVENT DISKUS 250, FLUTICASONE PROPIONATE
FLOVENT DISKUS 50, FLUTICASONE PROPIONATE

GLAXOSMITHKLINE

* GLAXOSMITHKLINE
ABREVA, DOCOSANOL (OTC)
AVODART, DUTASTERIDE
BECONASE AQ, BECLOMETHASONE DIPROPIONATE MONOHYDRATE
EPIVIR-HBV, LAMIVUDINE
IMITREX STATDOSE, SUMATRIPTAN SUCCINATE
IMITREX, SUMATRIPTAN
IMITREX, SUMATRIPTAN SUCCINATE
JALYN, DUTASTERIDE
MALARONE PEDIATRIC, ATOVAQUONE
MALARONE, ATOVAQUONE
NICORETTE (MINT), NICOTINE POLACRILEX (OTC)
NICORETTE, NICOTINE POLACRILEX (OTC)
RELENZA, ZANAMIVIR
VALTREX, VALACYCLOVIR HYDROCHLORIDE
WELLBUTRIN SR, BUPROPION HYDROCHLORIDE
ZYBAN, BUPROPION HYDROCHLORIDE

* GLAXOSMITHKLINE CONSUMER HEALTHCARE HOLDINGS (US) LLC
LAMISIL, TERBINAFFINE HYDROCHLORIDE (OTC)

* GLAXOSMITHKLINE INTELLECTUAL PROPERTY DEVELOPMENT LTD ENGLAND
ANORO ELLIPTA, UMECLIDINIUM BROMIDE
ARNUITY ELLIPTA, FLUTICASONE FUROATE
KRINTAFEL, TAFENOQUINE SUCCINATE
TRELEGY ELLIPTA, FLUTICASONE FUROATE

* GLAXOSMITHKLINE INTELLECTUAL PROPERTY LTD ENGLAND
SEREVENT, SALMETEROL XINAFOATE
VENTOLIN HFA, ALBUTEROL SULFATE

GLAXOSMITHKLINE CON

* GLAXOSMITHKLINE CONSUMER HEALTH
TRANSDERM SCOP, SCOPOLAMINE

GLAXOSMITHKLINE CONS

* GLAXOSMITHKLINE CONSUMER HEALTHCARE
ALLI, ORLISTAT (OTC)
EXCEDRIN (MIGRAINE), ACETAMINOPHEN (OTC)
FLONASE ALLERGY RELIEF, FLUTICASONE PROPIONATE (OTC)
FLONASE SENSIMIST ALLERGY RELIEF, FLUTICASONE FUROATE (OTC)
LAMISIL AT, TERBINAFFINE (OTC)
LAMISIL AT, TERBINAFFINE HYDROCHLORIDE (OTC)
NICORETTE, NICOTINE POLACRILEX (OTC)
PREVACID 24 HR, LANSOPRAZOLE (OTC)
VOLTAREN, DICLOFENAC SODIUM

GLAXOSMITHKLINE LLC

* GLAXOSMITHKLINE LLC
AMERGE, NARATRIPTAN HYDROCHLORIDE
DYAZIDE, HYDROCHLOROTHIAZIDE
FLOLAN, EPOPROSTENOL SODIUM
LAMICTAL CD, LAMOTRIGINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GLAXOSMITHKLINE LLC

LAMICTAL ODT, LAMOTRIGINE
 LAMICTAL XR, LAMOTRIGINE
 LAMICTAL, LAMOTRIGINE
 MEPRON, ATOVAQUONE
 REQUIP XL, ROPINIROLE HYDROCHLORIDE
 REQUIP, ROPINIROLE HYDROCHLORIDE
 RYTHMOL SR, PROPAFENONE HYDROCHLORIDE

GLENMARK

* GLENMARK THERAPEUTICS INC USA
 ECOZA, ECONAZOLE NITRATE

GLENMARK GENERICS

* GLENMARK GENERICS INC USA

ADAPALENE, ADAPALENE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 IMIQUIMOD, IMIQUIMOD
 MOMETASONE FUROATE, MOMETASONE FUROATE
 NIZATIDINE, NIZATIDINE
 ZONISAMIDE, ZONISAMIDE

* GLENMARK GENERICS LIMITED

BRIELLYN, ETHINYL ESTRADIOL
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE

* GLENMARK GENERICS LTD

ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
 ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
 ALYACEN 1/35, ETHINYL ESTRADIOL
 ALYACEN 7/7/7, ETHINYL ESTRADIOL
 ASHLYNA, ETHINYL ESTRADIOL
 ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
 CARVEDILOL, CARVEDILOL
 CICLOPIROX, CICLOPIROX
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 DESOXIMETASONE, DESOXIMETASONE
 ESZOPICLONE, ESZOPICLONE
 FELODIPINE, FELODIPINE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOCINONIDE, FLUOCINONIDE
 HEATHER, NORETHINDRONE
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 LAMOTRIGINE, LAMOTRIGINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LITHIUM CARBONATE, LITHIUM CARBONATE
 MARLISSA, ETHINYL ESTRADIOL
 MELOXICAM, MELOXICAM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NAPROXEN, NAPROXEN
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE, NORETHINDRONE
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 OMEPRAZOLE, OMEPRAZOLE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON, ONDANSETRON
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 THEOPHYLLINE, THEOPHYLLINE
 TOPIRAMATE, TOPIRAMATE
 TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE, TRANDOLAPRIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

- * GLENMARK GENERICS LTD
TROSPIMUM CHLORIDE, TROSPIMUM CHLORIDE
URSODIOL, URSODIOL
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
VIORELE, DESOGESTREL
ZOLMITRIPTAN, ZOLMITRIPTAN
- * GLENMARK GENERICS LTD INDIA
INDOMETHACIN, INDOMETHACIN
NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

GLENMARK PHARMS

- * GLENMARK PHARMACEUTICALS INC
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
TERIFLUNOMIDE, TERIFLUNOMIDE
- * GLENMARK PHARMACEUTICALS INC USA
CICLOPIROX, CICLOPIROX
CLOTRIMAZOLE, CLOTRIMAZOLE
MUPIROCIN, MUPIROCIN
- * GLENMARK PHARMACEUTICALS LTD
MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
- * GLENMARK PHARMACEUTICALS SA
ATOVAQUONE, ATOVAQUONE
AZELAIC ACID, AZELAIC ACID
CALCIPOTRIENE, CALCIPOTRIENE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
DESONIDE, DESONIDE
HAILEY 1.5/30, ETHINYL ESTRADIOL
HAILEY FE 1.5/30, ETHINYL ESTRADIOL
LINEZOLID, LINEZOLID
NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

GLENMARK PHARMS INC

- * GLENMARK PHARMACEUTICALS INC USA
CALCIPOTRIENE, CALCIPOTRIENE
LITHIUM CARBONATE, LITHIUM CARBONATE
MUPIROCIN, MUPIROCIN CALCIUM
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

GLENMARK PHARMS LTD

- * GLENMARK PHARMACEUTICALS LTD
ACYCLOVIR, ACYCLOVIR
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DESONIDE, DESONIDE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
ESTRADIOL, ESTRADIOL
EZETIMIBE, EZETIMIBE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FLUCINOLONE ACETONIDE, FLUCINOLONE ACETONIDE
FLUCINONIDE ACETONIDE, FLUCINOLONE ACETONIDE
FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
GABAPENTIN, GABAPENTIN
HAILEY FE 1/20, ETHINYL ESTRADIOL
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
INDOMETHACIN, INDOMETHACIN
LAMOTRIGINE, LAMOTRIGINE
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
LEVONORGESTREL, LEVONORGESTREL (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GLENMARK PHARMACEUTICALS LTD
 LIDOCAINE, LIDOCAINE
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OXCARBAZEPINE, OXCARBAZEPINE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RILUZOLE, RILUZOLE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 RUFINAMIDE, RUFINAMIDE
 TACROLIMUS, TACROLIMUS
 TELMISARTAN, TELMISARTAN
 TRETINOIN, TRETINOIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VORICONAZOLE, VORICONAZOLE

GLENMARK PHARMS SA

* GLENMARK PHARMACEUTICALS SA SWITZERLAND
 APREPITANT, APREPITANT
 NITROGLYCERIN, NITROGLYCERIN

GLOBAL ISOTOPES LLC

* GLOBAL ISOTOPES LLC DBA ZEVACOR MOLECULAR
 AMMONIA N 13, AMMONIA N-13
 CHOLINE C-11, CHOLINE C-11
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

GP-PHARM SA

* GP-PHARM SA
 LUTRATE DEPOT KIT, LEUPROLIDE ACETATE

GRANULES INDIA

* GRANULES INDIA LTD
 IBUPROFEN, IBUPROFEN (OTC)
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

GRANULES INDIA LTD

* GRANULES INDIA LTD
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHOCARBAMOL, METHOCARBAMOL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)

GRANULES PHARMS

* GRANULES PHARMACEUTICALS INC
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

GRAVITI PHARMS

* GRAVITI PHARMACEUTICALS PRIVATE LTD
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 FENOFIBRATE, FENOFIBRATE

GUARDIAN DRUG

* GUARDIAN DRUG CO
 GUAIFENESIN, GUAIFENESIN (OTC)
 IBUPROFEN, IBUPROFEN (OTC)

GUERBET

* GUERBET LLC
 DOTAREM, GADOTERATE MEGLUMINE
 LIPIODOL, ETHIODIZED OIL

GW RES LTD

* GW RESEARCH LTD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

* GW RESEARCH LTD
EPIDIOLEX, CANNABIDIOL

HANFORD GC

* GC HANFORD MANUFACTURING CO
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
AMPICILLIN SODIUM, AMPICILLIN SODIUM
PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM

POHL BOSKAMP

* G POHL BOSKAMP GMBH AND CO KG
GONITRO, NITROGLYCERIN

**** H ******HAEMONETICS**

* HAEMONETICS CORP
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

HAINAN POLY PHARM

* HAINAN POLY PHARMACEUTICAL CO LTD
AZITHROMYCIN, AZITHROMYCIN
GANCICLOVIR, GANCICLOVIR SODIUM
LEVETIRACETAM, LEVETIRACETAM
VORICONAZOLE, VORICONAZOLE

HALOCARBON PRODS

* HALOCARBON PRODUCTS CORP
ISOFLURANE, ISOFLURANE
SEVOFLURANE, SEVOFLURANE

HALOZYME THERAP

* HALOZYME THERAPEUTICS INC
HYLENEX RECOMBINANT, HYALURONIDASE RECOMBINANT HUMAN

HAMELN PHARMA PLUS

* HAMELN PHARMA PLUS GMBH
PENTETATE CALCIUM TRISODIUM, PENTETATE CALCIUM TRISODIUM
PENTETATE ZINC TRISODIUM, PENTETATE ZINC TRISODIUM

HANDA PHARMS LLC

* HANDA PHARMACEUTICALS LLC
LAMOTRIGINE, LAMOTRIGINE

HANGZHOU BINJIANG

* HANGZHOU MINSHENG BINJIANG PHARMACEUTICAL CO LTD
ALENDRONATE SODIUM, ALENDRONATE SODIUM

HANSAMED INC

* HANSAMED INC
ULTACAN FORTE, ARTICAINE HYDROCHLORIDE
ULTACAN, ARTICAINE HYDROCHLORIDE

HARRIS PHARM

* HARRIS PHARMACEUTICAL INC
FLUCONAZOLE, FLUCONAZOLE
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE

HEBEI CHANGSHAN

* HEBEI CHANGSHAN BIOCHEMICAL PHARMACEUTICAL CO LTD
AMLODIPINE BESYLATE, AMLDIPINE BESYLATE
SILDENAFIL CITRATE, SILDENAFIL CITRATE

HEC PHARM

* HEC PHARM USA INC
CLARITHROMYCIN, CLARITHROMYCIN
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
IBUPROFEN, IBUPROFEN
OLANZAPINE, OLANZAPINE
PRASUGREL, PRASUGREL HYDROCHLORIDE

HELSINN

* HELSINN BIREX PHARMACEUTICALS LTD
VALCHLOR, MECHLORETHAMINE HYDROCHLORIDE

HELSINN HLTHCARE

* HELSINN HEALTHCARE SA
AKYNZEO, FOSNETUPITANT CHLORIDE HYDROCHLORIDE
AKYNZEO, NETUPITANT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HELSINN HEALTHCARE SA
ALOXI, PALONOSETRON HYDROCHLORIDE

HERCON PHARM

* HERCON PHARMACEUTICAL LLC
NITROGLYCERIN, NITROGLYCERIN

HERITAGE LIFE

* HERITAGE LIFE SCIENCES BARBADOS INC
CLOZARIL, CLOZAPINE

HERITAGE PHARMA

* HERITAGE PHARMA LABS INC
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
ACETAZOLAMIDE, ACETAZOLAMIDE
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DIFLUNISAL, DIFLUNISAL
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
LITHIUM CARBONATE, LITHIUM CARBONATE
METHIMAZOLE, METHIMAZOLE
NIFEDIPINE, NIFEDIPINE

HERITAGE PHARMS INC

* HERITAGE PHARMACEUTICALS INC
ACETAZOLAMIDE, ACETAZOLAMIDE
ACHROMYCIN V, TETRACYCLINE HYDROCHLORIDE
ACYCLOVIR, ACYCLOVIR
ALPRAZOLAM, ALPRAZOLAM
CALCIUM ACETATE, CALCIUM ACETATE
CARISOPRODOL AND ASPIRIN, ASPIRIN
CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DOXYCYCLINE, DOXYCYCLINE
DUTASTERIDE, DUTASTERIDE
ETHOSUXIMIDE, ETHOSUXIMIDE
FELODIPINE, FELODIPINE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
GLYBURIDE, GLYBURIDE
GLYCOPYRROLATE, GLYCOPYRROLATE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
INDOMETHACIN, INDOMETHACIN
KETOPROFEN, KETOPROFEN
LEFLUNOMIDE, LEFLUNOMIDE
METRONIDAZOLE, METRONIDAZOLE
MODAFINIL, MODAFINIL
MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
NIMODIPINE, NIMODIPINE
NYSTATIN, NYSTATIN
PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

HERON THERAPS INC

* HERON THERAPEUTICS INC
CINVANTI, APREPITANT
SUSTOL, GRANISETRON

HETERO LABS LTD III

* HETERO LABS LTD UNIT III

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******* HETERO LABS LTD UNIT III**

ABACAVIR SULFATE, ABACAVIR SULFATE
 ATOVAQUONE, ATOVAQUONE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLOBAZAM, CLOBAZAM
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 EFAVIRENZ, EFAVIRENZ
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FENOFIBRATE, FENOFIBRATE
 FINASTERIDE, FINASTERIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 INDOMETHACIN, INDOMETHACIN
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 METHOCARBAMOL, METHOCARBAMOL
 NEVIRAPINE, NEVIRAPINE
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 RITONAVIR, RITONAVIR
 ROFLUMILAST, ROFLUMILAST
 SIMVASTATIN, SIMVASTATIN
 STAVUDINE, STAVUDINE
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TORSEMIDE, TORSEMIDE
 ZIDOVUDINE, ZIDOVUDINE

HETERO LABS LTD V*** HETERO LABS LTD UNIT V**

ACYCLOVIR, ACYCLOVIR
 ARIPIPRAZOLE, ARIPIPRAZOLE
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 ENTECAVIR, ENTECAVIR
 FAMCICLOVIR, FAMCICLOVIR
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 IRBESARTAN, IRBESARTAN
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LINEZOLID, LINEZOLID
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TELMISARTAN, TELMISARTAN
 TETRABENAZINE, TETRABENAZINE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
 VALSARTAN, VALSARTAN

HEYL CHEMISCH*** HEYL CHEMISCH PHARMAZEUTISCHE FABRIK**

RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE(II)

HI TECH PHARMA*** HI TECH PHARMACAL CO INC**

ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACYCLOVIR, ACYCLOVIR
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 CALCIPOTRIENE, CALCIPOTRIENE
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CICLOPIROX, CICLOPIROX

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HI TECH PHARMACAL CO INC
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 CORMAX, CLOBETASOL PROPIONATE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 EMBELINE E, CLOBETASOL PROPIONATE
 EMBELINE, CLOBETASOL PROPIONATE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 GABAPENTIN, GABAPENTIN
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 LACTULOSE, LACTULOSE
 LEVOCARNITINE, LEVOCARNITINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 NYSTATIN, NYSTATIN
 OFLOXACIN, OFLOXACIN
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 VOSOL HC, ACETIC ACID, GLACIAL
 VOSOL, ACETIC ACID, GLACIAL

HI TECH PHARMA CO

* HI TECH PHARMACAL CO INC
 FLUNISOLIDE, FLUNISOLIDE
 PREDNISOLONE, PREDNISOLONE

HI-TECH PHARMA CO

* HI-TECH PHARMACAL CO INC
 BIMATOPROST, BIMATOPROST
 FAMOTIDINE, FAMOTIDINE
 GATIFLOXACIN, GATIFLOXACIN
 LORAZEPAM, LORAZEPAM
 PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE

HI-TECH PHARMACAL

* HI-TECH PHARMACAL CO INC
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BIMATOPROST, BIMATOPROST
 BROMFENAC SODIUM, BROMFENAC SODIUM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DESONIDE, DESONIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 IBUPROFEN, IBUPROFEN
 LEVETIRACETAM, LEVETIRACETAM
 MEGESTROL ACETATE, MEGESTROL ACETATE
 MORPHINE SULFATE, MORPHINE SULFATE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE
 TRIFLURIDINE, TRIFLURIDINE

HIGH TECH PHARMA

* HIGH TECHNOLOGY PHARMACAL CO INC
 VALPROIC ACID, VALPROIC ACID

HIKMA

* HIKMA FARMACEUTICA LDA
 CEFOTAXIME, CEFOTAXIME SODIUM
 * HIKMA PHARMACEUTICALS
 AMOXICILLIN, AMOXICILLIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******* HIKMA PHARMACEUTICALS**

CEFACLOR, CEFACLOR
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEPHALEXIN, CEPHALEXIN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 GLYBURIDE (MICRONIZED), GLYBURIDE

HIKMA FARMACEUTICA

- * HIKMA FARMACEUTICA (PORTUGAL) SA**
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CEFOXITIN, CEFOXITIN SODIUM
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 CIPROFLOXACIN, CIPROFLOXACIN
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 ENALAPRILAT, ENALAPRILAT
 FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 FLUMAZENIL, FLUMAZENIL
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MILRINONE LACTATE IN PLASTIC CONTAINER, MILRINONE LACTATE
 MILRINONE LACTATE, MILRINONE LACTATE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PROGESTERONE, PROGESTERONE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 VALPROATE SODIUM, VALPROATE SODIUM
- * HIKMA FARMACEUTICA PORTUGAL LDA**
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFUROXIME SODIUM, CEFUROXIME SODIUM
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
- * HIKMA FARMACEUTICA PORTUGAL SA**
 CEFOTETAN, CEFOTETAN DISODIUM
 CEFTRIAZONE SODIUM, CEFTRIAZONE SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 ETOMIDATE, ETOMIDATE
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OXYTOCIN, OXYTOCIN
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
- * HIKMA FARMACEUTICA SA**
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

HIKMA INTL PHARMS

- * HIKMA INTERNATIONAL PHARMACEUTICALS LLC**
 BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
 CAPTOPRIL, CAPTOPRIL
 CARISOPRODOL, CARISOPRODOL
 CORTISONE ACETATE, CORTISONE ACETATE
 DIGOXIN, DIGOXIN
 DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROCORTISONE, HYDROCORTISONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HIKMA INTERNATIONAL PHARMACEUTICALS LLC
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINOPRIL, LISINOPRIL
 METHOCARBAMOL, METHOCARBAMOL
 MITIGARE, COLCHICINE
 PRIMIDONE, PRIMIDONE

HIKMA PHARM CO LTD

* HIKMA PHARM CO LTD
 ARGATROBAN, ARGATROBAN

HIKMA PHARMS

* HIKMA PHARMACEUTICALS
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 GEMFIBROZIL, GEMFIBROZIL
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 LETROZOLE, LETROZOLE
 MODAFINIL, MODAFINIL
 PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 RIFAMPIN, RIFAMPIN
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

* HIKMA PHARMACEUTICALS CO LTD
 PARICALCITOL, PARICALCITOL

* HIKMA PHARMACEUTICALS LLC
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DOXERCALCIFEROL, DOXERCALCIFEROL
 FOLIC ACID, FOLIC ACID
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 OLANZAPINE, OLANZAPINE
 PIROXICAM, PIROXICAM
 PREDNISONE, PREDNISONE
 ZALEPLON, ZALEPLON

HILL DERMAC

* HILL DERMACEUTICALS INC
 DERMA-SMOOTH/FS, FLUOCINOLONE ACETONIDE
 DERMOTIC, FLUOCINOLONE ACETONIDE

HILL DERMACEUTICALS

* HILL DERMACEUTICALS INC
 TOLAK, FLUOROURACIL

HISAMITSU PHARM CO

* HISAMITSU PHARMACEUTICAL CO INC
 SALONPAS, MENTHOL (OTC)

HISUN PHARM HANGZHOU

* HISUN PHARMACEUTICAL (HANGZHOU) CO LTD
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE

* HISUN PHARMACEUTICAL HANGZHOU CO LTD
 CAPREOMYCIN SULFATE, CAPREOMYCIN SULFATE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM

HOFFMANN LA ROCHE

* HOFFMANN LA ROCHE INC
 BONIVA, IBANDRONATE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HOFFMANN LA ROCHE INC
 VALCYTE, VALGANCICLOVIR HYDROCHLORIDE
 XELODA, CAPECITABINE
 ZELBORAF, VEMURAFENIB

HOFFMANN-LA ROCHE

* HOFFMANN-LA ROCHE INC
 ALECENSA, ALECTINIB HYDROCHLORIDE
 INVIRASE, SAQUINAVIR MESYLATE

HONG KONG

* HONG KONG KING-FRIEND INDUSTRIAL CO LTD
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 CYTARABINE, CYTARABINE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE

HOPE PHARMS

* HOPE PHARMACEUTICALS
 NITHIODE, SODIUM NITRITE
 SODIUM NITRITE, SODIUM NITRITE
 SODIUM THIOSULFATE, SODIUM THIOSULFATE

HORIZON

* HORIZON MEDICINES LLC
 DUEXIS, FAMOTIDINE
 VIMOVO, ESOMEPRAZOLE MAGNESIUM

HORIZON PHARMA INC

* HORIZON PHARMA INC
 BUPHENYL, SODIUM PHENYLBUTYRATE

HORIZON PHARMA USA

* HORIZON PHARMA USA INC
 PROCYSBI, CYSTEAMINE BITARTRATE
 RAYOS, PREDNISONE

HORIZON THERAPY INC

* HORIZON THERAPEUTICS INC
 RAVICTI, GLYCEROL PHENYLBUTYRATE

HOSPIRA

* HOSPIRA INC
 A-METHAPRED, METHYLPREDNISOLONE SODIUM SUCCINATE
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ALFENTANIL, ALFENTANIL HYDROCHLORIDE
 AMIDATE, ETOMIDATE
 AMINOCAPROIC ACID IN PLASTIC CONTAINER, AMINOCAPROIC ACID
 AMINOPHYLLINE, AMINOPHYLLINE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMMONIUM CHLORIDE IN PLASTIC CONTAINER, AMMONIUM CHLORIDE
 ARTICAIN HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, ARTICAIN HYDROCHLORIDE
 ATROPINE SULFATE ANSYR PLASTIC SYRINGE, ATROPINE SULFATE
 ATROPINE SULFATE LIFESHIELD ABBOJECT SYRINGE, ATROPINE SULFATE
 AZITHROMYCIN, AZITHROMYCIN
 BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BUMETANIDE, BUMETANIDE
 BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE W/EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 CARBOCAINE, MEPIVACAINE HYDROCHLORIDE
 CARBOPLATIN, CARBOPLATIN
 CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
 CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 CIPROFLOXACIN, CIPROFLOXACIN
 CORLOPAM, FENOLDOPAM MESYLATE
 CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE
 CYTARABINE, CYTARABINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******* HOSPIRA INC**

DACARBAZINE, DACARBAZINE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DEMEROL, MEPERIDINE HYDROCHLORIDE
 DEXTROSE 25%, DEXTROSE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 50%, DEXTROSE
 DIAZEPAM, DIAZEPAM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
 DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
 DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
 DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
 DROPERIDOL, DROPERIDOL
 ENALAPRILAT, ENALAPRILAT
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 ERYTHROCIN, ERYTHROMYCIN LACTOBIONATE
 FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
 FENTANYL CITRATE, FENTANYL CITRATE
 FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 FUROSEMIDE, FUROSEMIDE
 GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEVOPHED, NOREPINEPHRINE BITARTRATE
 LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE
 LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LORAZEPAM, LORAZEPAM
 M.V.I. ADULT (PHARMACY BULK PACKAGE), ASCORBIC ACID
 M.V.I. ADULT, ASCORBIC ACID
 M.V.I. PEDIATRIC, ASCORBIC ACID
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MANGANESE CHLORIDE IN PLASTIC CONTAINER, MANGANESE CHLORIDE
 MANNITOL 25%, MANNITOL
 MARCAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE HYDROCHLORIDE W/ EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 MARCAINE, BUPIVACAINE HYDROCHLORIDE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******* HOSPIRA INC**

METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
MORPHINE SULFATE, MORPHINE SULFATE
NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE
NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
PACLITAXEL, PACLITAXEL
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
PLEGISOL IN PLASTIC CONTAINER, CALCIUM CHLORIDE
POTASSIUM ACETATE, POTASSIUM ACETATE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PRECEDEX, DEXMEDETOMIDINE HYDROCHLORIDE
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
PROPOFOL, PROPOFOL
QUELICIN, SUCCINYLCHOLINE CHLORIDE
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
SODIUM ACETATE, SODIUM ACETATE
SODIUM BICARBONATE, SODIUM BICARBONATE
SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM LACTATE IN PLASTIC CONTAINER, SODIUM LACTATE
SODIUM PHOSPHATES IN PLASTIC CONTAINER, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
SUFENTANIL CITRATE, SUFENTANIL CITRATE
TAZICEF, CEFTAZIDIME
TOBRAMYCIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, TOBRAMYCIN SULFATE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
TPN ELECTROLYTES IN PLASTIC CONTAINER, CALCIUM CHLORIDE
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
VECURONIUM BROMIDE, VECURONIUM BROMIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
VINORELBINE TARTRATE, VINORELBINE TARTRATE
VITAMIN K1, PHYTONADIONE
ZINC CHLORIDE IN PLASTIC CONTAINER, ZINC CHLORIDE

*** HOSPIRA WORLDWIDE, INC**

DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
NITROPRESS, SODIUM NITROPRUSSIDE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE

HOSPIRA INC*** HOSPIRA INC**

ADENOSINE, ADENOSINE
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
AMPICILLIN SODIUM, AMPICILLIN SODIUM
ARGATROBAN, ARGATROBAN
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
BIVALIRUDIN, BIVALIRUDIN
BORTEZOMIB, BORTEZOMIB
BUSULFAN, BUSULFAN
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
CEFOTAXIME SODIUM, CEFOTAXIME SODIUM
CEFOXITIN, CEFOXITIN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******* HOSPIRA INC**

CEFTRIAZONE, CEFTRIAZONE SODIUM
 CEFUROXIME SODIUM, CEFUROXIME SODIUM
 CLOFARABINE, CLOFARABINE
 DAPTOMYCIN, DAPTOMYCIN
 DOCETAXEL, DOCETAXEL
 DOXERCALCIFEROL, DOXERCALCIFEROL
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 HEPARIN SODIUM, HEPARIN SODIUM
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
 INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LINEZOLID IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, LINEZOLID
 LINEZOLID, LINEZOLID
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MAXIPIME, CEFEPIME HYDROCHLORIDE
 MEROPENEM, MEROPENEM
 MILRINONE LACTATE, MILRINONE LACTATE
 MORPHINE SULFATE, MORPHINE SULFATE
 NIPENT, PENTOSTATIN
 OXACILLIN SODIUM, OXACILLIN SODIUM
 OXALIPLATIN, OXALIPLATIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PARICALCITOL, PARICALCITOL
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 SODIUM BICARBONATE, SODIUM BICARBONATE
 TACROLIMUS, TACROLIMUS
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

HOSPIRA WORLDWIDE

*** HOSPIRA WORLDWIDE PTY**
 OXALIPLATIN, OXALIPLATIN

HOT SHOTS NM LLC

*** HOT SHOTS NUCLEAR MEDICINE LLC**
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

HOUSTON CYCLOTRON

*** HOUSTON CYCLOTRON PARTNERS LP**
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

HQ SPCLT PHARMA

*** HQ SPECIALTY PHARMA CORP**
 CALCIUM GLUCONATE IN SODIUM CHLORIDE, CALCIUM GLUCONATE
 CISPLATIN, CISPLATIN
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 LINEZOLID, LINEZOLID
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 TAXOL, PACLITAXEL

HQ SPECIALITY PHARMA

*** HQ SPECIALITY PHARMA LLC**
 LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM

HUMANWELL PURACAP

*** HUMANWELL PURACAP PHARMACEUTICAL WUHAN CO LTD**
 DUTASTERIDE, DUTASTERIDE
 IBUPROFEN, IBUPROFEN (OTC)

HZNP

*** HZNP MEDICINES LLC**
 MIGERGOT, CAFFEINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

* HZNP MEDICINES LLC
PENNSAID, DICLOFENAC SODIUM

ROCHE

* HOFFMANN LA ROCHE INC
BONIVA, IBANDRONATE SODIUM
FUZEON, ENFUVIRTIDE
KLONOPIN, CLONAZEPAM
TAMIFLU, OSELTAMIVIR PHOSPHATE
VALIUM, DIAZEPAM

**** I ******IBSA INST BIO**

* IBSA INSTITUT BIOCHIMIQUE SA
LICART, DICLOFENAC EPOLAMINE

ICU MEDICAL INC

* ICU MEDICAL INC
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
AMINOSYN 10%, AMINO ACIDS
AMINOSYN 3.5% M, AMINO ACIDS
AMINOSYN 8.5% W/ELECTROLYTES, AMINO ACIDS
AMINOSYN 8.5%, AMINO ACIDS
AMINOSYN II 10% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 15% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN-PF 10%, AMINO ACIDS
AMINOSYN-PF 7%, AMINO ACIDS
DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 20% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 30% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 40% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL
MORPHINE SULFATE, MORPHINE SULFATE
NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
NORMOSOL-R IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PHYSIOSOL IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
POTASSIUM CHLORIDE 0.149% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, POTASSIUM
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******* ICU MEDICAL INC**

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER,
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SORBITOL-MANNITOL IN PLASTIC CONTAINER, MANNITOL
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION

IDENTI PHARMS INC*** IDENTI PHARMACEUTICALS INC**

FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE

IDT AUSTRALIA LTD*** IDT AUSTRALIA LTD**

TEMOZOLOMIDE, TEMOZOLOMIDE

IMPAX*** IMPAX LABORATORIES LLC**

ADRENACLICK, EPINEPHRINE

IMPAX LABS*** IMPAX LABORATORIES INC**

ACARBOSE, ACARBOSE
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 BACLOFEN, BACLOFEN
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DIGOXIN, DIGOXIN
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRATE, FENOFIBRATE
 FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 METHYLTESTOSTERONE, METHYLTESTOSTERONE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 NADOLOL AND BENDROFLUMETHIAZIDE, BENDROFLUMETHIAZIDE
 OMEPRAZOLE, OMEPRAZOLE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RILUZOLE, RILUZOLE
 RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE

IMPAX LABS INC*** IMPAX LABORATORIES INC**

ACITRETIN, ACITRETIN
 ALBENZA, ALBENDAZOLE
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 BUDESONIDE, BUDESONIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE
 COLESEVELAM HYDROCHLORIDE, COLESEVELAM HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

* IMPAX LABORATORIES INC
 DEXEDRINE, DEXTROAMPHETAMINE SULFATE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE
 EMVERM, MEBENDAZOLE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYBURIDE, GLYBURIDE
 HYDROCORTISONE, HYDROCORTISONE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 LAMOTRIGINE, LAMOTRIGINE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METHYLTESTOSTERONE, METHYLTESTOSTERONE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 MORPHINE SULFATE, MORPHINE SULFATE
 NABUMETONE, NABUMETONE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RYTARY, CARBIDOPA
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 URSODIOL, URSODIOL

IMPAX PHARMS

* IMPAX PHARMACEUTICALS
 GEMFIBROZIL, GEMFIBROZIL
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

INCYTE CORP

* INCYTE CORP
 JAKAFI, RUXOLITINIB PHOSPHATE

INDICUS PHARMA

* INDICUS PHARMA LLC
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 LETROZOLE, LETROZOLE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

INDIVIOR INC

* INDIVIOR INC
 BUPRENEX, BUPRENORPHINE HYDROCHLORIDE
 PERSERIS KIT, RISPERIDONE
 SUBLOCADE, BUPRENORPHINE
 SUBOXONE, BUPRENORPHINE HYDROCHLORIDE

INDOCO REMEDIES

* INDOCO REMEDIES LTD
 ALLOPURINOL, ALLOPURINOL
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 GLIMEPIRIDE, GLIMEPIRIDE

INFORLIFE

* INFORLIFE SA
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

INGENUS PHARMS LLC

* INGENUS PHARMACEUTICALS LLC
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 CABERGOLINE, CABERGOLINE
 CARBOPLATIN, CARBOPLATIN
 CICLOPIROX, CICLOPIROX
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

- * INGENUS PHARMACEUTICALS LLC
CLOFARABINE, CLOFARABINE
DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
DOCETAXEL, DOCETAXEL
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
OXALIPLATIN, OXALIPLATIN
TOLCAPONE, TOLCAPONE
- INGENUS PHARMS NJ**
- * INGENUS PHARMACEUTICALS NJ LLC
CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN
- INNOGENIX**
- * INNOGENIX LLC
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
METRONIDAZOLE, METRONIDAZOLE
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
- INSMED INC**
- * INSMED INC
ARIKAYCE KIT, AMIKACIN SULFATE
- INST BIOCHEM**
- * INSTITUT BIOCHEMIQUE SA
FLECTOR, DICLOFENAC EPOLAMINE
- INSTITUT BIOCHIMIQUE**
- * INSTITUT BIOCHIMIQUE SA (IBSA)
TIROSINT, LEVOTHYROXINE SODIUM
- INSYS DEV CO INC**
- * INSYS DEVELOPMENT CO INC
SUBSYS, FENTANYL
SYNDROS, DRONABINOL
- INTAS PHARMS USA**
- * INTAS PHARMACEUTICALS USA
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
- INTELLIPHARMACEUTICS**
- * INTELLIPHARMACEUTICS CORP
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
- INTERCEPT PHARMS INC**
- * INTERCEPT PHARMACEUTICALS INC
OCALIVA, OBETICHOLIC ACID
- INTERGEL PHARM**
- * INTERGEL PHARMACEUTICAL INC
NIFEDIPINE, NIFEDIPINE
- INTERGEL PHARMS INC**
- * INTERGEL PHARMACEUTICALS INC
DUTASTERIDE, DUTASTERIDE
- INTERPHARMA PRAHA AS**
- * INTERPHARMA PRAHA AS
ORALTAG, IOHEXOL
- INTERSECT ENT INC**
- * INTERSECT ENT INC
SINUVA, MOMETASONE FUROATE
- INTL MEDICATED**
- * INTERNATIONAL MEDICATED SYSTEMS LTD
MILRINONE LACTATE, MILRINONE LACTATE
- INTL MEDICATION**
- * INTERNATIONAL MEDICATION SYSTEM
LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
MANNITOL 25%, MANNITOL
NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

- * INTERNATIONAL MEDICATION SYSTEM
PHYTONADIONE, PHYTONADIONE
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
- * INTERNATIONAL MEDICATION SYSTEMS LTD
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE

INTL MEDICATION SYS

- * INTERNATIONAL MEDICATION SYSTEMS LTD
CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
LORAZEPAM, LORAZEPAM
SODIUM BICARBONATE, SODIUM BICARBONATE

INVAGEN PHARMS

- * INVAGEN PHARMACEUTICALS INC
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CALCIUM ACETATE, CALCIUM ACETATE
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
GABAPENTIN, GABAPENTIN
GEMFIBROZIL, GEMFIBROZIL
GLIMEPIRIDE, GLIMEPIRIDE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LISINOPRIL, LISINOPRIL
LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
MEPROBAMATE, MEPROBAMATE
NABUMETONE, NABUMETONE
NADOLOL, NADOLOL
NAPROXEN, NAPROXEN
OLANZAPINE, OLANZAPINE
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SEVELAMER CARBONATE, SEVELAMER CARBONATE
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
TOPIRAMATE, TOPIRAMATE
TROSPIUM CHLORIDE, TROSPIUM CHLORIDE
WARFARIN SODIUM, WARFARIN SODIUM
ZOLMITRIPTAN, ZOLMITRIPTAN
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
ZONISAMIDE, ZONISAMIDE

INVATECH PHARMA

- * INVATECH PHARMA SOLUTIONS LLC
CALCITRIOL, CALCITRIOL
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE

INVENTIA HLTHCARE

- * INVENTIA HEALTHCARE PRIVATE LTD
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
ILOPERIDONE, ILOPERIDONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

* INVENTIA HEALTHCARE PRIVATE LTD
 LANSOPRAZOLE, LANSOPRAZOLE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 TELMISARTAN, TELMISARTAN

IONETIX

* IONETIX CORP
 AMMONIA N 13, AMMONIA N-13

IPCA LABS LTD

* IPCA LABORATORIES LTD
 ALLOPURINOL, ALLOPURINOL
 ATENOLOL, ATENOLOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 FUROSEMIDE, FUROSEMIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM

IPR

* IPR PHARMACEUTICALS INC
 CRESTOR, ROSUVASTATIN CALCIUM
 ZOMIG, ZOLMITRIPTAN

IPSEN INC

* IPSEN BIOPHARMACEUTICALS INC
 INCRELEX, MECASERMIN RECOMBINANT
 ONIVYDE, IRINOTECAN HYDROCHLORIDE

IPSEN PHARMA

* IPSEN PHARMA BIOTECH SAS
 SOMATULINE DEPOT, LANREOTIDE ACETATE

IROKO PHARMS

* IROKO PHARMACEUTICALS LLC
 INDOCIN, INDOMETHACIN

IROKO PHARMS LLC

* IROKO PHARMACEUTICALS LLC
 TIVORBEX, INDOMETHACIN
 VIVLODEX, MELOXICAM
 ZORVOLEX, DICLOFENAC

IRONSHORE PHARMS

* IRONSHORE PHARMACEUTICALS AND DEVELOPMENT INC
 JORNAY PM, METHYLPHENIDATE HYDROCHLORIDE

IRONWOOD PHARMS INC

* IRONWOOD PHARMACEUTICALS INC
 DUZALLO, ALLOPURINOL
 ZURAMPIC, LESINURAD

ISO TEX

* ISO TEX DIAGNOSTICS INC
 JEANATOPE, ALBUMIN IODINATED I-125 SERUM
 MEGATOPE, ALBUMIN IODINATED I-131 SERUM

ISOTEX

* ISOTEX DIAGNOSTICS
 GLOFIL-125, IOTHALAMATE SODIUM I-125

ISTITUTO BIO ITA SPA

* ISTITUTO BIOCHIMICO ITALIANO SPA
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

* ISTITUTO BIOCHIMICO ITALIANO SPA
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PIPERACILLIN, PIPERACILLIN SODIUM

ITALFARMACO SPA

* ITALFARMACO SPA
 TIGLUTIK KIT, RILUZOLE

IVAX PHARMS

* IVAX PHARMACEUTICALS INC
 VALSARTAN, VALSARTAN

IVAX PHARMS INC

* IVAX PHARMACEUTICALS INC
 OLANZAPINE, OLANZAPINE

IVAX SUB TEVA PHARMS

* IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 BACLOFEN, BACLOFEN
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BUMETANIDE, BUMETANIDE
 CABERGOLINE, CABERGOLINE
 CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CIMETIDINE, CIMETIDINE (OTC)
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLOZAPINE, CLOZAPINE
 CYCLOSPORINE, CYCLOSPORINE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DIAZEPAM, DIAZEPAM
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FLUCONAZOLE, FLUCONAZOLE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FUROSEMIDE, FUROSEMIDE
 GABAPENTIN, GABAPENTIN
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 INDOMETHACIN, INDOMETHACIN
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINAPRIL, LISINAPRIL
 METHYLDOPA, METHYLDOPA
 MISOPROSTOL, MISOPROSTOL
 NADOLOL, NADOLOL
 OXAPROZIN, OXAPROZIN
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

**** J ******J AND J CONSUMER INC**

* JOHNSON AND JOHNSON CONSUMER INC MCNEIL CONSUMER HEALTHCARE DIVISION
 CHILDREN'S MOTRIN COLD, IBUPROFEN (OTC)
 CHILDREN'S MOTRIN, IBUPROFEN (OTC)
 CHILDREN'S ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S ZYRTEC HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 IMODIUM A-D, LOPERAMIDE HYDROCHLORIDE (OTC)
 IMODIUM MULTI-SYMPTOM RELIEF, LOPERAMIDE HYDROCHLORIDE (OTC)
 JUNIOR STRENGTH MOTRIN, IBUPROFEN (OTC)
 MOTRIN IB, IBUPROFEN (OTC)
 PEPCID AC, FAMOTIDINE (OTC)
 PEPCID COMPLETE, CALCIUM CARBONATE (OTC)
 SINE-AID IB, IBUPROFEN (OTC)
 SUDAFED 24 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ****

* JOHNSON AND JOHNSON CONSUMER INC MCNEIL CONSUMER HEALTHCARE DIVISION
 TYLENOL, ACETAMINOPHEN (OTC)
 ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 ZYRTEC-D 12 HOUR, CETIRIZINE HYDROCHLORIDE (OTC)

JACOBUS

* JACOBUS PHARMACEUTICAL CO
 DAPSONE, DAPSONE
 PASER, AMINOSALICYLIC ACID

JANSSEN BIOTECH

* JANSSEN BIOTECH INC
 ERLEADA, APALUTAMIDE
 ZYTIGA, ABIRATERONE ACETATE

JANSSEN PHARMS

* JANSSEN PHARMACEUTICALS INC
 AXERT, ALMOTRIPTAN MALATE
 CONCERTA, METHYLPHENIDATE HYDROCHLORIDE
 DITROPAN XL, OXYBUTYNIN CHLORIDE
 DURAGESIC-100, FENTANYL
 DURAGESIC-12, FENTANYL
 DURAGESIC-25, FENTANYL
 DURAGESIC-37, FENTANYL
 DURAGESIC-50, FENTANYL
 DURAGESIC-75, FENTANYL
 ELMIRON, PENTOSAN POLYSULFATE SODIUM
 HALDOL, HALOPERIDOL DECANOATE
 HALDOL, HALOPERIDOL LACTATE
 INVEGA SUSTENNA, PALIPERIDONE PALMITATE
 INVEGA TRINZA, PALIPERIDONE PALMITATE
 INVEGA, PALIPERIDONE
 INVOKAMET XR, CANAGLIFLOZIN
 INVOKAMET, CANAGLIFLOZIN
 INVOKANA, CANAGLIFLOZIN
 MICRONOR, NORETHINDRONE
 NIZORAL, KETOCONAZOLE
 ORTHO CYCLEN-28, ETHINYL ESTRADIOL
 ORTHO TRI-CYCLEN LO, ETHINYL ESTRADIOL
 ORTHO TRI-CYCLEN, ETHINYL ESTRADIOL
 ORTHO-NOVUM 1/35-28, ETHINYL ESTRADIOL
 ORTHO-NOVUM 7/7/7-28, ETHINYL ESTRADIOL
 RAZADYNE ER, GALANTAMINE HYDROBROMIDE
 RAZADYNE, GALANTAMINE HYDROBROMIDE
 RISPERDAL CONSTA, RISPERIDONE
 RISPERDAL, RISPERIDONE
 SPORANOX, ITRACONAZOLE
 TOPAMAX, TOPIRAMATE
 TYLENOL W/ CODEINE NO. 3, ACETAMINOPHEN
 TYLENOL W/ CODEINE NO. 4, ACETAMINOPHEN
 ULTRACET, ACETAMINOPHEN
 ULTRAM, TRAMADOL HYDROCHLORIDE
 XARELTO, RIVAROXABAN

JANSSEN PRODS

* JANSSEN PRODUCTS LP
 EDURANT, RILPIVIRINE HYDROCHLORIDE
 PREZCOBIX, COBICISTAT
 PREZISTA, DARUNAVIR ETHANOLATE
 SYMTUZA, COBICISTAT
 YONDELIS, TRABECTEDIN

JANSSEN R AND D

* JANSSEN RESEARCH AND DEVELOPMENT LLC
 INTELENCE, ETRAVIRINE

JANSSEN RES AND DEV

* JANSSEN RESEARCH AND DEVELOPMENT LLC
 DOXIL (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE

JANSSEN THERAP

* JANSSEN THERAPEUTICS DIV JANSSEN PRODUCTS LP

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ****

* JANSSEN THERAPEUTICS DIV JANSSEN PRODUCTS LP
SIRTURO, BEDAQUILINE FUMARATE

JAZZ PHARMS

* JAZZ PHARMACEUTICALS INC
XYREM, SODIUM OXYBATE

JAZZ PHARMS III

* JAZZ PHARMACEUTICALS III INTERNATIONAL LTD
FAZACLO ODT, CLOZAPINE

JAZZ PHARMS INC

* JAZZ PHARMACEUTICALS INC
DEFITELIO, DEFIBROTIDE SODIUM

JIANGSU HANSOH PHARM

* JIANGSU HANSOH PHARMACEUTICAL GROUP CO LTD
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
OLANZAPINE, OLANZAPINE
VINORELBINE TARTRATE, VINORELBINE TARTRATE

JIANGSU HENGRUI MED

* JIANGSU HENGRUI MEDICINE CO LTD
CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
DOCETAXEL, DOCETAXEL
FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
GABAPENTIN, GABAPENTIN
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
LETROZOLE, LETROZOLE
OXALIPLATIN, OXALIPLATIN
THIOTEPA, THIOTEPA

JOHNS HOPKINS UNIV

* JOHNS HOPKINS UNIV
AMMONIA N 13, AMMONIA N-13

JOHNSON AND JOHNSON

* JOHNSON AND JOHNSON CONSUMER INC
VISINE L.R., OXYMETAZOLINE HYDROCHLORIDE (OTC)
VISINE, NAPHAZOLINE HYDROCHLORIDE (OTC)
* JOHNSON AND JOHNSON GROUP CONSUMER COMPANIES
MEN'S ROGAINE, MINOXIDIL (OTC)
ROGAINE (FOR MEN), MINOXIDIL (OTC)
ROGAINE (FOR WOMEN), MINOXIDIL (OTC)
ROGAINE EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
WOMEN'S ROGAINE, MINOXIDIL (OTC)
* JOHNSON AND JOHNSON HEALTHCARE PRODUCTS DIV MCNEIL-PPC INC
NIZORAL A-D, KETOCONAZOLE (OTC)

JOURNEY

* JOURNEY MEDICAL CORP
EXELDERM, SULCONAZOLE NITRATE

JUBILANT CADISTA

* JUBILANT CADISTA PHARMACEUTICALS INC
ALENDRONATE SODIUM, ALENDRONATE SODIUM
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LAMOTRIGINE, LAMOTRIGINE
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
METHYLPREDNISOLONE, METHYLPREDNISOLONE
PREDNISONE, PREDNISONE
PROCOMP, PROCHLORPERAZINE MALEATE
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE

JUBILANT DRAXIMAGE

* JUBILANT DRAXIMAGE INC
DRAX EXAMETAZIME, TECHNETIUM TC-99M EXAMETAZIME KIT
DRAXIMAGE MDP-25, TECHNETIUM TC-99M MEDRONATE
DTPA, TECHNETIUM TC-99M PENTETATE KIT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ****

- * JUBILANT DRAXIMAGE INC
HICON, SODIUM IODIDE I-131
RUBY-FILL, RUBIDIUM CHLORIDE RB-82
SODIUM IODIDE I 131, SODIUM IODIDE I-131
- * JUBILANT DRAXIMAGE RADIOPHARMACIES INC
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18
- * JUBILANT DRAXIMAGE USA INC
TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT

JUBILANT GENERICS

- * JUBILANT GENERICS LTD
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CELECOXIB, CELECOXIB
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
FELODIPINE, FELODIPINE
INDOMETHACIN, INDOMETHACIN
IRBESARTAN, IRBESARTAN
ITRACONAZOLE, ITRACONAZOLE
LAMOTRIGINE, LAMOTRIGINE
LEVETIRACETAM, LEVETIRACETAM
LEVOFLOXACIN, LEVOFLOXACIN
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
MONTELUKAST SODIUM, MONTELUKAST SODIUM
NIACIN, NIACIN
OLANZAPINE, OLANZAPINE
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
RISPERIDONE, RISPERIDONE
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
SPIRONOLACTONE, SPIRONOLACTONE
TELMISARTAN, TELMISARTAN
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
VALSARTAN, VALSARTAN
ZOLMITRIPTAN, ZOLMITRIPTAN

JUBILANT HOLLISTRSTR

- * JUBILANT HOLLISTERSTIER LLC
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

STEVENS J

- * JEROME STEVENS PHARMACEUTICALS INC
BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
DIGOXIN, DIGOXIN
METHOCARBAMOL AND ASPIRIN, ASPIRIN
UNITHROID, LEVOTHYROXINE SODIUM **

**** K ******GRIFFEN**

- * KW GRIFFEN CO
BIOSCRUB, CHLORHEXIDINE GLUCONATE (OTC)

KADMON PHARMS LLC

- * KADMON PHARMACEUTICALS LLC
RIBASPHERE, RIBAVIRIN
RIBAVIRIN, RIBAVIRIN

KAI PHARMS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** K ****

* KAI PHARMACEUTICALS INC A WHOLLY OWNED SUBSIDIARY OF AMGEN INC
PARSABIV, ETELCALCETIDE

KALA PHARMS INC

* KALA PHARMACEUTICALS INC
INVELTYS, LOTEPIREDNOL ETABONATE

KALEO INC

* KALEO INC
AUVI-Q, EPINEPHRINE
EVZIO, NALOXONE HYDROCHLORIDE

KEMPHARM

* KEMPHARM INC
APADAZ, ACETAMINOPHEN

KERYX BIOPHARMS

* KERYX BIOPHARMACEUTICALS INC
AURYXIA, FERRIC CITRATE

KETTERING MEDCTR

* KETTERING MEDCTR
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

KING PHARMS

* KING PHARMACEUTICALS INC
SYNERCID, DALFOPRISTIN

* KING PHARMACEUTICALS RESEARCH AND DEVELOPMENT LLC
CYTOMEL, LIOETHYRONINE SODIUM
LEVOXYL, LEVOTHYROXINE SODIUM **
TUSSIGON, HOMATROPINE METHYLBROMIDE

* KING PHARMACEUTICALS RESEARCH AND DEVELOPMENT LLC A SUB OF PFIZER INC
SKELAXIN, METAXALONE

KING PHARMS LLC

* KING PHARMACEUTICALS LLC
ALTACE, RAMIPRIL
BICILLIN C-R 900/300, PENICILLIN G BENZATHINE
BICILLIN C-R, PENICILLIN G BENZATHINE
BICILLIN L-A, PENICILLIN G BENZATHINE
CORZIDE, BENDROFLUMETHIAZIDE
PENICILLIN G PROCAINE, PENICILLIN G PROCAINE
SILVADENE, SILVER SULFADIAZINE
TAPAZOLE, METHIMAZOLE
TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE

KITOV PHARMS LTD

* KITOV PHARMACEUTICALS LTD
CONSENSI, AMLODIPINE BESYLATE

KNIGHT THERAPS

* KNIGHT THERAPEUTICS USA INC
IMPAVIDO, MILTEFOSINE

KOWA CO

* KOWA CO LTD
LIVALO, PITAVASTATIN CALCIUM

KREITCHMAN PET CTR

* KREITCHMAN PET CENTER
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

KRKA TOVARNA ZDRAVIL

* KRKA TOVARNA ZDRAVIL DD NOVO MESTO
LANSOPRAZOLE, LANSOPRAZOLE

KVK TECH

* KVK TECH INC
BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
KALEXATE, SODIUM POLYSTYRENE SULFONATE
PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** K ****

* KVK TECH INC
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

KVK TECH INC

* KVK TECH INC
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

KYOWA KIRIN

* KYOWA KIRIN INC
FARESTON, TOREMIFENE CITRATE
SANCUSO, GRANISETRON

KYTHERA BIOPHARMS

* KYTHERA BIOPHARMACEUTICALS INC
KYBELLA, DEOXYCHOLIC ACID

**** L ******L PERRIGO CO**

* L PERRIGO CO
CIMETIDINE, CIMETIDINE (OTC)
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE (OTC)
IBUPROFEN, IBUPROFEN (OTC)
JUNIOR STRENGTH IBUPROFEN, IBUPROFEN (OTC)
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)

LA JOLLA PHARMA

* LA JOLLA PHARMA LLC
GIAPREZA, ANGIOTENSIN II ACETATE

LAB HRA PHARMA

* LABORATOIRE HRA PHARMA
ELLA, ULIPRISTAL ACETATE

LABORATOIRE HRA

* LABORATOIRE HRA PHARMA
LYSODREN, MITOTANE

LABORATORIE HRA

* LABORATORIE HRA PHARMA
METOPIRONE, METYRAPONE

LABORATORIOS GRIFOLS

* LABORATORIOS GRIFOLS SA
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

LABORATORIOS SALVAT

* LABORATORIOS SALVAT SA
OTOVEL, CIPROFLOXACIN HYDROCHLORIDE

LABS LEON FARMA

* LABORATORIOS LEON FARMA SA
ALTAVERA, ETHINYL ESTRADIOL
ELIFEMME, ETHINYL ESTRADIOL
ESTARYLLA, ETHINYL ESTRADIOL
INTROVALE, ETHINYL ESTRADIOL
ISIBLOOM, DESOGESTREL
JAIMIESS, ETHINYL ESTRADIOL
LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
LORYNA, DROSPIRENONE
SYEDA, DROSPIRENONE
TRI-ESTARYLLA, ETHINYL ESTRADIOL
TRI-LO-ESTARYLLA, ETHINYL ESTRADIOL
VIENVA, ETHINYL ESTRADIOL
VOLNEA, DESOGESTREL

LABS LICONSA

* LABORATORIOS LICONSA SA
LANSOPRAZOLE, LANSOPRAZOLE

LANDELA PHARM

* LANDELA PHARMACEUTICAL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LANDELA PHARMACEUTICAL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE

LANNETT

* LANNETT CO INC
 ACETAZOLAMIDE, ACETAZOLAMIDE
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 LANIAZID, ISONIAZID
 LANORINAL, ASPIRIN
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PRIMIDONE, PRIMIDONE
 PROBALAN, PROBENECID

LANNETT CO INC

* LANNETT CO INC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACETIC ACID, ACETIC ACID, GLACIAL
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BACLOFEN, BACLOFEN
 BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CODEINE SULFATE, CODEINE SULFATE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DANAZOL, DANAZOL
 DEXAMETHASONE, DEXAMETHASONE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DIAZEPAM, DIAZEPAM
 DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DRONABINOL, DRONABINOL
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 GLYCOLAX, POLYETHYLENE GLYCOL 3350
 GLYCOLAX, POLYETHYLENE GLYCOL 3350 (OTC)
 HALOPERIDOL, HALOPERIDOL LACTATE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCORTISONE, HYDROCORTISONE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 LACTULOSE, LACTULOSE
 LAMIVUDINE, LAMIVUDINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 LIDOCAINE HYDROCHLORIDE VISCOUS, LIDOCAINE HYDROCHLORIDE
 LOPINAVIR AND RITONAVIR, LOPINAVIR
 LORATADINE, LORATADINE (OTC)
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METADATE CD, METHYLPHENIDATE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******* LANNETT CO INC**

METADATE ER, METHYLPHENIDATE HYDROCHLORIDE
 METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
 METAXALONE, METAXALONE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MONOKET, ISOSORBIDE MONONITRATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MORPHINE SULFATE, MORPHINE SULFATE
 NEOMYCIN SULFATE, NEOMYCIN SULFATE
 NIACIN, NIACIN
 NYSTATIN, NYSTATIN
 OMEPRAZOLE, OMEPRAZOLE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREDNISOLONE, PREDNISOLONE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RIFAMPIN, RIFAMPIN
 RISPERIDONE, RISPERIDONE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SUMATRIPTAN, SUMATRIPTAN
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 THEOPHYLLINE, THEOPHYLLINE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 URSODIOL, URSODIOL
 VALPROIC ACID, VALPROIC ACID
 ZAROXOLYN, METOLAZONE

LANTHEUS MEDCL*** LANTHEUS MEDICAL IMAGING INC**

CARDIOLITE, TECHNETIUM TC-99M SESTAMIBI KIT
 DEFINITY, PERFLUTREN
 GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67
 NEUROLITE, TECHNETIUM TC-99M BICISATE KIT
 TECHNELITE, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
 XENON XE 133, XENON XE-133

LANTHEUS MEDICAL*** LANTHEUS MEDICAL IMAGING INC**

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 QUADRAMET, SAMARIUM SM-153 LEXIDRONAM PENTASODIUM

LARKEN LABS*** LARKEN LABORATORIES INC**

DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 OFLOXACIN, OFLOXACIN

LARKEN LABS INC*** LARKEN LABORATORIES INC**

ACETAMINOPHEN, CAFFEINE AND DIHYDROCODEINE BITARTRATE, ACETAMINOPHEN
 ALLZITAL, ACETAMINOPHEN
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 DEXAMETHASONE, DEXAMETHASONE

LAURUS LABS LTD*** LAURUS LABS LTD**

METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

LAVIPHARM LABS*** LAVIPHARM LABORATORIES INC**

FENTANYL-100, FENTANYL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LAVIPHARM LABORATORIES INC
 FENTANYL-25, FENTANYL
 FENTANYL-50, FENTANYL
 FENTANYL-75, FENTANYL

LEADIANT BIOSCI INC

* LEADIANT BIOSCIENCES INC
 ABELCET, AMPHOTERICIN B
 ADAGEN, PEGADEMASE BOVINE
 CARNITOR SF, LEVOCARNITINE
 CARNITOR, LEVOCARNITINE
 CYSTARAN, CYSTEAMINE HYDROCHLORIDE
 MATULANE, PROCARBAZINE HYDROCHLORIDE

LEADING PHARMA LLC

* LEADING PHARMA LLC
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 FOLIC ACID, FOLIC ACID
 FUROSEMIDE, FUROSEMIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 LORAZEPAM, LORAZEPAM
 NIFEDIPINE, NIFEDIPINE

LEO LABS

* LEO LABORATORIES LTD
 PICATO, INGENOL MEBUTATE

LEO PHARMA AS

* LEO PHARMA AS
 DESONATE, DESONIDE
 DOVONEX, CALCIPOTRIENE
 ENSTILAR, BETAMETHASONE DIPROPIONATE
 FINACEA, AZELAIC ACID
 PROTOPIC, TACROLIMUS
 TACLONEX, BETAMETHASONE DIPROPIONATE

LEXICON PHARMS INC

* LEXICON PHARMACEUTICALS INC
 XERMELO, TELOTRISTAT ETIPRATE

LG CHEM LTD

* LG CHEM LTD
 FACTIVE, GEMIFLOXACIN MESYLATE

LIEBEL-FLARSHEIM

* LIEBEL-FLARSHEIM CO LLC
 CONRAY 43, IOTHALAMATE MEGLUMINE
 CONRAY, IOTHALAMATE MEGLUMINE
 CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
 MD-GASTROVIEW, DIATRIZOATE MEGLUMINE
 OPTIRAY 240, IOVERSOL
 OPTIRAY 300, IOVERSOL
 OPTIRAY 320, IOVERSOL
 OPTIRAY 350, IOVERSOL
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

LIFE MOLECULAR

* LIFE MOLECULAR IMAGING SA
 NEURACEQ, FLORBETABEN F-18

LIFEPHARMA

* LIFEPHARMA FZE
 LACTULOSE, LACTULOSE

LNK

* LNK INTERNATIONAL INC
 DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)

LNK INTL INC

* LNK INTERNATIONAL INC
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******LOREAL USA**

- * LOREAL USA PRODUCTS INC
 - ANTHELIOS 20, AVOBENZONE (OTC)
 - ANTHELIOS 40, AVOBENZONE (OTC)
 - ANTHELIOS SX, AVOBENZONE (OTC)
 - CAPITAL SOLEIL 15, AVOBENZONE (OTC)

LOTUS PHARM CO LTD

- * LOTUS PHARMACEUTICAL CO LTD
 - ROWEEPR, LEVETIRACETAM
- * LOTUS PHARMACEUTICAL CO LTD NANTOU PLANT
 - CALCIUM ACETATE, CALCIUM ACETATE
 - LEVETIRACETAM, LEVETIRACETAM
 - LEVONORGESTREL, LEVONORGESTREL
 - LEVONORGESTREL, LEVONORGESTREL (OTC)
 - PARICALCITOL, PARICALCITOL

LOXO ONCOLOGY INC

- * LOXO ONCOLOGY INC
 - VITRAKVI, LAROTRECTINIB

LUITPOLD

- * LUITPOLD PHARMACEUTICALS INC
 - ACETYLCYSTEINE, ACETYLCYSTEINE
 - ADENOSINE, ADENOSINE
 - AMINOCAPROIC ACID, AMINOCAPROIC ACID
 - AMINOPHYLLINE, AMINOPHYLLINE
 - BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 - BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE, BETAMETHASONE ACETATE
 - BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 - BUSULFAN, BUSULFAN
 - CAFFEINE CITRATE, CAFFEINE CITRATE
 - CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE
 - CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 - CYANOCOBALAMIN, CYANOCOBALAMIN
 - CYCLOSPORINE, CYCLOSPORINE
 - DACTINOMYCIN, DACTINOMYCIN
 - DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 - DEXFERRUM, IRON DEXTRAN
 - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 - DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 - DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
 - DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 - DROPERIDOL, DROPERIDOL
 - EPINEPHRINE, EPINEPHRINE
 - ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 - ESTRADIOL VALERATE, ESTRADIOL VALERATE
 - ETOMIDATE, ETOMIDATE
 - FLOXURIDINE, FLOXURIDINE
 - FOMEPIZOLE, FOMEPIZOLE
 - FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 - GANCICLOVIR, GANCICLOVIR SODIUM
 - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 - GLYCOPYRROLATE, GLYCOPYRROLATE
 - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 - HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE
 - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 - IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
 - INJECTAFER, FERRIC CARBOXYMALTOS
 - LEVETIRACETAM, LEVETIRACETAM
 - LEVOCARNITINE, LEVOCARNITINE
 - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 - MANNITOL 25%, MANNITOL
 - METHOCARBAMOL, METHOCARBAMOL
 - METHYLDOPATE HYDROCHLORIDE, METHYLDOPATE HYDROCHLORIDE
 - METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LUITPOLD PHARMACEUTICALS INC
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 NANDROLONE DECANOATE, NANDROLONE DECANOATE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 NITROGLYCERIN, NITROGLYCERIN
 OLANZAPINE, OLANZAPINE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PHENYTOIN SODIUM, PHENYTOIN SODIUM
 PROGESTERONE, PROGESTERONE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRIMETHOBENZAMIDE HYDROCHLORIDE PRESERVATIVE FREE, TRIMETHOBENZAMIDE HYDROCHLORIDE
 TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
 VENOFER, IRON SUCROSE
 ZIDOVUDINE, ZIDOVUDINE

LUKARE MEDICAL LLC

* LUKARE MEDICAL LLC
 ELLIOTTS B SOLUTION, CALCIUM CHLORIDE

LUNDBECK NA LTD

* LUNDBECK NA LTD
 NORTHERA, DROXIDOPA

LUNDBECK PHARMS LLC

* LUNDBECK PHARMACEUTICALS LLC
 ONFI, CLOBAZAM
 SABRIL, VIGABATRIN

LUPIN

* LUPIN INC
 SOLOSEC, SECNIDAZOLE
 TOBRAMYCIN, TOBRAMYCIN

* LUPIN LTD
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 CARVEDILOL, CARVEDILOL
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFDINIR, CEFDINIR
 CEFPROZIL, CEFPROZIL
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CEPHALEXIN, CEPHALEXIN
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN, LEVOFLOXACIN
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINAPRIL, LISINAPRIL
 LOVASTATIN, LOVASTATIN
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 RAMIPRIL, RAMIPRIL
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SIMVASTATIN, SIMVASTATIN
 TOPIRAMATE, TOPIRAMATE
 TRANDOLAPRIL, TRANDOLAPRIL

LUPIN ATLANTIS

* LUPIN ATLANTIS HOLDINGS SA
 ANTARA (MICRONIZED), FENOFIBRATE
 BUDESONIDE, BUDESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 MIBELAS 24 FE, ETHINYL ESTRADIOL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

LUPIN LTD

* LUPIN LIMITED

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

* LUPIN LIMITED
 LEVETIRACETAM, LEVETIRACETAM
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

* LUPIN LTD
 ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
 ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE, ABACAVIR SULFATE
 AMABELZ, ESTRADIOL
 AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 ARMODAFINIL, ARMODAFINIL
 ATOVAQUONE, ATOVAQUONE
 AZITHROMYCIN, AZITHROMYCIN
 BEKYREE, DESOGESTREL
 BIMATOPROST, BIMATOPROST
 BLISOVI 24 FE, ETHINYL ESTRADIOL
 BLISOVI FE 1.5/30, ETHINYL ESTRADIOL
 BLISOVI FE 1/20, ETHINYL ESTRADIOL
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CALCIUM ACETATE, CALCIUM ACETATE
 CELECOXIB, CELECOXIB
 CIPROFLOXACIN, CIPROFLOXACIN
 CLARITHROMYCIN, CLARITHROMYCIN
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DAYSEE, ETHINYL ESTRADIOL
 DECITABINE, DECITABINE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENSKYCE, DESOGESTREL
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESZOPICLONE, ESZOPICLONE
 FALLBACK SOLO, LEVONORGESTREL (OTC)
 FAMOTIDINE, FAMOTIDINE
 FAYOSIM, ETHINYL ESTRADIOL
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FYAVOLV, ETHINYL ESTRADIOL
 GABAPENTIN, GABAPENTIN
 GATIFLOXACIN, GATIFLOXACIN
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 JENCYCLA, NORETHINDRONE
 KAITLIB FE, ETHINYL ESTRADIOL
 KURVELO, ETHINYL ESTRADIOL
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LORAZEPAM, LORAZEPAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******* LUPIN LTD**

MEFENAMIC ACID, MEFENAMIC ACID
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NABUMETONE, NABUMETONE
 NADOLOL, NADOLOL
 NIACIN, NIACIN
 NIKKI, DROSPIRENONE
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE, NORETHINDRONE
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OMEPRAZOLE, OMEPRAZOLE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PIRMELLA 1/35, ETHINYL ESTRADIOL
 PIRMELLA 7/7/7, ETHINYL ESTRADIOL
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUININE SULFATE, QUININE SULFATE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 RIFABUTIN, RIFABUTIN
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILODOSIN, SILODOSIN
 SUPRAX, CEFIXIME
 TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TESTOSTERONE, TESTOSTERONE
 TETRABENAZINE, TETRABENAZINE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TYDEMY, DROSPIRENONE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VYFEMLA, ETHINYL ESTRADIOL
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

LUPIN PHARMS*** LUPIN PHARMACEUTICALS INC**

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 DESLORATADINE, DESLORATADINE
 MELOXICAM, MELOXICAM
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 RIFAMPIN, RIFAMPIN
 SUPRAX, CEFIXIME
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

LYMOL MEDCL*** LYMOL MEDICAL CORP**

SCLEROSOL, TALC
 TALC, TALC

LYNE*** LYNE LABORATORIES INC**

CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DEXAMETHASONE, DEXAMETHASONE
 ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 LEVOCARNITINE, LEVOCARNITINE
 NYSTATIN, NYSTATIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******PERRIGO**

* L PERRIGO CO

ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
 CHILDREN'S IBUPROFEN, IBUPROFEN (OTC)
 CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
 DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
 FAMOTIDINE, FAMOTIDINE (OTC)
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN, IBUPROFEN (OTC)
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE, LORATADINE (OTC)
 MICONAZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 TAB-PROFEN, IBUPROFEN (OTC)
 TIOCONAZOLE, TIOCONAZOLE (OTC)

**** M ******MA GENERAL HOSP**

* MASSACHUSETTS GENERAL HOSP

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

MACLEODS PHARMS LTD

* MACLEODS PHARMACEUTICALS LTD

AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CELECOXIB, CELECOXIB
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DARIFENACIN, DARIFENACIN HYDROBROMIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTACAPONE, ENTACAPONE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESZOPICLONE, ESZOPICLONE
 FAMCICLOVIR, FAMCICLOVIR
 FLUOCINONIDE, FLUOCINONIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 NEVIRAPINE, NEVIRAPINE
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MACLEODS PHARMACEUTICALS LTD**

RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SILODOSIN, SILODOSIN
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRIAMCINOLONE ACETATE, TRIAMCINOLONE ACETONIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN

MAGNA PHARMS

*** MAGNA PHARMACEUTICALS INC**
 ZOLPIMIST, ZOLPIDEM TARTRATE

MAIA PHARMS INC

*** MAIA PHARMACEUTICALS INC**
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE

MAINPOINTE

*** MAINPOINTE PHARMACEUTICALS LLC**
 TUXARIN ER, CHLORPHENIRAMINE MALEATE

MALLINCKRODT ARD

*** MALLINCKRODT ARD INC**
 H.P. ACTHAR GEL, CORTICOTROPIN

MALLINCKRODT HOSP

*** MALLINCKRODT HOSP PRODUCTS IP LTD**
 INOMAX, NITRIC OXIDE
 OFIRMEV, ACETAMINOPHEN
 UVADEX, METHOXSALEN

MALLINKRODT NUCLEAR

*** MALLINCKRODT NUCLEAR MEDICINE LLC**
 GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67
 INDIUM IN 111 CHLORIDE, INDIUM IN-111 CHLORIDE
 OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT
 SODIUM IODIDE I 123, SODIUM IODIDE I-123
 TECHNISCAN MAG3, TECHNETIUM TC-99M MERTIATIDE KIT
 TECHNISCAN PYP KIT, TECHNETIUM TC-99M PYROPHOSPHATE KIT
 TECHNISCAN, TECHNETIUM TC-99M OXIDRONATE KIT
 TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
 THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
 ULTRA-TECHNEKOW FM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 ULTRATAG, TECHNETIUM TC-99M RED BLOOD CELL KIT
 XENON XE 133, XENON XE-133

MANNKIND

*** MANNKIND CORP**
 AFREZZA, INSULIN RECOMBINANT HUMAN

MARINA BIOTECH

*** MARINA BIOTECH INC**
 PRESTALIA, AMLODIPINE BESYLATE

MARKSANS PHARMA

*** MARKSANS PHARMA LTD**
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 DUTASTERIDE, DUTASTERIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 IBUPROFEN, IBUPROFEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MARKSANS PHARMA LTD**

IBUPROFEN, IBUPROFEN (OTC)
 LORATADINE, LORATADINE (OTC)
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NAPROXEN, NAPROXEN
 PARICALCITOL, PARICALCITOL

MARNEL PHARMS*** MARNEL PHARMACEUTICALS LLC**
CROTAN, CROTAMITON**MAYER LABS INC***** MAYER LABORATORIES INC**
TODAY, NONOXYNOL-9 (OTC)**MAYNE PHARMA***** MAYNE PHARMA INTERNATIONAL PTY LTD**
DORYX MPC, DOXYCYCLINE HYCLATE
DORYX, DOXYCYCLINE HYCLATE
ERYC, ERYTHROMYCIN*** MAYNE PHARMA LLC**

BUDESONIDE, BUDESONIDE
 CAMILA, NORETHINDRONE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CLARITHROMYCIN, CLARITHROMYCIN
 CLONIDINE, CLONIDINE
 CLOZAPINE, CLOZAPINE
 CYCLOSPORINE, CYCLOSPORINE
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DIAZEPAM, DIAZEPAM
 DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 ERRIN, NORETHINDRONE
 ESTAZOLAM, ESTAZOLAM
 ESTRADIOL, ESTRADIOL
 FABIOR, TAZAROTENE
 FENTANYL-100, FENTANYL
 FENTANYL-25, FENTANYL
 FENTANYL-50, FENTANYL
 FENTANYL-75, FENTANYL
 FLUOROURACIL, FLUOROURACIL
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVORA 0.15/30-28, ETHINYL ESTRADIOL
 LOW-OGESTREL-28, ETHINYL ESTRADIOL
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MICROGESTIN 1.5/30, ETHINYL ESTRADIOL
 MICROGESTIN 1/20, ETHINYL ESTRADIOL
 MICROGESTIN FE 1.5/30, ETHINYL ESTRADIOL
 MICROGESTIN FE 1/20, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 SORILUX, CALCIPOTRIENE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TRI-NORINYL 28-DAY, ETHINYL ESTRADIOL
 TRIMETHOPRIM, TRIMETHOPRIM
 TRIVORA-28, ETHINYL ESTRADIOL
 ZOVIA 1/35E-28, ETHINYL ESTRADIOL

MAYNE PHARMA INC*** MAYNE PHARMA INC**
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MAYNE PHARMA INC
 BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 DOFETILIDE, DOFETILIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 HYDROCODONE BITARTRATE AND PSEUDOEPHEDRINE HYDROCHLORIDE, HYDROCODONE BITARTRATE
 HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE,
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NYSTATIN, NYSTATIN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE AND ASPIRIN, ASPIRIN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

MAYNE PHARMA INTL

* MAYNE PHARMA INTERNATIONAL PTY LTD
 TOLSURA, ITRACONAZOLE

MCGUFF

* MCGUFF PHARMACEUTICALS INC
 ASCOR, ASCORBIC ACID

MCNEIL

* MCNEIL CONSUMER PRODUCTS CO DIV MCNEILAB INC
 IBUPROFEN, IBUPROFEN (OTC)

MCNEIL CONS

* MCNEIL CONSUMER HEALTHCARE
 SUDAFED 12 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)

MCPRF

* MAYO CLINIC PET RADIOCHEMISTRY FACILITY
 AMMONIA N 13, AMMONIA N-13
 CHOLINE C-11, CHOLINE C-11
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

MDGH

* MEDICINES DEVELOPMENT FOR GLOBAL HEALTH
 MOXIDECTIN, MOXIDECTIN

MEDAC PHARMA INC

* MEDAC PHARMA INC
 RASUVO, METHOTREXATE

MEDEFIL INC

* MEDEFIL INC
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE

MEDICINES360

* MEDICINES360
 LILETTA, LEVONORGESTREL

MEDICIS

* MEDICIS PHARMACEUTICAL CORP
 ALDARA, IMIQUIMOD
 AMMONUL, SODIUM BENZOATE
 CALCIUM DISODIUM VERSENATE, EDETATE CALCIUM DISODIUM
 LOPROX, CICLOPIROX
 LUZU, LULICONAZOLE
 METROGEL-VAGINAL, METRONIDAZOLE
 MINITRAN, NITROGLYCERIN
 SOLODYN, MINOCYCLINE HYDROCHLORIDE
 VANOS, FLUOCINONIDE
 ZIANA, CLINDAMYCIN PHOSPHATE
 ZYCLARA, IMIQUIMOD

MEDICURE

* MEDICURE INTERNATIONAL INC
 AGGRASTAT, TIROFIBAN HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE

MEDIMETRIKS PHARMS

* MEDIMETRIKS PHARMACEUTICALS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MEDIMETRIKS PHARMACEUTICALS INC
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 LOPROX, CICLOPIROX
 NEO-SYNALAR, FLUOCINOLONE ACETONIDE
 SYNALAR, FLUOCINOLONE ACETONIDE

MEDLINE INDUSTRIES

* MEDLINE INDUSTRIES INC
 READYPREP CHG, CHLORHEXIDINE GLUCONATE (OTC)

MEDTECH PRODUCTS

* MEDTECH PRODUCTS INC
 MONISTAT 1 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MONISTAT 3 COMBINATION PACK (PREFILLED), MICONAZOLE NITRATE (OTC)
 MONISTAT 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MONISTAT 3, MICONAZOLE NITRATE
 MONISTAT 3, MICONAZOLE NITRATE (OTC)
 MONISTAT 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 MONISTAT 7, MICONAZOLE NITRATE (OTC)
 NIX, PERMETHRIN (OTC)
 TAGAMET HB, CIMETIDINE (OTC)

MELINTA

* MELINTA SUBSIDIARY CORP
 BAXDELA, DELAFLOXACIN MEGLUMINE

MELINTA THERAP

* MELINTA THERAPEUTICS INC
 ORBACTIV, ORITAVANCIN DIPHOSPHATE

MEM SLOAN-KETTERING

* MEMORIAL SLOAN-KETTERING CANCER CENTER
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

MERCK

* MERCK AND CO INC
 CANCIDAS, CASPOFUNGIN ACETATE
 EMEND, APREPITANT
 FOSAMAX PLUS D, ALENDRONATE SODIUM
 MAXALT, RIZATRIPTAN BENZOATE
 MAXALT-MLT, RIZATRIPTAN BENZOATE
 PRIMAXIN, CILASTATIN SODIUM
 PROSCAR, FINASTERIDE
 ZOLINZA, VORINOSTAT

* MERCK RESEARCH LABORATORIES DIV MERCK CO INC
 PRINIVIL, LISINAPRIL
 PROPECIA, FINASTERIDE
 SINGULAIR, MONTELUKAST SODIUM
 TRUSOPT, DORZOLAMIDE HYDROCHLORIDE

MERCK AND CO INC

* MERCK AND CO INC
 EMEND, FOSAPREPITANT DIMEGLUMINE
 FOSAMAX, ALENDRONATE SODIUM

MERCK SHARP DOHME

* MERCK SHARP AND DOHME CORP
 ASMANEX HFA, MOMETASONE FUROATE
 ASMANEX TWISTHALER, MOMETASONE FUROATE
 BELSOMRA, SUVOREXANT
 CELESTONE SOLUSPAN, BETAMETHASONE ACETATE
 CLARINEX D 24 HOUR, DESLORATADINE
 CLARINEX, DESLORATADINE
 CLARINEX-D 12 HOUR, DESLORATADINE
 COZAAR, LOSARTAN POTASSIUM
 CRIXIVAN, INDINAVIR SULFATE
 DIPROLENE AF, BETAMETHASONE DIPROPIONATE
 DIPROLENE, BETAMETHASONE DIPROPIONATE
 DULERA, FORMOTEROL FUMARATE
 ELOCON, MOMETASONE FUROATE
 GUANIDINE HYDROCHLORIDE, GUANIDINE HYDROCHLORIDE
 HYZAAR, HYDROCHLOROTHIAZIDE
 INVANZ, ERTAPENEM SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MERCK SHARP AND DOHME CORP
 ISENTRESS HD, RALTEGRAVIR POTASSIUM
 ISENTRESS, RALTEGRAVIR POTASSIUM
 JANUMET XR, METFORMIN HYDROCHLORIDE
 JANUMET, METFORMIN HYDROCHLORIDE
 JANUVIA, SITAGLIPTIN PHOSPHATE
 LOTRISONE, BETAMETHASONE DIPROPIONATE
 NASONEX, MOMETASONE FUROATE
 NOXAFIL, POSACONAZOLE
 PREVYMIS, LETERMOVIR
 REBETOL, RIBAVIRIN
 SEGLUROMET, ERTUGLIFLOZIN
 SINEMET CR, CARBIDOPA
 SINEMET, CARBIDOPA
 STEGLATRO, ERTUGLIFLOZIN
 STEGLUJAN, ERTUGLIFLOZIN
 STROMECTOL, IVERMECTIN
 TEMODAR, TEMOZOLOMIDE
 ZEPATIER, ELBASVIR

MERIDIAN MEDCL

* MERIDIAN MEDICAL TECHNOLOGIES INC
 DUODOTE, ATROPINE

MERIDIAN MEDCL TECHN

* MERIDIAN MEDICAL TECHNOLOGIES INC
 ATROPEN, ATROPINE SULFATE
 MORPHINE SULFATE, MORPHINE SULFATE
 PRALIDOXIME CHLORIDE, PRALIDOXIME CHLORIDE
 SEIZALAM, MIDAZOLAM HYDROCHLORIDE

MERLION PHARMS GMBH

* MERLION PHARMACEUTICALS GMBH
 XTORO, FINAFLOXACIN

MERRO PHARM

* MERRO PHARMACEUTICAL CO LTD
 IBUPROFEN, IBUPROFEN (OTC)

MERZ PHARMS

* MERZ PHARMACEUTICALS LLC
 CUVPOSA, GLYCOPYRROLATE

METHAPHARM

* METHAPHARM INC
 PROVOCHOLINE, METHACHOLINE CHLORIDE

METUCHEN PHARMS

* METUCHEN PHARMACEUTICALS LLC
 STENDRA, AVANAFIL

MICRO LABS

* MICRO LABS LTD
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CELECOXIB, CELECOXIB
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 MEFENAMIC ACID, MEFENAMIC ACID
 METHAZOLAMIDE, METHAZOLAMIDE
 PIROXICAM, PIROXICAM
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 TELMISARTAN, TELMISARTAN
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

MICRO LABS LTD

* MICRO LABS LTD
 NEVIRAPINE, NEVIRAPINE

MICRO LABS LTD INDIA

* MICRO LABS LTD INDIA
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 CROMOLYN SODIUM, CROMOLYN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MICRO LABS LTD INDIA
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN

MIDATECH PHARMA US

* MIDATECH PHARMA US INC
 ORAVIG, MICONAZOLE
 SOLTAMOX, TAMOXIFEN CITRATE
 ZUPLENZ, ONDANSETRON

MIDWEST MEDCL

* MIDWEST MEDICAL ISOTOPES LLC CYCLOTRON DIV
 AMMONIA N 13, AMMONIA N-13
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

MIKART

* MIKART LLC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 BENZONATATE, BENZONATATE
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 BUTAPAP, ACETAMINOPHEN
 CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
 CHLORZOXAZONE, CHLORZOXAZONE
 ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 ISONIAZID, ISONIAZID
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 METHAZOLAMIDE, METHAZOLAMIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

MIKART INC

* MIKART INC
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN

MILLENNIUM PHARMS

* MILLENNIUM PHARMACEUTICALS INC
 NINLARO, IXAZOMIB CITRATE
 VELCADE, BORTEZOMIB

MILLICENT

* MILLICENT HOLDINGS LTD
 FEMRING, ESTRADIOL ACETATE

MIPS CRF

* MIPS CYCLOTRON AND RADIOCHEMISTRY FACILITY
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

MISSION PHARMA

* MISSION PHARMACAL CO
 BINOSTO, ALENDRONATE SODIUM
 LITHOSTAT, ACETOHYDROXAMIC ACID
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 TEXACORT, HYDROCORTISONE
 THIOLA, TIOPRONIN
 TINDAMAX, TINIDAZOLE
 UROCIT-K, POTASSIUM CITRATE

MISSION PHARMACAL CO

* MISSION PHARMACAL CO
 CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
 POTASSIUM IODIDE, POTASSIUM IODIDE (OTC)

MIST PHARMS LLC

* MIST PHARMACEUTICALS LLC
 NITROMIST, NITROGLYCERIN

MITSUBISHI TANABE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MITSUBISHI TANABE PHARMA CORP
RADICAVA, EDARAVONE

MOBERG PHARMA NORTH

* MOBERG PHARMA NORTH AMERICA LLC
CHILDREN'S ELIXSURE, IBUPROFEN (OTC)

MOBIUS THERAP

* MOBIUS THERAPEUTICS LLC
MITOSOL, MITOMYCIN

MOLNLYCKE HLTH

* MOLNLYCKE HEALTH CARE
HIBICLENS, CHLORHEXIDINE GLUCONATE (OTC)
HIBISTAT, CHLORHEXIDINE GLUCONATE (OTC)

MONARCH PHARMS

* MONARCH PHARMACEUTICALS LLC
CORTISPORIN, BACITRACIN ZINC
CORTISPORIN, HYDROCORTISONE ACETATE
MENEST, ESTROGENS, ESTERIFIED
NEOSPORIN G.U. IRRIGANT, NEOMYCIN SULFATE
NEOSPORIN, GRAMICIDIN
SEPTRA DS, SULFAMETHOXAZOLE
SEPTRA, SULFAMETHOXAZOLE
VIROPTIC, TRIFLURIDINE

MONTEREY PHARMS LLC

* MONTEREY PHARMACEUTICALS LLC
METHOCARBAMOL, METHOCARBAMOL

MOUNTAIN

* MOUNTAIN LLC
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
METAXALONE, METAXALONE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
POTASSIUM CITRATE, POTASSIUM CITRATE

MSD INTL

* MSD INTERNATIONAL GMBH
VYTORIN, EZETIMIBE

MSD INTL GMBH

* MSD INTERNATIONAL GMBH
ZETIA, EZETIMIBE

MSD MERCK CO

* MERCK SHARP AND DOHME CORP A SUB OF MERCK AND CO INC
DELSTRIGO, DORAVIRINE
EMEND, APREPITANT
PIFELTRO, DORAVIRINE
SINGULAIR, MONTELUKAST SODIUM
ZOCOR, SIMVASTATIN

MSN LABS PVT LTD

* MSN LABORATORIES PRIVATE LTD
CAPECITABINE, CAPECITABINE
CLOFARABINE, CLOFARABINE
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
SILODOSIN, SILODOSIN

MURTY PHARMS

* MURTY PHARMACEUTICALS INC
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
DIPYRIDAMOLE, DIPYRIDAMOLE

MUSTAFA NEVZAT ILAC

* MUSTAFA NEVZAT ILAC SANAYII AS (MN PHARMACEUTICALS)
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

MUTUAL PHARM

* MUTUAL PHARMACEUTICAL CO INC
PREDNISONE, PREDNISONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******MYLAN**

* MYLAN PHARMACEUTICALS
 FENOFIBRATE, FENOFIBRATE
 METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID

* MYLAN PHARMACEUTICALS INC
 ACARBOSE, ACARBOSE
 ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 ALLOPURINOL, ALLOPURINOL
 ALPRAZOLAM, ALPRAZOLAM
 AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 ANASTROZOLE, ANASTROZOLE
 ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 ATENOLOL, ATENOLOL
 AVITA, TRETINOIN
 AZATHIOPRINE, AZATHIOPRINE
 AZITHROMYCIN, AZITHROMYCIN
 BACLOFEN, BACLOFEN
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BUDESONIDE, BUDESONIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARVEDILOL, CARVEDILOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHLOROTHIAZIDE, CHLOROTHIAZIDE
 CHLORPROPAMIDE, CHLORPROPAMIDE
 CHLORTHALIDONE, CHLORTHALIDONE
 CIMETIDINE, CIMETIDINE
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 CLOZAPINE, CLOZAPINE
 CYSTAGON, CYSTEAMINE BITARTRATE
 DIAZEPAM, DIAZEPAM
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN PHARMACEUTICALS INC
 ESTRADIOL, ESTRADIOL
 ESTROPIPATE, ESTROPIPATE
 ETIDRONATE DISODIUM, ETIDRONATE DISODIUM
 ETOPOSIDE, ETOPOSIDE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 FAMCICLOVIR, FAMCICLOVIR
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FELODIPINE, FELODIPINE
 FENOFIBRATE, FENOFIBRATE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FINASTERIDE, FINASTERIDE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FLURBIPROFEN, FLURBIPROFEN
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 FUROSEMIDE, FUROSEMIDE
 GABAPENTIN, GABAPENTIN
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GLIMEPIRIDE, GLIMEPIRIDE
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 GLIPIZIDE, GLIPIZIDE
 GLYBURIDE (MICRONIZED), GLYBURIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HALOPERIDOL, HALOPERIDOL
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 INDAPAMIDE, INDAPAMIDE
 INDOMETHACIN, INDOMETHACIN
 KETOCONAZOLE, KETOCONAZOLE
 KETOPROFEN, KETOPROFEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LAMOTRIGINE, LAMOTRIGINE
 LATANOPROST, LATANOPROST
 LEVETIRACETAM, LEVETIRACETAM
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM **
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINAPRIL, LISINAPRIL
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 LORAZEPAM, LORAZEPAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LOVASTATIN, LOVASTATIN
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 MAPROTIline HYDROCHLORIDE, MAPROTIline HYDROCHLORIDE
 MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM
 MENTAX, BUTENAFINE HYDROCHLORIDE
 MERCAPTOPYRINE, MERCAPTOPYRINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHIMAZOLE, METHIMAZOLE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METHYLDOPA AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 METHYLDOPA, METHYLDOPA
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIRTAZAPINE, MIRTAZAPINE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN PHARMACEUTICALS INC
 NADOLOL, NADOLOL
 NAPROXEN, NAPROXEN
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NIFEDIPINE, NIFEDIPINE
 NISOLDIPINE, NISOLDIPINE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 OMEPRAZOLE, OMEPRAZOLE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON, ONDANSETRON
 OXYBUTYNYN CHLORIDE, OXYBUTYNYN CHLORIDE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PENTOXIFYLLINE, PENTOXIFYLLINE
 PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 PHENYTEK, PHENYTOIN SODIUM
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PROBENECID, PROBENECID
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 RISPERIDONE, RISPERIDONE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 STAVUDINE, STAVUDINE
 SULINDAC, SULINDAC
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TACROLIMUS, TACROLIMUS
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TEMAZEPAM, TEMAZEPAM
 THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
 THIOTHIXENE, THIOTHIXENE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOLMETIN SODIUM, TOLMETIN SODIUM
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 URSODIOL, URSODIOL
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
 ZONISAMIDE, ZONISAMIDE

MYLAN ASI

* MYLAN ASI LLC
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 ADENOSINE, ADENOSINE
 AZITHROMYCIN, AZITHROMYCIN
 BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MYLAN ASI LLC**

POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE

MYLAN INSTITUTIONAL*** MYLAN INSTITUTIONAL INC**

SULFAMYLON, MAFENIDE ACETATE

*** MYLAN INSTITUTIONAL LLC**

ACETYLCYSTEINE, ACETYLCYSTEINE
 ALOPRIM, ALLOPURINOL SODIUM
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 ARGATROBAN, ARGATROBAN
 AZACITIDINE, AZACITIDINE
 BIVALIRUDIN, BIVALIRUDIN
 CARBOPLATIN, CARBOPLATIN
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CIDOFOVIR, CIDOFOVIR
 COSYNTROPIN, COSYNTROPIN
 DANTROLENE SODIUM, DANTROLENE SODIUM
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 DIMETHYL SULFOXIDE, DIMETHYL SULFOXIDE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DURACLON, CLONIDINE HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 FOMEPIZOLE, FOMEPIZOLE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
 ISOSULFAN BLUE, ISOSULFAN BLUE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 MEFOXIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHOCARBAMOL, METHOCARBAMOL
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PENTOSTATIN, PENTOSTATIN
 RIMSO-50, DIMETHYL SULFOXIDE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 SOTRADECOL, SODIUM TETRADECYL SULFATE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 ULTIVA, REMIFENTANIL HYDROCHLORIDE

MYLAN IRELAND LTD*** MYLAN IRELAND LTD**

ARIXTRA, FONDAPARINUX SODIUM
 CARBAMAZEPINE, CARBAMAZEPINE
 MIACALCIN, CALCITONIN SALMON
 PIROXICAM, PIROXICAM
 THEOPHYLLINE, THEOPHYLLINE
 YUPELRI, REVEFENACIN

MYLAN LABS*** MYLAN LABORATORIES LTD**

NEVIRAPINE, NEVIRAPINE

MYLAN LABS LTD*** MYLAN LABORATORIES LTD**

ADENOSINE, ADENOSINE
 AMIFOSTINE, AMIFOSTINE
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MYLAN LABORATORIES LTD**

ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 AZITHROMYCIN, AZITHROMYCIN
 BACLOFEN, BACLOFEN
 BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
 BUSULFAN, BUSULFAN
 CAPREOMYCIN SULFATE, CAPREOMYCIN SULFATE
 CARBOPLATIN, CARBOPLATIN
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 CIMDUO, LAMIVUDINE
 CISPLATIN, CISPLATIN
 CLADRIBINE, CLADRIBINE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOFARABINE, CLOFARABINE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYTARABINE, CYTARABINE
 DACTINOMYCIN, DACTINOMYCIN
 DAPTOMYCIN, DAPTOMYCIN
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DOCETAXEL, DOCETAXEL
 DOCETAXEL, DOCETAXEL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE HYCLATE
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 EPTIFIBATIDE, EPTIFIBATIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 ETOMIDATE, ETOMIDATE
 ETOPOSIDE, ETOPOSIDE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUMAZENIL, FLUMAZENIL
 FLUOROURACIL, FLUOROURACIL
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 GANCICLOVIR, GANCICLOVIR SODIUM
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM, HEPARIN SODIUM
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
 IFOSFAMIDE, IFOSFAMIDE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN, LEVOFLOXACIN
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 LEVONORGESTREL, LEVONORGESTREL
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LINEZOLID, LINEZOLID
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
 MITOMYCIN, MITOMYCIN
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER, MOXIFLOXACIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MYLAN LABORATORIES LTD**

MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE, NORETHINDRONE
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXALIPLATIN, OXALIPLATIN
 PACLITAXEL, PACLITAXEL
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARICALCITOL, PARICALCITOL
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RIFAMPIN, RIFAMPIN
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 SYMFI, EFAVIRENZ
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

MYLAN PHARMS INC*** MYLAN PHARMACEUTICALS INC**

ABACAVIR SULFATE, ABACAVIR SULFATE
 ABIRATERONE ACETATE, ABIRATERONE ACETATE
 ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
 ACITRETIN, ACITRETIN
 ACYCLOVIR, ACYCLOVIR
 ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
 AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMNESTEEM, ISOTRETINOIN
 ARMODAFINIL, ARMODAFINIL
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
 AVITA, TRETINOIN
 BACLOFEN, BACLOFEN
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CABERGOLINE, CABERGOLINE
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CAPECITABINE, CAPECITABINE
 CAPTOPRIL, CAPTOPRIL
 CELECOXIB, CELECOXIB
 CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CLOZAPINE, CLOZAPINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN PHARMACEUTICALS INC
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DENAVIR, PENCICLOVIR
 DESLORATADINE, DESLORATADINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIGOXIN, DIGOXIN
 DISULFIRAM, DISULFIRAM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 ECONAZOLE NITRATE, ECONAZOLE NITRATE
 EFAVIRENZ, EFAVIRENZ
 ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
 ELIMITE, PERMETHRIN
 EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
 EPLERENONE, EPLERENONE
 EPROSARTAN MESYLATE, EPROSARTAN MESYLATE
 ERYGEL, ERYTHROMYCIN
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 ESTRADIOL, ESTRADIOL
 ESZOPICLONE, ESZOPICLONE
 EVOCLIN, CLINDAMYCIN PHOSPHATE
 EXEMESTANE, EXEMESTANE
 EXTINA, KETOCONAZOLE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRATE, FENOFIBRATE
 FENOFIBRIC ACID, CHOLINE FENOFIBRATE
 FLUOROURACIL, FLUOROURACIL
 FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM
 FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
 GABAPENTIN, GABAPENTIN
 GATIFLOXACIN, GATIFLOXACIN
 GLATIRAMER ACETATE, GLATIRAMER ACETATE
 GLIPIZIDE, GLIPIZIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 INDOMETHACIN, INDOMETHACIN
 ITRACONAZOLE, ITRACONAZOLE
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LANSOPRAZOLE, LANSOPRAZOLE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 LINEZOLID, LINEZOLID
 LITHIUM CARBONATE, LITHIUM CARBONATE
 LUXIQ, BETAMETHASONE VALERATE
 MAXZIDE, HYDROCHLOROTHIAZIDE
 MAXZIDE-25, HYDROCHLOROTHIAZIDE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 MESALAMINE, MESALAMINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHYLCLOTHIAZIDE, METHYLCLOTHIAZIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MYLAN PHARMACEUTICALS INC
MODAFINIL, MODAFINIL
MONTELUKAST SODIUM, MONTELUKAST SODIUM
MORPHINE SULFATE, MORPHINE SULFATE
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
NABUMETONE, NABUMETONE
NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
NEVIRAPINE, NEVIRAPINE
OLANZAPINE, OLANZAPINE
OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
OLUX E, CLOBETASOL PROPIONATE
OLUX, CLOBETASOL PROPIONATE
OMEPRAZOLE, OMEPRAZOLE
PALIPERIDONE, PALIPERIDONE
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
PERPHENAZINE, PERPHENAZINE
PHENYTOIN, PHENYTOIN
PINDOLOL, PINDOLOL
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PRASUGREL, PRASUGREL HYDROCHLORIDE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
PREDNISON, PREDNISON
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
QUININE SULFATE, QUININE SULFATE
RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
RASAGILINE MESYLATE, RASAGILINE MESYLATE
REPAGLINIDE, REPAGLINIDE
RILUZOLE, RILUZOLE
RISEDRONATE SODIUM, RISEDRONATE SODIUM
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
RUFINAMIDE, RUFINAMIDE
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SILDENAFIL CITRATE, SILDENAFIL CITRATE
SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
SYMFI LO, EFAVIRENZ
TADALAFIL, TADALAFIL
TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TELMISARTAN, TELMISARTAN
TENOFVIR DISOPROXIL FUMARATE, TENOFVIR DISOPROXIL FUMARATE
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TOBRAMYCIN, TOBRAMYCIN
TOLAZAMIDE, TOLAZAMIDE
TOLBUTAMIDE, TOLBUTAMIDE
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRAVOPROST, TRAVOPROST
TRETINOIN, TRETINOIN
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
TRIAZOLAM, TRIAZOLAM
TRILYTE, POLYETHYLENE GLYCOL 3350
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
VALSARTAN, VALSARTAN
VORICONAZOLE, VORICONAZOLE
VUSION, MICONAZOLE NITRATE
ZIDOVUDINE, ZIDOVUDINE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

- * MYLAN PHARMACEUTICALS INC
ZONALON, DOXEPIN HYDROCHLORIDE
ZOVIRAX, ACYCLOVIR
- * MYLAN PHARMACEUTICALS INC.
FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
NIZATIDINE, NIZATIDINE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

MYLAN SPECIALITY LP

- * MYLAN SPECIALTY LP
ACCUNEB, ALBUTEROL SULFATE
AEROSPAN HFA, FLUNISOLIDE
ANADROL-50, OXYMETHOLONE
ASTELIN, AZELASTINE HYDROCHLORIDE
ASTEPRO, AZELASTINE HYDROCHLORIDE
AVC, SULFANILAMIDE
BUTISOL SODIUM, BUTABARBITAL SODIUM
CESAMET, NABILONE
COLYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
CORTIFOAM, HYDROCORTISONE ACETATE
CROMOLYN SODIUM, CROMOLYN SODIUM
DEMADEX, TORSEMIDE
DEPEN, PENICILLAMINE
DIPENTUM, OLSALAZINE SODIUM
DYMISTA, AZELASTINE HYDROCHLORIDE
EDLUAR, ZOLPIDEM TARTRATE
ELESTRIN, ESTRADIOL
EPIFOAM, HYDROCORTISONE ACETATE
EPIPEN JR., EPINEPHRINE
EPIPEN, EPINEPHRINE
FELBATOL, FELBAMATE
GASTROCROM, CROMOLYN SODIUM
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
MUSE, ALPROSTADIL
PROCTOFOAM HC, HYDROCORTISONE ACETATE
ROWASA, MESALAMINE
SFROWASA, MESALAMINE
SOMA, CARISOPRODOL
TOBI PODHALER, TOBRAMYCIN
TOBI, TOBRAMYCIN

MYLAN SPECLT

- * MYLAN SPECIALTY LP
PERFOROMIST, FORMOTEROL FUMARATE

MYLAN TECHNOLOGIES

- * MYLAN TECHNOLOGIES INC
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
CLONIDINE, CLONIDINE
ESTRADIOL, ESTRADIOL
FENTANYL-100, FENTANYL
FENTANYL-12, FENTANYL
FENTANYL-25, FENTANYL
FENTANYL-37, FENTANYL
FENTANYL-50, FENTANYL
FENTANYL-62, FENTANYL
FENTANYL-75, FENTANYL
FENTANYL-87, FENTANYL
LIDOCAINE, LIDOCAINE
NITROGLYCERIN, NITROGLYCERIN
RIVASTIGMINE, RIVASTIGMINE
XULANE, ETHINYL ESTRADIOL

MYLAN TEORANTA

- * MYLAN TEORANTA
LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM

**** N ****

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ******NALPROPION**

* NALPROPION PHARMACEUTICALS INC
CONTRAVE, BUPROPION HYDROCHLORIDE

NAMIGEN LLC

* NAMIGEN LLC
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE

NANG KUANG PHARM CO

* NANG KUANG PHARMACEUTICAL CO LTD
LINEZOLID, LINEZOLID

NANJING KING-FRIEND

* NANJING KING-FRIEND BIOCHEMICAL PHARMACEUTICAL CO LTD
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
CARBOPLATIN, CARBOPLATIN
HEPARIN SODIUM, HEPARIN SODIUM

NAPO PHARMS INC

* NAPO PHARMACEUTICALS INC
MYTESI, CROFELEMER

NATCO PHARMA

* NATCO PHARMA LTD
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE

NATCO PHARMA LTD

* NATCO PHARMA LIMITED
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE

* NATCO PHARMA LTD
ALPRAZOLAM, ALPRAZOLAM
ANASTROZOLE, ANASTROZOLE
ARMODAFINIL, ARMODAFINIL
AZACITIDINE, AZACITIDINE
CARISOPRODOL, CARISOPRODOL
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
GLYCOPYRROLATE, GLYCOPYRROLATE
LANSOPRAZOLE, LANSOPRAZOLE
LANSOPRAZOLE, LANSOPRAZOLE (OTC)
LANTHANUM CARBONATE, LANTHANUM CARBONATE
LETROZOLE, LETROZOLE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

NAVINTA LLC

* NAVINTA LLC
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
CARMUSTINE, CARMUSTINE
FAMOTIDINE, FAMOTIDINE
FOMEPIZOLE, FOMEPIZOLE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
METHOCARBAMOL, METHOCARBAMOL
RIBAVIRIN, RIBAVIRIN
ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE
TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

NCM USA BRONX LLC

* NCM USA BRONX LLC
AMMONIA N 13, AMMONIA N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

NEOPHARMA

* NEOPHARMA INC
ALENDRONATE SODIUM, ALENDRONATE SODIUM
AMOXIL, AMOXICILLIN
ANASTROZOLE, ANASTROZOLE
AUGMENTIN '125', AMOXICILLIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ******* NEOPHARMA INC**

AUGMENTIN '250', AMOXICILLIN
 AUGMENTIN XR, AMOXICILLIN
 IRBESARTAN, IRBESARTAN
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LAROTID, AMOXICILLIN
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE

NEOS THERAP INC

* NEOS THERAPEUTICS INC
 HYDROCODONE POLISTIREX AND CHLORPHENIRAMNE POLISTIREX, CHLORPHENIRAMINE POLISTIREX

NEOS THERAPS

* NEOS THERAPEUTICS
 ADZENYS XR-ODT, AMPHETAMINE

NEOS THERAPS INC

* NEOS THERAPEUTICS INC
 ADZENYS ER, AMPHETAMINE
 COTEMPLA XR-ODT, METHYLPHENIDATE

NEPHRON

* NEPHRON CORP
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE

* NEPHRON PHARMACEUTICALS CORP
 ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE

NESHER PHARMS

* NESHER PHARMACEUTICALS USA LLC
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 MICRO-K 10, POTASSIUM CHLORIDE
 MICRO-K, POTASSIUM CHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NYSTATIN, NYSTATIN
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

NEUROCRINE

* NEUROCRINE BIOSCIENCES INC
 INGREZZA, VALBENZAZINE TOSYLATE

NEW RIVER

* NEW RIVER PHARMACEUTICALS INC
 PROFERDEX, IRON DEXTRAN

NEXGEN PHARMA

* NEXGEN PHARMA INC
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CHENODIOL, CHENODIOL
 GLYCOPYRROLATE, GLYCOPYRROLATE
 MECAMYLAMINE HYDROCHLORIDE, MECAMYLAMINE HYDROCHLORIDE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)

NEXGEN PHARMA INC

* NEXGEN PHARMA INC
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350

NEXTWAVE PHARMS

* NEXTWAVE PHARMACEUTICALS INC
 QUILLICHEW ER, METHYLPHENIDATE HYDROCHLORIDE
 QUILLIVANT XR, METHYLPHENIDATE HYDROCHLORIDE

NEXUS PHARMS

* NEXUS PHARMACEUTICALS INC
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ******* NEXUS PHARMACEUTICALS INC**

BUSULFAN, BUSULFAN
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE

NIAGARA PHARMS*** NIAGARA PHARMACEUTICALS INC**

PUR-WASH, PURIFIED WATER (OTC)

NODEN PHARMA*** NODEN PHARMA DAC**

TEKTURNA HCT, ALISKIREN HEMIFUMARATE
 TEKTURNA, ALISKIREN HEMIFUMARATE

NORTEC DEV ASSOC*** NORTEC DEVELOPMENT ASSOC INC**

PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

NORTHLAND*** NORTHLAND NUCLEAR MEDICINE LLC**

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

NORTHSTAR HLTHCARE*** NORTHSTAR HEALTHCARE HOLDINGS LTD**

ALLOPURINOL, ALLOPURINOL
 BACLOFEN, BACLOFEN
 GEMFIBROZIL, GEMFIBROZIL
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

NORTHSTAR MEDICAL*** NORTHSTAR MEDICAL RADIOISOTOPES LLC**

RADIOGENIX SYSTEM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

NORTON WATERFORD*** NORTON WATERFORD LTD**

QVAR REDIHALER, BECLOMETHASONE DIPROPIONATE

NOSTRUM LABS INC*** NOSTRUM LABORATORIES INC**

ACETAZOLAMIDE, ACETAZOLAMIDE
 CALCIUM ACETATE, CALCIUM ACETATE
 CARISOPRODOL, CARISOPRODOL
 DAPSONE, DAPSONE
 ELIXOPHYLLIN, THEOPHYLLINE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NITROFURANTOIN, NITROFURANTOIN
 PINDOLOL, PINDOLOL
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

NOSTRUM PHARMS LLC*** NOSTRUM PHARMACEUTICALS LLC**

METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 THEOCHRON, THEOPHYLLINE

NOVA LABS LTD*** NOVA LABORATORIES LTD**

PURIXAN, MERCAPTOPYRINE

NOVADAQ TECH*** NOVADAQ TECHNOLOGIES ULC**

SPY AGENT GREEN KIT, INDOCYANINE GREEN

NOVARTIS*** NOVARTIS PHARMACEUTICALS CORP**

AFINITOR, EVEROLIMUS
 COARTEM, ARTEMETHER
 DESFERAL, DEFEROXAMINE MESYLATE
 DIOVAN HCT, HYDROCHLOROTHIAZIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVARTIS PHARMACEUTICALS CORP
 DIOVAN, VALSARTAN
 EXELON, RIVASTIGMINE
 EXELON, RIVASTIGMINE TARTRATE
 EXFORGE HCT, AMLODIPINE BESYLATE
 EXFORGE, AMLODIPINE BESYLATE
 EXJADE, DEFERASIROX
 FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE
 GILENYA, FINGOLIMOD HYDROCHLORIDE
 GLEEVEC, IMATINIB MESYLATE
 LAMISIL, TERBINAFINE HYDROCHLORIDE
 LESCOL XL, FLUVASTATIN SODIUM
 LOPRESSOR, METOPROLOL TARTRATE
 LOTREL, AMLODIPINE BESYLATE
 MYFORTIC, MYCOPHENOLIC ACID
 NEORAL, CYCLOSPORINE
 RECLAST, ZOLEDRONIC ACID
 RITALIN LA, METHYLPHENIDATE HYDROCHLORIDE
 RITALIN, METHYLPHENIDATE HYDROCHLORIDE
 SANDIMMUNE, CYCLOSPORINE
 SANDOSTATIN LAR, OCTREOTIDE ACETATE
 SANDOSTATIN, OCTREOTIDE ACETATE
 SIGNIFOR, PASIREOTIDE DIASPARTATE
 STARLIX, NATEGLINIDE
 TASIGNA, NILOTINIB HYDROCHLORIDE
 TEGRETOL, CARBAMAZEPINE
 TEGRETOL-XR, CARBAMAZEPINE
 TRILEPTAL, OXCARBAZEPINE
 VIVELLE-DOT, ESTRADIOL
 VOLTAREN, DICLOFENAC SODIUM
 ZOMETA, ZOLEDRONIC ACID
 ZORTRESS, EVEROLIMUS

NOVARTIS PHARM

* NOVARTIS PHARMACEUTICAL CORP
 AFINITOR DISPERZ, EVEROLIMUS

NOVARTIS PHARMS

* NOVARTIS PHARMACEUTICALS CORP
 FEMARA, LETROZOLE

NOVARTIS PHARMS CORP

* NOVARTIS PHARMACEUTICALS CORP
 ALOMIDE, LODOXAMIDE TROMETHAMINE
 ARGATROBAN, ARGATROBAN
 ARRANON, NELARABINE
 AZOPT, BRINZOLAMIDE
 BETOPTIC S, BETAXOLOL HYDROCHLORIDE
 CILOXAN, CIPROFLOXACIN HYDROCHLORIDE
 CIPRO HC, CIPROFLOXACIN HYDROCHLORIDE
 CIPRODEX, CIPROFLOXACIN
 DUREZOL, DIFLUPREDNATE
 EMADINE, EMEDASTINE DIFUMARATE
 ENTRESTO, SACUBITRIL
 FARYDAK, PANOBINOSTAT LACTATE
 FLAREX, FLUOROMETHOLONE ACETATE
 HYCAMTIN, TOPOTECAN HYDROCHLORIDE
 ILEVRO, NEPAFENAC
 IOPIDINE, APRACLONIDINE HYDROCHLORIDE
 ISOPTO CARPINE, PILOCARPINE HYDROCHLORIDE
 JADENU SPRINKLE, DEFERASIROX
 JADENU, DEFERASIROX
 KISQALI FEMARA CO-PACK (COPACKAGED), LETROZOLE
 KISQALI, RIBOCICLIB SUCCINATE
 MAXIDEX, DEXAMETHASONE
 MAXITROL, DEXAMETHASONE
 MEKINIST, TRAMETINIB DIMETHYL SULFOXIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVARTIS PHARMACEUTICALS CORP
 MOXEZA, MOXIFLOXACIN HYDROCHLORIDE
 MYDRIACYL, TROPICAMIDE
 NATACYN, NATAMYCIN
 NEVANAC, NEPAFENAC
 OMNIPRED, PREDNISOLONE ACETATE
 PATADAY, OLOPATADINE HYDROCHLORIDE
 PATANASE, OLOPATADINE HYDROCHLORIDE
 PATANOL, OLOPATADINE HYDROCHLORIDE
 PAZEO, OLOPATADINE HYDROCHLORIDE
 PROMACTA KIT, ELTROMBOPAG OLAMINE
 PROMACTA, ELTROMBOPAG OLAMINE
 RYDAPT, MIDOSTAURIN
 SIGNIFOR LAR KIT, PASIREOTIDE PAMOATE
 SIMBRINZA, BRIMONIDINE TARTRATE
 TAFINLAR, DABRAFENIB MESYLATE
 TOBRADEX ST, DEXAMETHASONE
 TOBRADEX, DEXAMETHASONE
 TOBREX, TOBRAMYCIN
 TRAVATAN Z, TRAVOPROST
 TRISENCE, TRIAMCINOLONE ACETONIDE
 TYKERB, LAPATINIB DITOSYLATE
 VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE
 VOTRIENT, PAZOPANIB HYDROCHLORIDE
 ZOFRAN ODT, ONDANSETRON
 ZOFRAN, ONDANSETRON HYDROCHLORIDE
 ZYKADIA, CERITINIB

NOVAST LABS

* NOVAST LABORATORIES CHINA LTD
 NORETHINDRONE, NORETHINDRONE

* NOVAST LABORATORIES LTD
 ACETAZOLAMIDE, ACETAZOLAMIDE
 CARISOPRODOL AND ASPIRIN, ASPIRIN
 CARISOPRODOL, CARISOPRODOL
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 HER STYLE, LEVONORGESTREL (OTC)
 INDOMETHACIN, INDOMETHACIN
 LARIN 1.5/30, ETHINYL ESTRADIOL
 LARIN 1/20, ETHINYL ESTRADIOL
 LARIN 24 FE, ETHINYL ESTRADIOL
 LARIN FE 1.5/30, ETHINYL ESTRADIOL
 LARIN FE 1/20, ETHINYL ESTRADIOL
 LERIBANE, ETHINYL ESTRADIOL
 MAFENIDE ACETATE, MAFENIDE ACETATE
 MALMOREDE, ETHINYL ESTRADIOL
 MELAMISA, DROSPIRENONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NADOLOL, NADOLOL
 NIFEDIPINE, NIFEDIPINE
 NORETHINDRONE, NORETHINDRONE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PIMTREA, DESOGESTREL
 PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE
 PROBENECID AND COLCHICINE, COLCHICINE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUININE SULFATE, QUININE SULFATE
 SETLAKIN, ETHINYL ESTRADIOL
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 YAELA, DROSPIRENONE

NOVAST LABS LTD

* NOVAST LABORATORIES LTD
 DASETTA 1/35, ETHINYL ESTRADIOL
 DASETTA 7/7/7, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ******* NOVAST LABORATORIES LTD**

ELINEST, ETHINYL ESTRADIOL
 FALMINA, ETHINYL ESTRADIOL
 LEVONEST, ETHINYL ESTRADIOL
 MONO-LINYAH, ETHINYL ESTRADIOL
 PHILITH, ETHINYL ESTRADIOL
 TRI-LINYAH, ETHINYL ESTRADIOL
 WERA, ETHINYL ESTRADIOL

NOVATECH SA

* NOVATECH SA
 STERITALC, TALC

NOVEL LABS INC

* NOVEL LABORATORIES INC
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 CALCIPOTRIENE, CALCIPOTRIENE
 CARBIDOPA, CARBIDOPA
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DESOXIMETASONE, DESOXIMETASONE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 FAMOTIDINE, FAMOTIDINE
 FLUCYTOSINE, FLUCYTOSINE
 FLUOCINONIDE, FLUOCINONIDE
 HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 LEVONORGESTREL, LEVONORGESTREL (OTC)
 LINEZOLID, LINEZOLID
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MISOPROSTOL, MISOPROSTOL
 MORPHINE SULFATE, MORPHINE SULFATE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NITROFURANTOIN, NITROFURANTOIN
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PEG 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
 PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL
 PHENELZINE SULFATE, PHENELZINE SULFATE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE
 TEMAZEPAM, TEMAZEPAM
 TINIDAZOLE, TINIDAZOLE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIMETHOPRIM, TRIMETHOPRIM
 VORICONAZOLE, VORICONAZOLE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

NOVELGENIX THERAPS

* NOVELGENIX THERAPEUTICS PVT LTD
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

NOVEN

* NOVEN PHARMACEUTICALS INC
 MINIVELLE, ESTRADIOL

NOVEN PHARMS INC

* NOVEN PHARMACEUTICALS INC
 COMBIPATCH, ESTRADIOL
 DAYTRANA, METHYLPHENIDATE

NOVITIUM PHARMA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVITIUM PHARMA LLC
 DAPSONE, DAPSONE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 RANITIDINE, RANITIDINE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

NOVO

* NOVO NORDISK INC
 FIASP FLEXTOUCH, INSULIN ASPART
 FIASP, INSULIN ASPART
 MACRILEN, MACIMORELIN ACETATE
 OZEMPIC, SEMAGLUTIDE
 RYZODEG 70/30, INSULIN ASPART
 SAXENDA, LIRAGLUTIDE RECOMBINANT
 TRESIBA, INSULIN DEGLUDEC
 XULTOPHY 100/3.6, INSULIN DEGLUDEC

NOVO NORDISK

* NOVO NORDISK PHARMACEUTICALS INC
 GLUCAGEN, GLUCAGON HYDROCHLORIDE

NOVO NORDISK INC

* NOVO NORDISK INC
 LEVEMIR FLEXTOUCH, INSULIN DETEMIR RECOMBINANT
 LEVEMIR, INSULIN DETEMIR RECOMBINANT
 NORDITROPIN FLEXPRO, SOMATROPIN RECOMBINANT
 NOVOLIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
 NOVOLIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)
 NOVOLIN R, INSULIN RECOMBINANT HUMAN (OTC)
 NOVOLOG FLEXPEN, INSULIN ASPART RECOMBINANT
 NOVOLOG MIX 70/30 FLEXPEN, INSULIN ASPART PROTAMINE RECOMBINANT
 NOVOLOG MIX 70/30, INSULIN ASPART PROTAMINE RECOMBINANT
 NOVOLOG PENFILL, INSULIN ASPART RECOMBINANT
 NOVOLOG, INSULIN ASPART RECOMBINANT
 VAGIFEM, ESTRADIOL
 VICTOZA, LIRAGLUTIDE RECOMBINANT

NOVOCOL INC

* NOVOCOL INC
 DYCLOPRO, DYCLONINE HYDROCHLORIDE

NPS PHARMS INC

* NPS PHARMACEUTICALS INC
 GATTEX KIT, TEDUGLUTIDE RECOMBINANT

NUVO PHARM

* NUVO PHARMACEUTICAL INC
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

NUVO PHARMS INC

* NUVO PHAMACEUTICALS INC
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 FOLIC ACID, FOLIC ACID
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE

NXDC

* NX DEVELOPMENT CORP
 GLEOLAN, AMINOLEVULINIC ACID HYDROCHLORIDE

NYCOMED US

* NYCOMED US INC
 TERCONAZOLE, TERCONAZOLE

**** O ******OAK PHARMS**

* OAK PHARMACEUTICALS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** O **

* OAK PHARMACEUTICALS INC
NEMBUTAL SODIUM, PENTOBARBITAL SODIUM
XYLOCAINE, LIDOCAINE HYDROCHLORIDE

OAK PHARMS AKORN

* OAK PHARMACEUTICALS INC SUB AKORN INC
COGENTIN, BENZTROPINE MESYLATE
DIURIL, CHLOROTHIAZIDE SODIUM

OAK PHARMS INC

* OAK PHARMACEUTICALS INC
ZIOPTAN, TAFLUPROST
* OAK PHARMACEUTICALS INC SUBSIDIARY OF AKORN INC
AZASITE, AZITHROMYCIN
BETIMOL, TIMOLOL
COSOPT PF, DORZOLAMIDE HYDROCHLORIDE
COSOPT, DORZOLAMIDE HYDROCHLORIDE
XOPENEX, LEVALBUTEROL HYDROCHLORIDE

OC PHARMA

* OC PHARMA LLC
LEVONORGESTREL, LEVONORGESTREL (OTC)
NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

OCULAR THERAPEUTIX

* OCULAR THERAPEUTIX INC
DEXTENZA, DEXAMETHASONE

ODYSSEY PHARMS

* ODYSSEY PHARMACEUTICALS INC
ANTABUSE, DISULFIRAM
SURMONTIL, TRIMIPRAMINE MALEATE
URECHOLINE, BETHANECHOL CHLORIDE
VIVACTIL, PROTRIPTYLINE HYDROCHLORIDE

OHM

* OHM CORP
IBUPROFEN, IBUPROFEN (OTC)

OHM LABS

* OHM LABORATORIES INC
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)
IBUPROHM, IBUPROFEN (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)

OHM LABS INC

* OHM LABORATORIES INC
EZETIMIBE, EZETIMIBE
GUAIFENESIN, GUAIFENESIN (OTC)
VALSARTAN, VALSARTAN

OLTA PHARMS

* OLTA PHARMACEUTICALS CORP
LINDANE, LINDANE

OMEROS

* OMEROS CORP
OMIDRIA, KETOROLAC TROMETHAMINE

ONY

* ONY INC
INFASURF PRESERVATIVE FREE, CALFACTANT

ONYX THERAP

* ONYX THERAPEUTICS INC A WHOLLY OWNED SUB OF AMGEN INC
KYPROLIS, CARFILZOMIB

OPKO IRELAND GLOBAL

* OPKO IRELAND GLOBAL HOLDINGS LTD
RAYALDEE, CALCIFEDIOL

OPTINOSE US INC

* OPTINOSE US INC
XHANCE, FLUTICASONE PROPIONATE

ORAPHARMA

* ORAPHARMA INC
ARESTIN, MINOCYCLINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** O **

ORCHID HLTHCARE

* ORCHID HEALTHCARE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFDINIR, CEFDINIR
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CEFPROZIL, CEFPROZIL
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CEPHALEXIN, CEPHALEXIN
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 DESLORATADINE, DESLORATADINE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 ESZOPICLONE, ESZOPICLONE
 FELODIPINE, FELODIPINE
 GEMIFLOXACIN MESYLATE, GEMIFLOXACIN MESYLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN, LEVOFLOXACIN
 MODAFINIL, MODAFINIL
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZALEPLON, ZALEPLON

OREXO US INC

* OREXO US INC
 ZUBSOLV, BUPRENORPHINE HYDROCHLORIDE

ORGANON SUB MERCK

* ORGANON USA INC A SUB OF MERCK AND CO INC
 BRIDION, SUGAMMADEX SODIUM
 NUVARING, ETHINYL ESTRADIOL

ORGANON USA INC

* ORGANON USA INC
 DESOGEN, DESOGESTREL
 FOLLISTIM AQ, FOLLITROPIN ALFA/BETA
 GANIRELIX ACETATE, GANIRELIX ACETATE
 NEXPLANON, ETNOGESTREL
 PREGNYL, GONADOTROPIN, CHORIONIC
 REMERON SOLTAB, MIRTAZAPINE
 REMERON, MIRTAZAPINE

ORIENT PHARMA CO LTD

* ORIENT PHARMA CO LTD
 CARISOPRODOL, CARISOPRODOL
 MIGLITOL, MIGLITOL
 PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM

ORION PHARMA

* ORION PHARMA
 COMTAN, ENTACAPONE
 STALEVO 100, CARBIDOPA
 STALEVO 125, CARBIDOPA
 STALEVO 150, CARBIDOPA
 STALEVO 200, CARBIDOPA
 STALEVO 50, CARBIDOPA
 STALEVO 75, CARBIDOPA

ORIT LABS LLC

* ORIT LABORATORIES LLC
 BENZONATATE, BENZONATATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** O **

* ORIT LABORATORIES LLC
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 ERGOCALCIFEROL, ERGOCALCIFEROL
 GLYCOPYRROLATE, GLYCOPYRROLATE
 LEVETIRACETAM, LEVETIRACETAM
 METRONIDAZOLE, METRONIDAZOLE

ORPHAN EUROPE

* ORPHAN EUROPE
 CARBAGLU, CARGLUMIC ACID
 * ORPHAN EUROPE SARL
 CYSTADANE, BETAINE

OSI PHARMS

* OSI PHARMACEUTICALS INC
 TARCEVA, ERLOTINIB HYDROCHLORIDE

OSMOTICA

* OSMOTICA KERESKEDELMI ES SZOLGALTATO KFT
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

OSMOTICA PHARM

* OSMOTICA PHARMACEUTICAL
 OSMOLEX ER, AMANTADINE HYDROCHLORIDE
 * OSMOTICA PHARMACEUTICAL CORP
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

OSMOTICA PHARM CORP

* OSMOTICA PHARMACEUTICAL CORP
 KHEDEZLA, DESVENLAFAXINE

OSMOTICA PHARM US

* OSMOTICA PHARMACEUTICAL US LLC
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 NIFEDIPINE, NIFEDIPINE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

OTONOMY INC

* OTONOMY INC
 OTIPRIO, CIPROFLOXACIN

OTSUKA

* OTSUKA PHARMACEUTICAL CO LTD
 ABILIFY, ARIPIPIRAZOLE

OTSUKA AMERICA PHARM

* OTSUKA AMERICA PHARMACEUTICAL INC
 SAMSCA, TOLVAPTAN

OTSUKA PHARM

* OTSUKA PHARMACEUTICAL CO LTD
 BUSULFEX, BUSULFAN

OTSUKA PHARM CO LTD

* OTSUKA PHARMACEUTICAL CO LTD
 ABILIFY MAINTENA KIT, ARIPIPIRAZOLE
 ABILIFY MYCITE KIT, ARIPIPIRAZOLE
 DACOGEN, DECITABINE
 JYNARQUE, TOLVAPTAN
 REXULTI, BREXPIPIRAZOLE

OUTLOOK PHARMS

* OUTLOOK PHARMACEUTICALS INC
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE

OXFORD PHARMS

* OXFORD PHARMACEUTICALS LLC
 ALPRAZOLAM, ALPRAZOLAM
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 BACLOFEN, BACLOFEN
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** O ****

* OXFORD PHARMACEUTICALS LLC
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LORAZEPAM, LORAZEPAM
 METHOCARBAMOL, METHOCARBAMOL
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PRIMIDONE, PRIMIDONE
 RIMACTANE, RIFAMPIN
 RISPERIDONE, RISPERIDONE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SIMVASTATIN, SIMVASTATIN
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

**** P ******P AND L DEV LLC**

* P AND L DEVELOPMENT LLC DBA PLD DEVELOPMENTS LLC
 IBUPROFEN, IBUPROFEN (OTC)

PACIFIC PHARMA

* PACIFIC PHARMA
 TIMOLOL MALEATE, TIMOLOL MALEATE
 * PACIFIC PHARMA INC
 TIMOLOL MALEATE, TIMOLOL MALEATE

PACIRA PHARMS INC

* PACIRA PHARMACEUTICALS INC
 EXPAREL, BUPIVACAINE

PACK PHARMS LLC

* PACK PHARMACEUTICALS LLC
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE

PADDOCK LLC

* PADDOCK LABORATORIES LLC
 ATOVAQUONE, ATOVAQUONE
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CALCIUM ACETATE, CALCIUM ACETATE
 CICLOPIROX, CICLOPIROX
 CLINDA-DERM, CLINDAMYCIN PHOSPHATE
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 COLOCORT, HYDROCORTISONE
 COMPRO, PROCHLORPERAZINE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROCODONE BITARTRATE AND PSEUDOEPHEDRINE HYDROCHLORIDE, HYDROCODONE BITARTRATE
 HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE,
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 KIONEX, SODIUM POLYSTYRENE SULFONATE
 LAX-LYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
 MIDAMOR, AMILORIDE HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NYSTOP, NYSTATIN
 PODOFILOX, PODOFILOX
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 REPAGLINIDE, REPAGLINIDE
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PADDOCK LABORATORIES LLC
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TROSPIUM CHLORIDE, TROSPIUM CHLORIDE

PANACEA BIOTEC LTD

* PANACEA BIOTEC LTD
 PRASUGREL, PRASUGREL HYDROCHLORIDE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 TACROLIMUS, TACROLIMUS

PAR FORM

* PAR FORMULATIONS PRIVATE LTD
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 MAFENIDE ACETATE, MAFENIDE ACETATE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

PAR PHARM

* PAR PHARMACEUTICAL
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 TESTOSTERONE, TESTOSTERONE

* PAR PHARMACEUTICAL INC
 ALPRAZOLAM, ALPRAZOLAM
 AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 CABERGOLINE, CABERGOLINE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CLOMIPHENE CITRATE, CLOMIPHENE CITRATE
 CLONAZEPAM, CLONAZEPAM
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE
 ESTAZOLAM, ESTAZOLAM
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUTAMIDE, FLUTAMIDE
 GLIPIZIDE, GLIPIZIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROXYUREA, HYDROXYUREA
 IBUPROFEN, IBUPROFEN (OTC)
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 LAMOTRIGINE, LAMOTRIGINE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
 METRONIDAZOLE, METRONIDAZOLE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MINOXIDIL, MINOXIDIL
 NATEGLINIDE, NATEGLINIDE
 NIFEDIPINE, NIFEDIPINE
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
 OXANDROLONE, OXANDROLONE
 PIMOZIDE, PIMOZIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PAR PHARMACEUTICAL INC
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TORSEMIDE, TORSEMIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRANYLCPROMINE SULFATE, TRANYLCPROMINE SULFATE
 TRAVOPROST, TRAVOPROST
 URSODIOL, URSODIOL
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

PAR PHARM INC

* PAR PHARMACEUTICAL INC
 ACCOLATE, ZAFIRLUKAST
 ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 ANTIZOL, FOMEPIZOLE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
 COLCHICINE, COLCHICINE
 DEXLANSOPRAZOLE, DEXLANSOPRAZOLE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DOFETILIDE, DOFETILIDE
 DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
 ENTECAVIR, ENTECAVIR
 ETHACRYNIC ACID, ETHACRYNIC ACID
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 ITRACONAZOLE, ITRACONAZOLE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 OLMESARTAN MEDOXMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 PRAZIQUANTEL, PRAZIQUANTEL
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOLCAPONE, TOLCAPONE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VIGABATRIN, VIGABATRIN
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

PAR STERILE PRODUCTS

* PAR STERILE PRODUCTS LLC
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 ADRENALIN, EPINEPHRINE
 ARGATROBAN, ARGATROBAN
 BREVITAL SODIUM, METHOHEXITAL SODIUM
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 COLY-MYCIN M, COLISTIMETHATE SODIUM
 CORPHEDRA, EPHEDRINE SULFATE
 DANTRIUM, DANTROLENE SODIUM
 DELESTROGEN, ESTRADIOL VALERATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
 ETOMIDATE, ETOMIDATE
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 GANCICLOVIR, GANCICLOVIR SODIUM
 KETALAR, KETAMINE HYDROCHLORIDE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
 NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
 PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ******* PAR STERILE PRODUCTS LLC**

PITOCIN, OXYTOCIN
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE
 TRIOSTAT, LIOTHYRONINE SODIUM
 VASOSTRICT, VASOPRESSIN

PARAGON BIOTECK*** PARAGON BIOTECK INC**

PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE

PARAPRO LLC*** PARAPRO LLC**

NATROBA, SPINOSAD

PARATEK PHARMS INC*** PARATEK PHARMACEUTICALS INC**

NUZYRA, OMADACYCLINE TOSYLATE

PARKE DAVIS*** PARKE DAVIS DIV WARNER LAMBERT CO**

CELONTIN, METHSUXIMIDE
 CEREBYX, FOSPHENYTOIN SODIUM
 DILANTIN-125, PHENYTOIN
 NARDIL, PHENELZINE SULFATE
 NEURONTIN, GABAPENTIN
 ZARONTIN, ETHOSUXIMIDE

PARKE-DAVIS*** PARKE-DAVIS DIVISION OF PFIZER INC**

DILANTIN, PHENYTOIN SODIUM
 ZARONTIN, ETHOSUXIMIDE

PATRIN PHARMA INC*** PATRIN PHARMA INC**

FLAC, FLUOCINOLONE ACETONIDE

PERNIX IRELAND LTD*** PERNIX IRELAND LTD**

TREXIMET, NAPROXEN SODIUM

PERNIX IRELAND PAIN*** PERNIX IRELAND PAIN LIMITED**

ZOHYDRO ER, HYDROCODONE BITARTRATE

PERNIX THERAPS LLC*** PERNIX THERAPEUTICS LLC**

SILENOR, DOXEPIN HYDROCHLORIDE

PERRIGO*** PERRIGO CO**

MICONAZOLE NITRATE COMBINATION PACK, MICONAZOLE NITRATE (OTC)

PERRIGO CO*** PERRIGO CO OF TENNESSEE INC**

CICLOPIROX, CICLOPIROX
 CLINDETS, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 ERYTHROMYCIN, ERYTHROMYCIN
 STIE-CORT, HYDROCORTISONE

PERRIGO CO TENNESSEE*** PERRIGO CO TENNESSEE INC**

BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 BACITRACIN, BACITRACIN
 BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE, BACITRACIN ZINC
 ERYTHROMYCIN, ERYTHROMYCIN
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
 NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM

PERRIGO ISRAEL*** PERRIGO ISRAEL PHARMACEUTICALS LTD**

ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PERRIGO ISRAEL PHARMACEUTICALS LTD
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DESOXIMETASONE, DESOXIMETASONE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 GYNAZOLE-1, BUTOCONAZOLE NITRATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 IMIQUIMOD, IMIQUIMOD
 KETOCONAZOLE, KETOCONAZOLE
 MESALAMINE, MESALAMINE
 MINOXIDIL, MINOXIDIL (OTC)
 MOMETASONE FUROATE, MOMETASONE FUROATE
 NITROGLYCERIN, NITROGLYCERIN
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 TESTOSTERONE, TESTOSTERONE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE (OTC)

PERRIGO NEW YORK

* PERRIGO NEW YORK INC
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 AMMONIUM LACTATE, AMMONIUM LACTATE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CENTANY, MUPIROCIN
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 DESONIDE, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 ECONAZOLE NITRATE, ECONAZOLE NITRATE
 ERYTHROMYCIN, ERYTHROMYCIN
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 HYDROCORTISONE, HYDROCORTISONE
 KETOCONAZOLE, KETOCONAZOLE
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MUPIROCIN, MUPIROCIN
 NYSTATIN, NYSTATIN
 PERMETHRIN, PERMETHRIN
 PERMETHRIN, PERMETHRIN (OTC)
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 SELENIUM SULFIDE, SELENIUM SULFIDE
 TERCONAZOLE, TERCONAZOLE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

PERRIGO PHARMA INTL

* PERRIGO PHARMA INTERNATIONAL DAC
 CLINDESSE, CLINDAMYCIN PHOSPHATE
 ENTOCORT EC, BUDESONIDE
 EVAMIST, ESTRADIOL
 LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
 LORATADINE, LORATADINE (OTC)
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 TRETINOIN, TRETINOIN

PERRIGO PHARMS CO

* PERRIGO PHARMACEUTICALS CO
 SCOPOLAMINE, SCOPOLAMINE

PERRIGO R AND D

* PERRIGO R AND D CO
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ******* PERRIGO R AND D CO**

DES Loratadine, Desloratadine
 Esomeprazole Magnesium, Esomeprazole Magnesium (OTC)
 Famotidine, Calcium Carbonate, and Magnesium Hydroxide, Calcium Carbonate (OTC)
 Famotidine, Famotidine
 Famotidine, Famotidine (OTC)
 Guaifenesin and Dextromethorphan Hydrobromide, Dextromethorphan Hydrobromide (OTC)
 Guaifenesin, Guaifenesin (OTC)
 Ibuprofen and Diphenhydramine Citrate, Diphenhydramine Citrate (OTC)
 Ibuprofen and Phenylephrine Hydrochloride, Ibuprofen (OTC)
 Ibuprofen Sodium, Ibuprofen Sodium (OTC)
 Ibuprofen, Ibuprofen
 Ibuprofen, Ibuprofen (OTC)
 Lansoprazole, Lansoprazole (OTC)
 Levonorgestrel, Levonorgestrel
 Levonorgestrel, Levonorgestrel (OTC)
 Loperamide Hydrochloride and Simethicone, Loperamide Hydrochloride (OTC)
 Loperamide Hydrochloride, Loperamide Hydrochloride (OTC)
 Miconazole Nitrate, Miconazole Nitrate (OTC)
 Montelukast Sodium, Montelukast Sodium
 Naproxen, Naproxen
 Nicotine Polacrilex, Nicotine Polacrilex (OTC)
 Omeprazole and Sodium Bicarbonate, Omeprazole (OTC)
 Omeprazole Magnesium, Omeprazole Magnesium (OTC)
 Pantoprazole Sodium, Pantoprazole Sodium
 Polyethylene Glycol 3350, Polyethylene Glycol 3350 (OTC)
 Ranitidine Hydrochloride, Ranitidine Hydrochloride (OTC)

PERRIGO UK FINCO*** PERRIGO UK FINCO LTD PARTNERSHIP**

Betamethasone Valerate, Betamethasone Valerate
 Clindamycin Phosphate, Clindamycin Phosphate
 Diclofenac Sodium, Diclofenac Sodium
 Estradiol, Estradiol
 Flurandrenolide, Flurandrenolide
 Ingenol Mebutate, Ingenol Mebutate
 Nystatin and Triamcinolone Acetonide, Nystatin
 Testosterone, Testosterone
 Triamcinolone Acetonide, Triamcinolone Acetonide

PETNET*** PETNET SOLUTIONS INC**

Ammonia N 13, Ammonia N-13
 Fludeoxyglucose F18, Fludeoxyglucose F-18
 Sodium Fluoride F-18, Sodium Fluoride F-18

PF PRISM CV*** PF PRISM CV**

Bosulif, Bosutinib Monohydrate
 Chantix, Varenicline Tartrate
 Inlyta, Axitinib
 Lyrica CR, Pregabalin
 Lyrica, Pregabalin
 Pristiq, Desvenlafaxine Succinate
 Rapamune, Sirolimus
 Torisel, Temsirolimus
 Tygacil, Tigecycline
 Vfend, Voriconazole
 Xalkori, Crizotinib
 Xeljanz, Tofacitinib Citrate

PFIZER*** PFIZER CENTRAL RESEARCH**

Diflucan, Fluconazole
 Zithromax, Azithromycin

*** PFIZER CHEMICALS DIV PFIZER INC**

Diflucan, Fluconazole
 Zithromax, Azithromycin

*** PFIZER INC**

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

- * PFIZER INC
 ADVIL ALLERGY AND CONGESTION RELIEF, CHLORPHENIRAMINE MALEATE (OTC)
 ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
 ADVIL COLD AND SINUS, IBUPROFEN (OTC)
 ADVIL CONGESTION RELIEF, IBUPROFEN (OTC)
 ADVIL LIQUI-GELS, IBUPROFEN (OTC)
 ADVIL MIGRAINE LIQUI-GELS, IBUPROFEN (OTC)
 ADVIL MULTI-SYMPTOM COLD & FLU, CHLORPHENIRAMINE MALEATE (OTC)
 ADVIL PM, DIPHENHYDRAMINE CITRATE (OTC)
 ADVIL PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 ADVIL, IBUPROFEN (OTC)
 ALAVERT, LORATADINE (OTC)
 AXID AR, NIZATIDINE (OTC)
 CADUET, AMLODIPINE BESYLATE
 CALAN SR, VERAPAMIL HYDROCHLORIDE
 CARDURA XL, DOXAZOSIN MESYLATE
 CHILDREN'S ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
 CHILDREN'S ADVIL COLD, IBUPROFEN (OTC)
 CHILDREN'S ADVIL, IBUPROFEN (OTC)
 CHILDREN'S ADVIL-FLAVORED, IBUPROFEN (OTC)
 ELELYSO, TALIGLUCERASE ALFA
 GEODON, ZIPRASIDONE HYDROCHLORIDE
 GEODON, ZIPRASIDONE MESYLATE
 GLUCOTROL XL, GLIPIZIDE
 GLUCOTROL, GLIPIZIDE
 HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 HEPARIN SODIUM, HEPARIN SODIUM
 JUNIOR STRENGTH ADVIL, IBUPROFEN (OTC)
 LIPITOR, ATORVASTATIN CALCIUM
 LORATADINE, LORATADINE (OTC)
 MERREM, MEROPENEM
 NORVASC, AMLODIPINE BESYLATE
 PEDIATRIC ADVIL, IBUPROFEN (OTC)
 PROCARDIA, NIFEDIPINE
 REVATIO, SILDENAFIL CITRATE
 SONATA, ZALEPLON
 TESSALON, BENZONATATE
 TOVIAZ, FESOTERODINE FUMARATE
 UNASYN, AMPICILLIN SODIUM
 ZITHROMAX, AZITHROMYCIN
- * PFIZER LABORATORIES DIV PFIZER INC
 CARDURA, DOXAZOSIN MESYLATE
 FELDENE, PIROXICAM
 MINIPRESS, PRAZOSIN HYDROCHLORIDE
 PFIZERPEN, PENICILLIN G POTASSIUM
 PROCARDIA XL, NIFEDIPINE
 UNASYN, AMPICILLIN SODIUM
 VIBRAMYCIN, DOXYCYCLINE
 VIBRAMYCIN, DOXYCYCLINE CALCIUM
 VIBRAMYCIN, DOXYCYCLINE HYCLATE
 VISTARIL, HYDROXYZINE PAMOATE
- * PFIZER PHARMACEUTICALS INC
 DILANTIN, PHENYTOIN
 ZOLOFT, SERTRALINE HYDROCHLORIDE
- * PFIZER PHARMACEUTICALS PRODUCTION CORP LTD
 TIKOSYN, DOFETILIDE
- PFIZER CONS HLTHCARE**
- * PFIZER CONSUMER HEALTHCARE
 ADVIL, IBUPROFEN SODIUM (OTC)
- PFIZER INC**
- * PFIZER INC
 CAMPTOSAR, IRINOTECAN HYDROCHLORIDE
 DAURISMO, GLASDEGIB
 ELLENCE, EPIRUBICIN HYDROCHLORIDE
 FRAGMIN, DALTEPARIN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PFIZER INC
 IBRANCE, PALBOCICLIB
 LORBRENA, LORLATINIB
 NICOTROL, NICOTINE
 TALZENNA, TALAZOPARIB TOSYLATE
 VIAGRA, SILDENAFIL CITRATE
 VIZIMPRO, DACOMITINIB
 XELJANZ XR, TOFACITINIB CITRATE

PFIZER IRELAND

* PFIZER IRELAND PHARMACEUTICALS
 RELPAX, ELETRIPTAN HYDROBROMIDE

PFIZER PHARMS

* PFIZER PHARMACEUTICALS LTD
 ACCUPRIL, QUINAPRIL HYDROCHLORIDE
 ACCURETIC, HYDROCHLOROTHIAZIDE
 LOPID, GEMFIBROZIL
 NEURONTIN, GABAPENTIN
 NITROSTAT, NITROGLYCERIN

PHARM ASSOC

* PHARMACEUTICAL ASSOC INC
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 LORAZEPAM, LORAZEPAM
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE

* PHARMACEUTICAL ASSOCIATES INC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 LACTULOSE, LACTULOSE
 LEVETIRACETAM, LEVETIRACETAM
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PREDNISOLONE, PREDNISOLONE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE
 THEOPHYLLINE, THEOPHYLLINE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 VALPROIC ACID, VALPROIC ACID

PHARMA RES SOFTWARE

* PHARMA RESEARCH SOFTWARE SOLUTION LLC
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

PHARMACHEMIE BV

* PHARMACHEMIE BV
 CARBOPLATIN, CARBOPLATIN
 CISPLATIN, CISPLATIN
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM

PHARMACIA AND UPJOHN

* PHARMACIA AND UPJOHN
 XANAX XR, ALPRAZOLAM

* PHARMACIA AND UPJOHN CO
 AROMASIN, EXEMESTANE
 AZULFIDINE EN-TABS, SULFASALAZINE
 AZULFIDINE, SULFASALAZINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

- * PHARMACIA AND UPJOHN CO
 BACITRACIN, BACITRACIN
 CAVERJECT IMPULSE, ALPROSTADIL
 CAVERJECT, ALPROSTADIL
 CLEOCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 CLEOCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLEOCIN T, CLINDAMYCIN PHOSPHATE
 CLEOCIN, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLEOCIN, CLINDAMYCIN PHOSPHATE
 CORTEF, HYDROCORTISONE
 CORVERT, IBUTILIDE FUMARATE
 CYKLOKAPRON, TRANEXAMIC ACID
 DEPO-ESTRADIOL, ESTRADIOL CYPIONATE
 DEPO-MEDROL, METHYLPREDNISOLONE ACETATE
 DEPO-PROVERA, MEDROXYPROGESTERONE ACETATE
 DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE
 DETROL LA, TOLTERODINE TARTRATE
 DETROL, TOLTERODINE TARTRATE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 EMCYT, ESTRAMUSTINE PHOSPHATE SODIUM
 ESTRING, ESTRADIOL
 GENOTROPIN PRESERVATIVE FREE, SOMATROPIN RECOMBINANT
 GENOTROPIN, SOMATROPIN RECOMBINANT
 GLYNASE, GLYBURIDE
 GLYSET, MIGLITOL
 HALCION, TRIAZOLAM
 HEMABATE, CARBOPROST TROMETHAMINE
 IDAMYCIN PFS, IDARUBICIN HYDROCHLORIDE
 LINCOCIN, LINCOMYCIN HYDROCHLORIDE
 MEDROL, METHYLPREDNISOLONE
 MYCOBUTIN, RIFABUTIN
 NICOTROL, NICOTINE
 OGEN 5, ESTROPIPATE
 PREPIDIL, DINOPROSTONE
 PROSTIN E2, DINOPROSTONE
 PROSTIN VR PEDIATRIC, ALPROSTADIL
 PROVERA, MEDROXYPROGESTERONE ACETATE
 R-GENE 10, ARGININE HYDROCHLORIDE
 SOLU-CORTEF, HYDROCORTISONE SODIUM SUCCINATE
 SOLU-MEDROL, METHYLPREDNISOLONE SODIUM SUCCINATE
 SOMAVERT, PEGVISOMANT
 XALATAN, LATANOPROST
 XANAX, ALPRAZOLAM
 ZINECARD, DEXRAZOXANE HYDROCHLORIDE
 ZYVOX, LINEZOLID
- * PHARMACIA AND UPJOHN SUB PFIZER INC
 DEPO-SUBQ PROVERA 104, MEDROXYPROGESTERONE ACETATE
- PHARMACIA UPJOHN**
- * PHARMACIA AND UPJOHN CO A SUB OF PFIZER INC
 COLESTID, COLESTIPOL HYDROCHLORIDE
 FLAVORED COLESTID, COLESTIPOL HYDROCHLORIDE
- PHARMACYCLICS INC**
- * PHARMACYCLICS INC
 IMBRUVICA, IBRUTINIB
- PHARMADAX INC**
- * PHARMADAX INC
 GLYBURIDE, GLYBURIDE
 LEVETIRACETAM, LEVETIRACETAM
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
- PHARMALUCENCE**
- * PHARMALUCENCE INC
 AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT
 CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT
 CIS-PYRO, TECHNETIUM TC-99M PYROPHOSPHATE KIT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PHARMALUCENCE INC
 HEPATOLITE, TECHNETIUM TC-99M DISOFENIN KIT
 TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
 TECHNETIUM TC-99M MEBROFENIN, TECHNETIUM TC-99M MEBROFENIN KIT

PHARMASCIENCE INC

* PHARMASCIENCE INC
 BUSULFAN, BUSULFAN
 DECITABINE, DECITABINE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM

PHARMAXIS LTD

* PHARMAXIS LTD
 ARIDOL KIT, MANNITOL

PHARMTAK INC

* PHARMTAK INC
 LEVETIRACETAM, LEVETIRACETAM

PHOTOCURE ASA

* PHOTOCURE ASA
 CYSVIEW KIT, HEXAMINOLEVULINATE HYDROCHLORIDE

PIERRE FABRE

* PIERRE FABRE MEDICAMENT
 NAVELBINE, VINORELBINE TARTRATE

PIERRE FABRE DERMA

* PIERRE FABRE DERMATOLOGIE
 HEMANGEOL, PROPRANOLOL HYDROCHLORIDE

PIERREL

* PIERREL S.P.A.
 ORABLOC, ARTICAINA HYDROCHLORIDE

PII

* PHARMACEUTICS INTERNATIONAL INC
 BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
 DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 HYDROCORTISONE, HYDROCORTISONE
 PIROXICAM, PIROXICAM
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

PIRAMAL CRITICAL

* PIRAMAL CRITICAL CARE INC
 ISOFLURANE, ISOFLURANE
 SOJOURN, SEVOFLURANE
 * PIRAMAL CRITICAL CARE LTD
 GABLOFEN, BACLOFEN
 GLYCOPYRROLATE, GLYCOPYRROLATE
 LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 MITIGO, MORPHINE SULFATE

PIRAMAL ENT

* PIRAMAL ENTERPRISES LTD
 ISOFLURANE, ISOFLURANE

PIRAMAL HLTHCARE UK

* PIRAMAL HEALTHCARE UK LTD
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM

PLD ACQUISITIONS

* PLD ACQUISITIONS LLC DBA AVENA PHARMA SOLUTIONS
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE

PLD ACQUISITIONS LLC

* PLD ACQUISITIONS LLC
 LORATADINE, LORATADINE (OTC)
 ZOLMITRIPTAN, ZOLMITRIPTAN

PLIVA

* PLIVA INC
 AZITHROMYCIN, AZITHROMYCIN
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 CIMETIDINE, CIMETIDINE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ******* PLIVA INC**

CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 METRONIDAZOLE, METRONIDAZOLE
 NAPROXEN, NAPROXEN
 THEOPHYLLINE, THEOPHYLLINE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 WARFARIN SODIUM, WARFARIN SODIUM

PLIVA HRVATSKA DOO*** PLIVA HRVATSKA DOO**

ADAPALENE, ADAPALENE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE

PLIVA LACHEMA*** PLIVA LACHEMA AS**

CARBOPLATIN, CARBOPLATIN
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM

PLIVA PHARM IND*** PLIVA PHARMACEUTICAL INDUSTRY INC**

TORSEMIDE, TORSEMIDE

PLX PHARMA*** PLX PHARMA INC**

VAZALORE, ASPIRIN (OTC)

POHL BOSKAMP*** POHL BOSKAMP**

NITROLINGUAL PUMPSPRAY, NITROGLYCERIN

POLYGEN PHARMS*** POLYGEN PHARMACEUTICALS INC**

AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

POLYMEDICA*** POLYMEDICA INDUSTRIES INC**

NEOPAP, ACETAMINOPHEN (OTC)

PORTOLA PHARMS INC*** PORTOLA PHARMACEUTICALS INC**

BEVYXXA, BETRIXABAN

POWDER PHARMS*** POWDER PHARMACEUTICALS INC**

ZINGO, LIDOCAINE HYDROCHLORIDE

PRAGMA*** PRAGMA PHARMACEUTICALS LLC**

KEFLEX, CEPHALEXIN

PRAXAIR DISTRIBUTION*** PRAXAIR DISTRIBUTION INC**

NOXIVENT, NITRIC OXIDE

PRECISION DERMAT*** PRECISION DERMATOLOGY INC**

CLINDAGEL, CLINDAMYCIN PHOSPHATE
 LOCOID LIPOCREAM, HYDROCORTISONE BUTYRATE
 LOCOID, HYDROCORTISONE BUTYRATE
 MINOCIN, MINOCYCLINE HYDROCHLORIDE

PRECISION DOSE INC*** PRECISION DOSE INC**

PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE

PRECISION NUCLEAR*** PRECISION NUCLEAR LLC**

AMMONIA N 13, AMMONIA N-13

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PRECISION NUCLEAR LLC
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

PRINSTON INC

* PRINSTON PHARMACEUTICAL INC
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CAPTOPRIL, CAPTOPRIL
 CLONAZEPAM, CLONAZEPAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTECAVIR, ENTECAVIR
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 FENOFIBRATE, FENOFIBRATE
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 FUROSEMIDE, FUROSEMIDE
 GLIMEPIRIDE, GLIMEPIRIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 LEVETIRACETAM, LEVETIRACETAM
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINOPRIL, LISINOPRIL
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHOCARBAMOL, METHOCARBAMOL
 NEVIRAPINE, NEVIRAPINE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 PAROXETINE MESYLATE, PAROXETINE MESYLATE
 PAROXETINE, PAROXETINE HYDROCHLORIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMAZEPAM, TEMAZEPAM
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALSARTAN, VALSARTAN
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE

PROF DSPLS

* PROFESSIONAL DISPOSABLES INC
 PREVANTICS MAXI SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 PREVANTICS SWAB, CHLORHEXIDINE GLUCONATE (OTC)
 PREVANTICS SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)

PROGENICS PHARMS INC

* PROGENICS PHARMACEUTICALS INC
 AZEDRA, IOBENGUANE I-131

PROMIUS PHARMA LLC

* PROMIUS PHARMA LLC
 SERNIVO, BETAMETHASONE DIPROPIONATE

PROVELL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PROVELL PHARMACEUTICALS LLC
EUTHYROX, LEVOTHYROXINE SODIUM **

PROVENSIS

* PROVENSIS LTD
VARITHENA, POLIDOCANOL

PROVEPHARM SAS

* PROVEPHARM SAS
PROVAYBLUE, METHYLENE BLUE

PTC THERAP

* PTC THERAPEUTICS INC
EMFLAZA, DEFLAZACORT

PULMOFLOW INC

* PULMOFLOW INC
KITABIS PAK, TOBRAMYCIN

PUMA BIOTECH

* PUMA BIOTECHNOLOGY INC
NERLYNX, NERATINIB MALEATE

PURACAP PHARM

* PURACAP PHARMACEUTICAL LLC
MELOXICAM, MELOXICAM

PURACAP PHARM LLC

* PURACAP PHARMACEUTICAL LLC
BENZONATATE, BENZONATATE
ERGOCALCIFEROL, ERGOCALCIFEROL
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE

PURDUE GMP

* PURDUE GMP CENTER LLC DBA THE CHAO CENTER INDUSTRIAL PHARMACY
SEROMYCIN, CYCLOSERINE

PURDUE PHARMA

* PURDUE PHARMA PRODUCTS LP
INTERMEZZO, ZOLPIDEM TARTRATE

PURDUE PHARMA LP

* PURDUE PHARMA LP
BUTRANS, BUPRENORPHINE
HYSINGLA, HYDROCODONE BITARTRATE
MS CONTIN, MORPHINE SULFATE
OXYCONTIN, OXYCODONE HYDROCHLORIDE

**** Q ******QILU PHARM CO LTD**

* QILU PHARMACEUTICAL CO LTD
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
CEFTRIAZONE, CEFTRIAZONE SODIUM
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
OLANZAPINE, OLANZAPINE
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
OXALIPLATIN, OXALIPLATIN
PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
TENOFIVIR DISOPROXIL FUMARATE, TENOFIVIR DISOPROXIL FUMARATE

QILU TIANHE

* QILU TIANHE PHARMACEUTICAL CO LTD
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM

QINGDAO BAHEAL PHARM

* QINGDAO BAHEAL PHARMACEUTICAL CO LTD
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

QOL MEDCL

* QOL MEDICAL LLC
ETHAMOLIN, ETHANOLAMINE OLEATE
SUCRAID, SACROSIDASE

QUAGEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Q ****

* QUAGEN PHARMACEUTICALS LLC
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

QUEEN HAMAMATSU PET

* QUEEN HAMAMATSU PET IMAGING CENTER
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**** R ******R-PHARM US LLC**

* R-PHARM US LLC
IXEMPRA KIT, IXABEPILONE

R2 PHARMA LLC

* R2 PHARMA LLC
ESOMEPRAZOLE STRONTIUM, ESOMEPRAZOLE STRONTIUM

RADIUS HEALTH INC

* RADIUS HEALTH INC
TYMLOS, ABALOPARATIDE

RB HLTH

* RB HEALTH US LLC
DELSYM, DEXTROMETHORPHAN POLISTIREX (OTC)
MUCINEX D, GUAIFENESIN (OTC)
MUCINEX DM, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
MUCINEX, GUAIFENESIN (OTC)

RECIP

* RECIP AB
THYROSAFE, POTASSIUM IODIDE (OTC)

RECIPHARM

* RECIPHARM PHARMASERVICES PRIVATE LTD
FLUCYTOSINE, FLUCYTOSINE

RECKITT BENCKISER

* RECKITT BENCKISER LLC
LEVONORGESTREL, LEVONORGESTREL (OTC)

RECORDATI RARE

* RECORDATI RARE DISEASES INC
CHEMET, SUCCIMER
COSMEGEN, DACTINOMYCIN
DESOXYN, METHAMPHETAMINE HYDROCHLORIDE
INDOCIN, INDOMETHACIN SODIUM
NEOPROFEN, IBUPROFEN LYSINE
PEGANONE, ETHOTOIN
TRANXENE, CLORAZEPATE DIPOTASSIUM

RECRO GAINESVILLE

* RECRO GAINESVILLE LLC
VERELAN PM, VERAPAMIL HYDROCHLORIDE
VERELAN, VERAPAMIL HYDROCHLORIDE

RELYPSA INC

* RELYPSA INC
VELTASSA, PATIROMER SORBITEX CALCIUM

REMPEX PHARMS

* REMPEX PHARMACEUTICALS
MINOCIN, MINOCYCLINE HYDROCHLORIDE
* REMPEX PHARMACEUTICALS A WHOLLY OWNED SUB OF MELINTA THERAPEUTICS INC
VABOMERE, MEROPENEM

RENAISSANCE SSA LLC

* RENAISSANCE SSA LLC
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
METHOCARBAMOL, METHOCARBAMOL
MILRINONE LACTATE IN DEXTROSE 5%, MILRINONE LACTATE
NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
OXACILLIN SODIUM, OXACILLIN SODIUM
PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

* RENAISSANCE SSA LLC
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE

RENATA

* RENATA LTD
RISPERIDONE, RISPERIDONE

RHODES PHARMS

* RHODES PHARMACEUTICALS LP
APTENSIO XR, METHYLPHENIDATE HYDROCHLORIDE
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
DILAUDID, HYDROMORPHONE HYDROCHLORIDE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FENOFIBRATE, FENOFIBRATE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
MORPHINE SULFATE, MORPHINE SULFATE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
THEOPHYLLINE, THEOPHYLLINE

RICONPHARMA LLC

* RICONPHARMA LLC
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
CHLORTHALIDONE, CHLORTHALIDONE
DESOXIMETASONE, DESOXIMETASONE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
LIDOCAINE, LIDOCAINE
METHYLPREDNISOLONE, METHYLPREDNISOLONE

RIGEL PHARMS INC

* RIGEL PHARMACEUTICALS INC
TAVALISSE, FOSTAMATINIB DISODIUM

RISING PHARMS

* RISING PHARMACEUTICALS INC
ACETIC ACID, ACETIC ACID, GLACIAL
ALBUTEROL SULFATE, ALBUTEROL SULFATE
BUDESONIDE, BUDESONIDE
CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
CROMOLYN SODIUM, CROMOLYN SODIUM
DESOXIMETASONE, DESOXIMETASONE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
DOXERCALCIFEROL, DOXERCALCIFEROL
DUTASTERIDE, DUTASTERIDE
GLYCOPYRROLATE, GLYCOPYRROLATE
HYDROCORTISONE, HYDROCORTISONE
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN, AMOXICILLIN
LEVOFLOXACIN, LEVOFLOXACIN
METAXALONE, METAXALONE
METHIMAZOLE, METHIMAZOLE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
PARICALCITOL, PARICALCITOL
PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
TEMOZOLOMIDE, TEMOZOLOMIDE
TOREMIFENE CITRATE, TOREMIFENE CITRATE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
VORICONAZOLE, VORICONAZOLE
ZILEUTON, ZILEUTON

ROCHE PALO

* ROCHE PALO ALTO LLC
CELLCEPT, MYCOPHENOLATE MOFETIL
CELLCEPT, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
CYTOVENE, GANCICLOVIR SODIUM

ROCKWELL MEDICAL INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

* ROCKWELL MEDICAL INC
TRIFERIC, FERRIC PYROPHOSPHATE CITRATE

ROMARK

* ROMARK LABORATORIES
ALINIA, NITAZOXANIDE

ROUSES POINT PHARMS

* ROUSES POINT PHARMACEUTICALS LLC
LEVETIRACETAM, LEVETIRACETAM

RP SCHERER

* RP SCHERER TECHNOLOGIES LLC
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

RTRX

* RETROPHIN INC
CHOLBAM, CHOLIC ACID

RUBICON RES PVT LTD

* RUBICON RESEARCH PVT LTD
BACLOFEN, BACLOFEN
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
SILDENAFIL CITRATE, SILDENAFIL CITRATE

RXMTM THERAPS LLC

* RXMTM THERAPEUTICS LLC A WHOLLY OWNED SUB OF CUTISPHARMA INC
FIRVANQ KIT, VANCOMYCIN HYDROCHLORIDE

**** S ******SAGE PRODS**

* SAGE PRODUCTS INC
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

SAGENT PHARMS

* SAGENT PHARMACEUTICALS INC
ACETYLCYSTEINE, ACETYLCYSTEINE
AMIKACIN SULFATE, AMIKACIN SULFATE
AMPICILLIN SODIUM, AMPICILLIN SODIUM
CAFFEINE CITRATE, CAFFEINE CITRATE
CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
DAPTOMYCIN, DAPTOMYCIN
DECITABINE, DECITABINE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
EPTIFIBATIDE, EPTIFIBATIDE
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
FLUMAZENIL, FLUMAZENIL
FLUOROURACIL, FLUOROURACIL
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
GLYDO, LIDOCAINE HYDROCHLORIDE
HALOPERIDOL, HALOPERIDOL LACTATE
HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
HEPARIN SODIUM, HEPARIN SODIUM
IBANDRONATE SODIUM, IBANDRONATE SODIUM
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
LEVETIRACETAM, LEVETIRACETAM
LINEZOLID, LINEZOLID
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
MESNA, MESNA
METHOCARBAMOL, METHOCARBAMOL
METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
NAFCILLIN SODIUM, NAFCILLIN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SAGENT PHARMACEUTICALS INC
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXACILLIN SODIUM, OXACILLIN SODIUM
 OXYTOCIN, OXYTOCIN
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM
 PROPOFOL, PROPOFOL
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

SAGENT STRIDES

* SAGENT STRIDES LLC
 MIDOZALAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE

SALIX PHARMS

* SALIX PHARMACEUTICALS INC
 ANUSOL HC, HYDROCORTISONE
 DIURIL, CHLOROTHIAZIDE
 MOVIPREP, ASCORBIC ACID
 OSMOPREP, SODIUM PHOSPHATE, DIBASIC, ANHYDROUS
 PEPCID, FAMOTIDINE
 RELISTOR, METHYLNALTREXONE BROMIDE
 XIFAXAN, RIFAXIMIN

SALIX PHARMS INC

* SALIX PHARMACEUTICALS INC
 PLENVU, ASCORBIC ACID
 RELISTOR, METHYLNALTREXONE BROMIDE

SAMSON MEDCL

* SAMSON MEDICAL TECHNOLOGIES LLC
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFEPIME HYDROCHLORIDE IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE
 CEFIXITIN IN PLASTIC CONTAINER, CEFIXITIN SODIUM
 CEFTRIAXONE, CEFTRIAXONE SODIUM
 VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE

SANDOZ

* SANDOZ
 DOCETAXEL, DOCETAXEL

* SANDOZ INC
 ALPRAZOLAM, ALPRAZOLAM
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE
 APREPITANT, APREPITANT
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 ATENOLOL, ATENOLOL
 AZITHROMYCIN, AZITHROMYCIN
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BUMETANIDE, BUMETANIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CAFERGOT, CAFFEINE
 CARISOPRODOL AND ASPIRIN, ASPIRIN
 CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN
 CARVEDILOL, CARVEDILOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SANDOZ INC
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFDINIR, CEFDINIR
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CEFPROZIL, CEFPROZIL
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CLARITHROMYCIN, CLARITHROMYCIN
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 COSYNTROPIN, COSYNTROPIN
 CYCLOSPORINE, CYCLOSPORINE
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
 DESLORATADINE, DESLORATADINE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
 DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 EPLERENONE, EPLERENONE
 ETODOLAC, ETODOLAC
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FUROSEMIDE, FUROSEMIDE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GLIPIZIDE, GLIPIZIDE
 HALOPERIDOL, HALOPERIDOL
 HEPARIN SODIUM, HEPARIN SODIUM
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 INDOMETHACIN, INDOMETHACIN
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 ISONIAZID, ISONIAZID
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 ITRACONAZOLE, ITRACONAZOLE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVOFLOXACIN, LEVOFLOXACIN
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LORAZEPAM, LORAZEPAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LOVASTATIN, LOVASTATIN
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 METAXALONE, METAXALONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHAZOLAMIDE, METHAZOLAMIDE
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 METOLAZONE, METOLAZONE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NADOLOL, NADOLOL
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 NIZATIDINE, NIZATIDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SANDOZ INC
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OMEPRAZOLE, OMEPRAZOLE
 OMNITROPE, SOMATROPIN RECOMBINANT
 ONDANSETRON, ONDANSETRON
 ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE, ASPIRIN
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXALIPLATIN, OXALIPLATIN
 OXAPROZIN, OXAPROZIN
 OXAZEPAM, OXAZEPAM
 PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
 PENICILLIN G SODIUM, PENICILLIN G SODIUM
 PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
 PERPHENAZINE, PERPHENAZINE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE AND GLIMEPIRIDE, GLIMEPIRIDE
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUINIDINE SULFATE, QUINIDINE SULFATE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RIBAVIRIN, RIBAVIRIN
 RIFAMPIN, RIFAMPIN
 RISPERIDONE, RISPERIDONE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 SULFADIAZINE, SULFADIAZINE
 TACROLIMUS, TACROLIMUS
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TEMAZEPAM, TEMAZEPAM
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

SANDOZ INC

* SANDOZ INC
 ACETAMINOPHEN, ACETAMINOPHEN
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 ANECTINE, SUCCINYLCHOLINE CHLORIDE
 ANGIOMAX, BIVALIRUDIN
 ARISTOSPAN, TRIAMCINOLONE HEXACETONIDE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BETOPTIC, BETAXOLOL HYDROCHLORIDE
 BIMATOPROST, BIMATOPROST
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BUDESONIDE, BUDESONIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSULFAN, BUSULFAN
 CARBOPLATIN, CARBOPLATIN
 CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
 CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
 CEFIXIME, CEFIXIME
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
 CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
 CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 CROMOLYN SODIUM, CROMOLYN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SANDOZ INC**

DECITABINE, DECITABINE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIGOXIN, DIGOXIN
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
 EPHEDRINE SULFATE, EPHEDRINE SULFATE
 EZETIMIBE, EZETIMIBE
 FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
 FLUMAZENIL, FLUMAZENIL
 GATIFLOXACIN, GATIFLOXACIN
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GLATOPA, GLATIRAMER ACETATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 GRISEOFULVIN, ULTRAMICROSIZED, GRISEOFULVIN, ULTRAMICROSIZED
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE
 INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID
 INFUVITE PEDIATRIC, ASCORBIC ACID
 ISONIAZID, ISONIAZID
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN, AMOXICILLIN
 LATANOPROST, LATANOPROST
 LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 LINEZOLID, LINEZOLID
 MAXITROL, DEXAMETHASONE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METIPRANOLOL, METIPRANOLOL HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYDRIACYL, TROPICAMIDE
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 NEVIRAPINE, NEVIRAPINE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OFLOXACIN, OFLOXACIN
 OLANZAPINE, OLANZAPINE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PACLITAXEL, PACLITAXEL
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARICALCITOL, PARICALCITOL
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PROGESTERONE, PROGESTERONE
 QOLIANA, BRIMONIDINE TARTRATE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 REGONOL, PYRIDOSTIGMINE BROMIDE
 RIBAVIRIN, RIBAVIRIN
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILODOSIN, SILODOSIN
 SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TELMISARTAN, TELMISARTAN
 TERIFLUNOMIDE, TERIFLUNOMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** s ****

* SANDOZ INC
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TIGECYCLINE, TIGECYCLINE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 TOBREX, TOBRAMYCIN
 TREPROSTINIL, TREPROSTINIL
 TRIFLURIDINE, TRIFLURIDINE
 TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

SANOCHEMIA CORP USA

* SANOCHEMIA CORP USA
 SCANLUX-300, IOPAMIDOL
 SCANLUX-370, IOPAMIDOL

SANOFI

* SANOFI GENZYME
 HECTOROL, DOXERCALCIFEROL
 RENVELA, SEVELAMER CARBONATE

SANOFI AVENTIS US

* SANOFI AVENTIS US INC
 JEVTANA KIT, CABAZITAXEL

* SANOFI AVENTIS US LLC
 ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
 ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
 AMARYL, GLIMEPIRIDE
 AMBIEN CR, ZOLPIDEM TARTRATE
 AMBIEN, ZOLPIDEM TARTRATE
 APIDRA SOLOSTAR, INSULIN GLULISINE RECOMBINANT
 APIDRA, INSULIN GLULISINE RECOMBINANT
 ARAVA, LEFLUNOMIDE
 AUBAGIO, TERIFLUNOMIDE
 AVALIDE, HYDROCHLOROTHIAZIDE
 AVAPRO, IRBESARTAN
 CHILDREN'S ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 DIABETA, GLYBURIDE
 ELOXATIN, OXALIPLATIN
 FERRLECIT, SODIUM FERRIC GLUCONATE COMPLEX
 FLOMAX, TAMSULOSIN HYDROCHLORIDE
 GAVISCON, ALUMINUM HYDROXIDE (OTC)
 LANTUS SOLOSTAR, INSULIN GLARGINE RECOMBINANT
 LANTUS, INSULIN GLARGINE RECOMBINANT
 LOVENOX (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 LOVENOX, ENOXAPARIN SODIUM
 MULTAQ, DRONEDARONE HYDROCHLORIDE
 NASACORT ALLERGY 24 HOUR, TRIAMCINOLONE ACETONIDE (OTC)
 NICODERM CQ, NICOTINE (OTC)
 PLAVIX, CLOPIDOGREL BISULFATE
 PRIFTIN, RIFAPENTINE
 PRIMAQUINE, PRIMAQUINE PHOSPHATE
 RIFADIN, RIFAMPIN
 RIFAMATE, ISONIAZID
 RIFATER, ISONIAZID
 TAXOTERE, DOCETAXEL
 XYZAL ALLERGY 24HR, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
 XYZAL, LEVOCETIRIZINE DIHYDROCHLORIDE

SANOFI US

* SANOFI US
 ZANTAC 150, RANITIDINE HYDROCHLORIDE (OTC)
 ZANTAC 75, RANITIDINE HYDROCHLORIDE (OTC)

SANOFI US SERVICES

* SANOFI US SERVICES INC
 TOUJEO MAX SOLOSTAR, INSULIN GLARGINE RECOMBINANT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SANOFI US SERVICES INC
TOUJEO SOLOSTAR, INSULIN GLARGINE RECOMBINANT

SANOFI-AVENTIS US

* SANOFI-AVENTIS US LLC
ADLYXIN, LIXISENATIDE
ADMELOG SOLOSTAR, INSULIN LISPRO
ADMELOG, INSULIN LISPRO
SOLIQUA 100/33, INSULIN GLARGINE

SANTARUS INC

* SANTARUS INC
FENOGLIDE, FENOFIBRATE
GLUMETZA, METFORMIN HYDROCHLORIDE
ZEGERID, OMEPRAZOLE

SAOL THERAPS RES LTD

* SAOL THERAPEUTICS RESEARCH LTD
LIORESAL, BACLOFEN

SAREPTA THERAPS INC

* SAREPTA THERAPEUTICS INC
EXONDYS 51, ETEPLIRSEN

SAVIOR LIFETEC CORP

* SAVIOR LIFETEC CORP
MEROPENEM, MEROPENEM

SAWAI USA

* SAWAI USA INC
PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM

SB PHARMCO

* SB PHARMCO PUERTO RICO INC
AVANDIA, ROSIGLITAZONE MALEATE

SCHERING

* SCHERING CORP
INTEGRILIN, EPTIFIBATIDE
NOXAFIL, POSACONAZOLE
REBETOL, RIBAVIRIN

SCIECURE PHARMA INC

* SCIECURE PHARMA INC
BUDESONIDE, BUDESONIDE
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE

SCIEGEN PHARMS INC

* SCIEGEN PHARMACEUTICALS INC
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
ARIPIPRAZOLE, ARIPIPRAZOLE
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CARISOPRODOL, CARISOPRODOL
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
GABAPENTIN, GABAPENTIN
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
IRBESARTAN, IRBESARTAN
LAMOTRIGINE, LAMOTRIGINE
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
METAXALONE, METAXALONE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE

SCILEX PHARMS INC

* SCILEX PHARMACEUTICALS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SCILEX PHARMACEUTICALS

ZTLIDO, LIDOCAINE

SCINOPHARM TAIWAN

* SCINOPHARM TAIWAN LTD

FONDAPARINUX SODIUM, FONDAPARINUX SODIUM

SCIOS LLC

* SCIOS LLC

NATRECOR, NESIRITIDE RECOMBINANT

SEBELA IRELAND LTD

* SEBELA IRELAND LTD

BRISDELLE, PAROXETINE MESYLATE

IMURAN, AZATHIOPRINE

LOTRONEX, ALOSETRON HYDROCHLORIDE

MICORT-HC, HYDROCORTISONE ACETATE

MOTOFEN, ATROPINE SULFATE

NAFTIN, NAFTIFINE HYDROCHLORIDE

ONMEL, ITRACONAZOLE

PEXEVA, PAROXETINE MESYLATE

PRAMOSONE, HYDROCORTISONE ACETATE

RIDAUURA, AURANOFIN

SECAN PHARMS

* SECAN PHARMACEUTICALS INC

LEVETIRACETAM, LEVETIRACETAM

SENTYNL THERAPS INC

* SENTYNL THERAPEUTICS INC

ABSTRAL, FENTANYL CITRATE

LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE

SEPTODONT

* SEPTODONT INC

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE

SEPTODONT HOLDING

* SEPTODONT HOLDING SAS

ORAVVERSE, PHENTOLAMINE MESYLATE

SEPTODONT INC

* SEPTODONT INC

LIDOCAINE, LIDOCAINE

PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, EPINEPHRINE BITARTRATE

PRILOCAINE HYDROCHLORIDE, PRILOCAINE HYDROCHLORIDE

SERB SA

* SERB SA

CYANOKIT, HYDROXOCOBALAMIN

SETON PHARM

* SETON PHARMACEUTICAL LLC

PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE

SETON PHARMS

* SETON PHARMACEUTICALS LLC

PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE

SHANDONG XINHUA

* SHANDONG XINHUA PHARMACEUTICAL CO LTD

IBUPROFEN, IBUPROFEN

IBUPROFEN, IBUPROFEN (OTC)

SHANGHAI DESANO

* SHANGHAI DESANO BIO-PHARMACEUTICALS CO LTD

LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE

SHANGHAI HENGRUI

* SHANGHAI HENGRUI PHARMACEUTICAL CO LTD

DESFLURANE, DESFLURANE

SEVOFLURANE, SEVOFLURANE

SHENZHEN TECHDOW

* SHENZHEN TECHDOW PHARMACEUTICAL CO LTD

HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM

HEPARIN SODIUM, HEPARIN SODIUM

SHERTECH LABS LLC

* SHERTECH LABORATORIES LLC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SHERTECH LABORATORIES LLC
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

SHILPA MEDICARE

* SHILPA MEDICARE LTD
 AZACITIDINE, AZACITIDINE

SHILPA MEDICARE LTD

* SHILPA MEDICARE LTD
 CAPECITABINE, CAPECITABINE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE

SHIONOGI INC

* SHIONOGI INC
 MULPLETA, LUSUTROMBOPAG
 PONSTEL, MEFENAMIC ACID
 SYMPROIC, NALDEMEDINE TOSYLATE
 ULESFIA, BENZYL ALCOHOL

SHIRE

* SHIRE DEVELOPMENT INC
 ADDERALL XR 10, AMPHETAMINE ASPARTATE
 ADDERALL XR 15, AMPHETAMINE ASPARTATE
 ADDERALL XR 20, AMPHETAMINE ASPARTATE
 ADDERALL XR 25, AMPHETAMINE ASPARTATE
 ADDERALL XR 30, AMPHETAMINE ASPARTATE
 ADDERALL XR 5, AMPHETAMINE ASPARTATE
 CARBATROL, CARBAMAZEPINE
 INTUNIV, GUANFACINE HYDROCHLORIDE
 LIALDA, MESALAMINE
 PENTASA, MESALAMINE

SHIRE DEV LLC

* SHIRE DEVELOPMENT LLC
 FOSRENOL, LANTHANUM CARBONATE
 MOTEGRITY, PRUCALOPRIDE SUCCINATE
 MYDAYIS, AMPHETAMINE ASPARTATE
 VYVANSE, LISDEXAMFETAMINE DIMESYLATE
 XIIDRA, LIFITEGRAST

SHIRE DEVELOPMENT

* SHIRE DEVELOPMENT INC
 VYVANSE, LISDEXAMFETAMINE DIMESYLATE

SHIRE HUMAN GENETIC

* SHIRE HUMAN GENETIC THERAPIES INC
 VPRIV, VELAGLUCERASE ALFA

SHIRE LLC

* SHIRE DEVELOPMENT LLC
 AGRYLIN, ANAGRELIDE HYDROCHLORIDE
 FOSRENOL, LANTHANUM CARBONATE

SHIRE ORPHAN THERAP

* SHIRE ORPHAN THERAPIES INC
 FIRAZYR, ICATIBANT ACETATE

SIDMAK LABS INDIA

* SIDMAK LABORATORIES INDIA PVT LTD
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE

SIGA TECHNOLOGIES

* SIGA TECHNOLOGIES INC
 TPOXX, TECOVIRIMAT

SIGMAPHARM LABS LLC

* SIGMAPHARM LABORATORIES LLC
 ACITRETIN, ACITRETIN
 ADEFOVIR DIPIVOXIL, ADEFOVIR DIPIVOXIL
 AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
 ASENAPINE MALEATE, ASENAPINE MALEATE
 DISULFIRAM, DISULFIRAM
 DOFETILIDE, DOFETILIDE
 ERGOCALCIFEROL, ERGOCALCIFEROL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SIGMAPHARM LABORATORIES LLC
 FLUCYTOSINE, FLUCYTOSINE
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 GRISEOFULVIN,ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 NITROGLYCERIN, NITROGLYCERIN
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE

SILVERGATE PHARMS

* SILVERGATE PHARMACEUTICALS INC
 EPANED KIT, ENALAPRIL MALEATE
 EPANED, ENALAPRIL MALEATE
 QBRELIS, LISINOPRIL
 XATMEP, METHOTREXATE SODIUM

SINOTHERAPEUTICS INC

* SINOTHERAPEUTICS INC
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE

SKINMEDICA

* SKINMEDICA INC
 VANIQA, EFLORNITHINE HYDROCHLORIDE

SKYEPHARMA AG

* SKYEPHARMA AG
 TRIGLIDE, FENOFIBRATE

SLAYBACK PHARMA LLC

* SLAYBACK PHARMA LLC
 HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE

SMITHKLINE BEECHAM

* SMITHKLINE BEECHAM
 LOVAZA, OMEGA-3-ACID ETHYL ESTERS
 * SMITHKLINE BEECHAM (CORK) LTD IRELAND
 COREG CR, CARVEDILOL PHOSPHATE
 COREG, CARVEDILOL

SOAPCO

* SOAPCO INC
 BRIAN CARE, CHLORHEXIDINE GLUCONATE (OTC)

SOFGEN PHARMS

* SOFGEN PHARMACEUTICALS
 NIMODIPINE, NIMODIPINE
 * SOFGEN PHARMACEUTICALS LLC
 IBUPROFEN, IBUPROFEN (OTC)
 PROGESTERONE, PROGESTERONE

SOFIE

* SOFIE CO DBA SOFIE (FKA ZEVACOR PHARMA INC)
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

SOLARIS PHARMA CORP

* SOLARIS PHARMA CORP
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE

SOMERSET

* SOMERSET PHARMACEUTICALS INC
 EMSAM, SELEGILINE

SOMERSET THERAPS LLC

* SOMERSET THERAPEUTICS LLC
 CYANOCOBALAMIN, CYANOCOBALAMIN
 DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 METHOCARBAMOL, METHOCARBAMOL
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SOMERSET THERAPEUTICS LLC**

OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 TOBRAMYCIN, TOBRAMYCIN
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

SOVEREIGN PHARMS*** SOVEREIGN PHARMACEUTICALS LLC**

OBREDON, GUAIFENESIN

SPARC*** SUN PHARMA ADVANCED RESEARCH CO LTD**

ELEPSIA XR, LEVETIRACETAM

SPECGX LLC*** SPECGX LLC**

ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE
 ANEXSIA 5/325, ACETAMINOPHEN
 ANEXSIA 7.5/325, ACETAMINOPHEN
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 EXALGO, HYDROMORPHONE HYDROCHLORIDE
 FENTANYL CITRATE, FENTANYL CITRATE
 FENTANYL-100, FENTANYL
 FENTANYL-12, FENTANYL
 FENTANYL-25, FENTANYL
 FENTANYL-50, FENTANYL
 FENTANYL-75, FENTANYL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHADOSE, METHADONE HYDROCHLORIDE
 METHYLIN ER, METHYLPHENIDATE HYDROCHLORIDE
 METHYLIN, METHYLPHENIDATE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 OXYCET, ACETAMINOPHEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PAMELOR, NORTRIPTYLINE HYDROCHLORIDE
 RESTORIL, TEMAZEPAM
 ROXICODONE, OXYCODONE HYDROCHLORIDE
 TOFRANIL, IMIPRAMINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

SPECTRA MDCL DEVICES*** SPECTRA MEDICAL DEVICES INC**

LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE

SPECTRON MRC LLC*** SPECTRON MRC LLC**

AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

SPECTRUM PHARMS*** SPECTRUM PHARMACEUTICALS INC**

BELEODAQ, BELINOSTAT
 EVOMELA, MELPHALAN HYDROCHLORIDE
 FUSILEV, LEVOLEUCOVORIN CALCIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SPECTRUM PHARMACEUTICALS INC
KHAPZORY, LEVOLEUCOVORIN

SPIIL

* SUN PHARMA INDUSTRIES LTD
KAPSPARGO SPRINKLE, METOPROLOL SUCCINATE
OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)

SPROUT PHARMS

* SPROUT PHARMACEUTICALS INC
ADDYI, FLIBANSERIN

SQUARE PHARMS LTD

* SQUARE PHARMACEUTICALS LTD
VALSARTAN, VALSARTAN

ST RENATUS

* ST RENATUS LLC
KOVANAZE, OXYMETAZOLINE HYDROCHLORIDE

STAND HOMEOPATH

* STANDARD HOMEOPATHIC CO
IVY BLOCK, BENTOQUATAM (OTC)

STASON PHARMS

* STASON PHARMACEUTICALS INC
PURINETHOL, MERCAPTOPURINE

STERINOVA INC

* STERINOVA INC
HEPARIN SODIUM, HEPARIN SODIUM

STI PHARMA LLC

* STI PHARMA LLC
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
DEXAMETHASONE, DEXAMETHASONE
MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE
TRIAICIN-C, CODEINE PHOSPHATE

STIEFEL

* STIEFEL LABORATORIES INC
DUAC, BENZOYL PEROXIDE

STIEFEL LABS INC

* STIEFEL LABORATORIES INC
SORIATANE, ACITRETIN

STRIDES PHARMA

* STRIDES PHARMA GLOBAL PTE LTD
ABACAVIR SULFATE, ABACAVIR SULFATE
ACARBOSE, ACARBOSE
ACETAZOLAMIDE, ACETAZOLAMIDE
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
BENZONATATE, BENZONATATE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CALCITRIOL, CALCITRIOL
CARISOPRODOL, CARISOPRODOL
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
DUTASTERIDE, DUTASTERIDE
EFAVIRENZ, EFAVIRENZ
ERGOCALCIFEROL, ERGOCALCIFEROL
GABAPENTIN, GABAPENTIN
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
KETOCONAZOLE, KETOCONAZOLE
LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
LAMIVUDINE, LAMIVUDINE
LIDOCAINE, LIDOCAINE
MELOXICAM, MELOXICAM
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* STRIDES PHARMA GLOBAL PTE LTD
 METHOXSALEN, METHOXSALEN
 METRONIDAZOLE, METRONIDAZOLE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NEVIRAPINE, NEVIRAPINE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
 PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL
 PIROXICAM, PIROXICAM
 POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 TACROLIMUS, TACROLIMUS
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

STRIDES VIVIMED

* STRIDES VIVIMED PTE LTD
 ALBENDAZOLE, ALBENDAZOLE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 METRONIDAZOLE, METRONIDAZOLE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)

STRONGBRIDGE US

* STRONGBRIDGE US INC
 KEVEYIS, DICHLORPHENAMIDE

SUCAMPO PHARMA LLC

* SUCAMPO PHARMA AMERICAS LLC
 AMITIZA, LUBIPROSTONE

SUN PHARM INDS

* SUN PHARMACEUTICAL INDUSTRIES LTD
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 DESLORATADINE, DESLORATADINE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OLANZAPINE, OLANZAPINE
 ONDANSETRON, ONDANSETRON
 OXCARBAZEPINE, OXCARBAZEPINE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE

SUN PHARM INDS (IN)

* SUN PHARMACEUTICAL INDUSTRIES LTD
 CEPHALEXIN, CEPHALEXIN
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ZONISAMIDE, ZONISAMIDE

SUN PHARM INDS INC

* SUN PHARMACEUTICAL INDUSTRIES INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** s **

* SUN PHARMACEUTICAL INDUSTRIES INC
 ABSORICA, ISOTRETINOIN
 ALLOPURINOL, ALLOPURINOL
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ATENOLOL, ATENOLOL
 BACLOFEN, BACLOFEN
 BENZONATATE, BENZONATATE
 CARVEDILOL, CARVEDILOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLONAZEPAM, CLONAZEPAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CLOZAPINE, CLOZAPINE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DIGOXIN, DIGOXIN
 ERGOCALCIFEROL, ERGOCALCIFEROL
 EURAX, CROTAMITON
 FLUMADINE, RIMANTADINE HYDROCHLORIDE
 FLURBIPROFEN, FLURBIPROFEN
 GEMFIBROZIL, GEMFIBROZIL
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 GLIPIZIDE, GLIPIZIDE
 HALOG, HALCINONIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 INDOMETHACIN, INDOMETHACIN
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 KENALOG, TRIAMCINOLONE ACETONIDE
 LITHIUM CARBONATE, LITHIUM CARBONATE
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHIMAZOLE, METHIMAZOLE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 NIMODIPINE, NIMODIPINE
 OXAPROZIN, OXAPROZIN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 REPAGLINIDE, REPAGLINIDE
 RISPERIDONE, RISPERIDONE
 TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 ULTRAVATE, HALOBETASOL PROPIONATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

SUN PHARM INDS LTD

* SUN PHARMACEUTICAL INDUSTRIES LTD
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZITHROMYCIN, AZITHROMYCIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******* SUN PHARMACEUTICAL INDUSTRIES LTD**

BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CARVEDILOL, CARVEDILOL
 CERINTA, ETHINYL ESTRADIOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FAMOTIDINE, FAMOTIDINE (OTC)
 FELODIPINE, FELODIPINE
 FENOFIBRATE, FENOFIBRATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 GABAPENTIN, GABAPENTIN
 GANIRELIX ACETATE, GANIRELIX ACETATE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 INFUGEM, GEMCITABINE HYDROCHLORIDE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINOPRIL, LISINOPRIL
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
 LORATADINE REDIDOSE, LORATADINE (OTC)
 LORATADINE, LORATADINE (OTC)
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 OMEPRAZOLE, OMEPRAZOLE (OTC)
 ONDANSETRON, ONDANSETRON
 OPCICON ONE-STEP, LEVONORGESTREL (OTC)
 OXCARBAZEPINE, OXCARBAZEPINE
 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 RILUZOLE, RILUZOLE
 RIOMET, METFORMIN HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TOPIRAMATE, TOPIRAMATE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALPROIC ACID, VALPROIC ACID
 XIMINO, MINOCYCLINE HYDROCHLORIDE

SUN PHARM INDUSTRIES*** SUN PHARMACEUTICAL INDUSTRIES INC**

ACETAZOLAMIDE, ACETAZOLAMIDE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALLOPURINOL, ALLOPURINOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUN PHARMACEUTICAL INDUSTRIES INC
 ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 ATENOLOL, ATENOLOL
 BACTRIM DS, SULFAMETHOXAZOLE
 BACTRIM, SULFAMETHOXAZOLE
 CARISOPRODOL, CARISOPRODOL
 CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE
 CHLORTHALIDONE, CHLORTHALIDONE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ERGOLOID MESYLATES, ERGOLOID MESYLATES
 FELODIPINE, FELODIPINE
 IBANDRONATE SODIUM, IBANDRONATE SODIUM
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LOVASTATIN, LOVASTATIN
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MINOXIDIL, MINOXIDIL
 MORPHINE SULFATE, MORPHINE SULFATE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 NYSTATIN, NYSTATIN
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PINDOLOL, PINDOLOL
 PIROXICAM, PIROXICAM
 PREDNISON, PREDNISON
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 QUALAQUIN, QUININE SULFATE
 QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
 QUINIDINE SULFATE, QUINIDINE SULFATE
 SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SULINDAC, SULINDAC
 TEMAZEPAM, TEMAZEPAM
 THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
 ULTRAVATE, HALOBETASOL PROPIONATE

SUN PHARMA GLOBAL

* SUN PHARMA GLOBAL FZE
 ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 AMIFOSTINE, AMIFOSTINE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BROMSITE, BROMFENAC SODIUM
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CARBIDOPA AND LEVODOPA, CARBIDOPA
 CARBIDOPA, LEVODOPA AND ENTACAPONE, CARBIDOPA
 CEQUA, CYCLOSPORINE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DECITABINE, DECITABINE
 DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

- * SUN PHARMA GLOBAL FZE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DOFETILIDE, DOFETILIDE
 DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ENTACAPONE, ENTACAPONE
 ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
 ESZOPICLONE, ESZOPICLONE
 EZALLOR, ROSUVASTATIN CALCIUM
 FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
 FINASTERIDE, FINASTERIDE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 NIACIN, NIACIN
 ODOMZO, SONIDEGIB PHOSPHATE
 OXALIPLATIN, OXALIPLATIN
 PALIPERIDONE, PALIPERIDONE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SUMATRIPTAN AND NAPROXEN SODIUM, NAPROXEN SODIUM
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TETRABENAZINE, TETRABENAZINE
 TOPIRAMATE, TOPIRAMATE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 XELPROS, LATANOPROST
 YONSA, ABIRATERONE ACETATE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
- * SUN PHARMA GLOBAL INC
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 ALPRAZOLAM, ALPRAZOLAM
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 CARBOPLATIN, CARBOPLATIN
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
- SUNGEN PHARMA**
- * SUNGEN PHARMA LLC
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
- SUNNY PHARMTECH INC**
- * SUNNY PHARMTECH INC
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
- SUNOVION**
- * SUNOVION PHARMACEUTICALS INC
 BROVANA, ARFORMOTEROL TARTRATE
 XOPENEX HFA, LEVALBUTEROL TARTRATE
- SUNOVION PHARMS INC**
- * SUNOVION PHARMACEUTICALS INC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SUNOVION PHARMACEUTICALS INC
 APTIOM, ESLICARBAZEPINE ACETATE
 ARCAPTA NEOHALER, INDACATEROL MALEATE
 LATUDA, LURASIDONE HYDROCHLORIDE
 LUNESTA, ESZOPICLONE
 SEEBRI, GLYCOPYRROLATE
 UTIBRON, GLYCOPYRROLATE
 ZONEGRAN, ZONISAMIDE

SUNOVION RESP

* SUNOVION RESPIRATORY DEVELOPMENT INC
 LONHALA MAGNAIR KIT, GLYCOPYRROLATE

SUNSHINE LAKE

* SUNSHINE LAKE PHARMA CO LTD
 AZITHROMYCIN, AZITHROMYCIN
 CLARITHROMYCIN, CLARITHROMYCIN
 ENTACAPONE, ENTACAPONE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE

SUNSTAR AMERICAS

* SUNSTAR AMERICAS INC
 PAROEX, CHLORHEXIDINE GLUCONATE

SUPERNUS PHARMS

* SUPERNUS PHARMACEUTICALS INC
 OXTELLAR XR, OXCARBAZEPINE
 TROKENDI XR, TOPIRAMATE

SUVEN LIFE

* SUVEN LIFE SCIENCES LTD
 MALATHION, MALATHION

SVC PHARMA

* SVC PHARMA LP
 DRONABINOL, DRONABINOL

SWEDISH ORPHAN

* SWEDISH ORPHAN BIOVITRUM AB PUBL
 ORFADIN, NITISINONE

SYNERGY PHARMS

* SYNERGY PHARMACEUTICALS INC
 TRULANCE, PLECANATIDE

SYNTHON PHARMS

* SYNTHON PHARMACEUTICALS INC
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE

**** T ******ACME LABS**

* THE ACME LABORATORIES LTD
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

GEN HOSP

* THE GENERAL HOSPITAL CORP
 AMMONIA N 13, AMMONIA N-13

METHODIST HOSP RES

* THE METHODIST HOSP RESEARCH INSTITUTE
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

RITEDOSE CORP

* THE RITEDOSE CORP
 ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

TAIHO ONCOLOGY

* TAIHO ONCOLOGY INC
 LONSURF, TIPIRACIL HYDROCHLORIDE

TAKEDA PHARMS USA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TAKEDA PHARMACEUTICALS USA INC
 ACTOPLUS MET XR, METFORMIN HYDROCHLORIDE
 ACTOPLUS MET, METFORMIN HYDROCHLORIDE
 ACTOS, PIOGLITAZONE HYDROCHLORIDE
 COLCRYS, COLCHICINE
 DEXILANT, DEXLANSOPRAZOLE
 DUETACT, GLIMEPIRIDE
 KAZANO, ALOGLIPTIN BENZOATE
 NESINA, ALOGLIPTIN BENZOATE
 OSENI, ALOGLIPTIN BENZOATE
 PREVACID, LANSOPRAZOLE
 ROZEREM, RAMELTEON
 TRINTELLIX, VORTIOXETINE HYDROBROMIDE
 ULORIC, FEBUXOSTAT

TALON THERAP

* TALON THERAPEUTICS INC
 MARQIBO KIT, VINCRISTINE SULFATE

TAMARANG

* TAMARANG SA
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE

TARO

* TARO PHARMACEUTICAL INDUSTRIES LTD
 ACETAZOLAMIDE, ACETAZOLAMIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CARVEDILOL, CARVEDILOL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ETODOLAC, ETODOLAC
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 FLUCONAZOLE, FLUCONAZOLE
 FLUOROURACIL, FLUOROURACIL
 GABAPENTIN, GABAPENTIN
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 IMIQUIMOD, IMIQUIMOD
 KETOCONAZOLE, KETOCONAZOLE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LORATADINE, LORATADINE (OTC)
 MELOXICAM, MELOXICAM
 METRONIDAZOLE, METRONIDAZOLE
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 OXCARBAZEPINE, OXCARBAZEPINE
 PHENYTOIN, PHENYTOIN

* TARO PHARMACEUTICALS USA INC
 ACETIC ACID, ACETIC ACID, GLACIAL
 ACYCLOVIR, ACYCLOVIR
 ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
 ADAPALENE, ADAPALENE
 ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
 AMMONIUM LACTATE, AMMONIUM LACTATE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
 DAPSONE, DAPSONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TARO PHARMACEUTICALS USA INC
 DERMABET, BETAMETHASONE VALERATE
 DESONIDE, DESONIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 ECONAZOLE NITRATE, ECONAZOLE NITRATE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE, FLUOCINONIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 HYDROCORTISONE, HYDROCORTISONE
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 KETOZOLE, KETOCONAZOLE
 LIDOCAINE, LIDOCAINE
 LORATADINE, LORATADINE (OTC)
 MICONAZOLE 3, MICONAZOLE NITRATE (OTC)
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MUPIROCI, MUPIROCI
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 PHENYTOIN, PHENYTOIN
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE (OTC)
 TERCONAZOLE, TERCONAZOLE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)
 U-CORT, HYDROCORTISONE ACETATE

TARO PHARM

* TARO PHARMACEUTICAL INDUSTRIES LTD
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 DESLORATADINE, DESLORATADINE
 DESONIDE, DESONIDE
 FELBAMATE, FELBAMATE
 FLUOROURACIL, FLUOROURACIL
 GABAPENTIN, GABAPENTIN
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 METRONIDAZOLE, METRONIDAZOLE
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 TERIL, CARBAMAZEPINE
 WARFARIN SODIUM, WARFARIN SODIUM

TARO PHARM INDS

* TARO PHARMACEUTICAL INDUSTRIES LTD
 AMCINONIDE, AMCINONIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 ETODOLAC, ETODOLAC

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TARO PHARMACEUTICAL INDUSTRIES LTD
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 LAMOTRIGINE, LAMOTRIGINE

TARO PHARM INDS LTD

* TARO PHARMACEUTICAL INDUSTRIES LTD
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 DESLORATADINE, DESLORATADINE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
 HYDROCORTISONE, HYDROCORTISONE
 INFANTS' FEVERALL, ACETAMINOPHEN (OTC)
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 OVIDE, MALATHION
 TOPICORT, DESOXIMETASONE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

TARO PHARMS

* TARO PHARMACEUTICALS INC
 BUTENAFINE HYDROCHLORIDE, BUTENAFINE HYDROCHLORIDE (OTC)
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE
 OXICONAZOLE NITRATE, OXICONAZOLE NITRATE
 PLIAGLIS, LIDOCAINE
 TAZAROTENE, TAZAROTENE
 TOPICORT, DESOXIMETASONE

TASMAN PHARMA

* TASMAN PHARMA INC
 VERSACLOZ, CLOZAPINE

TCG FLUENT PHARMA

* TCG FLUENT PHARMA INVESTORS LP
 FLOLIPID, SIMVASTATIN

TEIKOKU PHARMA USA

* TEIKOKU PHARMA USA INC
 LIDODERM, LIDOCAINE

TELIGENT

* TELIGENT OU
 CEFOTAN, CEFOTETAN DISODIUM
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 FORTAZ, CEFTAZIDIME
 ZANTAC, RANTIDINE HYDROCHLORIDE
 ZINACEF, CEFUROXIME SODIUM

TELIGENT PHARMA INC

* TELIGENT PHARMA INC
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 CICLOPIROX, CICLOPIROX
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DESONIDE, DESONIDE
 DESOXIMETASONE, DESOXIMETASONE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 ECONAZOLE NITRATE, ECONAZOLE NITRATE
 ERYTHROMYCIN, ERYTHROMYCIN
 FLUOCINONIDE, FLUOCINONIDE
 FLURANDRENOLIDE, FLURANDRENOLIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 HYDROCORTISONE, HYDROCORTISONE
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE, LIDOCAINE
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ******TERSERA THERAPS LLC**

* TERSERA THERAPEUTICS LLC
 ERGOMAR, ERGOTAMINE TARTRATE
 PRIALT, ZICONOTIDE ACETATE
 QMIIZ ODT, MELOXICAM
 VARUBI, ROLAPITANT HYDROCHLORIDE
 ZOLADEX, GOSERELIN ACETATE

TESARO INC

* TESARO INC
 ZEJULA, NIRAPARIB TOSYLATE

TETRAPHASE PHARMS

* TETRAPHASE PHARMACEUTICALS INC
 XERAVAL, ERAVACYCLINE DIHYDROCHLORIDE

TEVA

* TEVA NEUROSCIENCE INC
 AZILECT, RASAGILINE MESYLATE

* TEVA PHARMACEUTICALS USA INC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACYCLOVIR, ACYCLOVIR
 ADIPEX-P, PHENTERMINE HYDROCHLORIDE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN PEDIATRIC, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 ATENOLOL, ATENOLOL
 AZITHROMYCIN, AZITHROMYCIN
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BICALUTAMIDE, BICALUTAMIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CALCITRIOL, CALCITRIOL
 CAPTOPRIL, CAPTOPRIL
 CARVEDILOL, CARVEDILOL
 CEFACLOR, CEFACLOR
 CEFPROZIL, CEFPROZIL
 CELECOXIB, CELECOXIB
 CEPHALEXIN, CEPHALEXIN
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CILOSTAZOL, CILOSTAZOL
 CIMETIDINE, CIMETIDINE
 CLARITHROMYCIN, CLARITHROMYCIN
 CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
 DIFLUNISAL, DIFLUNISAL
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 EPITOL, CARBAMAZEPINE
 ESZOPICLONE, ESZOPICLONE
 ETODOLAC, ETODOLAC
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA PHARMACEUTICALS USA INC
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FINASTERIDE, FINASTERIDE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLURBIPROFEN, FLURBIPROFEN
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 GALZIN, ZINC ACETATE
 GEMFIBROZIL, GEMFIBROZIL
 GLIMEPIRIDE, GLIMEPIRIDE
 GLYBURIDE (MICRONIZED), GLYBURIDE
 GLYBURIDE, GLYBURIDE
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 KETOCONAZOLE, KETOCONAZOLE
 KETOPROFEN, KETOPROFEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LAMOTRIGINE, LAMOTRIGINE
 LEVOFLOXACIN, LEVOFLOXACIN
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 LOVASTATIN, LOVASTATIN
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MIRTAZAPINE, MIRTAZAPINE
 MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 MUPIROCIN, MUPIROCIN
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NAPROXEN, NAPROXEN
 NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
 NEOMYCIN SULFATE, NEOMYCIN SULFATE
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 NYSTATIN, NYSTATIN
 OFLOXACIN, OFLOXACIN
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON, ONDANSETRON
 ORAP, PIMOZIDE
 OXAPROZIN, OXAPROZIN
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PENICILLIN-VK, PENICILLIN V POTASSIUM
 PIROXICAM, PIROXICAM
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PRELONE, PREDNISOLONE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RIBAVIRIN, RIBAVIRIN
 RISPERIDONE, RISPERIDONE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SUCRALFATE, SUCRALFATE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH, SULFAMETHOXAZOLE
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ******* TEVA PHARMACEUTICALS USA INC**

TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOLMETIN SODIUM, TOLMETIN SODIUM
 TOPIRAMATE, TOPIRAMATE
 TORSEMIDE, TORSEMIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

TEVA BRANDED PHARM*** TEVA BRANDED PHARMACEUTICAL PRODUCTS R AND D INC**

AUSTEDO, DEUTETRABENAZINE
 DIAMOX, ACETAZOLAMIDE
 LOSEASONIQUE, ETHINYL ESTRADIOL
 PROAIR HFA, ALBUTEROL SULFATE
 PROAIR RESPICLICK, ALBUTEROL SULFATE
 PROGLYCEM, DIAZOXIDE
 QNASL, BECLOMETHASONE DIPROPIONATE
 QUARTETTE, ETHINYL ESTRADIOL
 SEASONALE, ETHINYL ESTRADIOL
 SEASONIQUE, ETHINYL ESTRADIOL
 TENUATE DOSPAN, DIETHYLPROPION HYDROCHLORIDE
 TENUATE, DIETHYLPROPION HYDROCHLORIDE
 ZIAC, BISOPROLOL FUMARATE

TEVA PARENTERAL*** TEVA PARENTERAL MEDICINES INC**

LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

TEVA PHARM*** TEVA PHARMACEUTICAL INDUSTRIES LTD**

AIRDUO RESPICLICK, FLUTICASONE PROPIONATE
 ARMONAIR RESPICLICK, FLUTICASONE PROPIONATE

TEVA PHARMS*** TEVA PHARMACEUTICALS USA**

ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 ANASTROZOLE, ANASTROZOLE
 AZITHROMYCIN, AZITHROMYCIN
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 BUDESONIDE, BUDESONIDE
 CARBAMAZEPINE, CARBAMAZEPINE
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFDINIR, CEFDINIR
 CEFPROZIL, CEFPROZIL
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 FAMCICLOVIR, FAMCICLOVIR
 FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 GABAPENTIN, GABAPENTIN
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HYDROCORTISONE, HYDROCORTISONE
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 IRBESARTAN, IRBESARTAN
 LANSOPRAZOLE, LANSOPRAZOLE
 LEFLUNOMIDE, LEFLUNOMIDE
 LETROZOLE, LETROZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ******* TEVA PHARMACEUTICALS USA**

LEVETIRACETAM, LEVETIRACETAM
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
 LINEZOLID, LINEZOLID
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 MELOXICAM, MELOXICAM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 OXALIPLATIN, OXALIPLATIN
 PACLITAXEL, PACLITAXEL
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 QUININE SULFATE, QUININE SULFATE
 RAMIPRIL, RAMIPRIL
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH, SULFAMETHOXAZOLE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TRANDOLAPRIL, TRANDOLAPRIL
 URSODIOL, URSODIOL
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VANDAZOLE, METRONIDAZOLE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZALEPLON, ZALEPLON

TEVA PHARMS INTL

*** TEVA PHARMACEUTICALS INTERNATIONAL GMBH**
 AMRIX, CYCLOBENZAPRINE HYDROCHLORIDE
 SYNRIPO, OMACETAXINE MEPESUCCINATE

TEVA PHARMS USA

*** TEVA PHARMACEUTICALS USA**
 ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
 ACITRETIN, ACITRETIN
 ADENOSINE, ADENOSINE
 ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
 ALPROSTADIL, ALPROSTADIL
 AMIKACIN SULFATE, AMIKACIN SULFATE
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 ARGATROBAN IN 0.9% SODIUM CHLORIDE, ARGATROBAN
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BUDESONIDE, BUDESONIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 CARBOPLATIN, CARBOPLATIN
 CLARAVIS, ISOTRETINOIN
 CLOZAPINE, CLOZAPINE
 COPAXONE, GLATIRAMER ACETATE
 DACARBAZINE, DACARBAZINE
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TEVA PHARMACEUTICALS USA
 DOCETAXEL, DOCETAXEL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 EFAVIRENZ, EMTRICITABINE, AND TENOFOVIR DISOPROXIL FUMARATE, EFAVIRENZ
 ENALAPRILAT, ENALAPRILAT
 ENTECAVIR, ENTECAVIR
 EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM
 EPTIFIBATIDE, EPTIFIBATIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESTRADIOL, ESTRADIOL
 ETOPOSIDE, ETOPOSIDE
 EZETIMIBE, EZETIMIBE
 FLUOROURACIL, FLUOROURACIL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
 GABAPENTIN, GABAPENTIN
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 IDARUBICIN HYDROCHLORIDE PFS, IDARUBICIN HYDROCHLORIDE
 IFOSFAMIDE, IFOSFAMIDE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 LANSOPRAZOLE, LANSOPRAZOLE
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
 LINEZOLID, LINEZOLID
 LOGILIA, ULIPRISTAL ACETATE
 MESNA, MESNA
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
 OMEPRAZOLE, OMEPRAZOLE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
 PARICALCITOL, PARICALCITOL
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TADALAFIL, TADALAFIL
 TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TOBRAMYCIN, TOBRAMYCIN
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VIGABATRIN, VIGABATRIN
 VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TEVA PHARMACEUTICALS USA
VINORELBINE TARTRATE, VINORELBINE TARTRATE
ZANOSAR, STREPTOZOCIN
ZOLMITRIPTAN, ZOLMITRIPTAN
- * TEVA PHARMACEUTICALS USA INC
ABIRATERONE ACETATE, ABIRATERONE ACETATE
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
CAPECITABINE, CAPECITABINE
CASSIPA, BUPRENORPHINE HYDROCHLORIDE
DAPTOMYCIN, DAPTOMYCIN
DARUNAVIR ETHANOLATE, DARUNAVIR ETHANOLATE
ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
EPINEPHRINE (AUTOINJECTOR), EPINEPHRINE
ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
ESTRADIOL, ESTRADIOL
LIDOCAINE, LIDOCAINE
LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
METRONIDAZOLE, METRONIDAZOLE
OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
SELFEMRA, FLUOXETINE HYDROCHLORIDE
TERIFLUNOMIDE, TERIFLUNOMIDE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
- TEVA PHARMS USA INC**
- * TEVA PHARMACEUTICALS USA INC
POTASSIUM CITRATE, POTASSIUM CITRATE
- THE FEINSTEIN INST**
- * THE FEINSTEIN INSTITUTE FOR MEDICAL RESEARCH
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18
- THEPHARMANETWORK LLC**
- * THEPHARMANETWORK LLC
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
BENZONATATE, BENZONATATE
ISONIAZID, ISONIAZID
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
NIMODIPINE, NIMODIPINE
THERMAZENE, SILVER SULFADIAZINE
- THERAPEUTICSMD INC**
- * THERAPEUTICSMD INC
ANNOVERA, ETHINYL ESTRADIOL
BIJUVA, ESTRADIOL
IMVEXXY, ESTRADIOL
- THERATECHNOLOGIES**
- * THERATECHNOLOGIES INC
EGRIFTA, TESAMORELIN ACETATE
- TIME-CAP LABS INC**
- * TIME-CAP LABORATORIES INC
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
- TITAN PHARMS**
- * TITAN PHARMACEUTICALS INC
PROBUPHINE, BUPRENORPHINE HYDROCHLORIDE
- TOLMAR**
- * TOLMAR INC
ACYCLOVIR, ACYCLOVIR
ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE
ADAPALENE, ADAPALENE
ATRIDOX, DOXYCYCLINE HYCLATE
AZELAIC ACID, AZELAIC ACID
CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CALCIPOTRIENE, CALCIPOTRIENE
CICLOPIROX, CICLOPIROX

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TOLMAR INC
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 KETOCONAZOLE, KETOCONAZOLE
 LEVETIRACETAM, LEVETIRACETAM
 LIDOCAINE AND PRILOCAINE, LIDOCAINE
 METRONIDAZOLE, METRONIDAZOLE
 NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE

TOLMAR THERAP

* TOLMAR THERAPEUTICS INC
 ELIGARD, LEUPROLIDE ACETATE

TORPHARM

* TORPHARM INC
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE

TORRENT PHARMA INC

* TORRENT PHARMA INC
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE

TORRENT PHARMS

* TORRENT PHARMACEUTICALS LIMITED
 LEVOFLOXACIN, LEVOFLOXACIN

* TORRENT PHARMACEUTICALS LTD
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 CARBAMAZEPINE, CARBAMAZEPINE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

* TORRENT PHARMACEUTICALS LTD.
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE

TORRENT PHARMS LLC

* TORRENT PHARMACEUTICALS LLC
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 OLANZAPINE, OLANZAPINE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

TORRENT PHARMS LTD

* TORRENT PHARMACEUTICALS LTD
 ACYCLOVIR, ACYCLOVIR
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CELECOXIB, CELECOXIB
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
 DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
 ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
 ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
 FELODIPINE, FELODIPINE
 FENOFIBRATE (MICRONIZED), FENOFIBRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ******* TORRENT PHARMACEUTICALS LTD**

FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 ITRACONAZOLE, ITRACONAZOLE
 LAMOTRIGINE, LAMOTRIGINE
 LEVETIRACETAM, LEVETIRACETAM
 LURASIDONE HYDROCHLORIDE, LURASIDONE HYDROCHLORIDE
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MOMETASONE FUROATE, MOMETASONE FUROATE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 OLANZAPINE, OLANZAPINE
 OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
 OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
 PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
 ROFLUMILAST, ROFLUMILAST
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 VALSARTAN, VALSARTAN

TRIS PHARMA INC*** TRIS PHARMA INC**

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)
 DYANAVAL XR, AMPHETAMINE
 GABAPENTIN, GABAPENTIN
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX, CHLORPHENIRAMINE POLISTIREX
 IBUPROFEN, IBUPROFEN (OTC)
 KARBINAL ER, CARBINOXAMINE MALEATE
 LEVETIRACETAM, LEVETIRACETAM
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 THEOPHYLLINE, THEOPHYLLINE

TRUSTEES UNIV PA*** TRUSTEES OF THE UNIV OF PENNSYLVANIA**

FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

TULEX PHARMS INC*** TULEX PHARMACEUTICALS INC**

OXYBUTYNYN CHLORIDE, OXYBUTYNYN CHLORIDE

TWI PHARMS*** TWI PHARMACEUTICALS INC**

BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

* TWI PHARMACEUTICALS INC
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NIFEDIPINE, NIFEDIPINE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 ZOLMITRIPTAN, ZOLMITRIPTAN

**** U ******UCB INC**

* UCB INC
 BRIVIACT, BRIVARACETAM
 KEPPRA XR, LEVETIRACETAM
 KEPPRA, LEVETIRACETAM
 NEUPRO, ROTIGOTINE
 VIMPAT, LACOSAMIDE

UCLA BIOMEDICAL

* UCLA BIOMEDICAL CYCLOTRON
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UCSF RADIOPHARM

* UCSF RADIOPHARMACEUTICAL FACILITY
 AMMONIA N 13, AMMONIA N-13
 CHOLINE C-11, CHOLINE C-11
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

UIHC PET IMAGING

* UNIV IOWA HOSPS AND CLINICS PET IMAGING CENTER
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UMEDICA LABS PVT LTD

* UMEDICA LABORATORIES PRIVATE LTD
 CHLORTHALIDONE, CHLORTHALIDONE

UNICHEM

* UNICHEM LABORATORIES LTD
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 MELOXICAM, MELOXICAM
 ZALEPLON, ZALEPLON

UNICHEM LABS LTD

* UNICHEM LABORATORIES LIMITED
 DIVALPROEX SODIUM, DIVALPROEX SODIUM

* UNICHEM LABORATORIES LTD
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 LAMOTRIGINE, LAMOTRIGINE
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 METRONIDAZOLE, METRONIDAZOLE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 PIROXICAM, PIROXICAM
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 VALSARTAN, VALSARTAN

UNICHEM PHARMS (USA)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ****

* UNICHEM PHARMACEUTICALS (USA) INC
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE

UNIMARK REMEDIES LTD

* UNIMARK REMEDIES LTD
MONTELUKAST SODIUM, MONTELUKAST SODIUM

UNIQUE PHARM LABS

* UNIQUE PHARMACEUTICAL LABORATORIES A DIVISION OF J.B. CHEMICALS AND PHARMACEUTICALS LTD
ATENOLOL, ATENOLOL
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
FLUCONAZOLE, FLUCONAZOLE
GLIPIZIDE, GLIPIZIDE
LITHIUM CARBONATE, LITHIUM CARBONATE
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
TINIDAZOLE, TINIDAZOLE

UNITED BIOMEDCL

* UNITED BIOMEDICAL INC
TERBUTALINE SULFATE, TERBUTALINE SULFATE

UNITED GUARDIAN

* UNITED GUARDIAN INC
RENACIDIN, CITRIC ACID

UNITED THERAP

* UNITED THERAPEUTICS CORP
ORENITRAM, TREPROSTINIL DIOLAMINE
REMODULIN, TREPROSTINIL
TYVASO, TREPROSTINIL

UNIV MICHIGAN

* UNIV MICHIGAN PET RADIOPHARMACEUTICAL PRODUCTION PROGRAM
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UNIV TX MD ANDERSON

* UNIV TEXAS MD ANDERSON CANCER CENTER
CHOLINE C-11, CHOLINE C-11
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

UNIV UTAH CYCLOTRON

* UNIV UTAH CYCLOTRON RADIOCHEMISTRY LAB
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

UPSHER SMITH LABS

* UPSHER SMITH LABORATORIES LLC
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
BEXAROTENE, BEXAROTENE
BUMETANIDE, BUMETANIDE
CLOBAZAM, CLOBAZAM
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
EXEMESTANE, EXEMESTANE
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
Klor-CON M10, POTASSIUM CHLORIDE
Klor-CON M15, POTASSIUM CHLORIDE
Klor-CON M20, POTASSIUM CHLORIDE
Klor-CON, POTASSIUM CHLORIDE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
MORPHINE SULFATE, MORPHINE SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ****

* UPSHER SMITH LABORATORIES LLC
 NYSTATIN, NYSTATIN
 ORVATEN, MIDODRINE HYDROCHLORIDE
 OXANDROLONE, OXANDROLONE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PACERONE, AMIODARONE HYDROCHLORIDE
 PENTOXIL, PENTOXIFYLLINE
 PREVALITE, CHOLESTYRAMINE
 QUDEXY XR, TOPIRAMATE
 SORINE, SOTALOL HYDROCHLORIDE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 VOGELXO, TESTOSTERONE

US PHARM HOLDINGS

* US PHARMACEUTICAL HOLDINGS II LLC
 DEMEROL, MEPERIDINE HYDROCHLORIDE
 DRISDOL, ERGOCALCIFEROL
 HIPREX, METHENAMINE HIPPURATE
 LASIX, FUROSEMIDE
 NORPRAMIN, DESIPRAMINE HYDROCHLORIDE

US PHARMS HOLDINGS I

* US PHARMACEUTICALS HOLDINGS I LLC
 LOPRESSOR HCT, HYDROCHLOROTHIAZIDE
 LOPRESSOR, METOPROLOL TARTRATE
 LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
 LOTENSIN, BENAZEPRIL HYDROCHLORIDE
 PARLODEL, BROMOCRIPTINE MESYLATE

US WORLDMEDS

* US WORLDMEDS LLC
 APOKYN, APOMORPHINE HYDROCHLORIDE
 REVONTO, DANTROLENE SODIUM

US WORLDMEDS LLC

* US WORLDMEDS LLC
 CORGARD, NADOLOL
 LUCEMYRA, LOFEXIDINE HYDROCHLORIDE
 XADAGO, SAFINAMIDE MESYLATE

USL PHARMA

* USL PHARMA LLC
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 BACLOFEN, BACLOFEN
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 FLUOXYMESTERONE, FLUOXYMESTERONE
 JANTOVEN, WARFARIN SODIUM

USPHARMA

* USPHARMA LTD
 NITRO-DUR, NITROGLYCERIN

USPHARMA WINDLAS

* USPHARMA WINDLAS LLC
 AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
 PRASUGREL, PRASUGREL HYDROCHLORIDE

USV NORTH AMERICA

* USV NORTH AMERICA INC
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

**** V ******VALEANT**

* VALEANT PHARMACEUTICALS INTERNATIONAL
 ANCOBON, FLUCYTOSINE
 BONTRIL PDM, PHENDIMETRAZINE TARTRATE
 D.H.E. 45, DIHYDROERGOTAMINE MESYLATE
 MIGRANAL, DIHYDROERGOTAMINE MESYLATE
 MYSOLINE, PRIMIDONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** V **

VALEANT BERMUDA

- * VALEANT INTERNATIONAL BERMUDA
 - BENZACLIN, BENZOYL PEROXIDE
 - DERMATOP E EMOLLIENT, PREDNICARBATE
 - ELIDEL, PIMECROLIMUS
 - PENLAC, CICLOPIROX
 - RETIN-A, TRETINOIN
 - XERESE, ACYCLOVIR
 - ZOVIRAX, ACYCLOVIR

VALEANT INTL

- * VALEANT INTERNATIONAL BARBADOS SRL
 - ATIVAN, LORAZEPAM
 - CARDIZEM CD, DILTIAZEM HYDROCHLORIDE
 - CARDIZEM LA, DILTIAZEM HYDROCHLORIDE
 - CARDIZEM, DILTIAZEM HYDROCHLORIDE
 - RETIN-A MICRO, TRETINOIN
 - RETIN-A, TRETINOIN
 - RETIN-A-MICRO, TRETINOIN
 - VASERETIC, ENALAPRIL MALEATE
 - WELLBUTRIN XL, BUPROPION HYDROCHLORIDE
- * VALEANT INTERNATIONAL SRL
 - BENZAMYCIN, BENZOYL PEROXIDE

VALEANT LUXEMBOURG

- * VALEANT PHARMACEUTICALS LUXEMBOURG SARL
 - ERTACZO, SERTACONAZOLE NITRATE
 - TARGRETIN, BEXAROTENE
 - VISUDYNE, VERTEPORFIN

VALEANT PHARM INTL

- * VALEANT PHARMACEUTICALS INTERNATIONAL
 - ANDROID 25, METHYLTESTOSTERONE
 - EFUDEX, FLUOROURACIL
 - LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE
 - MESTINON, PYRIDOSTIGMINE BROMIDE
 - TESTRED, METHYLTESTOSTERONE
 - VIRAZOLE, RIBAVIRIN
 - ZELAPAR, SELEGILINE HYDROCHLORIDE

VALEANT PHARMS

- * VALEANT PHARMACEUTICALS NORTH AMERICA
 - MEPHYTON, PHYTONADIONE
- * VALEANT PHARMACEUTICALS NORTH AMERICA LLC
 - LIBRAX, CHLORDIAZEPOXIDE HYDROCHLORIDE
 - MESTINON, PYRIDOSTIGMINE BROMIDE
 - MINITRAN, NITROGLYCERIN
 - PENTOXIFYLLINE, PENTOXIFYLLINE

VALEANT PHARMS INC

- * VALEANT PHARMACEUTICALS INTERNATIONAL INC
 - GRIS-PEG, GRISEOFULVIN, ULTRAMICROSIZED

VALEANT PHARMS INTL

- * VALEANT PHARMACEUTICALS INTERNATIONAL
 - APRISO, MESALAMINE
 - COLAZAL, BALSALAZIDE DISODIUM
 - GIAZO, BALSALAZIDE DISODIUM
 - JUBLIA, EFINACONAZOLE
 - UCERIS, BUDESONIDE

VALEANT PHARMS LLC

- * VALEANT PHARMACEUTICALS NORTH AMERICA LLC
 - MACUGEN, PEGAPTANIB SODIUM
 - MESTINON, PYRIDOSTIGMINE BROMIDE
 - TASMAR, TOLCAPONE
 - TIMOPTIC-XE, TIMOLOL MALEATE

VALEANT PHARMS NORTH

- * VALEANT PHARMACEUTICALS NORTH AMERICA LLC
 - APLENZIN, BUPROPION HYDROBROMIDE
 - CARAC, FLUOROURACIL
 - DERMATOP, PREDNICARBATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** v ****

* VALEANT PHARMACEUTICALS NORTH AMERICA LLC
 DIASTAT ACUDIAL, DIAZEPAM
 DIASTAT, DIAZEPAM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 FENOFIBRATE, FENOFIBRATE
 ISORDIL, ISOSORBIDE DINITRATE
 ISUPREL, ISOPROTERENOL HYDROCHLORIDE
 KLARON, SULFACETAMIDE SODIUM
 MINITRAN, NITROGLYCERIN
 NIFEDIPINE, NIFEDIPINE
 NORITATE, METRONIDAZOLE
 PEPCID, FAMOTIDINE
 RENOVA, TRETINOIN
 RETIN-A, TRETINOIN
 SECONAL SODIUM, SECOBARBITAL SODIUM
 TIAZAC, DILTIAZEM HYDROCHLORIDE
 VASOTEC, ENALAPRIL MALEATE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 XENAZINE, TETRABENAZINE

VALIDUS PHARMS

* VALIDUS PHARMACEUTICALS LLC
 BUMEX, BUMETANIDE
 EQUETRO, CARBAMAZEPINE
 ROCALTROL, CALCITRIOL

VALIDUS PHARMS INC

* VALIDUS PHARMACEUTICALS INC
 MARPLAN, ISOCARBOXAZID

VANDA PHARMS INC

* VANDA PHARMACEUTICALS INC
 FANAPT, ILOPERIDONE
 HETLIOZ, TASIMELTEON

VELOXIS PHARMS INC

* VELOXIS PHARMACEUTICALS INC
 ENVARUS XR, TACROLIMUS

VERASTEM INC

* VERASTEM INC
 COPIKTRA, DUVELISIB

VEROSCIENCE

* VEROSCIENCE LLC
 CYCLOSET, BROMOCRIPTINE MESYLATE

VERTEX PHARMS

* VERTEX PHARMACEUTICALS INC
 KALYDECO, IVACAFTOR

VERTEX PHARMS INC

* VERTEX PHARMACEUTICALS INC
 KALYDECO, IVACAFTOR
 ORKAMBI, IVACAFTOR
 SYMDEKO (COPACKAGED), IVACAFTOR

VERTICAL PHARMS LLC

* VERTICAL PHARMACEUTICALS LLC
 DIVIGEL, ESTRADIOL

VIB

* VALEANT INTERNATIONAL BERMUDA
 ZOVIRAX, ACYCLOVIR

VICURON

* VICURON PHARMACEUTICALS INC
 ERAXIS, ANIDULAFUNGIN

VIFOR FRESENIUS

* VIFOR FRESENIUS MEDICAL CARE RENAL PHARMA FRANCE
 VELPHORO, SUCROFERRIC OXYHYDROXIDE

VIIV HLTHCARE

* VIIV HEALTHCARE CO
 COMBIVIR, LAMIVUDINE
 EPIVIR, LAMIVUDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VIIV HEALTHCARE CO
 EPZICOM, ABACAVIR SULFATE
 JULUCA, DOLUTEGRAVIR SODIUM
 LEXIVA, FOSAMPRENAVIR CALCIUM
 RESCRIPTOR, DELAVIRDINE MESYLATE
 RETROVIR, ZIDOVUDINE
 SELZENTRY, MARAVIROC
 TIVICAY, DOLUTEGRAVIR SODIUM
 TRIUMEQ, ABACAVIR SULFATE
 TRIZIVIR, ABACAVIR SULFATE
 ZIAGEN, ABACAVIR SULFATE

VINTAGE

* VINTAGE PHARMACEUTICALS LLC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 FOLIC ACID, FOLIC ACID
 HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
 HYDROCORTISONE, HYDROCORTISONE
 NYSTATIN, NYSTATIN
 PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
 PROMETHAZINE DM, DEXTROMETHORPHAN HYDROBROMIDE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROMETHAZINE WITH CODEINE, CODEINE PHOSPHATE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

VINTAGE PHARMS

* VINTAGE PHARMACEUTICALS
 ALPRAZOLAM, ALPRAZOLAM
 BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
 CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CYCLAFEM 0.5/35, ETHINYL ESTRADIOL
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 GILDAGIA, ETHINYL ESTRADIOL
 GILDESS 24 FE, ETHINYL ESTRADIOL
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 KIMIDESS, DESOGESTREL
 METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

* VINTAGE PHARMACEUTICALS INC
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ALLOPURINOL, ALLOPURINOL
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 BACLOFEN, BACLOFEN
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
 CARISOPRODOL, CARISOPRODOL
 DIAZEPAM, DIAZEPAM
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 LEVETIRACETAM, LEVETIRACETAM
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 OXYBUTYNYN CHLORIDE, OXYBUTYNYN CHLORIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PERPHENAZINE, PERPHENAZINE
 PREDNISON, PREDNISON

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VINTAGE PHARMACEUTICALS INC
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 SULFASALAZINE, SULFASALAZINE
 TORSEMIDE, TORSEMIDE

VINTAGE PHARMS LLC

* VINTAGE PHARMACEUTICALS LLC
 CYCLAFEM 1/35, ETHINYL ESTRADIOL
 CYCLAFEM 7/7/7, ETHINYL ESTRADIOL
 DUTASTERIDE, DUTASTERIDE
 EMOQUETTE, DESOGESTREL
 FELODIPINE, FELODIPINE
 GILDESS 1.5/30, ETHINYL ESTRADIOL
 GILDESS 1/20, ETHINYL ESTRADIOL
 GILDESS FE 1.5/30, ETHINYL ESTRADIOL
 GILDESS FE 1/20, ETHINYL ESTRADIOL
 LETROZOLE, LETROZOLE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MORPHINE SULFATE, MORPHINE SULFATE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYZILRA, ETHINYL ESTRADIOL
 ORSYTHIA, ETHINYL ESTRADIOL
 PERCOCET, ACETAMINOPHEN
 PREVIFEM, ETHINYL ESTRADIOL
 TRI-PREVIFEM, ETHINYL ESTRADIOL

VIRTUS PHARM

* VIRTUS PHARMACEUTICAL INC
 ACARBOSE, ACARBOSE
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE

VIRTUS PHARMS

* VIRTUS PHARMACEUTICALS LLC
 DAPSONE, DAPSONE
 LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
 PROMETRIUM, PROGESTERONE
 TRANEXAMIC ACID, TRANEXAMIC ACID

VISTA PHARMS

* VISTA PHARMACEUTICALS INC
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

VISTAPHARM

* VISTAPHARM INC
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 LACTULOSE, LACTULOSE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MORPHINE SULFATE, MORPHINE SULFATE
 NYSTATIN, NYSTATIN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PHENYTOIN, PHENYTOIN
 PREDNISOLONE, PREDNISOLONE
 VALPROIC ACID, VALPROIC ACID

VITRUVIAS THERAP

* VITRUVIAS THERAPEUTICS
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 * VITRUVIAS THERAPEUTICS LLC
 CYANOCOBALAMIN, CYANOCOBALAMIN
 LIDOCAINE, LIDOCAINE

VIVA HLTHCARE

* VIVA HEALTHCARE FZ LLC
 GLIMEPIRIDE, GLIMEPIRIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 SIMVASTATIN, SIMVASTATIN
 TRANEXAMIC ACID, TRANEXAMIC ACID

VIVIMED GLOBAL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VIVIMED GLOBAL GENERICS PTE LTD
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

VIVUS

* VIVUS INC
QSYMIA, PHENTERMINE HYDROCHLORIDE

VIVUS INC

* VIVUS INC
PANCREAZE, PANCRELIPASE (AMYLASE)

VPNA

* VALEANT PHARMACEUTICALS NORTH AMERICA
DICLOFENAC SODIUM, DICLOFENAC SODIUM

VYERA PHARMS LLC

* VYERA PHARMACEUTICALS LLC
DARAPRIM, PYRIMETHAMINE

**** W ******WA UNIV SCH MED**

* WASHINGTON UNIV SCHOOL MEDICINE
AMMONIA N 13, AMMONIA N-13
CHOLINE C-11, CHOLINE C-11

WATSON LABS

* WATSON LABORATORIES
FOLIC ACID, FOLIC ACID
PROPAPENONE HYDROCHLORIDE, PROPAPENONE HYDROCHLORIDE

* WATSON LABORATORIES INC
ACARBOSE, ACARBOSE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ALLOPURINOL, ALLOPURINOL
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
AMOXAPINE, AMOXAPINE
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
CAPTOPRIL, CAPTOPRIL
CARISOPRODOL, CARISOPRODOL
CHLORZOXAZONE, CHLORZOXAZONE
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CLONAZEPAM, CLONAZEPAM
COL-PROBENECID, COLCHICINE
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
ESTAZOLAM, ESTAZOLAM
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
GLIPIZIDE, GLIPIZIDE
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LAMOTRIGINE, LAMOTRIGINE
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
LISINAPRIL, LISINAPRIL
LORAZEPAM, LORAZEPAM
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
MEPROBAMATE, MEPROBAMATE
METHOCARBAMOL, METHOCARBAMOL
METHYLDOPA, METHYLDOPA
METHYLPREDNISOLONE, METHYLPREDNISOLONE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
METRONIDAZOLE, METRONIDAZOLE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MINOXIDIL, MINOXIDIL
MIRTAZAPINE, MIRTAZAPINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WATSON LABORATORIES INC
 NABUMETONE, NABUMETONE
 NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NATEGLINIDE, NATEGLINIDE
 NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 NIZATIDINE, NIZATIDINE
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 OGESTREL 0.5/50-28, ETHINYL ESTRADIOL
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PREDNISOLONE, PREDNISOLONE
 PREDNISON, PREDNISON
 PRIMIDONE, PRIMIDONE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 QUASENSE, ETHINYL ESTRADIOL
 QUINIDINE SULFATE, QUINIDINE SULFATE
 RAMIPRIL, RAMIPRIL
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 SULFASALAZINE, SULFASALAZINE
 SULINDAC, SULINDAC
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TELMISARTAN, TELMISARTAN
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 TRIMETHOPRIM, TRIMETHOPRIM
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 ZOVIA 1/50E-28, ETHINYL ESTRADIOL

* WATSON LABS INC
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

WATSON LABS INC

* WATSON LABORATORIES INC
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 AMMONIUM LACTATE, AMMONIUM LACTATE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CELECOXIB, CELECOXIB
 CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONE
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE
 EZETIMIBE, EZETIMIBE
 METRONIDAZOLE, METRONIDAZOLE
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 MODAFINIL, MODAFINIL
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 PERPHENAZINE, PERPHENAZINE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 PROPOFOL, PROPOFOL
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
 RASAGILINE MESYLATE, RASAGILINE MESYLATE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
 SILDENAFIL CITRATE, SILDENAFIL CITRATE
 TICAGRELOR, TICAGRELOR
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ******WATSON LABS TEVA**

- * WATSON LABORATORIES INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
 - BICALUTAMIDE, BICALUTAMIDE
 - BUPRENORPHINE, BUPRENORPHINE
 - CINACALCET HYDROCHLORIDE, CINACALCET HYDROCHLORIDE
 - EZETIMIBE AND ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 - GLIPIZIDE, GLIPIZIDE
 - IBANDRONATE SODIUM, IBANDRONATE SODIUM
 - ISRADIPINE, ISRADIPINE
 - LEVOFLOXACIN, LEVOFLOXACIN
 - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 - NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 - NORETHINDRONE AND ETHINYL ESTRADIOL (10/11), ETHINYL ESTRADIOL
 - NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 - PROBENECID, PROBENECID
 - SIMVASTATIN, SIMVASTATIN
 - TEMOZOLOMIDE, TEMOZOLOMIDE
 - TERIFLUNOMIDE, TERIFLUNOMIDE
 - TRIENTINE HYDROCHLORIDE, TRIENTINE HYDROCHLORIDE

WATSON PHARMS INC

- * WATSON PHARMACEUTICALS INC
 - TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 - TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE

WATSON PHARMS TEVA

- * WATSON PHARMACEUTICALS INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
 - RIFAMPIN, RIFAMPIN

WELLSTAT THERAP

- * WELLSTAT THERAPEUTICS CORP
 - VISTOGARD, URIDINE TRIACETATE
 - XURIDEN, URIDINE TRIACETATE

WES PHARMA INC

- * WES PHARMA INC
 - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 - OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

WEST WARD

- * WEST WARD PHARMACEUTICAL CORP
 - DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

WEST WARD PHARM CORP

- * WEST WARD PHARMACEUTICAL CORP
 - PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 - ROCURONIUM BROMIDE, ROCURONIUM BROMIDE

WEST-WARD PHARM CORP

- * WEST-WARD PHARMACEUTICAL CORP
 - CEFOTETAN, CEFOTETAN DISODIUM

WEST-WARD PHARMS INT

- * WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
 - ACARBOSE, ACARBOSE
 - ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 - ADENOSINE, ADENOSINE
 - ALENDRONATE SODIUM, ALENDRONATE SODIUM
 - ALLOPURINOL SODIUM, ALLOPURINOL SODIUM
 - ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
 - ALPRAZOLAM, ALPRAZOLAM
 - ALPROSTADIL, ALPROSTADIL
 - AMIKACIN SULFATE, AMIKACIN SULFATE
 - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 - AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 - AMRINONE LACTATE, INAMRINONE LACTATE
 - ANASTROZOLE, ANASTROZOLE
 - ATIVAN, LORAZEPAM
 - ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 - ATRACURIUM BESYLATE, ATRACURIUM BESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
 AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BUMETANIDE, BUMETANIDE
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CAFECIT, CAFFEINE CITRATE
 CALCITRIOL, CALCITRIOL
 CALCIUM ACETATE, CALCIUM ACETATE
 CAPECITABINE, CAPECITABINE
 CARBOPLATIN, CARBOPLATIN
 CEFOXITIN, CEFOXITIN SODIUM
 CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
 CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 CILOSTAZOL, CILOSTAZOL
 CISPLATIN, CISPLATIN
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLADRIBINE, CLADRIBINE
 CLARITHROMYCIN, CLARITHROMYCIN
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBAZAM, CLOBAZAM
 CLOTRIMAZOLE, CLOTRIMAZOLE
 CODEINE SULFATE, CODEINE SULFATE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 CYCLOSPORINE, CYCLOSPORINE
 CYTARABINE, CYTARABINE
 DACARBAZINE, DACARBAZINE
 DALFAMPRIDINE, DALFAMPRIDINE
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DEXAMETHASONE INTENSOL, DEXAMETHASONE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DEXAMETHASONE, DEXAMETHASONE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 DIAZEPAM INTENSOL, DIAZEPAM
 DIAZEPAM, DIAZEPAM
 DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE), DICYCLOMINE HYDROCHLORIDE
 DIGOXIN, DIGOXIN
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DISULFIRAM, DISULFIRAM
 DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
 DOLOPHINE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 DOPRAM, DOXAPRAM HYDROCHLORIDE
 DOXERCALCIFEROL, DOXERCALCIFEROL
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE HYCLATE
 DURAMORPH PF, MORPHINE SULFATE
 DUTASTERIDE, DUTASTERIDE
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 ESZOPICLONE, ESZOPICLONE
 ETHACRYNIC ACID, ETHACRYNIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** W **

* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
 ETOMIDATE, ETOMIDATE
 ETOPOSIDE, ETOPOSIDE
 EVEROLIMUS, EVEROLIMUS
 EXEMESTANE, EXEMESTANE
 FAMCICLOVIR, FAMCICLOVIR
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
 FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLOXURIDINE, FLOXURIDINE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
 FLUCONAZOLE, FLUCONAZOLE
 FLUCYTOSINE, FLUCYTOSINE
 FLUMAZENIL, FLUMAZENIL
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FUROSEMIDE, FUROSEMIDE
 GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HEPARIN SODIUM, HEPARIN SODIUM
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
 IFOSFAMIDE, IFOSFAMIDE
 IMATINIB MESYLATE, IMATINIB MESYLATE
 IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
 INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
 INFUMORPH, MORPHINE SULFATE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 IRBESARTAN, IRBESARTAN
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LACTULOSE, LACTULOSE
 LETROZOLE, LETROZOLE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LEVOCARNITINE, LEVOCARNITINE
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE
 LINEZOLID, LINEZOLID
 LITHIUM CARBONATE, LITHIUM CARBONATE
 LITHIUM CITRATE, LITHIUM CITRATE
 LORAZEPAM INTENSOL, LORAZEPAM
 LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 MERCAPTOPYRINE, MERCAPTOPYRINE
 MESNA, MESNA
 METHADONE HYDROCHLORIDE INTENSOL, METHADONE HYDROCHLORIDE
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
 METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 MILRINONE LACTATE, MILRINONE LACTATE
 MITOMYCIN, MITOMYCIN
 MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 MONTELUKAST SODIUM, MONTELUKAST SODIUM
 MORPHINE SULFATE, MORPHINE SULFATE
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 NALOXONE, NALOXONE HYDROCHLORIDE
 NAPROXEN, NAPROXEN
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXCARBAZEPINE, OXCARBAZEPINE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 OXYTOCIN, OXYTOCIN
 PACLITAXEL, PACLITAXEL
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PENTOSTATIN, PENTOSTATIN
 PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
 PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
 PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE
 PHENYTOIN SODIUM, PHENYTOIN SODIUM
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 PREDNISONE INTENSOL, PREDNISONE
 PREDNISONE, PREDNISONE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPANTHELINE BROMIDE, PROPANTHELINE BROMIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
 RAMIPRIL, RAMIPRIL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RIFAMPIN, RIFAMPIN
 RISPERIDONE, RISPERIDONE
 RITONAVIR, RITONAVIR
 ROBAXIN, METHOCARBAMOL
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROXICET, ACETAMINOPHEN
 RUFINAMIDE, RUFINAMIDE
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
 SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE, SODIUM FERRIC GLUCONATE COMPLEX
 SODIUM OXYBATE, SODIUM OXYBATE
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
 SUFENTANIL CITRATE, SUFENTANIL CITRATE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 THIOTEPA, THIOTEPA
 TINIDAZOLE, TINIDAZOLE
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 TORSEMIDE, TORSEMIDE
 TRIAZOLAM, TRIAZOLAM
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VINBLASTINE SULFATE, VINBLASTINE SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
 VINOELBINE TARTRATE, VINOELBINE TARTRATE
 ZALEPLON, ZALEPLON
 ZIDOVUDINE, ZIDOVUDINE

WI MEDCL CYCLOTRON

* WISCONSIN MEDICAL CYCLOTRON LLC
 AMMONIA N 13, AMMONIA N-13
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

WILSHIRE PHARMS INC

* WILSHIRE PHARMACEUTICALS INC
 CARISOPRODOL, CARISOPRODOL
 NATEGLINIDE, NATEGLINIDE
 PERPHENAZINE, PERPHENAZINE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 SEVELAMER CARBONATE, SEVELAMER CARBONATE
 TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE

WOCKHARDT

* WOCKHARDT LTD
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AZITHROMYCIN, AZITHROMYCIN
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 CEFOTAXIME SODIUM, CEFOTAXIME SODIUM
 CEFOTAXIME, CEFOTAXIME SODIUM
 CEFPROZIL, CEFPROZIL
 CEFTAZIDIME, CEFTAZIDIME
 CEFTRIAZONE, CEFTRIAZONE SODIUM
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 CLARITHROMYCIN, CLARITHROMYCIN
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 FUROSEMIDE, FUROSEMIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEVETIRACETAM, LEVETIRACETAM
 LEVOFLOXACIN, LEVOFLOXACIN
 LISINAPRIL, LISINAPRIL
 METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 NIACIN, NIACIN
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 RISPERIDONE, RISPERIDONE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TIMOLOL MALEATE, TIMOLOL MALEATE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
 ZONISAMIDE, ZONISAMIDE

WOCKHARDT BIO AG

* WOCKHARDT BIO AG
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACETIC ACID, ACETIC ACID, GLACIAL
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 BROMFED-DM, BROMPHENIRAMINE MALEATE
 CARBAMAZEPINE, CARBAMAZEPINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ******* WOCKHARDT BIO AG**

CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 CYCLOSPORINE, CYCLOSPORINE
 DEXAMETHASONE, DEXAMETHASONE
 DEXCHLORPHENIRAMINE MALEATE, DEXCHLORPHENIRAMINE MALEATE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 ERYTHROMYCIN, ERYTHROMYCIN
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 FUROSEMIDE, FUROSEMIDE
 GENERLAC, LACTULOSE
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 LACTULOSE, LACTULOSE
 LEVETIRACETAM, LEVETIRACETAM
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 LINDANE, LINDANE
 LITHIUM CITRATE, LITHIUM CITRATE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE, LORATADINE (OTC)
 MEGESTROL ACETATE, MEGESTROL ACETATE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 NYSTATIN, NYSTATIN
 OXACILLIN SODIUM, OXACILLIN SODIUM
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PHENYTOIN, PHENYTOIN
 PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PREDNISOLONE, PREDNISOLONE
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE PLAIN, PROMETHAZINE HYDROCHLORIDE
 PROMETHAZINE W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE
 SELENIUM SULFIDE, SELENIUM SULFIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 VALPROIC ACID, VALPROIC ACID

WOCKHARDT LTD*** WOCKHARDT LTD**

BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 CAPTOPRIL, CAPTOPRIL
 CARBIDOPA, LEVODOPA AND ENTACAPONE, CARBIDOPA
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ENTACAPONE, ENTACAPONE
 FAMOTIDINE, FAMOTIDINE
 FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 LAMOTRIGINE, LAMOTRIGINE
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

WOCKHARDT USA*** WOCKHARDT USA INC**

GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE

*** WOCKHARDT USA LLC**

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

* WOCKHARDT USA LLC
LANSOPRAZOLE, LANSOPRAZOLE

WRASER PHARMS

* WRASER PHARMACEUTICALS LLC
CETRAXAL, CIPROFLOXACIN HYDROCHLORIDE

WRASER PHARMS LLC

* WRASER PHARMACEUTICALS LLC
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
TREZIX, ACETAMINOPHEN

WUSM CYCLOTRON

* WASHINGTON UNIV SCH MEDICINE CYCLOTRON FACILITY
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

WYETH PHARMS

* WYETH PHARMACEUTICALS LLC
DUAVEE, BAZEDOXIFENE ACETATE
EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE
PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE
PREMARIN, ESTROGENS, CONJUGATED
PREMPHASE 14/14, ESTROGENS, CONJUGATED
PREMPRO, ESTROGENS, CONJUGATED
PROTONIX IV, PANTOPRAZOLE SODIUM
PROTONIX, PANTOPRAZOLE SODIUM
TRECATOR, ETHIONAMIDE
ZOSYN IN PLASTIC CONTAINER, PIPERACILLIN SODIUM
ZOSYN, PIPERACILLIN SODIUM

**** X ******X GEN PHARMS**

* X GEN PHARMACEUTICALS INC
ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
AMPHOTERICIN B, AMPHOTERICIN B
BACIIM, BACITRACIN
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
LEVETIRACETAM, LEVETIRACETAM
LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
NEOMYCIN SULFATE, NEOMYCIN SULFATE
NYSTATIN, NYSTATIN
POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE

X-GEN PHARMS

* X-GEN PHARMACEUTICALS INC
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

X-GEN PHARMS INC

* X-GEN PHARMACEUTICALS INC
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
IBUPROFEN LYSINE, IBUPROFEN LYSINE
LINCOMYCIN, LINCOMYCIN HYDROCHLORIDE
TRANEXAMIC ACID, TRANEXAMIC ACID

XELLIA PHARMS APS

* XELLIA PHARMACEUTICALS APS
BACITRACIN, BACITRACIN
CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
DAPTOMYCIN, DAPTOMYCIN
POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
VORICONAZOLE, VORICONAZOLE

XIAMEN LP PHARM CO

* XIAMEN LP PHARMACUETICAL CO LTD
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** X ******XSPIRE PHARMA**

- * XSPIRE PHARMA
NALFON, FENOPROFEN CALCIUM
- * XSPIRE PHARMA LLC
DEXAMETHASONE, DEXAMETHASONE
FENOPROFEN CALCIUM, FENOPROFEN CALCIUM

XTTRIUM

- * XTTRIUM LABORATORIES INC
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)

XYLOPIA

- * XYLOPIA
ACETAZOLAMIDE, ACETAZOLAMIDE

**** Y ******YABAO PHARM**

- * YABAO PHARMACEUTICAL CO LTD BEIJING
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE

YAOPHARMA CO LTD

- * YAOPHARMA CO LTD
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

YICHANG HUMANWELL

- * YICHANG HUMANWELL PHARMACEUTICAL CO LTD
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

YILING PHARM LTD

- * YILING PHARMACEUTICAL LTD
ACYCLOVIR, ACYCLOVIR
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
FELODIPINE, FELODIPINE

YUNG SHIN PHARM

- * YUNG SHIN PHARMACEUTICAL INDUSTRIAL CO LTD
CEFACTOR, CEFACOR
CEPHALEXIN, CEPHALEXIN
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
MELOXICAM, MELOXICAM
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**** Z ******ZAMBON SPA**

- * ZAMBON SPA ITALY
MONUROL, FOSFOMYCIN TROMETHAMINE

ZHEJIANG HISUN PHARM

- * ZHEJIANG HISUN PHARMACEUTICAL CO LTD
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL

ZO SKIN HEALTH

- * ZO SKIN HEALTH
TRETINOIN, TRETINOIN

ZYDUS HLTHCARE

- * ZYDUS HEALTHCARE USA LLC
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
LANSOPRAZOLE, LANSOPRAZOLE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

ZYDUS PHARMS USA

- * ZYDUS PHARMACEUTICALS USA INC
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
ATENOLOL, ATENOLOL
AZATHIOPRINE, AZATHIOPRINE
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BENZONATATE, BENZONATATE
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ******* ZYDUS PHARMACEUTICALS USA INC**

HALOPERIDOL, HALOPERIDOL
 LAMOTRIGINE, LAMOTRIGINE
 MELOXICAM, MELOXICAM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 NAPROXEN, NAPROXEN
 PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 RAMIPRIL, RAMIPRIL
 RIBAVIRIN, RIBAVIRIN
 RISPERIDONE, RISPERIDONE
 SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 SIMVASTATIN, SIMVASTATIN
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 WARFARIN SODIUM, WARFARIN SODIUM
 ZONISAMIDE, ZONISAMIDE

ZYDUS PHARMS USA INC*** ZYDUS PHARMACEUTICALS USA INC**

ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
 ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 ACYCLOVIR, ACYCLOVIR
 ALBENDAZOLE, ALBENDAZOLE
 ALLOPURINOL, ALLOPURINOL
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
 AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
 ANASTROZOLE, ANASTROZOLE
 ARIPIPIRAZOLE, ARIPIPIRAZOLE
 ARSENIC TRIOXIDE, ARSENIC TRIOXIDE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BICALUTAMIDE, BICALUTAMIDE
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BUDESONIDE, BUDESONIDE
 BUMETANIDE, BUMETANIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
 CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
 CARBIDOPA, CARBIDOPA
 CARVEDILOL, CARVEDILOL
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLOBAZAM, CLOBAZAM
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 DESOXIMETASONE, DESOXIMETASONE
 DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE
 DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DIFLUNISAL, DIFLUNISAL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ****

* ZYDUS PHARMACEUTICALS USA INC
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DIPYRIDAMOLE, DIPYRIDAMOLE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DOXYCYCLINE, DOXYCYCLINE
DOXYCYCLINE, DOXYCYCLINE HYCLATE
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
DUTASTERIDE, DUTASTERIDE
ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
ENTECAVIR, ENTECAVIR
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
ETODOLAC, ETODOLAC
ETOMIDATE, ETOMIDATE
EXEMESTANE, EXEMESTANE
EZETIMIBE, EZETIMIBE
FELBAMATE, FELBAMATE
FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
FINASTERIDE, FINASTERIDE
FLUCONAZOLE, FLUCONAZOLE
FLUOCINONIDE, FLUOCINONIDE
GABAPENTIN, GABAPENTIN
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
GEMFIBROZIL, GEMFIBROZIL
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
GLIPIZIDE, GLIPIZIDE
GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
GLYBURIDE, GLYBURIDE
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
INDOMETHACIN, INDOMETHACIN
IRBESARTAN, IRBESARTAN
ITRACONAZOLE, ITRACONAZOLE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LAMOTRIGINE, LAMOTRIGINE
LANSOPRAZOLE, LANSOPRAZOLE
LEVETIRACETAM, LEVETIRACETAM
LEVOFLOXACIN, LEVOFLOXACIN
LINEZOLID, LINEZOLID
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
MESALAMINE, MESALAMINE
METHOTREXATE SODIUM, METHOTREXATE SODIUM
METHYLPREDNISOLONE, METHYLPREDNISOLONE
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
METRONIDAZOLE, METRONIDAZOLE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
MODAFINIL, MODAFINIL
MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
NADOLOL, NADOLOL
NATEGLINIDE, NATEGLINIDE
NIFEDIPINE, NIFEDIPINE
NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
NYSTATIN, NYSTATIN
OLANZAPINE, OLANZAPINE
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
OMEPRAZOLE, OMEPRAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ****

* ZYDUS PHARMACEUTICALS USA INC
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PINDOLOL, PINDOLOL
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
 PIROXICAM, PIROXICAM
 POTASSIUM CITRATE, POTASSIUM CITRATE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RISPERIDONE, RISPERIDONE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 SIROLIMUS, SIROLIMUS
 SPIRONOLACTONE, SPIRONOLACTONE
 SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TELMISARTAN, TELMISARTAN
 TEMOZOLOMIDE, TEMOZOLOMIDE
 TERIFLUNOMIDE, TERIFLUNOMIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRANEXAMIC ACID, TRANEXAMIC ACID
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VARDENAFIL HYDROCHLORIDE, VARDENAFIL HYDROCHLORIDE
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 VORICONAZOLE, VORICONAZOLE
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
 ZOLMITRIPTAN, ZOLMITRIPTAN
 ZYPITAMAG, PITAVASTATIN MAGNESIUM

ZYDUS WORLDWIDE

* ZYDUS WORLDWIDE DMCC
 AZITHROMYCIN, AZITHROMYCIN
 BACLOFEN, BACLOFEN
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 URSODIOL, URSODIOL

39TH EDITION - 2019 - APPROVED DRUG PRODUCTS LIST

APPENDIX C**UNIFORM TERMS*****DOSAGE FORMS***

| | |
|----------------------------------|----------------------------------------------------|
| AEROSOL, FOAM | OINTMENT |
| AEROSOL, METERED | OINTMENT, AUGMENTED |
| CAPSULE | PASTE |
| CAPSULE, DELAYED REL PELLETS | PATCH |
| CAPSULE, DELAYED RELEASE | PELLET |
| CAPSULE, EXTENDED RELEASE | POWDER |
| CAPSULE, PELLET | POWDER, EXTENDED RELEASE |
| CLOTH | POWDER, METERED |
| CONCENTRATE | RING |
| CREAM | SHAMPOO |
| CREAM, AUGMENTED | SOLUTION |
| ELIXIR | SOLUTION FOR SLUSH |
| EMULSION | SOLUTION, EXTENDED RELEASE |
| ENEMA | SOLUTION, GEL FORMING/DROPS |
| FILM | SOLUTION, METERED |
| FILM, EXTENDED RELEASE | SOLUTION/DROPS |
| FOR SOLUTION | SPONGE |
| FOR SUSPENSION | SPRAY |
| FOR SUSPENSION, DELAYED RELEASE | SPRAY, METERED |
| FOR SUSPENSION, EXTENDED RELEASE | SUPPOSITORY |
| GAS | SUSPENSION |
| GEL | SUSPENSION, EXTENDED RELEASE |
| GEL, AUGMENTED | SUSPENSION, LIPOSOMAL |
| GEL, METERED | SUSPENSION/DROPS |
| GRANULE | SWAB |
| GRANULE, DELAYED RELEASE | SYRUP |
| GUM, CHEWING | SYSTEM |
| IMPLANT | SYSTEM, EXTENDED RELEASE |
| INHALANT | TABLET |
| INJECTABLE | TABLET, CHEWABLE |
| INJECTABLE, LIPID COMPLEX | TABLET, DELAYED RELEASE |
| INJECTABLE, LIPOSOMAL | TABLET, EFFERVESCENT |
| INJECTION, EXTENDED RELEASE | TABLET, EXTENDED RELEASE |
| INSERT | TABLET, EXTENDED RELEASE, CHEWABLE |
| INSERT, EXTENDED RELEASE | TABLET, FOR SUSPENSION |
| INTRAUTERINE DEVICE | TABLET, ORALLY DISINTEGRATING |
| JELLY | TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE |
| LIQUID | TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE |
| LOTION | TAPE |
| LOTION, AUGMENTED | TROCHE/LOZENGE |
| LOTION/SHAMPOO | |
| OIL | |
| OIL/DROPS | |

Note: Terms comprise currently marketed products

APPENDIX C

UNIFORM TERMS

ROUTES OF ADMINISTRATION

| | |
|-------------------|--------------------|
| BUCCAL | INTRAVITREAL |
| DENTAL | IONTOPHORESIS |
| ENDOCERVICAL | IRRIGATION |
| ENDOTRACHEAL | IV (INFUSION) |
| ENTERAL | N/A |
| IMPLANTATION | NASAL |
| INHALATION | OPHTHALMIC |
| INJECTION | ORAL |
| INTRA-ANAL | ORAL-21 |
| INTRA-ARTERIAL | ORAL-28 |
| INTRA-ARTICULAR | OTIC |
| INTRACRANIAL | PERFUSION, CARDIAC |
| INTRADERMAL | PERIODONTAL |
| INTRAMUSCULAR | RECTAL |
| INTRAOCULAR | SPINAL |
| INTRAOSSEOUS | SUBCUTANEOUS |
| INTRAPERITONEAL | SUBLINGUAL |
| INTRAPLEURAL | TOPICAL |
| INTRATHECAL | TRANSDERMAL |
| INTRATRACHEAL | TRANSMUCOSAL |
| INTRAUTERINE | URETHRAL |
| INTRAVENOUS | VAGINAL |
| INTRAVENOUS BOLUS | |
| INTRAVESICAL | |

Note: Terms comprise currently marketed products

APPENDIX C**UNIFORM TERMS*****ABBREVIATIONS***

| | |
|---------|----------------------------|
| AMP | AMPULE |
| AMPICIL | AMPICILLIN |
| APPROX | APPROXIMATELY |
| BOT | BOTTLE |
| CI | CURIE |
| CSR | CAROTID SINUS REFLEX |
| CU | CLINICAL UNITS |
| DIPROP | DIPROPIONATE |
| ELECT | ELECTROLYTE |
| EQ | EQUIVALENT TO |
| ER | EXTENDED RELEASE |
| GM | GRAM |
| HBR | HYDROBROMIDE |
| HCL | HYDROCHLORIDE |
| HR | HOUR |
| IM | INTRAMUSCULAR |
| INH | INHALATION |
| IU | INTERNATIONAL UNITS |
| IV | INTRAVENOUS |
| KIU | KALLIKREIN INHIBITOR UNITS |
| MCG | MICROGRAM |
| mCi | MILLICURIE |
| MEQ | MILLIEQUIVALENT |
| MG | MILLIGRAM |
| ML | MILLILITER |
| N/A | NOT APPLICABLE |
| PPM | PARTS PER MILLION |
| REL | RELEASE |
| SQ CM | SQUARE CENTIMETER |
| SC | SUBCUTANEOUS |
| U | UNITS |
| uCi | MICROCURIE |
| UMOLAR | MICROMOLAR |
| USP | UNITED STATES PHARMACOPEIA |

PATENT AND EXCLUSIVITY INFORMATION ADDENDUM

This *Addendum* identifies drugs that qualify under the Federal Food, Drug, and Cosmetic Act (FD&C Act) for periods of exclusivity and provides patent information that has been submitted to the Food and Drug Administration (FDA) concerning the listed drug products.

Exclusivity

During relevant exclusivity periods, certain abbreviated new drug applications (ANDAs) and applications described in Section 505(b)(2) of the FD&C Act (505(b)(2) applications) may not be submitted or approved as described below. This *Addendum* identifies drugs approved under section 505(c) of the FD&C Act that qualify under the Drug Price Competition and Patent Term Restoration Act of 1984 (Hatch-Waxman Amendments) for five-year and three-year periods of exclusivity pursuant to Section 505(c)(3)(E) and Section 505(j)(5)(F) and certain generic drugs approved under section 505(j) of the FD&C Act that have qualified for 180-day exclusivity pursuant to Section 505(j)(5)(B)(iv). This *Addendum* also identifies those drugs that have qualified for Orphan Drug Exclusivity pursuant to Section 527 of the FD&C Act, those drugs that have qualified for Pediatric Exclusivity pursuant to Section 505A of the FD&C Act, those drugs that have qualified for Generating Antibiotics Incentives Now (GAIN) exclusivity pursuant to Section 505E of the FD&C Act, and those generic drugs that have qualified for Competitive Generic Therapy (CGT) exclusivity pursuant to Section 505(j)(5)(B)(v) of the FD&C Act. This section is arranged in alphabetical order by active ingredient name followed by the proprietary name. Active ingredient headings for multiple ingredient fixed-combination drug products are arranged alphabetically.

For an explanation of the codes used in the *Addendum*, see the *Patent and Exclusivity Terms* Section. The exclusivity codes are general shorthand descriptions and do not necessarily identify, with specificity, the actual scope of exclusivity. Please note that beginning with the publication of the 38th edition of the Orange Book, individual descriptions of the protected use have been added to each Orphan Drug Exclusivity entry listed in the Orange Book. In previous editions of the Orange Book, Orphan Drug Exclusivity was not described with any specificity.

Exclusivity does not prevent the submission or approval of an application submitted pursuant to Section 505(b)(1) of the FD&C Act that would otherwise be blocked if it had been submitted pursuant to Section 505(b)(2) or 505(j), except in the case of Orphan Drug Exclusivity. Drugs approved under section 505(c) of the FD&C Act that may qualify for periods of exclusivity include:

- (1) A new chemical entity, submitted in a new drug application under Section 505(b) of the FD&C Act and approved after September 24, 1984. A new chemical entity is an active ingredient that contains "no active ingredient (including any ester or salt of the active ingredient)" that has been approved by FDA in any other application submitted under Section 505(b) of the FD&C Act. An active ingredient would be eligible for 5-year exclusivity for a new chemical entity, provided that it meets the definition of a new chemical entity, regardless of whether that active ingredient is approved in a single-ingredient drug product, in a fixed-combination with another active ingredient that contains no other previously approved active moiety, or in a fixed-combination with

another active ingredient that contains a previously approved active moiety.¹ No subsequent ANDA or 505(b)(2) application for a drug that contains the same active moiety may be *submitted* for a period of *five years* from the date of approval of the original application, except that such an application may be *submitted* after *four years* if it contains a certification that a patent claiming the drug is invalid or will not be infringed by the product for which approval is sought. See Sections 505(j)(5)(F)(ii) and 505(c)(3)(E)(ii) of the FD&C Act.

- (2) A new drug application approved after September 24, 1984, for a drug product containing "an active ingredient (including any ester or salt of the active ingredient)" that has been approved in an earlier new drug application and that includes reports of new clinical investigations (other than bioavailability studies). Such investigations must have been conducted or sponsored by the applicant and must have been essential to approval of the application. If these requirements are met, a subsequent ANDA or a 505(b)(2) application may not be approved for the exclusivity-protected "conditions of approval of such drug" before the expiration of *three years* from the date of approval of the original application. See Sections 505(j)(5)(F)(iii) and 505(c)(3)(E)(iii) of the FD&C Act. If an NDA has exclusivity only for a new indication or use, this exclusivity generally does not preclude the approval of an ANDA or 505(b)(2) application for indications and uses not covered by the exclusivity, assuming the proposed drug product will be safe and effective as labeled.
- (3) A supplement to a new drug application for a drug containing a previously approved "active ingredient (including any ester or salt of the active ingredient)" approved after September 24, 1984, that contains reports of new clinical investigations (other than bioavailability studies) essential to the approval of the supplement and conducted or sponsored by the applicant. See Sections 505(j)(5)(F)(iv) and 505(c)(3)(E)(iv) of the FD&C Act. A subsequent ANDA or 505(b)(2) application may not be approved for an exclusivity-protected change approved in the supplement for *three years* from the date of approval of the supplement.

Patent Information

The FD&C Act requires that patent information be filed with all newly submitted Section 505(b) drug applications. No NDA may be approved after September 24, 1984, without the submission of patent information to the Agency. Effective August 18, 2003, this information must be filed using Form FDA 3542a "Patent Information Submitted with the Filing of an NDA, Amendment or Supplement".

Effective August 18, 2003, upon approval of an application, patent information for purposes of listing in the Orange Book must be submitted to the Agency within 30 days of the date of approval on Form FDA 3542 "Patent

¹ For more information on exclusivity for fixed-dose combination drug products that include new chemical entities, see FDA's guidance on *New Chemical Entity Exclusivity Determinations for Certain Fixed-Dose Combination Drug Products* at <https://www.fda.gov/ucm/groups/fdagov-public/@fdagov-drugs-gen/documents/document/ucm386685.pdf>.

Information Submitted Upon and After Approval of an NDA or Supplement."² In November 2017, the Agency began including in the Orange Book the patent submission date (i.e., the date on which the FDA receives patent information from the NDA holder) for each newly listed patent to facilitate assessments of whether patent information is untimely filed with respect to a pending 505(b)(2) application or ANDA.³ Patent information on unapproved applications or on patents beyond the scope of the FD&C Act (i.e., process or manufacturing patents) will not be published. Form FDA 3542 will be the only form used for purposes of this publication.

The patents that FDA regards as covered by the statutory provisions for submission of patent information are: patents that claim the active ingredient(s); drug product patents, which include formulation/composition patents; method-of-use patents that claim one or more approved methods of using the approved drug product; and certain other patents as detailed in the regulations and on Form FDA 3542.⁴ This information, as provided by the sponsor on Form FDA 3542, will be published as described above. As of December 5, 2016, an NDA holder submitting information on a patent that claims both the drug substance and the drug product (and is eligible for listing on either basis) is required only to specify that the patent claims either the drug substance or the drug product.

A requirement for submission of patent information to FDA for certain old antibiotics became effective October 7, 2008 under Section 4(b)(1) of the QI Program Supplemental Funding Act (Public Law 110-379) (QI Act).

Upon approval of an NDA or at such time as patent information is updated for the drug product, FDA adds to the Orange Book the patent number, expiration date, type of patent, and submission date... The Addendum lists patent and exclusivity information up to January of the Edition year. The monthly Cumulative Supplements to the annual edition list patent and exclusivity information changes since the Annual Edition Addendum. Since all parts of this publication are subject to changes, additions, or deletions, the Orange Book, updated daily, should be consulted for the most recent patent and exclusivity information.

² Please note that the date of approval for an NDA for a drug for which FDA intends to recommend controls under the Controlled Substances Act is the later of the date on the approval letter for the NDA or the date of issuance of the interim final rule controlling the drug (see Section 505(x)(1) and (2) of the FD&C Act).

³ See 21 CFR 314.50(i)(4) and 314.94(a)(12)(vi). The submission date for patent information is determined in accordance with 21 CFR 314.53(d)(5).

⁴ See 21 CFR 314.53(c)(2)(ii)(M), (N)(2) and (N)(3).

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ABACAVIR SULFATE - ZIAGEN</u> | | | | | | |
| N 020978 | 001 6641843 | Feb 04, 2019 | DP | | | |
| <u>ABACAVIR SULFATE; DOLUTEGRAVIR SODIUM; LAMIVUDINE - TRIUMEQ</u> | | | | | | |
| N 205551 | 001 8129385 | Oct 05, 2027 | DS DP | | | |
| | 9242986 | Dec 08, 2029 | DS DP | | | |
| <u>ABALOPARATIDE - TYMLOS</u> | | | | | | |
| N 208743 | 001 7803770 | Mar 26, 2028 | | U-2009 | NCE | Apr 28, 2022 |
| | 8148333 | Nov 08, 2027 | DP | | | |
| | 8748382 | Oct 03, 2027 | | U-2009 | | |
| <u>ABEMACICLIB - VERZENIO</u> | | | | | | |
| N 208716 | 001 7855211 | Dec 15, 2029 | DS DP U-2132 | | I-768 | Feb 26, 2021 |
| | 7855211 | Dec 15, 2029 | DS DP U-2135 | | NCE | Sep 28, 2022 |
| | 7855211 | Dec 15, 2029 | DS DP U-2251 | | | |
| <u>ABEMACICLIB - VERZENIO</u> | | | | | | |
| N 208716 | 002 7855211 | Dec 15, 2029 | DS DP U-2132 | | I-768 | Feb 26, 2021 |
| | 7855211 | Dec 15, 2029 | DS DP U-2135 | | NCE | Sep 28, 2022 |
| | 7855211 | Dec 15, 2029 | DS DP U-2251 | | | |
| <u>ABEMACICLIB - VERZENIO</u> | | | | | | |
| N 208716 | 003 7855211 | Dec 15, 2029 | DS DP U-2132 | | I-768 | Feb 26, 2021 |
| | 7855211 | Dec 15, 2029 | DS DP U-2135 | | NCE | Sep 28, 2022 |
| | 7855211 | Dec 15, 2029 | DS DP U-2251 | | | |
| <u>ABEMACICLIB - VERZENIO</u> | | | | | | |
| N 208716 | 004 7855211 | Dec 15, 2029 | DS DP U-1981 | | I-768 | Feb 26, 2021 |
| | 7855211 | Dec 15, 2029 | DS DP U-2132 | | NCE | Sep 28, 2022 |
| | 7855211 | Dec 15, 2029 | DS DP U-2135 | | | |
| | 7855211 | Dec 15, 2029 | DS DP U-2251 | | | |
| <u>ABIRATERONE ACETATE - ZYTIGA</u> | | | | | | |
| N 202379 | 001 8822438 | Aug 24, 2027 | | U-1579 | I-765 | Feb 07, 2021 |
| | 8822438 | Aug 24, 2027 | | U-1580 | | |
| | 8822438 | Aug 24, 2027 | | U-2235 | | |
| <u>ABIRATERONE ACETATE - ZYTIGA</u> | | | | | | |
| N 202379 | 002 8822438 | Aug 24, 2027 | | U-1579 | I-765 | Feb 07, 2021 |
| | 8822438 | Aug 24, 2027 | | U-1580 | | |
| | 8822438 | Aug 24, 2027 | | U-2235 | | |
| <u>ABIRATERONE ACETATE - YONSA</u> | | | | | | |
| N 210308 | 001 9889144 | Mar 17, 2034 | DP | | | |
| <u>ACALABRUTINIB - CALOQUENCE</u> | | | | | | |
| N 210259 | 001 7459554 | Nov 24, 2026 | DS | | NCE | Oct 31, 2022 |
| | 9290504 | Jul 11, 2032 | DS DP | | ODE-175 | Oct 31, 2024 |
| | 9758524 | Jul 11, 2032 | | U-2145 | | |
| | 9796721 | Jul 01, 2036 | DS DP U-2145 | | | |
| <u>ACETAMINOPHEN - OFIRMEV</u> | | | | | | |
| N 022450 | 001 6992218 | Jun 06, 2021 | DP | | M-196 | Jan 27, 2020 |
| | 6992218*PED | Dec 06, 2021 | | | PED | Jul 27, 2020 |
| | 9399012 | Sep 11, 2031 | | U-2261 | | |
| | 9399012 | Sep 11, 2031 | | U-2262 | | |
| | 9399012*PED | Mar 11, 2032 | | | | |
| | 9610265 | Nov 13, 2028 | | U-2263 | | |
| | 9610265*PED | May 13, 2029 | | | | |
| | 9987238 | Nov 13, 2028 | | U-2261 | | |
| | 9987238*PED | May 13, 2029 | | | | |
| <u>ACETAMINOPHEN; BENZHYDROCODONE HYDROCHLORIDE - APADAZ</u> | | | | | | |
| N 208653 | 001 8461137 | Feb 22, 2031 | DS DP | | | |
| | 8748413 | Jul 01, 2030 | DS DP | | | |
| | 8828978 | Jul 01, 2030 | DP | | | |
| | 9132125 | Jul 01, 2030 | DS DP U-2249 | | | |
| | 9549923 | Jul 01, 2030 | DS DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE - XARTEMIS XR</u> | | | | | | |
| N 204031 | 001 | 6488962 | Jun 20, 2020 | DP | | |
| | | 7976870 | Jun 01, 2027 | DP | U-1498 | |
| | | 8372432 | Mar 11, 2029 | DP | U-1499 | |
| | | 8377453 | Nov 19, 2029 | DP | U-1499 | |
| | | 8394408 | Mar 11, 2029 | DP | | |
| | | 8597681 | Dec 21, 2030 | DP | | |
| | | 8658631 | May 16, 2032 | DP | | |
| | | 8668929 | Mar 11, 2029 | DP | U-1499 | |
| | | 8741885 | May 16, 2032 | DP | U-1499 | |
| | | 8980319 | Dec 21, 2030 | DP | | |
| | | 8992975 | May 16, 2032 | DP | | |
| | | 9050335 | May 16, 2032 | DP | | |
| | | 9468636 | May 16, 2032 | DP | U-1499 | |
| <u>ACETYLCYSTEINE - ACETADOTE</u> | | | | | | |
| N 021539 | 001 | 8148356 | May 21, 2026 | DP | | |
| | | 8399445 | Aug 24, 2025 | DP | U-1373 | |
| | | 8653061 | Aug 24, 2025 | DP | U-1373 | |
| | | 8722738 | Apr 06, 2032 | DP | U-1373 | |
| | | 9327028 | Jul 21, 2031 | DP | U-1839 | |
| <u>ACETYLCYSTEINE - CETYLEV</u> | | | | | | |
| N 207916 | 001 | 8747894 | May 08, 2032 | DP | U-1373 | |
| | | 9427421 | May 08, 2032 | DP | | |
| | | 9561204 | May 08, 2032 | DP | U-1373 | |
| <u>ACETYLCYSTEINE - CETYLEV</u> | | | | | | |
| N 207916 | 002 | 8747894 | May 08, 2032 | DP | U-1373 | |
| | | 9427421 | May 08, 2032 | DP | | |
| | | 9561204 | May 08, 2032 | DP | U-1373 | |
| <u>ACLIDINIUM BROMIDE - TUDORZA PRESSAIR</u> | | | | | | |
| N 202450 | 001 | 10034867 | Jul 07, 2020 | DP | U-2431 | |
| | | 10034867 | Jul 07, 2020 | DP | U-2432 | |
| | | 10085974 | Mar 13, 2029 | DP | U-1263 | |
| | | 6681768 | Aug 07, 2022 | DP | | |
| | | 7078412 | Jul 07, 2020 | DS DP | U-2431 | |
| | | 8051851 | Apr 22, 2027 | DP | | |
| | | 9056100 | Jul 07, 2020 | DP | U-1263 | |
| | | 9333195 | Jul 07, 2020 | DP | U-1263 | |
| | | RE46417 | Feb 10, 2025 | DS DP | U-1263 | |
| <u>ACYCLOVIR - SITAVIG</u> | | | | | | |
| N 203791 | 001 | 8592434 | Jun 16, 2030 | DP | U-1460 | |
| | | 8747896 | Jun 03, 2027 | DP | U-1460 | |
| | | 8791127 | Mar 23, 2027 | DP | U-1460 | |
| <u>ACYCLOVIR; HYDROCORTISONE - XERESE</u> | | | | | | |
| N 022436 | 001 | 7223387 | Nov 13, 2022 | DP | U-1006 | |
| | | 7223387 | Nov 13, 2022 | DP | U-1484 | |
| <u>ADAPALENE - DIFFERIN</u> | | | | | | |
| N 020380 | 002 | | | | RTO | Jul 08, 2019 |
| <u>ADAPALENE - DIFFERIN</u> | | | | | | |
| N 021753 | 001 | 7579377 | Feb 23, 2025 | DP | U-818 | |
| | | 7737181 | Aug 29, 2024 | DP | | |
| | | 7834060 | Mar 12, 2023 | DP | U-1078 | |
| | | 7838558 | Mar 12, 2023 | DP | | |
| | | 7868044 | Mar 12, 2023 | DP | U-1078 | |
| | | 8703820 | Mar 12, 2023 | DP | U-1078 | |
| <u>ADAPALENE - DIFFERIN</u> | | | | | | |
| N 022502 | 001 | 7998467 | May 31, 2028 | DP | U-1078 | |
| | | 8435502 | Sep 15, 2026 | DP | U-1078 | |
| | | 8709392 | Sep 15, 2026 | DP | U-1078 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ADAPALENE; BENZOYL PEROXIDE - EPIDUO</u> | | | | | | |
| N 022320 | 001 | 7820186 | Nov 23, 2025 | DP | | |
| | | 7964202 | Sep 01, 2024 | DP | U-1078 | |
| | | 8071644 | Jul 18, 2027 | DP | U-1078 | |
| | | 8080537 | Jul 18, 2027 | | U-1078 | |
| | | 8105618 | Dec 23, 2022 | | U-1078 | |
| | | 8129362 | Jul 18, 2027 | | U-1078 | |
| | | 8241649 | Dec 23, 2022 | DP | | |
| | | 8445543 | Jul 12, 2027 | | U-1078 | |
| | | 8809305 | Dec 23, 2022 | | U-1078 | |
| | | 8936800 | Dec 23, 2022 | DP | U-1078 | |
| <u>ADAPALENE; BENZOYL PEROXIDE - EPIDUO FORTE</u> | | | | | | |
| N 207917 | 001 | 8445543 | Dec 23, 2022 | | U-1078 | |
| | | 8703820 | Mar 12, 2023 | | U-1078 | |
| | | 8729127 | Mar 12, 2023 | | U-1078 | |
| | | 8785420 | Dec 23, 2022 | | U-1078 | |
| | | 8809305 | Dec 23, 2022 | | U-1078 | |
| | | 8936800 | Dec 23, 2022 | DP | U-1078 | |
| | | 9381179 | Mar 12, 2023 | | U-1078 | |
| | | 9387187 | Mar 12, 2023 | | U-1078 | |
| | | 9814690 | Dec 23, 2022 | DP | U-1078 | |
| <u>AFATINIB DIMALEATE - GILOTRIF</u> | | | | | | |
| N 201292 | 001 | 10004743 | Jul 05, 2030 | DP | I-763 | Jan 12, 2021 |
| | | 8426586 | Oct 10, 2029 | DS | ODE-115 | Apr 15, 2023 |
| | | 8545884 | Dec 19, 2029 | DP | ODE-50 | Jul 12, 2020 |
| | | 9539258 | Nov 09, 2026 | | U-1950 | |
| | | RE43431 | Jan 22, 2022 | DS | | |
| <u>AFATINIB DIMALEATE - GILOTRIF</u> | | | | | | |
| N 201292 | 002 | 10004743 | Jul 05, 2030 | DP | I-763 | Jan 12, 2021 |
| | | 8426586 | Oct 10, 2029 | DS | ODE-115 | Apr 15, 2023 |
| | | 8545884 | Dec 19, 2029 | DP | ODE-50 | Jul 12, 2020 |
| | | 9539258 | Nov 09, 2026 | | U-1950 | |
| | | RE43431 | Jan 22, 2022 | DS | | |
| <u>AFATINIB DIMALEATE - GILOTRIF</u> | | | | | | |
| N 201292 | 003 | 10004743 | Jul 05, 2030 | DP | I-730 | Apr 15, 2019 |
| | | 8426586 | Oct 10, 2029 | DS | I-763 | Jan 12, 2021 |
| | | 8545884 | Dec 19, 2029 | DP | ODE-115 | Apr 15, 2023 |
| | | 9539258 | Nov 09, 2026 | | U-1950 | Jul 12, 2020 |
| | | RE43431 | Jan 22, 2022 | DS | | |
| <u>ALBUMIN HUMAN - OPTISON</u> | | | | | | |
| N 020899 | 001 | 6723303 | Apr 20, 2021 | DP | | |
| <u>ALBUTEROL SULFATE - ACCUNEE</u> | | | | | | |
| N 020949 | 001 | 6702997 | Dec 28, 2021 | | U-558 | |
| <u>ALBUTEROL SULFATE - ACCUNEE</u> | | | | | | |
| N 020949 | 002 | 6702997 | Dec 28, 2021 | | U-558 | |
| <u>ALBUTEROL SULFATE - VENTOLIN HFA</u> | | | | | | |
| N 020983 | 001 | 7500444 | Feb 26, 2026 | DP | | |
| | | 7500444*PED | Aug 26, 2026 | | | |
| | | 7832351 | Jun 19, 2023 | DP | | |
| | | 9861771 | Oct 11, 2020 | DP | | |
| <u>ALBUTEROL SULFATE - PROAIR HFA</u> | | | | | | |
| N 021457 | 001 | 10022509 | May 18, 2031 | DP | | |
| | | 10022510 | May 18, 2031 | DP | | |
| | | 10086156 | May 18, 2031 | DP | | |
| | | 7105152 | Sep 12, 2023 | DP | | |
| | | 8132712 | Sep 07, 2028 | DP | | |
| | | 9463289 | May 18, 2031 | DP | | |
| | | 9808587 | May 18, 2031 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ALBUTEROL SULFATE - PROAIR RESPICLICK</u> | | | | | | |
| N 205636 | 001 | 10022510 | May 18, 2031 | DP | NPP | Apr 28, 2019 |
| | | 10124131 | May 18, 2031 | DP | | |
| | | 6701917 | Jun 23, 2021 | DP | | |
| | | 6718972 | Jun 23, 2021 | DP | | |
| | | 6748947 | Jun 23, 2021 | DP | | |
| | | 6871646 | Jun 23, 2021 | DP | | |
| | | 7540282 | May 06, 2023 | DP | | |
| | | 8006690 | Jun 23, 2021 | DP | | |
| | | 8651103 | Mar 26, 2028 | DP | | |
| | | 8978966 | Jan 13, 2032 | DP | | |
| | | 9216260 | Jun 28, 2031 | DP | | |
| | | 9463288 | May 19, 2025 | DP | | |
| | | 9731087 | May 18, 2031 | DP | | |
| | | 9782550 | Aug 28, 2035 | DP | | |
| | | 9782551 | Aug 28, 2035 | DP | | |
| <u>ALBUTEROL SULFATE; IPRATROPIUM BROMIDE - DUONEB</u> | | | | | | |
| N 020950 | 001 | 6632842 | Dec 28, 2021 | U-532 | | |
| <u>ALBUTEROL SULFATE; IPRATROPIUM BROMIDE - COMBIVENT RESPIMAT</u> | | | | | | |
| N 021747 | 001 | 6988496 | Feb 23, 2020 | DP | | |
| | | 7284474 | Aug 26, 2024 | DP | | |
| | | 7396341 | Oct 10, 2026 | DP | | |
| | | 7802568 | Feb 26, 2019 | DP | | |
| | | 7837235 | Mar 13, 2028 | DP | | |
| | | 7896264 | May 26, 2025 | DP | | |
| | | 7988001 | Aug 04, 2021 | DP | | |
| | | 8733341 | Oct 16, 2030 | DP | | |
| | | 9027967 | Mar 31, 2027 | DP | | |
| <u>ALCAFTADINE - LASTACAFT</u> | | | | | | |
| N 022134 | 001 | 8664215 | Dec 23, 2027 | U-1493 | | |
| <u>ALCOHOL - ABLYSINOL</u> | | | | | | |
| N 207987 | 001 | | | | ODE-192 | Jun 21, 2025 |
| <u>ALCOHOL - ABLYSINOL</u> | | | | | | |
| N 207987 | 002 | | | | ODE-192 | Jun 21, 2025 |
| <u>ALECTINIB HYDROCHLORIDE - ALECENSA</u> | | | | | | |
| N 208434 | 001 | 9126931 | May 29, 2031 | DS | I-756 | Nov 06, 2020 |
| | | 9365514 | Mar 04, 2032 | DP | NCE | Dec 11, 2020 |
| | | 9440922 | Jun 09, 2030 | DP | ODE-105 | Dec 11, 2022 |
| | | | | | ODE-159 | Nov 06, 2024 |
| <u>ALENDRONATE SODIUM - BINOSTO</u> | | | | | | |
| N 202344 | 001 | 7488496 | Aug 11, 2023 | DS DP | | |
| | | 7964212 | Mar 06, 2023 | DS DP | | |
| <u>ALISKIREN HEMIFUMARATE - TEKTURNA</u> | | | | | | |
| N 021985 | 001 | 5559111*PED | Jan 21, 2019 | | | |
| | | 8617595 | Feb 19, 2026 | DP | | |
| | | 8617595*PED | Aug 19, 2026 | | | |
| <u>ALISKIREN HEMIFUMARATE - TEKTURNA</u> | | | | | | |
| N 021985 | 002 | 5559111*PED | Jan 21, 2019 | | | |
| | | 8617595 | Feb 19, 2026 | DP | | |
| | | 8617595*PED | Aug 19, 2026 | | | |
| <u>ALISKIREN HEMIFUMARATE - TEKTURNA</u> | | | | | | |
| N 210709 | 001 | | | | NP PED | Nov 14, 2020 May 14, 2021 |
| <u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u> | | | | | | |
| N 022545 | 001 | 8613949 | Dec 21, 2029 | DP | | |
| <u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u> | | | | | | |
| N 022545 | 002 | 8613949 | Dec 21, 2029 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u> | | | | | | |
| N 022545 | 003 | 8613949 | Dec 21, 2029 | DP | | |
| <u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u> | | | | | | |
| N 022545 | 004 | 8613949 | Dec 21, 2029 | DP | | |
| <u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u> | | | | | | |
| N 200045 | 001 | 8183295 | May 16, 2023 | DP | | |
| | | 8618174 | Nov 15, 2021 | DP | | |
| <u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u> | | | | | | |
| N 200045 | 002 | 8183295 | May 16, 2023 | DP | | |
| | | 8618174 | Nov 15, 2021 | DP | | |
| <u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u> | | | | | | |
| N 200045 | 003 | 8183295 | May 16, 2023 | DP | | |
| | | 8618174 | Nov 15, 2021 | DP | | |
| <u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u> | | | | | | |
| N 200045 | 004 | 8183295 | May 16, 2023 | DP | | |
| | | 8618174 | Nov 15, 2021 | DP | | |
| <u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u> | | | | | | |
| N 200045 | 005 | 8183295 | May 16, 2023 | DP | | |
| | | 8618174 | Nov 15, 2021 | DP | | |
| <u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u> | | | | | | |
| N 022107 | 001 | 5559111*PED | Jan 21, 2019 | | | |
| | | 8618172 | Jul 13, 2028 | DP | | |
| | | 9023893 | Mar 03, 2022 | DP | | |
| <u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u> | | | | | | |
| N 022107 | 002 | 5559111*PED | Jan 21, 2019 | | | |
| | | 8618172 | Jul 13, 2028 | DP | | |
| | | 9023893 | Mar 03, 2022 | DP | | |
| <u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u> | | | | | | |
| N 022107 | 003 | 5559111*PED | Jan 21, 2019 | | | |
| | | 8618172 | Jul 13, 2028 | DP | | |
| | | 9023893 | Mar 03, 2022 | DP | | |
| <u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u> | | | | | | |
| N 022107 | 004 | 5559111*PED | Jan 21, 2019 | | | |
| | | 8618172 | Jul 13, 2028 | DP | | |
| | | 9023893 | Mar 03, 2022 | DP | | |
| <u>ALISKIREN HEMIFUMARATE; VALSARTAN - VALTURNA</u> | | | | | | |
| N 022217 | 001 | 8168616 | Jul 03, 2026 | DP | | |
| <u>ALISKIREN HEMIFUMARATE; VALSARTAN - VALTURNA</u> | | | | | | |
| N 022217 | 002 | 8168616 | Jul 03, 2026 | DP | | |
| <u>ALLOPURINOL; LESINURAD - DUZALLO</u> | | | | | | |
| N 209203 | 001 | 8003681 | Aug 25, 2025 | DS | | |
| | | 8084483 | Aug 17, 2029 | | U-2104 | |
| | | 8283369 | Nov 26, 2028 | | U-2104 | |
| | | 8357713 | Nov 26, 2028 | DP | U-2104 | |
| | | 8546436 | Feb 29, 2032 | DS | | |
| | | 8546437 | Apr 29, 2029 | | U-2104 | |
| | | 9216179 | Aug 01, 2031 | | U-2104 | |
| | | 9956205 | Dec 28, 2031 | | U-2104 | |
| <u>ALLOPURINOL; LESINURAD - DUZALLO</u> | | | | | | |
| N 209203 | 002 | 8003681 | Aug 25, 2025 | DS | | |
| | | 8084483 | Aug 17, 2029 | | U-2104 | |
| | | 8283369 | Nov 26, 2028 | | U-2104 | |
| | | 8357713 | Nov 26, 2028 | DP | U-2104 | |
| | | 8546436 | Feb 29, 2032 | DS | | |
| | | 8546437 | Apr 29, 2029 | | U-2104 | |
| | | 9216179 | Aug 01, 2031 | | U-2104 | |
| | | 9956205 | Dec 28, 2031 | | U-2104 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ALLOPURINOL; LESINURAD - DUZALLO</u> | | | | | | |
| N 209203 | 002 | 8003681 | Aug 25, 2025 | DS | | |
| | | 8084483 | Aug 17, 2029 | | U-2104 | |
| | | 8283369 | Nov 26, 2028 | | U-2104 | |
| | | 8357713 | Nov 26, 2028 | | DP U-2104 | |
| | | 8546436 | Feb 29, 2032 | DS | | |
| | | 8546437 | Apr 29, 2029 | | U-2104 | |
| | | 9216179 | Aug 01, 2031 | | U-2104 | |
| | | 9956205 | Dec 28, 2031 | | U-2104 | |
| <u>ALOGLIPTIN BENZOATE - NESINA</u> | | | | | | |
| N 022271 | 001 | 6890898 | Feb 02, 2019 | | U-1335 | M-177 Apr 05, 2019 |
| | | 7078381 | Feb 02, 2019 | | U-1335 | |
| | | 7459428 | Feb 02, 2019 | | U-1336 | |
| | | 7807689 | Jun 27, 2028 | DS DP | U-1337 | |
| | | 8173663 | Dec 02, 2025 | | U-1338 | |
| | | 8288539 | Mar 15, 2025 | DS | | |
| | | 8697125 | Jun 16, 2029 | | DP | |
| <u>ALOGLIPTIN BENZOATE - NESINA</u> | | | | | | |
| N 022271 | 002 | 6890898 | Feb 02, 2019 | | U-1335 | M-177 Apr 05, 2019 |
| | | 7078381 | Feb 02, 2019 | | U-1335 | |
| | | 7459428 | Feb 02, 2019 | | U-1336 | |
| | | 7807689 | Jun 27, 2028 | DS DP | U-1337 | |
| | | 8173663 | Dec 02, 2025 | | U-1338 | |
| | | 8288539 | Mar 15, 2025 | DS | | |
| | | 8697125 | Jun 16, 2029 | | DP | |
| <u>ALOGLIPTIN BENZOATE - NESINA</u> | | | | | | |
| N 022271 | 003 | 6890898 | Feb 02, 2019 | | U-1335 | M-177 Apr 05, 2019 |
| | | 7078381 | Feb 02, 2019 | | U-1335 | |
| | | 7459428 | Feb 02, 2019 | | U-1336 | |
| | | 7807689 | Jun 27, 2028 | DS DP | U-1337 | |
| | | 8173663 | Dec 02, 2025 | | U-1338 | |
| | | 8288539 | Mar 15, 2025 | DS | | |
| | | 8697125 | Jun 16, 2029 | | DP | |
| <u>ALOGLIPTIN BENZOATE; METFORMIN HYDROCHLORIDE - KAZANO</u> | | | | | | |
| N 203414 | 001 | 6890898 | Feb 02, 2019 | | U-1335 | M-177 Apr 05, 2019 |
| | | 7078381 | Feb 02, 2019 | | U-1335 | |
| | | 7459428 | Feb 02, 2019 | | U-1336 | |
| | | 7807689 | Jun 27, 2028 | DS DP | U-1337 | |
| | | 8173663 | Mar 15, 2025 | | U-1338 | |
| | | 8288539 | Jun 24, 2025 | DS | | |
| | | 8900638 | May 24, 2029 | | DP | |
| <u>ALOGLIPTIN BENZOATE; METFORMIN HYDROCHLORIDE - KAZANO</u> | | | | | | |
| N 203414 | 002 | 6890898 | Feb 02, 2019 | | U-1335 | M-177 Apr 05, 2019 |
| | | 7078381 | Feb 02, 2019 | | U-1335 | |
| | | 7459428 | Feb 02, 2019 | | U-1336 | |
| | | 7807689 | Jun 27, 2028 | DS DP | U-1337 | |
| | | 8173663 | Mar 15, 2025 | | U-1338 | |
| | | 8288539 | Jun 24, 2025 | DS | | |
| | | 8900638 | May 24, 2029 | | DP | |
| <u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u> | | | | | | |
| N 022426 | 001 | 6329404 | Jun 19, 2021 | DP | U-1334 | M-177 Apr 05, 2019 |
| | | 6890898 | Feb 02, 2019 | | U-1335 | |
| | | 7078381 | Feb 02, 2019 | | U-1335 | |
| | | 7459428 | Feb 02, 2019 | | U-1336 | |
| | | 7807689 | Jun 27, 2028 | DS DP | U-1337 | |
| | | 8173663 | Mar 15, 2025 | | U-1338 | |
| | | 8288539 | Mar 15, 2025 | DS | | |
| | | 8637079 | Jun 04, 2029 | | DP | |
| <u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u> | | | | | | |
| N 022426 | 002 | 6329404 | Jun 19, 2021 | DP | U-1334 | M-177 Apr 05, 2019 |
| | | 6890898 | Feb 02, 2019 | | U-1335 | |
| | | 7078381 | Feb 02, 2019 | | U-1335 | |
| | | 7459428 | Feb 02, 2019 | | U-1336 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u> | | | | | | |
| N 022426 002 | 7807689 | Jun 27, 2028 | DS DP U-1337 | | | |
| | 8173663 | Mar 15, 2025 | U-1338 | | | |
| | 8288539 | Mar 15, 2025 | DS | | | |
| | 8637079 | Jun 04, 2029 | DP | | | |
| <u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u> | | | | | | |
| N 022426 003 | 6329404 | Jun 19, 2021 | DP U-1334 | | M-177 | Apr 05, 2019 |
| | 6890898 | Feb 02, 2019 | U-1335 | | | |
| | 7078381 | Feb 02, 2019 | U-1335 | | | |
| | 7459428 | Feb 02, 2019 | U-1336 | | | |
| | 7807689 | Jun 27, 2028 | DS DP U-1337 | | | |
| | 8173663 | Mar 15, 2025 | U-1338 | | | |
| | 8288539 | Mar 15, 2025 | DS | | | |
| | 8637079 | Jun 04, 2029 | DP | | | |
| <u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u> | | | | | | |
| N 022426 004 | 6329404 | Jun 19, 2021 | DP U-1334 | | M-177 | Apr 05, 2019 |
| | 6890898 | Feb 02, 2019 | U-1335 | | | |
| | 7078381 | Feb 02, 2019 | U-1335 | | | |
| | 7459428 | Feb 02, 2019 | U-1336 | | | |
| | 7807689 | Jun 27, 2028 | DS DP U-1337 | | | |
| | 8173663 | Mar 15, 2025 | U-1338 | | | |
| | 8288539 | Mar 15, 2025 | DS | | | |
| | 8637079 | Jun 04, 2029 | DP | | | |
| <u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u> | | | | | | |
| N 022426 005 | 6329404 | Jun 19, 2021 | DP U-1334 | | M-177 | Apr 05, 2019 |
| | 6890898 | Feb 02, 2019 | U-1335 | | | |
| | 7078381 | Feb 02, 2019 | U-1335 | | | |
| | 7459428 | Feb 02, 2019 | U-1336 | | | |
| | 7807689 | Jun 27, 2028 | DS DP U-1337 | | | |
| | 8173663 | Mar 15, 2025 | U-1338 | | | |
| | 8288539 | Mar 15, 2025 | DS | | | |
| | 8637079 | Jun 04, 2029 | DP | | | |
| <u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u> | | | | | | |
| N 022426 006 | 6890898 | Feb 02, 2019 | U-1335 | | M-177 | Apr 05, 2019 |
| | 7078381 | Feb 02, 2019 | U-1335 | | | |
| | 7459428 | Feb 02, 2019 | U-1336 | | | |
| | 7807689 | Jun 27, 2028 | DS DP U-1337 | | | |
| | 8173663 | Mar 15, 2025 | U-1338 | | | |
| | 8288539 | Mar 15, 2025 | DS | | | |
| | 8637079 | Jun 04, 2029 | DP | | | |
| <u>ALVIMOPAN - ENTEREG</u> | | | | | | |
| N 021775 001 | 6469030 | Nov 29, 2020 | U-879 | | | |
| | 8112290 | Jul 31, 2030 | U-1443 | Y | | |
| | 8645160 | Jun 18, 2029 | U-1485 | Y | | |
| | 8946262 | Feb 12, 2030 | U-1655 | | | |
| <u>AMANTADINE HYDROCHLORIDE - GOCOVRI</u> | | | | | | |
| N 208944 001 | 10154971 | Dec 04, 2034 | U-2459 | | I-769 | Aug 24, 2020 |
| | 8389578 | Jan 22, 2028 | U-2105 | | ODE-153 | Aug 24, 2024 |
| | 8741343 | Dec 02, 2030 | U-2106 | | | |
| | 8796337 | Nov 23, 2025 | U-2106 | | | |
| | 8889740 | Nov 23, 2025 | DP | | | |
| | 8895614 | Nov 23, 2025 | DP | | | |
| | 8895615 | Nov 23, 2025 | U-2106 | | | |
| | 8895616 | Nov 23, 2025 | U-2106 | | | |
| | 8895617 | Nov 23, 2025 | U-2106 | | | |
| | 8895618 | Nov 23, 2025 | DP | | | |
| | 9867791 | Dec 02, 2030 | U-2106 | | | |
| | 9867792 | Dec 02, 2030 | U-2106 | | | |
| | 9867793 | Dec 02, 2030 | U-2106 | | | |
| | 9877933 | Dec 02, 2030 | U-2224 | | | |
| <u>AMANTADINE HYDROCHLORIDE - GOCOVRI</u> | | | | | | |
| N 208944 002 | 10154971 | Dec 04, 2034 | U-2459 | | I-769 | Aug 24, 2020 |
| | 8389578 | Jan 22, 2028 | U-2105 | | ODE-153 | Aug 24, 2024 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>AMANTADINE HYDROCHLORIDE - GOCOVRI</u> | | | | | | |
| N 208944 | 002 | 8741343 | Dec 02, 2030 | U-2106 | | |
| | | 8796337 | Nov 23, 2025 | U-2106 | | |
| | | 8889740 | Nov 23, 2025 | DP | | |
| | | 8895614 | Nov 23, 2025 | DP | | |
| | | 8895615 | Nov 23, 2025 | U-2106 | | |
| | | 8895616 | Nov 23, 2025 | U-2106 | | |
| | | 8895617 | Nov 23, 2025 | U-2106 | | |
| | | 8895618 | Nov 23, 2025 | DP | | |
| | | 9867791 | Dec 02, 2030 | U-2106 | | |
| | | 9867792 | Dec 02, 2030 | U-2106 | | |
| | | 9867793 | Dec 02, 2030 | U-2106 | | |
| | | 9877933 | Dec 02, 2030 | U-2224 | | |
| <u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u> | | | | | | |
| N 209410 | 001 | 8252331 | Mar 13, 2030 | DP | | |
| | | 8574626 | Nov 28, 2025 | DP U-20 | | |
| <u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u> | | | | | | |
| N 209410 | 002 | 8252331 | Mar 13, 2030 | DP | | |
| | | 8574626 | Nov 28, 2025 | DP U-20 | | |
| <u>AMANTADINE HYDROCHLORIDE - OSMOLEX ER</u> | | | | | | |
| N 209410 | 003 | 8252331 | Mar 13, 2030 | DP | | |
| | | 8574626 | Nov 28, 2025 | DP U-20 | | |
| <u>AMBRISENTAN - LETAIRIS</u> | | | | | | |
| N 022081 | 001 | 8377933 | Dec 11, 2027 | U-1754 | | |
| | | 9474752 | Dec 11, 2027 | U-1754 | | |
| | | 9549926 | Oct 14, 2031 | U-1965 | | |
| <u>AMBRISENTAN - LETAIRIS</u> | | | | | | |
| N 022081 | 002 | 8377933 | Dec 11, 2027 | U-1754 | | |
| | | 9474752 | Dec 11, 2027 | U-1754 | | |
| | | 9549926 | Oct 14, 2031 | U-1965 | | |
| <u>AMIFAMPRIDINE PHOSPHATE - FIRDAPSE</u> | | | | | | |
| N 208078 | 001 | | | | NCE ODE-223 | Nov 28, 2023 Nov 28, 2025 |
| <u>AMIKACIN SULFATE - ARIKAYCE KIT</u> | | | | | | |
| N 207356 | 001 | 7718189 | Jun 06, 2025 | DP U-2415 | ODE-214 | Sep 28, 2025 |
| | | 8226975 | Aug 15, 2028 | DP | | |
| | | 8632804 | Dec 05, 2026 | U-2416 | | |
| | | 8642075 | Dec 05, 2026 | DP | | |
| | | 8679532 | Dec 05, 2026 | U-2415 | | |
| | | 8802137 | Apr 08, 2024 | DP U-2414 | | |
| | | 9566234 | Jan 18, 2034 | DP U-2415 | | |
| | | 9827317 | Apr 08, 2024 | DP U-2415 | | |
| | | 9895385 | May 15, 2035 | U-2417 | | |
| <u>AMINOLEVULINIC ACID HYDROCHLORIDE - LEVULAN</u> | | | | | | |
| N 020965 | 001 | 7723910 | Jun 17, 2019 | U-289 | I-766 | Mar 09, 2021 |
| <u>AMINOLEVULINIC ACID HYDROCHLORIDE - AMELUZ</u> | | | | | | |
| N 208081 | 001 | 6559183 | Nov 12, 2019 | DP U-804 | NP | May 10, 2019 |
| <u>AMINOLEVULINIC ACID HYDROCHLORIDE - GLEOLAN</u> | | | | | | |
| N 208630 | 001 | | | | ODE-146 | Jun 06, 2024 |
| <u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u> | | | | | | |
| N 022325 | 001 | 6869939 | May 04, 2022 | DP | | |
| | | 7635773 | Mar 13, 2029 | DP | | |
| <u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u> | | | | | | |
| N 022325 | 002 | 6869939 | May 04, 2022 | DP | | |
| | | 7635773 | Mar 13, 2029 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u> | | | | | | |
| N 022325 | 003 6869939 | May 04, 2022 | DP | | | |
| | 7635773 | Mar 13, 2029 | DP | | | |
| <u>AMLODIPINE BESYLATE - AMLODIPINE BESYLATE</u> | | | | | | |
| N 022026 | 001 6828339 | Nov 20, 2022 | DS | | | |
| <u>AMLODIPINE BESYLATE - AMLODIPINE BESYLATE</u> | | | | | | |
| N 022026 | 002 6828339 | Nov 20, 2022 | DS | | | |
| <u>AMLODIPINE BESYLATE - AMLODIPINE BESYLATE</u> | | | | | | |
| N 022026 | 003 6828339 | Nov 20, 2022 | DS | | | |
| <u>AMLODIPINE BESYLATE; CELECOXIB - CONSENSI</u> | | | | | | |
| N 210045 | 001 9662315 | Feb 28, 2030 | DP U-2410 | | NC | May 31, 2021 |
| <u>AMLODIPINE BESYLATE; CELECOXIB - CONSENSI</u> | | | | | | |
| N 210045 | 002 9662315 | Feb 28, 2030 | DP U-2410 | | NC | May 31, 2021 |
| <u>AMLODIPINE BESYLATE; CELECOXIB - CONSENSI</u> | | | | | | |
| N 210045 | 003 9662315 | Feb 28, 2030 | DP U-2410 | | NC | May 31, 2021 |
| <u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u> | | | | | | |
| N 022314 | 001 8101599 | May 16, 2023 | DP | | | |
| | 8475839 | May 16, 2023 | DP | | | |
| | 8475839*PED | Nov 16, 2023 | | | | |
| <u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u> | | | | | | |
| N 022314 | 002 8101599 | May 16, 2023 | DP | | | |
| | 8475839 | May 16, 2023 | DP | | | |
| | 8475839*PED | Nov 16, 2023 | | | | |
| <u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u> | | | | | | |
| N 022314 | 003 8101599 | May 16, 2023 | DP | | | |
| | 8475839 | May 16, 2023 | DP | | | |
| | 8475839*PED | Nov 16, 2023 | | | | |
| <u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u> | | | | | | |
| N 022314 | 004 8101599 | May 16, 2023 | DP | | | |
| | 8475839 | May 16, 2023 | DP | | | |
| | 8475839*PED | Nov 16, 2023 | | | | |
| <u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u> | | | | | | |
| N 022314 | 005 8101599 | May 16, 2023 | DP | | | |
| | 8475839 | May 16, 2023 | DP | | | |
| | 8475839*PED | Nov 16, 2023 | | | | |
| <u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u> | | | | | | |
| N 205003 | 001 6696481 | Apr 15, 2023 | DS DP U-3 | | | |
| | 7846961 | Oct 05, 2029 | DS DP U-3 | | | |
| <u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u> | | | | | | |
| N 205003 | 002 6696481 | Apr 15, 2023 | DS DP U-3 | | | |
| | 7846961 | Oct 05, 2029 | DS DP U-3 | | | |
| <u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u> | | | | | | |
| N 205003 | 003 6696481 | Apr 15, 2023 | DS DP U-3 | | | |
| | 7846961 | Oct 05, 2029 | DS DP U-3 | | | |
| <u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u> | | | | | | |
| N 021990 | 002 6395728 | Jul 08, 2019 | DP | | | |
| <u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u> | | | | | | |
| N 021990 | 003 6395728 | Jul 08, 2019 | DP | | | |
| <u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u> | | | | | | |
| N 021990 | 004 6395728 | Jul 08, 2019 | DP | | | |
| <u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u> | | | | | | |
| N 021990 | 005 6395728 | Jul 08, 2019 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>AMOXICILLIN - MOXATAG</u> | | | | | | |
| N 050813 | 001 | 6544555 | Oct 13, 2020 | DS DP U-897 | | |
| | | 6669948 | Oct 13, 2020 | DS DP U-897 | | |
| | | 6723341 | Oct 13, 2020 | DS DP U-897 | | |
| | | 8299052 | May 07, 2027 | | U-1304 | |
| | | 8357394 | Dec 08, 2026 | DP | | |
| | | 8778924 | Dec 08, 2026 | DS DP U-897 | | |
| <u>AMOXICILLIN; CLAVULANATE POTASSIUM - AUGMENTIN XR</u> | | | | | | |
| N 050785 | 001 | 6746692 | Apr 04, 2020 | DP | | |
| | | 6783773 | Apr 04, 2020 | DP | | |
| | | 6878386 | Apr 04, 2020 | | U-926 | |
| | | 7217430 | Apr 04, 2020 | DP | U-926 | |
| | | 7250176 | Apr 04, 2020 | | U-926 | |
| <u>AMPHETAMINE - ADZENYS ER</u> | | | | | | |
| N 204325 | 001 | 8709491 | Jun 28, 2032 | DP | | |
| | | 9017731 | Jun 28, 2032 | DP | | |
| | | 9265737 | Jun 28, 2032 | DP | | |
| <u>AMPHETAMINE - ADZENYS XR-ODT</u> | | | | | | |
| N 204326 | 001 | 8709491 | Jun 28, 2032 | DP | | |
| | | 8840924 | Apr 09, 2026 | DP | | |
| | | 9017731 | Jun 28, 2032 | DP | | |
| | | 9265737 | Jun 28, 2032 | DP | | |
| <u>AMPHETAMINE - ADZENYS XR-ODT</u> | | | | | | |
| N 204326 | 002 | 8709491 | Jun 28, 2032 | DP | | |
| | | 8840924 | Apr 09, 2026 | DP | | |
| | | 9017731 | Jun 28, 2032 | DP | | |
| | | 9265737 | Jun 28, 2032 | DP | | |
| <u>AMPHETAMINE - ADZENYS XR-ODT</u> | | | | | | |
| N 204326 | 003 | 8709491 | Jun 28, 2032 | DP | | |
| | | 8840924 | Apr 09, 2026 | DP | | |
| | | 9017731 | Jun 28, 2032 | DP | | |
| | | 9265737 | Jun 28, 2032 | DP | | |
| <u>AMPHETAMINE - ADZENYS XR-ODT</u> | | | | | | |
| N 204326 | 004 | 8709491 | Jun 28, 2032 | DP | | |
| | | 8840924 | Apr 09, 2026 | DP | | |
| | | 9017731 | Jun 28, 2032 | DP | | |
| | | 9265737 | Jun 28, 2032 | DP | | |
| <u>AMPHETAMINE - ADZENYS XR-ODT</u> | | | | | | |
| N 204326 | 005 | 8709491 | Jun 28, 2032 | DP | | |
| | | 8840924 | Apr 09, 2026 | DP | | |
| | | 9017731 | Jun 28, 2032 | DP | | |
| | | 9265737 | Jun 28, 2032 | DP | | |
| <u>AMPHETAMINE - ADZENYS XR-ODT</u> | | | | | | |
| N 204326 | 006 | 8709491 | Jun 28, 2032 | DP | | |
| | | 8840924 | Apr 09, 2026 | DP | | |
| | | 9017731 | Jun 28, 2032 | DP | | |
| | | 9265737 | Jun 28, 2032 | DP | | |
| <u>AMPHETAMINE - DYANA VEL XR</u> | | | | | | |
| N 208147 | 001 | 10086087 | Mar 15, 2027 | DP | | |
| | | 8062667 | Mar 29, 2029 | DP | | |
| | | 8597684 | Mar 15, 2027 | DP | | |
| | | 8747902 | Mar 15, 2027 | DP | | |
| | | 8883217 | Mar 15, 2027 | DP | | |
| | | 9675703 | Mar 15, 2027 | DP | | |
| <u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 10</u> | | | | | | |
| N 011522 | 007 | 6384020 | Jul 06, 2020 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 20</u> | | | | | | |
| N 011522 | 008 | 6384020 | | | | |
| <u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 5</u> | | | | | | |
| N 011522 | 009 | 6384020 | | | | |
| <u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 30</u> | | | | | | |
| N 011522 | 010 | 6384020 | | | | |
| <u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 7.5</u> | | | | | | |
| N 011522 | 011 | 6384020 | | | | |
| <u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 12.5</u> | | | | | | |
| N 011522 | 012 | 6384020 | | | | |
| <u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 15</u> | | | | | | |
| N 011522 | 013 | 6384020 | | | | |
| <u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u> | | | | | | |
| N 022063 | 001 | 6913768 | | | | |
| | | 8846100 | | DP U-2025 | NP | Jun 20, 2020 |
| | | 9173857 | | DP | | |
| | | | | U-2025 | | |
| <u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u> | | | | | | |
| N 022063 | 002 | 6913768 | | | | |
| | | 8846100 | | DP U-2025 | NP | Jun 20, 2020 |
| | | 9173857 | | DP | | |
| | | | | U-2025 | | |
| <u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u> | | | | | | |
| N 022063 | 003 | 6913768 | | | | |
| | | 8846100 | | DP U-2025 | NP | Jun 20, 2020 |
| | | 9173857 | | DP | | |
| | | | | U-2025 | | |
| <u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u> | | | | | | |
| N 022063 | 004 | 6913768 | | | | |
| | | 8846100 | | DP U-2025 | NP | Jun 20, 2020 |
| | | 9173857 | | DP | | |
| | | | | U-2025 | | |
| <u>AMPHETAMINE SULFATE - AMPHETAMINE SULFATE</u> | | | | | | |
| A 211139 | 001 | | | | CGT | Mar 26, 2019 |
| <u>AMPHETAMINE SULFATE - AMPHETAMINE SULFATE</u> | | | | | | |
| A 211139 | 002 | | | | CGT | Mar 30, 2019 |
| <u>AMPHOTERICIN B - ABELCET</u> | | | | | | |
| N 050724 | 001 | 6406713 | | DS | | |
| <u>ANGIOTENSIN II ACETATE - GIAPREZA</u> | | | | | | |
| N 209360 | 001 | 10028995 | | | | |
| | | 9220745 | | U-2338 | NCE | Dec 21, 2022 |
| | | 9220745 | | U-2217 | | |
| | | 9220745 | | U-2218 | | |
| | | 9572856 | | U-2221 | | |
| | | 9867863 | | U-2231 | | |
| <u>ANGIOTENSIN II ACETATE - GIAPREZA</u> | | | | | | |
| N 209360 | 002 | 10028995 | | | | |
| | | 9220745 | | U-2338 | NCE | Dec 21, 2022 |
| | | 9220745 | | U-2217 | | |
| | | 9220745 | | U-2218 | | |
| | | 9572856 | | U-2221 | | |
| | | 9867863 | | U-2231 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------|----------------------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ANIDULAFUNGIN - ERAXIS</u> | | | | | | |
| N 021632 001 | 5965525 | Feb 17, 2020 | DS DP U-540 | | | |
| | 6960564 | Apr 12, 2021 | DP U-540 | | | |
| | 7709444 | Apr 12, 2021 | DP U-540 | | | |
| <u>ANIDULAFUNGIN - ERAXIS</u> | | | | | | |
| N 021632 002 | 5965525 | Feb 17, 2020 | DS DP U-540 | | | |
| | 6960564 | Apr 12, 2021 | DP U-540 | | | |
| | 7709444 | Apr 12, 2021 | DP U-540 | | | |
| <u>APALUTAMIDE - ERLEADA</u> | | | | | | |
| N 210951 001 | 10052314 | Sep 23, 2033 | | U-2381 | NCE | Feb 14, 2023 |
| | 10052314 | Sep 23, 2033 | | U-2382 | | |
| | 8445507 | Sep 15, 2030 | DS DP | U-2237 | | |
| | 8802689 | Mar 27, 2027 | | U-2237 | | |
| | 9388159 | Mar 27, 2027 | DS DP | | | |
| | 9481663 | Jun 04, 2033 | DS DP | U-2237 | | |
| | 9884054 | Sep 23, 2033 | | U-2237 | | |
| | 9987261 | Mar 27, 2027 | DP | | | |
| <u>APIXABAN - ELIQUIS</u> | | | | | | |
| N 202155 001 | 6413980 | Dec 22, 2019 | DS DP | U-1200 | | |
| | 6413980 | Dec 22, 2019 | DS DP | U-1301 | | |
| | 6413980 | Dec 22, 2019 | DS DP | U-1302 | | |
| | 6413980 | Dec 22, 2019 | DS DP | U-1501 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1167 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1200 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1301 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1302 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1323 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1501 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1502 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1729 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1730 | | |
| | 9326945 | Feb 24, 2031 | DP | | | |
| | <u>APIXABAN - ELIQUIS</u> | | | | | |
| N 202155 002 | 6413980 | Dec 22, 2019 | DS DP | U-1200 | | |
| | 6413980 | Dec 22, 2019 | DS DP | U-1301 | | |
| | 6413980 | Dec 22, 2019 | DS DP | U-1302 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1200 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1301 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1302 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1323 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1502 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1729 | | |
| | 6967208 | Nov 21, 2026 | DS DP | U-1730 | | |
| 9326945 | Feb 24, 2031 | DP | | | | |
| <u>APREMILAST - OTEZLA</u> | | | | | | |
| N 205437 001 | 10092541 | May 29, 2034 | | U-2403 | NCE | Mar 21, 2019 |
| | 6962940 | Mar 19, 2023 | | U-1504 | | |
| | 7208516 | Mar 19, 2023 | | U-1505 | | |
| | 7427638 | Feb 16, 2028 | DS DP | | | |
| | 7659302 | Mar 19, 2023 | | U-1505 | | |
| | 7659302 | Mar 19, 2023 | | U-1595 | | |
| | 7893101 | Dec 09, 2023 | DS DP | | | |
| | 8455536 | Mar 19, 2023 | | U-1505 | | |
| | 8455536 | Mar 19, 2023 | | U-1595 | | |
| | 8802717 | Mar 19, 2023 | | U-1561 | | |
| | 9018243 | Mar 19, 2023 | | U-1505 | | |
| | 9018243 | Mar 19, 2023 | | U-1595 | | |
| | 9724330 | Mar 19, 2023 | | U-1561 | | |
| | 9724330 | Mar 19, 2023 | | U-1595 | | |
| | 9872854 | May 29, 2034 | | U-2232 | | |
| | 9872854 | May 29, 2034 | | U-2233 | | |
| | <u>APREMILAST - OTEZLA</u> | | | | | |
| N 205437 002 | 10092541 | May 29, 2034 | | U-2403 | NCE | Mar 21, 2019 |
| | 6962940 | Mar 19, 2023 | | U-1504 | | |
| | 7208516 | Mar 19, 2023 | | U-1505 | | |
| | 7427638 | Feb 16, 2028 | DS DP | | | |
| | 7659302 | Mar 19, 2023 | | U-1505 | | |
| | 7659302 | Mar 19, 2023 | | U-1595 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>APREMILAST - OTEZLA</u> | | | | | | |
| N 205437 | 002 | 7659302 | | | | |
| | | 7893101 | | DS DP | | |
| | | 8455536 | | | U-1505 | |
| | | 8455536 | | | U-1595 | |
| | | 8802717 | | | U-1561 | |
| | | 9018243 | | | U-1505 | |
| | | 9018243 | | | U-1595 | |
| | | 9724330 | | | U-1561 | |
| | | 9724330 | | | U-1595 | |
| | | 9872854 | | | U-2232 | |
| | | 9872854 | | | U-2233 | |
| <u>APREMILAST - OTEZLA</u> | | | | | | |
| N 205437 | 003 | 10092541 | | | | |
| | | 6962940 | | | | |
| | | 7208516 | | | | |
| | | 7427638 | | DS DP | | |
| | | 7659302 | | | U-1505 | |
| | | 7659302 | | | U-1595 | |
| | | 7893101 | | DS DP | | |
| | | 8455536 | | | U-1505 | |
| | | 8455536 | | | U-1595 | |
| | | 8802717 | | | U-1561 | |
| | | 9018243 | | | U-1505 | |
| | | 9018243 | | | U-1595 | |
| | | 9724330 | | | U-1561 | |
| | | 9724330 | | | U-1595 | |
| | | 9872854 | | | U-2232 | |
| | | 9872854 | | | U-2233 | |
| <u>APREPITANT - EMEND</u> | | | | | | |
| N 021549 | 001 | 8258132 | | | | |
| | | 8258132 | | DP | U-1743 | |
| | | | | DP | U-901 | |
| <u>APREPITANT - EMEND</u> | | | | | | |
| N 021549 | 002 | 8258132 | | | | |
| | | 8258132 | | DP | U-1743 | |
| | | | | DP | U-901 | |
| <u>APREPITANT - EMEND</u> | | | | | | |
| N 021549 | 003 | 8258132 | | | | |
| | | 8258132 | | DP | U-1743 | |
| | | | | DP | U-901 | |
| <u>APREPITANT - EMEND</u> | | | | | | |
| N 207865 | 001 | 8258132 | | | | |
| | | | | DP | U-1916 | |
| <u>APREPITANT - CINVANTI</u> | | | | | | |
| N 209296 | 001 | 9561229 | | | | |
| | | 9808465 | | | | |
| | | 9974742 | | | | |
| | | 9974793 | | | | |
| | | 9974794 | | | | |
| | | | | DP | U-2161 | |
| | | | | | U-2161 | |
| | | | | DP | | |
| | | | | DP | | |
| | | | | DP | U-2161 | |
| <u>ARFORMOTEROL TARTRATE - BROVANA</u> | | | | | | |
| N 021912 | 001 | 6472563 | | | | |
| | | 6667344 | | DS | | |
| | | 6720453 | | | | |
| | | 6814953 | | DS | | |
| | | 7145036 | | | U-793 | |
| | | 7145036 | | DS | | |
| | | 7348362 | | | U-793 | |
| | | 7462645 | | | U-793 | |
| | | 7465756 | | | | |
| | | 7465756 | | DP | | |
| | | 7473710 | | | U-793 | |
| | | 7541385 | | | U-793 | |
| | | 8110706 | | | | |
| | | | | DP | | |
| <u>ARGATROBAN - ARGATROBAN IN SODIUM CHLORIDE</u> | | | | | | |
| N 022434 | 001 | 7589106 | | | | |
| | | 7687516 | | | | |
| | | | | DP | U-1163 | |
| | | | | DP | U-1164 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ARIPIRAZOLE - ABILIFY</u> | | | | | | |
| N 021436 001 | 7053092 | Jan 28, 2022 | | U-839 | ODE-80 | Dec 12, 2021 |
| | 8017615 | Jun 16, 2024 | | DP | | |
| | 8017615*PED | Dec 16, 2024 | | | | |
| | 8580796 | Sep 25, 2022 | DS | | | |
| | 8580796*PED | Mar 25, 2023 | | | | |
| | 8642600 | Jan 28, 2022 | | U-1492 | | |
| | 8642600*PED | Jul 28, 2022 | | | | |
| | 8642760 | Sep 25, 2022 | DS | | | |
| | 8642760*PED | Mar 25, 2023 | | | | |
| | 8759350 | Mar 02, 2027 | | U-1529 | | |
| | 9089567 | Jan 28, 2022 | | U-543 | | |
| | 9125939 | Jul 28, 2026 | | U-1749 | | |
| | 9359302 | Sep 25, 2022 | DS DP | U-1859 | | |
| | 9387182 | Dec 25, 2023 | | U-1529 | | |
| <u>ARIPIRAZOLE - ABILIFY</u> | | | | | | |
| N 021436 002 | 7053092 | Jan 28, 2022 | | U-839 | ODE-80 | Dec 12, 2021 |
| | 8017615 | Jun 16, 2024 | | DP | | |
| | 8017615*PED | Dec 16, 2024 | | | | |
| | 8580796 | Sep 25, 2022 | DS | | | |
| | 8580796*PED | Mar 25, 2023 | | | | |
| | 8642600 | Jan 28, 2022 | | U-1492 | | |
| | 8642600*PED | Jul 28, 2022 | | | | |
| | 8642760 | Sep 25, 2022 | DS | | | |
| | 8642760*PED | Mar 25, 2023 | | | | |
| | 8759350 | Mar 02, 2027 | | U-1529 | | |
| | 9089567 | Jan 28, 2022 | | U-543 | | |
| | 9125939 | Jul 28, 2026 | | U-1749 | | |
| | 9359302 | Sep 25, 2022 | DS DP | U-1859 | | |
| | 9387182 | Dec 25, 2023 | | U-1529 | | |
| <u>ARIPIRAZOLE - ABILIFY</u> | | | | | | |
| N 021436 003 | 7053092 | Jan 28, 2022 | | U-839 | ODE-80 | Dec 12, 2021 |
| | 8017615 | Jun 16, 2024 | | DP | | |
| | 8017615*PED | Dec 16, 2024 | | | | |
| | 8580796 | Sep 25, 2022 | DS | | | |
| | 8580796*PED | Mar 25, 2023 | | | | |
| | 8642600 | Jan 28, 2022 | | U-1492 | | |
| | 8642600*PED | Jul 28, 2022 | | | | |
| | 8642760 | Sep 25, 2022 | DS | | | |
| | 8642760*PED | Mar 25, 2023 | | | | |
| | 8759350 | Mar 02, 2027 | | U-1529 | | |
| | 9089567 | Jan 28, 2022 | | U-543 | | |
| | 9125939 | Jul 28, 2026 | | U-1749 | | |
| | 9359302 | Sep 25, 2022 | DS DP | U-1859 | | |
| | 9387182 | Dec 25, 2023 | | U-1529 | | |
| <u>ARIPIRAZOLE - ABILIFY</u> | | | | | | |
| N 021436 004 | 7053092 | Jan 28, 2022 | | U-839 | ODE-80 | Dec 12, 2021 |
| | 8017615 | Jun 16, 2024 | | DP | | |
| | 8017615*PED | Dec 16, 2024 | | | | |
| | 8580796 | Sep 25, 2022 | DS | | | |
| | 8580796*PED | Mar 25, 2023 | | | | |
| | 8642600 | Jan 28, 2022 | | U-1492 | | |
| | 8642600*PED | Jul 28, 2022 | | | | |
| | 8642760 | Sep 25, 2022 | DS | | | |
| | 8642760*PED | Mar 25, 2023 | | | | |
| | 8759350 | Mar 02, 2027 | | U-1529 | | |
| | 9089567 | Jan 28, 2022 | | U-543 | | |
| | 9125939 | Jul 28, 2026 | | U-1749 | | |
| | 9359302 | Sep 25, 2022 | DS DP | U-1859 | | |
| | 9387182 | Dec 25, 2023 | | U-1529 | | |
| <u>ARIPIRAZOLE - ABILIFY</u> | | | | | | |
| N 021436 005 | 7053092 | Jan 28, 2022 | | U-839 | ODE-80 | Dec 12, 2021 |
| | 8017615 | Jun 16, 2024 | | DP | | |
| | 8017615*PED | Dec 16, 2024 | | | | |
| | 8580796 | Sep 25, 2022 | DS | | | |
| | 8580796*PED | Mar 25, 2023 | | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ARIPIRAZOLE - ABILIFY</u> | | | | | | |
| N 021436 005 | 8642600 | Jan 28, 2022 | | U-1492 | | |
| | 8642600*PED | Jul 28, 2022 | | | | |
| | 8642760 | Sep 25, 2022 | DS | | | |
| | 8642760*PED | Mar 25, 2023 | | | | |
| | 8759350 | Mar 02, 2027 | | U-1529 | | |
| | 9089567 | Jan 28, 2022 | | U-543 | | |
| | 9125939 | Jul 28, 2026 | | U-1749 | | |
| | 9359302 | Sep 25, 2022 | DS DP | U-1859 | | |
| | 9387182 | Dec 25, 2023 | | U-1529 | | |
| <u>ARIPIRAZOLE - ABILIFY</u> | | | | | | |
| N 021436 006 | 7053092 | Jan 28, 2022 | | U-839 | ODE-80 | Dec 12, 2021 |
| | 8017615 | Jun 16, 2024 | DP | | | |
| | 8017615*PED | Dec 16, 2024 | | | | |
| | 8580796 | Sep 25, 2022 | DS | | | |
| | 8580796*PED | Mar 25, 2023 | | | | |
| | 8642600 | Jan 28, 2022 | | U-1492 | | |
| | 8642600*PED | Jul 28, 2022 | | | | |
| | 8642760 | Sep 25, 2022 | DS | | | |
| | 8642760*PED | Mar 25, 2023 | | | | |
| | 8759350 | Mar 02, 2027 | | U-1529 | | |
| | 9089567 | Jan 28, 2022 | | U-543 | | |
| | 9125939 | Jul 28, 2026 | | U-1749 | | |
| | 9359302 | Sep 25, 2022 | DS DP | U-1859 | | |
| | 9387182 | Dec 25, 2023 | | U-1529 | | |
| <u>ARIPIRAZOLE - ABILIFY</u> | | | | | | |
| N 021713 001 | 6977257 | Apr 24, 2022 | | DP | ODE-80 | Dec 12, 2021 |
| | 6977257*PED | Oct 24, 2022 | | | | |
| | 7053092 | Jan 28, 2022 | | U-839 | | |
| | 8642600 | Jan 28, 2022 | | U-1492 | | |
| | 8642600*PED | Jul 28, 2022 | | | | |
| | 8759350 | Mar 02, 2027 | | U-1529 | | |
| | 9387182 | Dec 25, 2023 | | U-1529 | | |
| <u>ARIPIRAZOLE - ABILIFY</u> | | | | | | |
| N 021729 002 | 7053092 | Jan 28, 2022 | | U-839 | ODE-80 | Dec 12, 2021 |
| | 8017615 | Jun 16, 2024 | DP | | | |
| | 8017615*PED | Dec 16, 2024 | | | | |
| | 8518421 | Jan 24, 2021 | DP | | | |
| | 8518421*PED | Jul 24, 2021 | | | | |
| | 8580796 | Sep 25, 2022 | DS | | | |
| | 8580796*PED | Mar 25, 2023 | | | | |
| | 8642600 | Jan 28, 2022 | | U-1492 | | |
| | 8642600*PED | Jul 28, 2022 | | | | |
| | 8642760 | Sep 25, 2022 | DS | | | |
| | 8642760*PED | Mar 25, 2023 | | | | |
| | 8759350 | Mar 02, 2027 | | U-1529 | | |
| | 9089567 | Jan 28, 2022 | | U-543 | | |
| | 9125939 | Jul 28, 2026 | | U-1749 | | |
| | 9358207 | Apr 12, 2020 | | DP | | |
| | 9359302 | Sep 25, 2022 | DS DP | U-1859 | | |
| | 9387182 | Dec 25, 2023 | | U-1529 | | |
| <u>ARIPIRAZOLE - ABILIFY</u> | | | | | | |
| N 021729 003 | 7053092 | Jan 28, 2022 | | U-839 | ODE-80 | Dec 12, 2021 |
| | 8017615 | Jun 16, 2024 | DP | | | |
| | 8017615*PED | Dec 16, 2024 | | | | |
| | 8518421 | Jan 24, 2021 | DP | | | |
| | 8518421*PED | Jul 24, 2021 | | | | |
| | 8580796 | Sep 25, 2022 | DS | | | |
| | 8580796*PED | Mar 25, 2023 | | | | |
| | 8642600 | Jan 28, 2022 | | U-1492 | | |
| | 8642600*PED | Jul 28, 2022 | | | | |
| | 8642760 | Sep 25, 2022 | DS | | | |
| | 8642760*PED | Mar 25, 2023 | | | | |
| | 8759350 | Mar 02, 2027 | | U-1529 | | |
| | 9089567 | Jan 28, 2022 | | U-543 | | |
| | 9125939 | Jul 28, 2026 | | U-1749 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ARIPIRAZOLE - ABILIFY</u> | | | | | | |
| N 021729 | 003 | 9358207 | Apr 12, 2020 | DP | | |
| | | 9359302 | Sep 25, 2022 | DS DP | U-1859 | |
| | | 9387182 | Dec 25, 2023 | | U-1529 | |
| <u>ARIPIRAZOLE - ABILIFY</u> | | | | | | |
| N 021729 | 004 | 7053092 | Jan 28, 2022 | | U-839 | ODE-80 Dec 12, 2021 |
| | | 8017615 | Jun 16, 2024 | DP | | |
| | | 8017615*PED | Dec 16, 2024 | | | |
| | | 8518421 | Jan 24, 2021 | DP | | |
| | | 8518421*PED | Jul 24, 2021 | | | |
| | | 8580796 | Sep 25, 2022 | DS | | |
| | | 8580796*PED | Mar 25, 2023 | | | |
| | | 8642600 | Jan 28, 2022 | | U-1492 | |
| | | 8642600*PED | Jul 28, 2022 | | | |
| | | 8642760 | Sep 25, 2022 | DS | | |
| | | 8642760*PED | Mar 25, 2023 | | | |
| | | 9358207 | Apr 12, 2020 | DP | | |
| | | 9359302 | Sep 25, 2022 | DS DP | U-1859 | |
| | | 9387182 | Dec 25, 2023 | | U-1529 | |
| <u>ARIPIRAZOLE - ABILIFY</u> | | | | | | |
| N 021729 | 005 | 7053092 | Jan 28, 2022 | | U-839 | ODE-80 Dec 12, 2021 |
| | | 8017615 | Jun 16, 2024 | DP | | |
| | | 8017615*PED | Dec 16, 2024 | | | |
| | | 8518421 | Jan 24, 2021 | DP | | |
| | | 8518421*PED | Jul 24, 2021 | | | |
| | | 8580796 | Sep 25, 2022 | DS | | |
| | | 8580796*PED | Mar 25, 2023 | | | |
| | | 8642600 | Jan 28, 2022 | | U-1492 | |
| | | 8642600*PED | Jul 28, 2022 | | | |
| | | 8642760 | Sep 25, 2022 | DS | | |
| | | 8642760*PED | Mar 25, 2023 | | | |
| | | 9358207 | Apr 12, 2020 | DP | | |
| | | 9359302 | Sep 25, 2022 | DS DP | U-1859 | |
| | | 9387182 | Dec 25, 2023 | | U-1529 | |
| <u>ARIPIRAZOLE - ABILIFY</u> | | | | | | |
| N 021866 | 001 | 7115587 | Jul 21, 2024 | DP | U-764 | ODE-80 Dec 12, 2021 |
| | | 7115587*PED | Jan 21, 2025 | | | |
| | | 7550445 | Jul 21, 2024 | DP | | |
| <u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u> | | | | | | |
| N 202971 | 001 | 7807680 | Oct 19, 2024 | DP | | I-746 Jul 27, 2020 |
| | | 8030313 | Oct 19, 2024 | | U-1632 | |
| | | 8030313 | Oct 19, 2024 | | U-543 | |
| | | 8338427 | Mar 15, 2025 | DP | U-1633 | |
| | | 8338427 | Mar 15, 2025 | DP | U-543 | |
| | | 8338428 | Aug 06, 2023 | DP | U-1633 | |
| | | 8338428 | Aug 06, 2023 | DP | U-543 | |
| | | 8399469 | Jun 29, 2025 | DS | | |
| | | 8722679 | Oct 19, 2024 | DP | | |
| | | 8759351 | Aug 06, 2023 | DP | U-1530 | |
| | | 8759351 | Aug 06, 2023 | DP | U-1633 | |
| | | 8993761 | Sep 25, 2022 | DS | | |
| | | 9089567 | Jan 28, 2022 | | U-543 | |
| <u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u> | | | | | | |
| N 202971 | 002 | 7807680 | Oct 19, 2024 | DP | | I-746 Jul 27, 2020 |
| | | 8030313 | Oct 19, 2024 | | U-1632 | |
| | | 8030313 | Oct 19, 2024 | | U-543 | |
| | | 8338427 | Mar 15, 2025 | DP | U-1633 | |
| | | 8338427 | Mar 15, 2025 | DP | U-543 | |
| | | 8338428 | Aug 06, 2023 | DP | U-1633 | |
| | | 8338428 | Aug 06, 2023 | DP | U-543 | |
| | | 8399469 | Jun 29, 2025 | DS | | |
| | | 8722679 | Oct 19, 2024 | DP | | |
| | | 8759351 | Aug 06, 2023 | DP | U-1530 | |
| | | 8759351 | Aug 06, 2023 | DP | U-1633 | |
| | | 8993761 | Sep 25, 2022 | DS | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u> | | | | | | |
| N 202971 | 002 | 9089567 | Jan 28, 2022 | U-543 | | |
| <u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u> | | | | | | |
| N 202971 | 003 | 7807680 | Oct 19, 2024 | DP | I-746 | Jul 27, 2020 |
| | | 8030313 | Oct 19, 2024 | U-1632 | | |
| | | 8030313 | Oct 19, 2024 | U-543 | | |
| | | 8338427 | Mar 15, 2025 | DP U-1633 | | |
| | | 8338427 | Mar 15, 2025 | DP U-543 | | |
| | | 8338428 | Aug 06, 2023 | DP U-1633 | | |
| | | 8338428 | Aug 06, 2023 | DP U-543 | | |
| | | 8399469 | Jun 29, 2025 | DS | | |
| | | 8722679 | Oct 19, 2024 | DP | | |
| | | 8759351 | Aug 06, 2023 | DP U-1530 | | |
| | | 8759351 | Aug 06, 2023 | DP U-1633 | | |
| | | 8993761 | Sep 25, 2022 | DS | | |
| | | 9089567 | Jan 28, 2022 | U-543 | | |
| <u>ARIPIRAZOLE - ABILIFY MAINTENA KIT</u> | | | | | | |
| N 202971 | 004 | 7807680 | Oct 19, 2024 | DP | I-746 | Jul 27, 2020 |
| | | 8030313 | Oct 19, 2024 | U-1632 | | |
| | | 8030313 | Oct 19, 2024 | U-543 | | |
| | | 8338427 | Mar 15, 2025 | DP U-1633 | | |
| | | 8338427 | Mar 15, 2025 | DP U-543 | | |
| | | 8338428 | Aug 06, 2023 | DP U-1633 | | |
| | | 8338428 | Aug 06, 2023 | DP U-543 | | |
| | | 8399469 | Jun 29, 2025 | DS | | |
| | | 8722679 | Oct 19, 2024 | DP | | |
| | | 8759351 | Aug 06, 2023 | DP U-1530 | | |
| | | 8759351 | Aug 06, 2023 | DP U-1633 | | |
| | | 8993761 | Sep 25, 2022 | DS | | |
| | | 9089567 | Jan 28, 2022 | U-543 | | |
| <u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u> | | | | | | |
| N 207202 | 001 | 10097388 | Sep 19, 2034 | U-2169 | I-746 | Jul 27, 2020 |
| | | 7053092 | Jan 28, 2022 | U-1529 | | |
| | | 7978064 | Sep 14, 2026 | DP | | |
| | | 8017615 | Jun 16, 2024 | DP | | |
| | | 8114021 | Nov 02, 2026 | DP | | |
| | | 8258962 | Nov 25, 2030 | DP | | |
| | | 8545402 | Apr 27, 2030 | DP | | |
| | | 8547248 | Dec 18, 2030 | DP U-2167 | | |
| | | 8580796 | Sep 25, 2022 | DS | | |
| | | 8642760 | Sep 25, 2022 | DS | | |
| | | 8674825 | Apr 09, 2029 | DP U-2170 | | |
| | | 8718193 | Dec 05, 2029 | DP | | |
| | | 8759350 | Mar 02, 2027 | U-1529 | | |
| | | 8847766 | Mar 29, 2030 | DP U-2167 | | |
| | | 8945005 | Aug 19, 2029 | DP U-2167 | | |
| | | 8956288 | Jul 06, 2029 | DP U-2167 | | |
| | | 8961412 | Nov 17, 2030 | DP | | |
| | | 9060708 | Mar 05, 2029 | DP | | |
| | | 9089567 | Jan 28, 2022 | U-543 | | |
| | | 9119554 | Dec 16, 2028 | DP | | |
| | | 9125939 | Jul 28, 2026 | U-1749 | | |
| | | 9149577 | Dec 15, 2029 | DP | | |
| | | 9258035 | Mar 05, 2029 | DP | | |
| | | 9268909 | Oct 15, 2033 | DP U-2168 | | |
| | | 9270503 | Sep 19, 2034 | DP U-2169 | | |
| | | 9320455 | Dec 15, 2031 | DP | | |
| | | 9359302 | Sep 25, 2022 | DS DP U-1529 | | |
| | | 9359302 | Sep 25, 2022 | DS DP U-1749 | | |
| | | 9359302 | Sep 25, 2022 | DS DP U-543 | | |
| | | 9387182 | Dec 25, 2023 | U-1529 | | |
| | | 9433371 | Sep 15, 2029 | DP | | |
| | | 9444503 | Nov 19, 2027 | DP U-2169 | | |
| | | 9577864 | Oct 03, 2033 | DP U-2169 | | |
| | | 9787511 | Sep 19, 2034 | DP U-2169 | | |
| | | 9941931 | Nov 04, 2030 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u> | | | | | | |
| N 207202 002 | 10097388 | Sep 19, 2034 | | U-2169 | I-746 | Jul 27, 2020 |
| | 7053092 | Jan 28, 2022 | | U-1529 | | |
| | 7978064 | Sep 14, 2026 | DP | | | |
| | 8017615 | Jun 16, 2024 | DP | | | |
| | 8114021 | Nov 02, 2026 | DP | | | |
| | 8258962 | Nov 25, 2030 | DP | | | |
| | 8545402 | Apr 27, 2030 | DP | | | |
| | 8547248 | Dec 18, 2030 | DP | U-2167 | | |
| | 8580796 | Sep 25, 2022 | DS | | | |
| | 8642760 | Sep 25, 2022 | DS | | | |
| | 8674825 | Apr 09, 2029 | DP | U-2170 | | |
| | 8718193 | Dec 05, 2029 | DP | | | |
| | 8759350 | Mar 02, 2027 | | U-1529 | | |
| | 8847766 | Mar 29, 2030 | DP | U-2167 | | |
| | 8945005 | Aug 19, 2029 | DP | U-2167 | | |
| | 8956288 | Jul 06, 2029 | DP | U-2167 | | |
| | 8961412 | Nov 17, 2030 | DP | | | |
| | 9060708 | Mar 05, 2029 | DP | | | |
| | 9089567 | Jan 28, 2022 | | U-543 | | |
| | 9119554 | Dec 16, 2028 | DP | | | |
| | 9125939 | Jul 28, 2026 | | U-1749 | | |
| | 9149577 | Dec 15, 2029 | DP | | | |
| | 9258035 | Mar 05, 2029 | DP | | | |
| | 9268909 | Oct 15, 2033 | DP | U-2168 | | |
| | 9270503 | Sep 19, 2034 | DP | U-2169 | | |
| | 9320455 | Dec 15, 2031 | DP | | | |
| | 9359302 | Sep 25, 2022 | DS DP | U-1529 | | |
| | 9359302 | Sep 25, 2022 | DS DP | U-1749 | | |
| | 9359302 | Sep 25, 2022 | DS DP | U-543 | | |
| | 9387182 | Dec 25, 2023 | | U-1529 | | |
| | 9433371 | Sep 15, 2029 | DP | | | |
| | 9444503 | Nov 19, 2027 | DP | U-2169 | | |
| | 9577864 | Oct 03, 2033 | DP | U-2169 | | |
| | 9787511 | Sep 19, 2034 | DP | U-2169 | | |
| | 9941931 | Nov 04, 2030 | DP | | | |
| <u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u> | | | | | | |
| N 207202 003 | 10097388 | Sep 19, 2034 | | U-2169 | I-746 | Jul 27, 2020 |
| | 7053092 | Jan 28, 2022 | | U-1529 | | |
| | 7978064 | Sep 14, 2026 | DP | | | |
| | 8017615 | Jun 16, 2024 | DP | | | |
| | 8114021 | Nov 02, 2026 | DP | | | |
| | 8258962 | Nov 25, 2030 | DP | | | |
| | 8545402 | Apr 27, 2030 | DP | | | |
| | 8547248 | Dec 18, 2030 | DP | U-2167 | | |
| | 8580796 | Sep 25, 2022 | DS | | | |
| | 8642760 | Sep 25, 2022 | DS | | | |
| | 8674825 | Apr 09, 2029 | DP | U-2170 | | |
| | 8718193 | Dec 05, 2029 | DP | | | |
| | 8759350 | Mar 02, 2027 | | U-1529 | | |
| | 8847766 | Mar 29, 2030 | DP | U-2167 | | |
| | 8945005 | Aug 19, 2029 | DP | U-2167 | | |
| | 8956288 | Jul 06, 2029 | DP | U-2167 | | |
| | 8961412 | Nov 17, 2030 | DP | | | |
| | 9060708 | Mar 05, 2029 | DP | | | |
| | 9089567 | Jan 28, 2022 | | U-543 | | |
| | 9119554 | Dec 16, 2028 | DP | | | |
| | 9125939 | Jul 28, 2026 | | U-1749 | | |
| | 9149577 | Dec 15, 2029 | DP | | | |
| | 9258035 | Mar 05, 2029 | DP | | | |
| | 9268909 | Oct 15, 2033 | DP | U-2168 | | |
| | 9270503 | Sep 19, 2034 | DP | U-2169 | | |
| | 9320455 | Dec 15, 2031 | DP | | | |
| | 9359302 | Sep 25, 2022 | DS DP | U-1529 | | |
| | 9359302 | Sep 25, 2022 | DS DP | U-1749 | | |
| | 9359302 | Sep 25, 2022 | DS DP | U-543 | | |
| | 9387182 | Dec 25, 2023 | | U-1529 | | |
| | 9433371 | Sep 15, 2029 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u> | | | | | | |
| N 207202 003 | 9444503 | Nov 19, 2027 | DP U-2169 | | | |
| | 9577864 | Oct 03, 2033 | DP U-2169 | | | |
| | 9787511 | Sep 19, 2034 | DP U-2169 | | | |
| | 9941931 | Nov 04, 2030 | DP | | | |
| <u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u> | | | | | | |
| N 207202 004 | 10097388 | Sep 19, 2034 | U-2169 | | I-746 | Jul 27, 2020 |
| | 7053092 | Jan 28, 2022 | U-1529 | | | |
| | 7978064 | Sep 14, 2026 | DP | | | |
| | 8017615 | Jun 16, 2024 | DP | | | |
| | 8114021 | Nov 02, 2026 | DP | | | |
| | 8258962 | Nov 25, 2030 | DP | | | |
| | 8545402 | Apr 27, 2030 | DP | | | |
| | 8547248 | Dec 18, 2030 | DP U-2167 | | | |
| | 8580796 | Sep 25, 2022 | DS | | | |
| | 8642760 | Sep 25, 2022 | DS | | | |
| | 8674825 | Apr 09, 2029 | DP U-2170 | | | |
| | 8718193 | Dec 05, 2029 | DP | | | |
| | 8759350 | Mar 02, 2027 | U-1529 | | | |
| | 8847766 | Mar 29, 2030 | DP U-2167 | | | |
| | 8945005 | Aug 19, 2029 | DP U-2167 | | | |
| | 8956288 | Jul 06, 2029 | DP U-2167 | | | |
| | 8961412 | Nov 17, 2030 | DP | | | |
| | 9060708 | Mar 05, 2029 | DP | | | |
| | 9089567 | Jan 28, 2022 | U-543 | | | |
| | 9119554 | Dec 16, 2028 | DP | | | |
| | 9125939 | Jul 28, 2026 | U-1749 | | | |
| | 9149577 | Dec 15, 2029 | DP | | | |
| | 9258035 | Mar 05, 2029 | DP | | | |
| | 9268909 | Oct 15, 2033 | DP U-2168 | | | |
| | 9270503 | Sep 19, 2034 | DP U-2169 | | | |
| | 9320455 | Dec 15, 2031 | DP | | | |
| | 9359302 | Sep 25, 2022 | DS DP U-1529 | | | |
| | 9359302 | Sep 25, 2022 | DS DP U-1749 | | | |
| | 9359302 | Sep 25, 2022 | DS DP U-543 | | | |
| | 9387182 | Dec 25, 2023 | U-1529 | | | |
| | 9433371 | Sep 15, 2029 | DP | | | |
| | 9444503 | Nov 19, 2027 | DP U-2169 | | | |
| | 9577864 | Oct 03, 2033 | DP U-2169 | | | |
| | 9787511 | Sep 19, 2034 | DP U-2169 | | | |
| | 9941931 | Nov 04, 2030 | DP | | | |
| <u>ARIPIRAZOLE - ABILIFY MYCITE KIT</u> | | | | | | |
| N 207202 005 | 10097388 | Sep 19, 2034 | U-2169 | | I-746 | Jul 27, 2020 |
| | 7053092 | Jan 28, 2022 | U-1529 | | | |
| | 7978064 | Sep 14, 2026 | DP | | | |
| | 8017615 | Jun 16, 2024 | DP | | | |
| | 8114021 | Nov 02, 2026 | DP | | | |
| | 8258962 | Nov 25, 2030 | DP | | | |
| | 8545402 | Apr 27, 2030 | DP | | | |
| | 8547248 | Dec 18, 2030 | DP U-2167 | | | |
| | 8580796 | Sep 25, 2022 | DS | | | |
| | 8642760 | Sep 25, 2022 | DS | | | |
| | 8674825 | Apr 09, 2029 | DP U-2170 | | | |
| | 8718193 | Dec 05, 2029 | DP | | | |
| | 8759350 | Mar 02, 2027 | U-1529 | | | |
| | 8847766 | Mar 29, 2030 | DP U-2167 | | | |
| | 8945005 | Aug 19, 2029 | DP U-2167 | | | |
| | 8956288 | Jul 06, 2029 | DP U-2167 | | | |
| | 8961412 | Nov 17, 2030 | DP | | | |
| | 9060708 | Mar 05, 2029 | DP | | | |
| | 9089567 | Jan 28, 2022 | U-543 | | | |
| | 9119554 | Dec 16, 2028 | DP | | | |
| | 9125939 | Jul 28, 2026 | U-1749 | | | |
| | 9149577 | Dec 15, 2029 | DP | | | |
| | 9258035 | Mar 05, 2029 | DP | | | |
| | 9268909 | Oct 15, 2033 | DP U-2168 | | | |
| | 9270503 | Sep 19, 2034 | DP U-2169 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ARIPIPIRAZOLE - ABILIFY MYCITE KIT</u> | | | | | | |
| N 207202 005 | 9320455 | Dec 15, 2031 | DP | | | |
| | 9359302 | Sep 25, 2022 | DS DP U-1529 | | | |
| | 9359302 | Sep 25, 2022 | DS DP U-1749 | | | |
| | 9359302 | Sep 25, 2022 | DS DP U-543 | | | |
| | 9387182 | Dec 25, 2023 | U-1529 | | | |
| | 9433371 | Sep 15, 2029 | DP | | | |
| | 9444503 | Nov 19, 2027 | DP U-2169 | | | |
| | 9577864 | Oct 03, 2033 | DP U-2169 | | | |
| | 9787511 | Sep 19, 2034 | DP U-2169 | | | |
| | 9941931 | Nov 04, 2030 | DP | | | |
| <u>ARIPIPIRAZOLE - ABILIFY MYCITE KIT</u> | | | | | | |
| N 207202 006 | 10097388 | Sep 19, 2034 | U-2169 | | I-746 | Jul 27, 2020 |
| | 7053092 | Jan 28, 2022 | U-1529 | | | |
| | 7978064 | Sep 14, 2026 | DP | | | |
| | 8017615 | Jun 16, 2024 | DP | | | |
| | 8114021 | Nov 02, 2026 | DP | | | |
| | 8258962 | Nov 25, 2030 | DP | | | |
| | 8545402 | Apr 27, 2030 | DP | | | |
| | 8547248 | Dec 18, 2030 | DP U-2167 | | | |
| | 8580796 | Sep 25, 2022 | DS | | | |
| | 8642760 | Sep 25, 2022 | DS | | | |
| | 8674825 | Apr 09, 2029 | DP U-2170 | | | |
| | 8718193 | Dec 05, 2029 | DP | | | |
| | 8759350 | Mar 02, 2027 | U-1529 | | | |
| | 8847766 | Mar 29, 2030 | DP U-2167 | | | |
| | 8945005 | Aug 19, 2029 | DP U-2167 | | | |
| | 8956288 | Jul 06, 2029 | DP U-2167 | | | |
| | 8961412 | Nov 17, 2030 | DP | | | |
| | 9060708 | Mar 05, 2029 | DP | | | |
| | 9089567 | Jan 28, 2022 | U-543 | | | |
| | 9119554 | Dec 16, 2028 | DP | | | |
| | 9125939 | Jul 28, 2026 | U-1749 | | | |
| | 9149577 | Dec 15, 2029 | DP | | | |
| | 9258035 | Mar 05, 2029 | DP | | | |
| | 9268909 | Oct 15, 2033 | DP U-2168 | | | |
| | 9270503 | Sep 19, 2034 | DP U-2169 | | | |
| | 9320455 | Dec 15, 2031 | DP | | | |
| | 9359302 | Sep 25, 2022 | DS DP U-1529 | | | |
| | 9359302 | Sep 25, 2022 | DS DP U-1749 | | | |
| | 9359302 | Sep 25, 2022 | DS DP U-543 | | | |
| | 9387182 | Dec 25, 2023 | U-1529 | | | |
| | 9433371 | Sep 15, 2029 | DP | | | |
| | 9444503 | Nov 19, 2027 | DP U-2169 | | | |
| | 9577864 | Oct 03, 2033 | DP U-2169 | | | |
| | 9787511 | Sep 19, 2034 | DP U-2169 | | | |
| | 9941931 | Nov 04, 2030 | DP | | | |
| <u>ARIPIPIRAZOLE LAUROXIL - ARISTADA</u> | | | | | | |
| N 207533 001 | 10112903 | Jun 24, 2030 | DS U-543 | | NCE | Oct 05, 2020 |
| | 8431576 | Oct 26, 2030 | DS | | | |
| | 8796276 | Jun 24, 2030 | U-543 | | | |
| | 9034867 | Nov 07, 2032 | DP U-543 | | | |
| | 9193685 | Oct 24, 2033 | DP U-543 | | | |
| | 9452131 | Mar 19, 2035 | U-2402 | | | |
| <u>ARIPIPIRAZOLE LAUROXIL - ARISTADA</u> | | | | | | |
| N 207533 002 | 10112903 | Jun 24, 2030 | DS U-543 | | NCE | Oct 05, 2020 |
| | 8431576 | Oct 26, 2030 | DS | | | |
| | 8796276 | Jun 24, 2030 | U-543 | | | |
| | 9034867 | Nov 07, 2032 | DP U-543 | | | |
| | 9193685 | Oct 24, 2033 | DP U-543 | | | |
| | 9452131 | Mar 19, 2035 | U-2402 | | | |
| | 9526726 | Mar 19, 2035 | DP | | | |
| <u>ARIPIPIRAZOLE LAUROXIL - ARISTADA</u> | | | | | | |
| N 207533 003 | 10112903 | Jun 24, 2030 | DS U-543 | | NCE | Oct 05, 2020 |
| | 8431576 | Oct 26, 2030 | DS | | | |
| | 8796276 | Jun 24, 2030 | U-543 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ARIPIPIRAZOLE LAUROXIL - ARISTADA</u> | | | | | | |
| N 207533 | 003 | 9034867 | Nov 07, 2032 | DP U-543 | | |
| | | 9193685 | Oct 24, 2033 | DP U-543 | | |
| | | 9452131 | Mar 19, 2035 | U-2402 | | |
| | | 9526726 | Mar 19, 2035 | DP | | |
| <u>ARIPIPIRAZOLE LAUROXIL - ARISTADA</u> | | | | | | |
| N 207533 | 004 | 10112903 | Jun 24, 2030 | DS U-543 | NCE | Oct 05, 2020 |
| | | 8431576 | Oct 26, 2030 | DS | | |
| | | 8796276 | Jun 24, 2030 | U-543 | | |
| | | 9034867 | Nov 07, 2032 | DP U-543 | | |
| | | 9193685 | Oct 24, 2033 | DP U-543 | | |
| | | 9452131 | Mar 19, 2035 | U-2402 | | |
| <u>ARIPIPIRAZOLE LAUROXIL - ARISTADA INITIO KIT</u> | | | | | | |
| N 209830 | 001 | 10016415 | Sep 08, 2035 | DP | NCE | Oct 05, 2020 |
| | | 10112903 | Jun 24, 2030 | DS U-543 | | |
| | | 8431576 | Oct 26, 2030 | DS | | |
| | | 8796276 | Jun 24, 2030 | U-543 | | |
| <u>ARMODAFINIL - NUVIGIL</u> | | | | | | |
| N 021875 | 001 | 7132570 | Dec 18, 2023 | DS DP | | |
| | | 7297346 | Nov 29, 2023 | DP | | |
| <u>ARMODAFINIL - NUVIGIL</u> | | | | | | |
| N 021875 | 002 | 7132570 | Dec 18, 2023 | DS DP | | |
| | | 7297346 | Nov 29, 2023 | DP | | |
| <u>ARMODAFINIL - NUVIGIL</u> | | | | | | |
| N 021875 | 003 | 7132570 | Dec 18, 2023 | DS DP | | |
| | | 7297346 | Nov 29, 2023 | DP | | |
| <u>ARMODAFINIL - NUVIGIL</u> | | | | | | |
| N 021875 | 004 | 7132570 | Dec 18, 2023 | DS DP | | |
| | | 7297346 | Nov 29, 2023 | DP | | |
| <u>ARMODAFINIL - NUVIGIL</u> | | | | | | |
| N 021875 | 005 | 7132570 | Dec 18, 2023 | DS DP | | |
| | | 7297346 | Nov 29, 2023 | DP | | |
| <u>ARSENIC TRIOXIDE - TRISENOX</u> | | | | | | |
| N 021248 | 001 | | | | ODE-167 | Jan 12, 2025 |
| <u>ARSENIC TRIOXIDE - TRISENOX</u> | | | | | | |
| N 021248 | 002 | | | | ODE-167 | Jan 12, 2025 |
| <u>ASCORBIC ACID - ASCOR</u> | | | | | | |
| N 209112 | 001 | | | | ODE-160 | Oct 02, 2024 |
| <u>ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - MOVIPREP</u> | | | | | | |
| N 021881 | 001 | 7169381 | Sep 01, 2024 | DS DP | | |
| | | 7658914 | Sep 01, 2024 | DS DP | | |
| <u>ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - PLENUV</u> | | | | | | |
| N 209381 | 001 | 10016504 | Sep 10, 2033 | DP | NP | May 04, 2021 |
| | | 8999313 | Sep 10, 2033 | DP | | |
| | | 9326969 | Sep 10, 2033 | U-2310 | | |
| | | 9592252 | Aug 11, 2032 | DP U-2310 | | |
| | | 9707297 | Sep 10, 2033 | DP | | |
| <u>ASENAPINE MALEATE - SAPHRIS</u> | | | | | | |
| N 022117 | 001 | 5763476 | Jun 09, 2020 | DP U-1960 | D-166 | Jan 13, 2020 |
| | | 5763476 | Jun 09, 2020 | DP U-1961 | I-597 | Jan 13, 2020 |
| | | 5763476 | Jun 09, 2020 | DP U-1962 | | |
| | | 5763476 | Jun 09, 2020 | DP U-1963 | | |
| | | 5763476 | Jun 09, 2020 | DP U-326 | | |
| | | 5763476*PED | Dec 09, 2020 | | | |
| | | 7741358 | Apr 06, 2026 | DS DP U-1064 | | |
| | | 7741358 | Apr 06, 2026 | DS DP U-1960 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ASENAPINE MALEATE - SAPHRIS</u> | | | | | | |
| N 022117 001 | 7741358 | Apr 06, 2026 | DS DP U-1961 | | | |
| | 7741358 | Apr 06, 2026 | DS DP U-1962 | | | |
| | 7741358 | Apr 06, 2026 | DS DP U-1963 | | | |
| | 7741358*PED | Oct 06, 2026 | | | | |
| | 8022228 | Apr 06, 2026 | DS DP | | | |
| | 8022228*PED | Oct 06, 2026 | | | | |
| <u>ASENAPINE MALEATE - SAPHRIS</u> | | | | | | |
| N 022117 002 | 5763476 | Jun 09, 2020 | DP U-1960 | | D-166 | Jan 13, 2020 |
| | 5763476 | Jun 09, 2020 | DP U-1961 | | I-597 | Jan 13, 2020 |
| | 5763476 | Jun 09, 2020 | DP U-1962 | | | |
| | 5763476 | Jun 09, 2020 | DP U-1963 | | | |
| | 5763476 | Jun 09, 2020 | DP U-326 | | | |
| | 5763476*PED | Dec 09, 2020 | | | | |
| | 7741358 | Apr 06, 2026 | DS DP U-1064 | | | |
| | 7741358 | Apr 06, 2026 | DS DP U-1960 | | | |
| | 7741358 | Apr 06, 2026 | DS DP U-1961 | | | |
| | 7741358 | Apr 06, 2026 | DS DP U-1962 | | | |
| | 7741358 | Apr 06, 2026 | DS DP U-1963 | | | |
| | 7741358*PED | Oct 06, 2026 | | | | |
| | 8022228 | Apr 06, 2026 | DS DP | | | |
| | 8022228*PED | Oct 06, 2026 | | | | |
| <u>ASENAPINE MALEATE - SAPHRIS</u> | | | | | | |
| N 022117 003 | 5763476 | Jun 09, 2020 | DP U-1893 | | D-166 | Jan 13, 2020 |
| | 5763476 | Jun 09, 2020 | DP U-1966 | | I-597 | Jan 13, 2020 |
| | 5763476*PED | Dec 09, 2020 | | | | |
| | 7741358 | Apr 06, 2026 | DS DP U-1893 | | | |
| | 7741358 | Apr 06, 2026 | DS DP U-1966 | | | |
| | 7741358*PED | Oct 06, 2026 | | | | |
| | 8022228 | Apr 06, 2026 | DS DP | | | |
| | 8022228*PED | Oct 06, 2026 | | | | |
| <u>ASPIRIN - VAZALORE</u> | | | | | | |
| N 203697 001 | 8865187 | Mar 23, 2022 | DP | | | |
| | 9101637 | Mar 23, 2022 | U-1731 | | | |
| | 9101637 | Mar 23, 2022 | U-1732 | | | |
| | 9101637 | Mar 23, 2022 | U-1733 | | | |
| | 9216150 | Sep 29, 2032 | DP | | | |
| | 9226892 | Sep 29, 2032 | U-1731 | | | |
| | 9226892 | Sep 29, 2032 | U-1732 | | | |
| | 9226892 | Sep 29, 2032 | U-1733 | | | |
| | 9351984 | Dec 19, 2021 | DP | | | |
| <u>ASPIRIN; OMEPRAZOLE - YOSPRALA</u> | | | | | | |
| N 205103 001 | 6926907 | Feb 28, 2023 | DP U-1902 | | NC | Sep 14, 2019 |
| | 8206741 | Feb 28, 2023 | DP U-1902 | | | |
| | 9364439 | May 31, 2022 | DP U-1902 | | | |
| | 9539214 | Mar 13, 2033 | U-1902 | | | |
| | 9987231 | Jan 02, 2033 | U-2324 | | | |
| <u>ASPIRIN; OMEPRAZOLE - YOSPRALA</u> | | | | | | |
| N 205103 002 | 6926907 | Feb 28, 2023 | DP U-1902 | | NC | Sep 14, 2019 |
| | 8206741 | Feb 28, 2023 | DP U-1902 | | | |
| | 9364439 | May 31, 2022 | DP U-1902 | | | |
| | 9539214 | Mar 13, 2033 | U-1902 | | | |
| | 9987231 | Jan 02, 2033 | U-2324 | | | |
| <u>ATAZANAVIR SULFATE - REYATAZ</u> | | | | | | |
| N 021567 001 | 6087383*PED | Jun 21, 2019 | | | | |
| <u>ATAZANAVIR SULFATE - REYATAZ</u> | | | | | | |
| N 021567 002 | 6087383*PED | Jun 21, 2019 | | | | |
| <u>ATAZANAVIR SULFATE - REYATAZ</u> | | | | | | |
| N 021567 003 | 6087383*PED | Jun 21, 2019 | | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ATAZANAVIR SULFATE - REYATAZ</u> | | | | | | |
| N 021567 | 004 | 6087383*PED | Jun 21, 2019 | | | |
| <u>ATAZANAVIR SULFATE - REYATAZ</u> | | | | | | |
| N 206352 | 001 | 6087383*PED | Jun 21, 2019 | | | |
| <u>ATAZANAVIR SULFATE; COBICISTAT - EVOTAZ</u> | | | | | | |
| N 206353 | 001 | 10039718 | Oct 04, 2032 | DP | | |
| | | 6087383*PED | Jun 21, 2019 | | | |
| | | 8148374 | Sep 03, 2029 | DS DP | U-1279 | |
| <u>ATORVASTATIN CALCIUM - LIPITOR</u> | | | | | | |
| N 020702 | 001 | | | | M-204 | Jun 23, 2020 |
| <u>ATORVASTATIN CALCIUM - LIPITOR</u> | | | | | | |
| N 020702 | 002 | | | | M-204 | Jun 23, 2020 |
| <u>AVANAFIL - STENDRA</u> | | | | | | |
| N 202276 | 001 | 6656935 | Apr 27, 2025 | DS DP | U-155 | |
| | | 7501409 | May 05, 2023 | DP | | |
| <u>AVANAFIL - STENDRA</u> | | | | | | |
| N 202276 | 002 | 6656935 | Apr 27, 2025 | DS DP | U-155 | |
| | | 7501409 | May 05, 2023 | DP | | |
| <u>AVANAFIL - STENDRA</u> | | | | | | |
| N 202276 | 003 | 6656935 | Apr 27, 2025 | DS DP | U-155 | |
| | | 7501409 | May 05, 2023 | DP | | |
| <u>AVATROMBOPAG MALEATE - DOPTELET</u> | | | | | | |
| N 210238 | 001 | 7638536 | May 05, 2025 | DS DP | | NCE |
| | | 8765764 | Jan 15, 2023 | | U-2314 | May 21, 2023 |
| <u>AVIBACTAM SODIUM; CEFTAZIDIME - AVYCAZ</u> | | | | | | |
| N 206494 | 001 | 7112592 | Feb 24, 2022 | DS DP | U-2244 | |
| | | 7112592 | Feb 24, 2022 | DS DP | U-282 | |
| | | 7612087 | Nov 12, 2026 | DP | | |
| | | 8178554 | Jul 24, 2021 | DS DP | U-2245 | |
| | | 8178554 | Jul 24, 2021 | DS DP | U-282 | |
| | | 8471025 | Aug 12, 2031 | DS | | |
| | | 8835455 | Oct 08, 2030 | DP | | |
| | | 8969566 | Jun 15, 2032 | DS | | |
| | | 9284314 | Jun 15, 2032 | DS | | |
| | | 9695122 | Jun 15, 2032 | DS | | |
| <u>AXITINIB - INLYTA</u> | | | | | | |
| N 202324 | 001 | 6534524 | Apr 29, 2025 | DS DP | | |
| | | 7141581 | Jun 30, 2020 | | U-1220 | |
| | | 8791140 | Dec 14, 2030 | DS | | |
| <u>AXITINIB - INLYTA</u> | | | | | | |
| N 202324 | 002 | 6534524 | Apr 29, 2025 | DS DP | | |
| | | 7141581 | Jun 30, 2020 | | U-1220 | |
| | | 8791140 | Dec 14, 2030 | DS | | |
| <u>AZELAIC ACID - FINACEA</u> | | | | | | |
| N 207071 | 001 | 10117812 | Oct 18, 2027 | DP | U-1796 | |
| | | 6730288 | Sep 08, 2019 | DP | | |
| | | 7700076 | Sep 18, 2027 | DP | | |
| | | 8435498 | Mar 01, 2024 | | U-1727 | |
| | | 8722021 | Oct 24, 2023 | DP | | |
| | | 8900554 | Oct 24, 2023 | DP | | |
| | | 9211259 | Feb 28, 2029 | | U-1796 | |
| | | 9265725 | Dec 08, 2027 | DP | | |
| <u>AZELASTINE HYDROCHLORIDE - ASTEPRO</u> | | | | | | |
| N 022203 | 001 | 8071073 | Jun 04, 2028 | DP | | |
| | | 8518919 | Nov 22, 2025 | | U-1430 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>AZELASTINE HYDROCHLORIDE - ASTEPRO</u> | | | | | | |
| N 022203 | 002 | 8071073 | Jun 04, 2028 | DP | | |
| | | 8518919 | Nov 22, 2025 | | U-1430 | |
| | | 9919050 | Nov 22, 2025 | DP | | |
| <u>AZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE - DYMISTA</u> | | | | | | |
| N 202236 | 001 | 8163723 | Aug 29, 2023 | | U-1667 | |
| | | 8163723 | Aug 29, 2023 | | U-644 | |
| | | 8163723 | Aug 29, 2023 | | U-707 | |
| | | 8163723 | Aug 29, 2023 | | U-77 | |
| | | 8163723 | Aug 29, 2023 | | U-81 | |
| | | 8163723*PED | Feb 29, 2024 | | | |
| | | 8168620 | Feb 24, 2026 | DP | | |
| | | 9259428 | Jun 13, 2023 | | U-644 | |
| | | 9259428*PED | Dec 13, 2023 | | | |
| | | 9901585 | Jun 13, 2023 | DP | | |
| <u>AZILSARTAN KAMEDOXOMIL - EDARBI</u> | | | | | | |
| N 200796 | 001 | 7157584 | May 22, 2025 | DS | | |
| | | 7572920 | Jan 07, 2025 | | DP U-3 | |
| | | 9066936 | Mar 26, 2028 | DP | | |
| <u>AZILSARTAN KAMEDOXOMIL - EDARBI</u> | | | | | | |
| N 200796 | 002 | 7157584 | May 22, 2025 | DS | | |
| | | 7572920 | Jan 07, 2025 | | DP U-3 | |
| | | 9066936 | Mar 26, 2028 | DP | | |
| <u>AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE - EDARBYCLOR</u> | | | | | | |
| N 202331 | 001 | 7157584 | May 22, 2025 | DS | | |
| | | 7572920 | Jan 07, 2025 | | DP U-3 | |
| | | 9066936 | Mar 26, 2028 | DP | | |
| | | 9169238 | Feb 04, 2030 | DP | | |
| <u>AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE - EDARBYCLOR</u> | | | | | | |
| N 202331 | 002 | 7157584 | May 22, 2025 | DS | | |
| | | 7572920 | Jan 07, 2025 | | DP U-3 | |
| | | 9066936 | Mar 26, 2028 | DP | | |
| | | 9169238 | Feb 04, 2030 | DP | | |
| <u>AZITHROMYCIN - ZMAX</u> | | | | | | |
| N 050797 | 001 | 6984403 | Feb 14, 2024 | | DP U-282 | |
| | | 7887844 | Feb 14, 2024 | DP | | |
| <u>AZITHROMYCIN - AZASITE</u> | | | | | | |
| N 050810 | 001 | 6239113 | Mar 31, 2019 | | U-709 | |
| | | 6569443 | Mar 31, 2019 | | DP U-709 | |
| | | 7056893 | Mar 31, 2019 | | DP U-709 | |
| <u>AZTREONAM - CAYSTON</u> | | | | | | |
| N 050814 | 001 | 7208141 | Dec 20, 2021 | | DP U-1031 | |
| | | 7214364 | Dec 20, 2021 | | DP | |
| | | 7427633 | Dec 20, 2021 | | DP U-1031 | |
| | | 8399496 | Dec 20, 2021 | | DP U-1377 | |
| <u>BALOXAVIR MARBOXIL - XOFLUZA</u> | | | | | | |
| N 210854 | 001 | 8927710 | May 05, 2031 | | DP | |
| | | 8987441 | Sep 21, 2031 | DS | DP | |
| | | 9815835 | Jun 14, 2030 | | DP | |
| <u>BALOXAVIR MARBOXIL - XOFLUZA</u> | | | | | | |
| N 210854 | 002 | 8927710 | May 05, 2031 | | DP | |
| | | 8987441 | Sep 21, 2031 | DS | DP | |
| | | 9815835 | Jun 14, 2030 | | DP | |
| <u>BALSALAZIDE DISODIUM - COLAZAL</u> | | | | | | |
| N 020610 | 001 | 7452872 | Aug 24, 2026 | | U-141 | |
| | | 7625884 | Aug 24, 2026 | | U-141 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>BALSALAZIDE DISODIUM - GIAZO</u> | | | | | | |
| N 022205 | 001 | 7452872 | Aug 24, 2026 | U-1229 | | |
| | | 7625884 | Aug 24, 2026 | U-1229 | | |
| | | 8497256 | Jun 23, 2031 | U-1229 | | |
| | | 9192616 | Aug 02, 2026 | U-1229 | | |
| <u>BARICITINIB - OLUMIANT</u> | | | | | | |
| N 207924 | 001 | 8158616 | Jun 08, 2030 | DS DP | NCE | May 31, 2023 |
| | | 8420629 | Mar 10, 2029 | U-247 | | |
| <u>BAZEDOXIFENE ACETATE; ESTROGENS, CONJUGATED - DUAVEE</u> | | | | | | |
| N 022247 | 001 | 5998402 | Apr 04, 2019 | DS DP | U-594 | |
| | | 6479535 | May 06, 2019 | DP | U-594 | |
| | | 6479535 | May 06, 2019 | DP | U-904 | |
| | | 7683051 | Mar 10, 2027 | DS DP | U-594 | |
| | | 7683051 | Mar 10, 2027 | DS DP | U-904 | |
| | | 8815934 | May 06, 2019 | DP | | |
| <u>BECLOMETHASONE DIPROPIONATE - QVAR 80</u> | | | | | | |
| N 020911 | 001 | 10022509 | May 18, 2031 | DP | | |
| | | 10022510 | May 18, 2031 | DP | | |
| | | 10086156 | May 18, 2031 | DP | | |
| | | 9463289 | May 18, 2031 | DP | | |
| | | 9808587 | May 18, 2031 | DP | | |
| <u>BECLOMETHASONE DIPROPIONATE - QVAR 40</u> | | | | | | |
| N 020911 | 002 | 10022509 | May 18, 2031 | DP | | |
| | | 10022510 | May 18, 2031 | DP | | |
| | | 10086156 | May 18, 2031 | DP | | |
| | | 9463289 | May 18, 2031 | DP | | |
| | | 9808587 | May 18, 2031 | DP | | |
| <u>BECLOMETHASONE DIPROPIONATE - QNASL</u> | | | | | | |
| N 202813 | 001 | 7780038 | Jan 24, 2027 | DP | | |
| <u>BECLOMETHASONE DIPROPIONATE - QNASL</u> | | | | | | |
| N 202813 | 002 | 7780038 | Jan 24, 2027 | DP | | |
| <u>BECLOMETHASONE DIPROPIONATE - QVAR REDIHALER</u> | | | | | | |
| N 207921 | 001 | 10022509 | May 18, 2031 | DP | | |
| | | 10022510 | May 18, 2031 | DP | | |
| | | 10086156 | May 18, 2031 | DP | | |
| | | 7637260 | Aug 25, 2020 | DP | | |
| | | 8132712 | Sep 07, 2028 | DP | | |
| | | 8931476 | Jul 17, 2031 | DP | | |
| <u>BECLOMETHASONE DIPROPIONATE - QVAR REDIHALER</u> | | | | | | |
| N 207921 | 002 | 10022509 | May 18, 2031 | DP | | |
| | | 10022510 | May 18, 2031 | DP | | |
| | | 10086156 | May 18, 2031 | DP | | |
| | | 7637260 | Aug 25, 2020 | DP | | |
| | | 8132712 | Sep 07, 2028 | DP | | |
| | | 8931476 | Jul 17, 2031 | DP | | |
| <u>BEDAQUILINE FUMARATE - SIRTURO</u> | | | | | | |
| N 204384 | 001 | 7498343 | Dec 01, 2026 | DS DP | U-1321 | ODE-38 Dec 28, 2019 |
| | | 8546428 | Mar 19, 2029 | DS DP | U-1321 | |
| <u>BELINOSTAT - BELEODAQ</u> | | | | | | |
| N 206256 | 001 | 6888027 | Sep 27, 2021 | DS DP | U-1544 | NCE Jul 03, 2019 |
| | | 8835501 | Oct 27, 2027 | DP | ODE-68 | Jul 03, 2021 |
| <u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u> | | | | | | |
| N 022249 | 001 | 8436190 | Oct 26, 2030 | DP | | |
| | | 8436190*PED | Apr 26, 2031 | | | |
| | | 8445524 | Mar 26, 2029 | DS DP | U-1402 | |
| | | 8445524*PED | Sep 26, 2029 | | | |
| | | 8609863 | Jan 12, 2026 | DP | | |
| | | 8609863*PED | Jul 12, 2026 | | | |
| | | 8669279 | Mar 26, 2029 | DP | U-1402 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u> | | | | | | |
| N 022249 001 | 8669279*PED | Sep 26, 2029 | | | | |
| | 8791270 | Jan 12, 2026 | DP U-1542 | | | |
| | 8791270*PED | Jul 12, 2026 | | | | |
| | 8883836 | Mar 26, 2029 | DP U-1402 | | | |
| | 8883836*PED | Sep 26, 2029 | | | | |
| | 8895756 | Jan 12, 2026 | DP | | | |
| | 8895756*PED | Jul 12, 2026 | | | | |
| | 9533955 | Mar 26, 2029 | DP U-1949 | | | |
| | 9533955 | Mar 26, 2029 | DP U-1952 | | | |
| <u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u> | | | | | | |
| N 022249 002 | 8436190 | Oct 26, 2030 | DP | | | |
| | 8436190*PED | Apr 26, 2031 | | | | |
| | 8445524 | Mar 26, 2029 | DS DP U-1402 | | | |
| | 8445524*PED | Sep 26, 2029 | | | | |
| | 8609863 | Jan 12, 2026 | DP | | | |
| | 8609863*PED | Jul 12, 2026 | | | | |
| | 8669279 | Mar 26, 2029 | DP U-1402 | | | |
| | 8669279*PED | Sep 26, 2029 | | | | |
| | 8791270 | Jan 12, 2026 | DP U-1542 | | | |
| | 8791270*PED | Jul 12, 2026 | | | | |
| | 8883836 | Mar 26, 2029 | DP U-1402 | | | |
| | 8883836*PED | Sep 26, 2029 | | | | |
| | 8895756 | Jan 12, 2026 | DP | | | |
| | 8895756*PED | Jul 12, 2026 | | | | |
| | 9533955 | Mar 26, 2029 | DP U-1949 | | | |
| | 9533955 | Mar 26, 2029 | DP U-1952 | | | |
| <u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u> | | | | | | |
| N 022249 003 | 8344006 | Sep 23, 2029 | DP U-1402 | | | |
| | 8344006*PED | Mar 23, 2030 | | | | |
| | 8445524 | Mar 26, 2029 | DS | | | |
| | 8445524*PED | Sep 26, 2029 | | | | |
| | 8791270 | Jan 12, 2026 | DP U-1542 | | | |
| | 8791270*PED | Jul 12, 2026 | | | | |
| <u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u> | | | | | | |
| N 022249 004 | 8344006 | Sep 23, 2029 | DP U-1402 | | | |
| | 8344006*PED | Mar 23, 2030 | | | | |
| | 8445524 | Mar 26, 2029 | DS | | | |
| | 8445524*PED | Sep 26, 2029 | | | | |
| | 8791270 | Jan 12, 2026 | DP U-1542 | | | |
| | 8791270*PED | Jul 12, 2026 | | | | |
| <u>BENDAMUSTINE HYDROCHLORIDE - BELRAPZO</u> | | | | | | |
| N 205580 001 | 10010533 | Jan 28, 2031 | DP | | | |
| | 8609707 | Aug 11, 2031 | DP U-1971 | | | |
| | 8609707 | Aug 11, 2031 | DP U-1972 | | | |
| | 8791270 | Jan 12, 2026 | DP U-1971 | | | |
| | 8791270 | Jan 12, 2026 | DP U-1972 | | | |
| | 9265831 | Jan 28, 2031 | DP | | | |
| | 9572796 | Jan 28, 2031 | DP U-1971 | | | |
| | 9572796 | Jan 28, 2031 | DP U-1972 | | | |
| | 9572797 | Jan 28, 2031 | U-1971 | | | |
| | 9572797 | Jan 28, 2031 | U-1972 | | | |
| <u>BENDAMUSTINE HYDROCHLORIDE - BENDEKA</u> | | | | | | |
| N 208194 001 | 10010533 | Jan 28, 2031 | DP | | ODE-179 | Dec 07, 2022 |
| | 10052385 | Mar 15, 2033 | U-1971 | | | |
| | 10052385 | Mar 15, 2033 | U-1972 | | | |
| | 8609707 | Aug 11, 2031 | DP U-1542 | | | |
| | 8791270 | Jan 12, 2026 | DP U-1790 | | | |
| | 9000021 | Mar 15, 2033 | U-1542 | | | |
| | 9034908 | Mar 15, 2033 | U-1542 | | | |
| | 9144568 | Mar 15, 2033 | U-1542 | | | |
| | 9265831 | Jan 28, 2031 | DP | | | |
| | 9572796 | Jan 28, 2031 | DP U-1971 | | | |
| | 9572796 | Jan 28, 2031 | DP U-1972 | | | |
| | 9572797 | Jan 28, 2031 | U-1971 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>BENDAMUSTINE HYDROCHLORIDE - BENDEKA</u> | | | | | | |
| N 208194 001 | 9572797 | Jan 28, 2031 | U-1972 | | | |
| | 9572887 | Mar 15, 2033 | U-1971 | | | |
| | 9572887 | Mar 15, 2033 | U-1972 | | | |
| | 9579384 | Mar 15, 2033 | U-1971 | | | |
| | 9579384 | Mar 15, 2033 | U-1972 | | | |
| | 9597397 | Mar 15, 2033 | U-1971 | | | |
| | 9597397 | Mar 15, 2033 | U-1972 | | | |
| | 9597398 | Mar 15, 2033 | U-1971 | | | |
| | 9597399 | Mar 15, 2033 | U-1971 | | | |
| | 9597399 | Mar 15, 2033 | U-1972 | | | |
| <u>BENZNIDAZOLE - BENZNIDAZOLE</u> | | | | | | |
| N 209570 001 | | | | | NCE ODE-154 | Aug 29, 2022 Aug 29, 2024 |
| <u>BENZNIDAZOLE - BENZNIDAZOLE</u> | | | | | | |
| N 209570 002 | | | | | NCE ODE-154 | Aug 29, 2022 Aug 29, 2024 |
| <u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - ACANYA</u> | | | | | | |
| N 050819 001 | 8288434 | Aug 05, 2029 | DP U-124 | | | |
| | 8663699 | Jun 03, 2029 | U-124 | | | |
| | 8895070 | Jun 03, 2029 | U-124 | | | |
| | 9078870 | Jun 03, 2029 | DP | | | |
| <u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - ONEXTON</u> | | | | | | |
| N 050819 002 | 10137142 | Jun 03, 2029 | DP U-916 | | | |
| | 8288434 | Aug 05, 2029 | DP U-1033 | | | |
| | 8288434 | Aug 05, 2029 | DP U-124 | | | |
| | 8288434 | Aug 05, 2029 | DP U-134 | | | |
| | 8288434 | Aug 05, 2029 | DP U-818 | | | |
| | 8288434 | Aug 05, 2029 | DP U-916 | | | |
| | 8288434 | Aug 05, 2029 | DP U-921 | | | |
| | 9504704 | Jun 03, 2029 | DP U-124 | | | |
| | 9504704 | Jun 03, 2029 | DP U-134 | | | |
| | 9504704 | Jun 03, 2029 | DP U-818 | | | |
| | 9504704 | Jun 03, 2029 | DP U-916 | | | |
| | 9561208 | Jun 03, 2029 | DP U-916 | | | |
| <u>BENZYL ALCOHOL - ULESFIA</u> | | | | | | |
| N 022129 001 | 6793931 | Jul 11, 2022 | DP U-970 | | | |
| | 7294342 | May 19, 2024 | U-970 | | | |
| <u>BEPOTASTINE BESILATE - BEPREVE</u> | | | | | | |
| N 022288 001 | 6780877 | Sep 19, 2019 | DS DP | | | |
| | 8784789 | Sep 05, 2024 | DP | | | |
| | 8877168 | Jul 30, 2023 | DP | | | |
| <u>BESIFLOXACIN HYDROCHLORIDE - BESIVANCE</u> | | | | | | |
| N 022308 001 | 6685958 | Jun 29, 2021 | DP U-80 | | | |
| | 6699492 | Mar 31, 2019 | DP U-80 | | | |
| | 8415342 | Nov 07, 2030 | U-80 | | | |
| | 8481526 | Jan 09, 2031 | DS | | | |
| | 8604020 | Mar 12, 2030 | DP | | | |
| | 8937062 | Nov 13, 2029 | U-80 | | | |
| <u>BETAMETHASONE DIPROPIONATE - SERNIVO</u> | | | | | | |
| N 208079 001 | 9364485 | Aug 31, 2030 | DP U-1858 | | NDF | Feb 05, 2019 |
| | 9433630 | Aug 31, 2030 | DP U-1858 | | | |
| | 9439911 | Aug 31, 2030 | DP U-1858 | | | |
| | 9655907 | Aug 31, 2030 | DP U-1858 | | | |
| | 9775851 | Aug 31, 2030 | DP U-1858 | | | |
| | 9877974 | Aug 31, 2030 | DP U-1858 | | | |
| <u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE - ENSTILAR</u> | | | | | | |
| N 207589 001 | 10130640 | Jun 10, 2031 | DP | | | |
| | 6753013 | Jan 27, 2020 | DP U-1761 | | | |
| | 9119781 | Jun 10, 2031 | DP U-1761 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE - TACLONEX</u> | | | | | | |
| N 021852 | 001 | 6753013 | Jan 27, 2020 | DP U-193 | | |
| | | 6753013 | Jan 27, 2020 | DP U-88 | | |
| <u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE - TACLONEX</u> | | | | | | |
| N 022185 | 001 | 6753013 | Jan 27, 2020 | DP U-1761 | | |
| | | 6753013 | Jan 27, 2020 | DP U-193 | | |
| | | 6753013 | Jan 27, 2020 | DP U-88 | | |
| | | 6787529 | Jan 27, 2020 | DP U-1761 | | |
| | | 6787529 | Jan 27, 2020 | DP U-193 | | |
| | | 6787529 | Jan 27, 2020 | DP U-88 | | |
| <u>BETRIXABAN - BEVYXXA</u> | | | | | | |
| N 208383 | 001 | 6376515 | Sep 15, 2020 | DS DP U-1167 | NCE | Jun 23, 2022 |
| | | 6376515 | Sep 15, 2020 | DS DP U-1502 | | |
| | | 6376515 | Sep 15, 2020 | DS DP U-2029 | | |
| | | 6376515 | Sep 15, 2020 | DS DP U-2030 | | |
| | | 6835739 | Sep 15, 2020 | DS DP | | |
| | | 7598276 | Nov 08, 2026 | DS | | |
| | | 8404724 | Mar 29, 2031 | DP U-2034 | | |
| | | 8518977 | Sep 15, 2020 | DS | | |
| | | 8557852 | Sep 08, 2028 | U-1167 | | |
| | | 8557852 | Sep 08, 2028 | U-2030 | | |
| | | 8691847 | Sep 15, 2020 | DS DP U-2029 | | |
| | | 8691847 | Sep 15, 2020 | DS DP U-2035 | | |
| | | 8987463 | Dec 28, 2030 | DP | | |
| | | 9555023 | Nov 07, 2026 | U-1502 | | |
| | | 9629831 | Sep 15, 2020 | U-1167 | | |
| | | 9629831 | Sep 15, 2020 | U-1502 | | |
| | | 9629831 | Sep 15, 2020 | U-2030 | | |
| | | 9629831 | Sep 15, 2020 | U-2035 | | |
| <u>BETRIXABAN - BEVYXXA</u> | | | | | | |
| N 208383 | 002 | 6376515 | Sep 15, 2020 | DS DP U-1167 | NCE | Jun 23, 2022 |
| | | 6376515 | Sep 15, 2020 | DS DP U-1502 | | |
| | | 6376515 | Sep 15, 2020 | DS DP U-2029 | | |
| | | 6376515 | Sep 15, 2020 | DS DP U-2030 | | |
| | | 6835739 | Sep 15, 2020 | DS DP | | |
| | | 7598276 | Nov 08, 2026 | DS | | |
| | | 8404724 | Mar 29, 2031 | DP U-2034 | | |
| | | 8518977 | Sep 15, 2020 | DS | | |
| | | 8557852 | Sep 08, 2028 | U-1167 | | |
| | | 8557852 | Sep 08, 2028 | U-2030 | | |
| | | 8691847 | Sep 15, 2020 | DS DP U-2029 | | |
| | | 8691847 | Sep 15, 2020 | DS DP U-2035 | | |
| | | 8987463 | Dec 28, 2030 | DP | | |
| | | 9555023 | Nov 07, 2026 | U-1502 | | |
| | | 9629831 | Sep 15, 2020 | U-1167 | | |
| | | 9629831 | Sep 15, 2020 | U-1502 | | |
| | | 9629831 | Sep 15, 2020 | U-2030 | | |
| | | 9629831 | Sep 15, 2020 | U-2035 | | |
| <u>BICTEGRAVIR SODIUM; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - BIKTARVY</u> | | | | | | |
| N 210251 | 001 | 6642245 | Nov 04, 2020 | U-257 | NCE | Feb 07, 2023 |
| | | 6703396 | Mar 09, 2021 | DS DP | | |
| | | 7390791 | May 07, 2022 | DS DP | | |
| | | 7803788 | Feb 02, 2022 | U-257 | | |
| | | 8754065 | Aug 15, 2032 | DS DP U-257 | | |
| | | 9216996 | Dec 19, 2033 | DS DP | | |
| | | 9296769 | Aug 15, 2032 | DS DP U-257 | | |
| | | 9708342 | Jun 19, 2035 | DS DP | | |
| | | 9732092 | Dec 19, 2033 | DS DP | | |
| <u>BIMATOPROST - LUMIGAN</u> | | | | | | |
| N 022184 | 001 | 7851504 | Jun 13, 2027 | DS DP | | |
| | | 8278353 | Mar 16, 2025 | DP | | |
| | | 8299118 | Mar 16, 2025 | U-1295 | | |
| | | 8309605 | Mar 16, 2025 | U-1293 | | |
| | | 8309605 | Mar 16, 2025 | U-1294 | | |
| | | 8338479 | Mar 16, 2025 | DP U-1295 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>BIMATOPROST - LUMIGAN</u> | | | | | | |
| N 022184 | 001 | 8524777 | Mar 16, 2025 | U-1235 | | |
| | | 8586630 | Mar 16, 2025 | U-1458 | | |
| | | 8772338 | Mar 16, 2025 | DP U-1528 | | |
| | | 8933120 | Mar 16, 2025 | DP | | |
| | | 8933127 | Mar 16, 2025 | DP | | |
| | | 9155716 | Mar 16, 2025 | DP U-1528 | | |
| | | 9241918 | Mar 16, 2025 | DP U-1814 | | |
| <u>BIMATOPROST - LATISSE</u> | | | | | | |
| N 022369 | 001 | 8038988 | Aug 25, 2023 | DS DP U-1208 | | |
| | | 8101161 | May 25, 2024 | U-1217 | | |
| | | 8101161 | May 25, 2024 | U-1218 | | |
| | | 8263054 | Jan 15, 2023 | U-1277 | | |
| | | 8541466 | Jan 31, 2021 | U-1217 | | |
| | | 8632760 | Jan 15, 2023 | U-1487 | | |
| | | 8758733 | Jan 15, 2023 | U-1487 | | |
| | | 8906962 | Jan 31, 2021 | U-1217 | | |
| | | 8986715 | Jan 15, 2023 | U-1217 | | |
| | | 9216183 | Jan 15, 2023 | U-1487 | | |
| | | 9226931 | Jan 15, 2023 | U-1799 | | |
| | | 9579270 | Jan 31, 2021 | U-1975 | | |
| <u>BINIMETINIB - MEKTOVI</u> | | | | | | |
| N 210498 | 001 | 10005761 | Aug 27, 2030 | U-2331 | NCE | Jun 27, 2023 |
| | | 7777050 | Mar 13, 2023 | DS DP | ODE-194 | Jun 27, 2025 |
| | | 8178693 | Mar 13, 2023 | DS DP | | |
| | | 8193229 | Mar 13, 2023 | U-2330 | | |
| | | 8513293 | Mar 13, 2023 | U-2331 | | |
| | | 9314464 | Jul 04, 2031 | U-2332 | | |
| | | 9562016 | Oct 18, 2033 | DS DP | | |
| | | 9593100 | Aug 27, 2030 | DP | | |
| | | 9598376 | Oct 18, 2033 | U-2330 | | |
| | | 9850229 | Aug 27, 2030 | U-2333 | | |
| | | 9980944 | Oct 18, 2033 | U-2334 | | |
| <u>BISACODYL; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE - HALFLYTELY</u> | | | | | | |
| N 021551 | 003 | 7291324 | Oct 22, 2022 | U-837 | | |
| <u>BIVALIRUDIN - ANGIOMAX</u> | | | | | | |
| N 020873 | 001 | 7582727 | Jul 27, 2028 | DP | | |
| | | 7598343 | Jul 27, 2028 | DP | | |
| <u>BOCEPREVIR - VICTRELIS</u> | | | | | | |
| N 202258 | 001 | 7772178 | Nov 11, 2027 | DP U-1128 | | |
| | | 8119602 | Mar 17, 2027 | U-1233 | | |
| | | RE43298 | Dec 22, 2024 | DS DP U-1128 | | |
| <u>BORTEZOMIB - VELCADE</u> | | | | | | |
| N 021602 | 001 | 6713446 | Jan 25, 2022 | DS DP | ODE-76 | Oct 08, 2021 |
| | | 6713446*PED | Jul 25, 2022 | | PED | Apr 08, 2022 |
| | | 6958319 | Jan 25, 2022 | DS DP | | |
| | | 6958319*PED | Jul 25, 2022 | | | |
| <u>BORTEZOMIB - BORTEZOMIB</u> | | | | | | |
| N 205004 | 001 | 8962572 | Nov 03, 2032 | DP | | |
| <u>BOSENTAN - TRACLEER</u> | | | | | | |
| N 021290 | 001 | | | | NPP | Sep 05, 2020 |
| <u>BOSENTAN - TRACLEER</u> | | | | | | |
| N 021290 | 002 | | | | NPP | Sep 05, 2020 |
| <u>BOSENTAN - TRACLEER</u> | | | | | | |
| N 209279 | 001 | 7959945 | Dec 28, 2027 | DP | NP | Sep 05, 2020 |
| | | 8309126 | May 15, 2026 | DP | ODE-161 | Sep 05, 2024 |
| <u>BOSUTINIB MONOHYDRATE - BOSULIF</u> | | | | | | |
| N 203341 | 001 | 7417148 | Jan 23, 2026 | U-1283 | I-759 | Dec 19, 2020 |
| | | 7767678 | Nov 23, 2026 | DS DP | ODE-163 | Dec 19, 2024 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>BOSUTINIB MONOHYDRATE - BOSULIF</u> | | | | | | |
| N 203341 001 | 7919625 | Dec 11, 2025 | DP | | ODE-30 | Sep 04, 2019 |
| | RE42376 | Apr 13, 2024 | DS | | | |
| <u>BOSUTINIB MONOHYDRATE - BOSULIF</u> | | | | | | |
| N 203341 002 | 7417148 | Jan 23, 2026 | | U-1283 | I-759 | Dec 19, 2020 |
| | 7767678 | Nov 23, 2026 | DS DP | | ODE-163 | Dec 19, 2024 |
| | 7919625 | Dec 11, 2025 | DP | | ODE-30 | Sep 04, 2019 |
| | RE42376 | Apr 13, 2024 | DS | | | |
| <u>BOSUTINIB MONOHYDRATE - BOSULIF</u> | | | | | | |
| N 203341 003 | 7417148 | Jan 23, 2026 | | U-1283 | I-759 | Dec 19, 2020 |
| | 7767678 | Nov 23, 2026 | DS DP | | ODE-163 | Dec 19, 2024 |
| | 7919625 | Dec 11, 2025 | DP | | ODE-30 | Sep 04, 2019 |
| | RE42376 | Apr 13, 2024 | DS | | | |
| <u>BREXPIRAZOLE - REXULTI</u> | | | | | | |
| N 205422 001 | 7888362 | Feb 23, 2027 | DS | | M-186 | Sep 23, 2019 |
| | 8349840 | Apr 12, 2026 | DP U-1529 | | NCE | Jul 10, 2020 |
| | 8618109 | Apr 12, 2026 | | U-543 | | |
| | 9839637 | Apr 12, 2026 | DP U-1529 | | | |
| | 9839637 | Apr 12, 2026 | DP U-543 | | | |
| <u>BREXPIRAZOLE - REXULTI</u> | | | | | | |
| N 205422 002 | 7888362 | Feb 23, 2027 | DS | | M-186 | Sep 23, 2019 |
| | 8349840 | Apr 12, 2026 | DP U-1529 | | NCE | Jul 10, 2020 |
| | 8618109 | Apr 12, 2026 | | U-543 | | |
| | 9839637 | Apr 12, 2026 | DP U-1529 | | | |
| | 9839637 | Apr 12, 2026 | DP U-543 | | | |
| <u>BREXPIRAZOLE - REXULTI</u> | | | | | | |
| N 205422 003 | 7888362 | Feb 23, 2027 | DS | | M-186 | Sep 23, 2019 |
| | 8349840 | Apr 12, 2026 | DP U-1529 | | NCE | Jul 10, 2020 |
| | 8618109 | Apr 12, 2026 | | U-543 | | |
| | 9839637 | Apr 12, 2026 | DP U-1529 | | | |
| | 9839637 | Apr 12, 2026 | DP U-543 | | | |
| <u>BREXPIRAZOLE - REXULTI</u> | | | | | | |
| N 205422 004 | 7888362 | Feb 23, 2027 | DS | | M-186 | Sep 23, 2019 |
| | 8349840 | Apr 12, 2026 | DP U-1529 | | NCE | Jul 10, 2020 |
| | 8618109 | Apr 12, 2026 | | U-543 | | |
| | 9839637 | Apr 12, 2026 | DP U-1529 | | | |
| | 9839637 | Apr 12, 2026 | DP U-543 | | | |
| <u>BREXPIRAZOLE - REXULTI</u> | | | | | | |
| N 205422 005 | 7888362 | Feb 23, 2027 | DS | | M-186 | Sep 23, 2019 |
| | 8349840 | Apr 12, 2026 | DP U-1529 | | NCE | Jul 10, 2020 |
| | 8618109 | Apr 12, 2026 | | U-543 | | |
| | 9839637 | Apr 12, 2026 | DP U-1529 | | | |
| | 9839637 | Apr 12, 2026 | DP U-543 | | | |
| <u>BREXPIRAZOLE - REXULTI</u> | | | | | | |
| N 205422 006 | 7888362 | Feb 23, 2027 | DS | | M-186 | Sep 23, 2019 |
| | 8349840 | Apr 12, 2026 | DP U-1529 | | NCE | Jul 10, 2020 |
| | 8618109 | Apr 12, 2026 | | U-543 | | |
| | 9839637 | Apr 12, 2026 | DP U-1529 | | | |
| | 9839637 | Apr 12, 2026 | DP U-543 | | | |
| <u>BRIGATINIB - ALUNBRIG</u> | | | | | | |
| N 208772 001 | 9012462 | Feb 06, 2031 | DS | | NCE | Apr 28, 2022 |
| | 9273077 | May 21, 2029 | | U-1927 | ODE-142 | Apr 28, 2024 |
| | 9611283 | Apr 10, 2034 | | U-1927 | | |
| <u>BRIGATINIB - ALUNBRIG</u> | | | | | | |
| N 208772 002 | 9012462 | Feb 06, 2031 | DS | | NCE | Apr 28, 2022 |
| | 9273077 | May 21, 2029 | | U-1927 | ODE-142 | Apr 28, 2024 |
| | 9611283 | Apr 10, 2034 | | U-1927 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>BRIGATINIB - ALUNBRIG</u> | | | | | | |
| N 208772 | 003 | 9012462 | Feb 06, 2031 | DS | NCE | Apr 28, 2022 |
| | | 9273077 | May 21, 2029 | | ODE-142 | Apr 28, 2024 |
| | | 9611283 | Apr 10, 2034 | | | |
| <u>BRIMONIDINE TARTRATE - ALPHAGAN P</u> | | | | | | |
| N 021262 | 001 | 6562873 | Jul 10, 2021 | | | |
| | | 6627210 | Jul 18, 2021 | DP | | |
| | | 6641834 | Jul 28, 2021 | DP | | |
| | | 6673337 | Jul 26, 2021 | DP | | |
| | | 9295641 | Jul 10, 2021 | | U-1833 | |
| | | 9295641*PED | Jan 10, 2022 | | | |
| <u>BRIMONIDINE TARTRATE - QOLIANA</u> | | | | | | |
| N 021764 | 001 | 7265117 | Aug 19, 2025 | DP | | |
| <u>BRIMONIDINE TARTRATE - ALPHAGAN P</u> | | | | | | |
| N 021770 | 001 | 6562873 | Jul 10, 2021 | DP | | |
| | | 6627210 | Jul 18, 2021 | DP | | |
| | | 6641834 | Jul 28, 2021 | DP | | |
| | | 6673337 | Jul 26, 2021 | DP | | |
| | | 8858961 | Sep 02, 2023 | DP | | |
| | | 8858961*PED | Mar 02, 2024 | | | |
| | | 9295641 | Jul 10, 2021 | | U-1833 | |
| | | 9295641*PED | Jan 10, 2022 | | | |
| | | 9687443 | Jul 10, 2021 | DP | | |
| | | 9687443*PED | Jan 10, 2022 | | | |
| <u>BRIMONIDINE TARTRATE - MIRVASO</u> | | | | | | |
| N 204708 | 001 | 7439241 | Aug 25, 2025 | | U-1428 | |
| | | 8053427 | Jun 13, 2031 | DP | U-1428 | |
| | | 8163725 | Jun 13, 2031 | DP | | |
| | | 8231885 | May 24, 2025 | DP | | |
| | | 8410102 | May 24, 2025 | | U-1428 | |
| | | 8426410 | May 24, 2025 | | U-1428 | |
| | | 8513247 | Mar 25, 2031 | DP | U-1428 | |
| | | 8513249 | Mar 25, 2031 | DP | U-1428 | |
| | | 8859551 | May 25, 2024 | | U-1428 | |
| | | 9861631 | Mar 25, 2031 | | U-1428 | |
| | | 9861632 | Mar 25, 2031 | | U-1428 | |
| <u>BRIMONIDINE TARTRATE - LUMIFY</u> | | | | | | |
| N 208144 | 001 | 8293742 | Jul 14, 2030 | | NP | Dec 22, 2020 |
| <u>BRIMONIDINE TARTRATE; BRINZOLAMIDE - SIMBRINZA</u> | | | | | | |
| N 204251 | 001 | 6316441 | Dec 07, 2019 | | U-778 | |
| | | 9044484 | Oct 30, 2030 | DP | | |
| | | 9421265 | Jun 17, 2030 | DP | | |
| <u>BRIMONIDINE TARTRATE; TIMOLOL MALEATE - COMBIGAN</u> | | | | | | |
| N 021398 | 001 | 7030149 | Apr 19, 2022 | | U-849 | |
| | | 7320976 | Apr 19, 2022 | | U-849 | |
| | | 7323463 | Jan 19, 2023 | DP | | |
| | | 7642258 | Apr 19, 2022 | DS DP | U-1024 | |
| | | 8133890 | Apr 19, 2022 | | U-1235 | |
| | | 8354409 | Apr 19, 2022 | DP | U-1371 | |
| | | 8748425 | Apr 19, 2022 | DP | U-1524 | |
| | | 9474751 | Apr 19, 2022 | DP | U-1524 | |
| | | 9770453 | Apr 19, 2022 | DP | U-2131 | |
| | | 9907801 | Apr 19, 2022 | DP | U-2239 | |
| | | 9907802 | Apr 19, 2022 | DP | U-2240 | |
| <u>BRIVARACETAM - BRIVIACT</u> | | | | | | |
| N 205836 | 001 | 6784197 | Feb 21, 2021 | DS DP | U-2295 | NCE |
| | | 6911461 | Feb 21, 2021 | DS DP | U-2295 | |
| | | 8492416 | Feb 21, 2021 | | U-2295 | |
| <u>BRIVARACETAM - BRIVIACT</u> | | | | | | |
| N 205836 | 002 | 6784197 | Feb 21, 2021 | DS DP | U-2295 | NCE |
| | | 6911461 | Feb 21, 2021 | DS DP | U-2295 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>BRIVARACETAM - BRIVIACT</u> | | | | | | |
| N 205836 002 | 8492416 | Feb 21, 2021 | U-2295 | | | |
| <u>BRIVARACETAM - BRIVIACT</u> | | | | | | |
| N 205836 003 | 6784197 | Feb 21, 2021 | DS DP U-2295 | | NCE | May 12, 2021 |
| | 6911461 | Feb 21, 2021 | DS DP U-2295 | | | |
| | 8492416 | Feb 21, 2021 | U-2295 | | | |
| <u>BRIVARACETAM - BRIVIACT</u> | | | | | | |
| N 205836 004 | 6784197 | Feb 21, 2021 | DS DP U-2295 | | NCE | May 12, 2021 |
| | 6911461 | Feb 21, 2021 | DS DP U-2295 | | | |
| | 8492416 | Feb 21, 2021 | U-2295 | | | |
| <u>BRIVARACETAM - BRIVIACT</u> | | | | | | |
| N 205836 005 | 6784197 | Feb 21, 2021 | DS DP U-2295 | | NCE | May 12, 2021 |
| | 6911461 | Feb 21, 2021 | DS DP U-2295 | | | |
| | 8492416 | Feb 21, 2021 | U-2295 | | | |
| <u>BRIVARACETAM - BRIVIACT</u> | | | | | | |
| N 205837 001 | 6784197 | Feb 21, 2021 | DS DP U-1815 | | NCE | May 12, 2021 |
| | 6784197 | Feb 21, 2021 | DS DP U-2130 | | | |
| | 6911461 | Feb 21, 2021 | DS DP U-1815 | | | |
| | 6911461 | Feb 21, 2021 | DS DP U-2130 | | | |
| | 8492416 | Feb 21, 2021 | U-1815 | | | |
| | 8492416 | Feb 21, 2021 | U-2130 | | | |
| <u>BRIVARACETAM - BRIVIACT</u> | | | | | | |
| N 205838 001 | 6784197 | Feb 21, 2021 | DS DP U-2295 | | NCE | May 12, 2021 |
| | 6911461 | Feb 21, 2021 | DS DP U-2295 | | | |
| | 8492416 | Feb 21, 2021 | U-2295 | | | |
| <u>BROMFENAC SODIUM - PROLENSA</u> | | | | | | |
| N 203168 001 | 10085958 | Nov 19, 2032 | DP | | | |
| | 8129431 | Sep 11, 2025 | DS DP | | | |
| | 8669290 | Jan 16, 2024 | DP | | | |
| | 8754131 | Jan 16, 2024 | DP | | | |
| | 8871813 | Jan 16, 2024 | DP | | | |
| | 8927606 | Jan 16, 2024 | U-100 | | | |
| | 8927606 | Jan 16, 2024 | U-1095 | | | |
| | 8927606 | Jan 16, 2024 | U-810 | | | |
| | 9144609 | Jan 16, 2024 | DP | | | |
| | 9517220 | Nov 11, 2033 | U-1933 | | | |
| | 9561277 | Jan 16, 2024 | U-1933 | | | |
| <u>BROMFENAC SODIUM - BROMSITE</u> | | | | | | |
| N 206911 001 | 8778999 | Aug 07, 2029 | DP U-1834 | | NP | Apr 08, 2019 |
| <u>BROMOCRIPTINE MESYLATE - CYCLOSET</u> | | | | | | |
| N 020866 001 | 7888310 | Jul 25, 2023 | U-1433 | | | |
| | 8137992 | Jul 25, 2023 | U-1433 | | | |
| | 8137993 | Jul 25, 2023 | U-1433 | | | |
| | 8137994 | Jul 25, 2023 | U-1433 | | | |
| | 8431155 | Apr 30, 2032 | DP U-976 | | | |
| | 8613947 | Apr 30, 2032 | DP U-976 | | | |
| | 8877708 | Jun 07, 2030 | DP U-1706 | | | |
| | 9192576 | Apr 30, 2032 | DP U-976 | | | |
| | 9352025 | Jun 07, 2030 | U-2111 | | | |
| | 9352025 | Jun 07, 2030 | U-2112 | | | |
| | 9352025 | Jun 07, 2030 | U-2113 | | | |
| | 9352025 | Jun 07, 2030 | U-2114 | | | |
| | 9352025 | Jun 07, 2030 | U-2115 | | | |
| | 9352025 | Jun 07, 2030 | U-2116 | | | |
| | 9352025 | Jun 07, 2030 | U-2117 | | | |
| | 9352025 | Jun 07, 2030 | U-2118 | | | |
| | 9352025 | Jun 07, 2030 | U-2119 | | | |
| | 9522117 | Apr 30, 2032 | DP U-1939 | | | |
| | 9522117 | Apr 30, 2032 | DP U-976 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2183 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2184 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2185 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>BROMOCRIPTINE MESYLATE - CYCLOSET</u> | | | | | | |
| N 020866 001 | 9700555 | Apr 30, 2032 | DP U-2186 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2187 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2188 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2189 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2190 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2191 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2192 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2193 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2194 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2195 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2196 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2197 | | | |
| | 9700555 | Apr 30, 2032 | DP U-2198 | | | |
| | 9895422 | Jun 07, 2030 | U-2114 | | | |
| | 9895422 | Jun 07, 2030 | U-2116 | | | |
| | 9895422 | Jun 07, 2030 | U-2281 | | | |
| | 9895422 | Jun 07, 2030 | U-2282 | | | |
| | 9895422 | Jun 07, 2030 | U-2283 | | | |
| | 9895422 | Jun 07, 2030 | U-2284 | | | |
| | 9895422 | Jun 07, 2030 | U-2285 | | | |
| | 9895422 | Jun 07, 2030 | U-2286 | | | |
| | 9895422 | Jun 07, 2030 | U-2287 | | | |
| | 9993474 | Apr 30, 2032 | U-2384 | | | |
| | 9993474 | Apr 30, 2032 | U-2385 | | | |
| | 9993474 | Apr 30, 2032 | U-2386 | | | |
| | 9993474 | Apr 30, 2032 | U-2387 | | | |
| | 9993474 | Apr 30, 2032 | U-2388 | | | |
| | 9993474 | Apr 30, 2032 | U-2389 | | | |
| | 9993474 | Apr 30, 2032 | U-2390 | | | |
| | 9993474 | Apr 30, 2032 | U-2391 | | | |
| | 9993474 | Apr 30, 2032 | U-2392 | | | |
| | 9993474 | Apr 30, 2032 | U-2393 | | | |
| <u>BUDESONIDE - ENTOCORT EC</u> | | | | | | |
| N 021324 001 | | | | | M-178 | Apr 29, 2019 |
| | | | | | NPP | Apr 29, 2019 |
| <u>BUDESONIDE - UCERIS</u> | | | | | | |
| N 203634 001 | 10064878 | Jun 09, 2020 | DP U-1325 | | | |
| | 10105374 | Jun 09, 2020 | DP U-1325 | | | |
| | 10143698 | Jun 09, 2020 | DP U-1325 | | | |
| | 7410651 | Jun 09, 2020 | DP U-1325 | | | |
| | 7431943 | Jun 09, 2020 | DP | | | |
| | 8293273 | Jun 09, 2020 | DP | | | |
| | 8784888 | Jun 09, 2020 | DP | | | |
| | 8895064 | Sep 07, 2031 | DP | | | |
| | 9132093 | Sep 07, 2031 | DP | | | |
| | 9192581 | Sep 07, 2031 | DP U-1325 | | | |
| | 9320716 | Jun 09, 2020 | DP U-1325 | | | |
| | 9532954 | Jun 09, 2020 | DP U-1325 | | | |
| | RE43799 | Jun 09, 2020 | DP U-1325 | | | |
| <u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u> | | | | | | |
| N 021929 001 | 10166247 | Jan 29, 2023 | DP U-2001 | | M-210 | Sep 11, 2020 |
| | 10166247 | Jan 29, 2023 | DP U-2002 | | M-214 | Dec 20, 2020 |
| | 10166247 | Jan 29, 2023 | DP U-2122 | | NPP | Jan 27, 2020 |
| | 7367333*PED | May 11, 2019 | | | PED | Jul 27, 2020 |
| | 7587988 | Apr 10, 2026 | DP | | | |
| | 7587988*PED | Oct 10, 2026 | | | | |
| | 7759328 | Jan 29, 2023 | DP U-2001 | | | |
| | 7759328 | Jan 29, 2023 | DP U-2002 | | | |
| | 7759328 | Jan 29, 2023 | DP U-2122 | | | |
| | 7759328*PED | Jul 29, 2023 | | | | |
| | 7967011 | Aug 11, 2021 | DP | | | |
| | 7967011*PED | Feb 11, 2022 | | | | |
| | 8143239 | Jan 29, 2023 | DP U-2001 | | | |
| | 8143239 | Jan 29, 2023 | DP U-2002 | | | |
| | 8143239 | Jan 29, 2023 | DP U-2122 | | | |
| | 8143239*PED | Jul 29, 2023 | | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u> | | | | | | |
| N 021929 001 | 8387615 | Mar 26, 2027 | DP | | | |
| | 8387615*PED | Sep 26, 2027 | | | | |
| | 8528545 | Oct 16, 2028 | DP | | | |
| | 8528545*PED | Apr 16, 2029 | | | | |
| | 8575137 | Jan 29, 2023 | DP U-2001 | | | |
| | 8575137 | Jan 29, 2023 | DP U-2002 | | | |
| | 8575137 | Jan 29, 2023 | DP U-2122 | | | |
| | 8575137*PED | Jul 29, 2023 | | | | |
| | 8616196 | Apr 07, 2029 | DP | | | |
| | 8616196*PED | Oct 07, 2029 | | | | |
| | 8875699 | Nov 10, 2024 | DP | | | |
| | 8875699*PED | May 10, 2025 | | | | |
| <u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u> | | | | | | |
| N 021929 002 | 10166247 | Jan 29, 2023 | DP U-2001 | | M-210 | Sep 11, 2020 |
| | 10166247 | Jan 29, 2023 | DP U-2002 | | M-214 | Dec 20, 2020 |
| | 10166247 | Jan 29, 2023 | DP U-2122 | | | |
| | 7367333*PED | May 11, 2019 | | | | |
| | 7587988 | Apr 10, 2026 | DP | | | |
| | 7587988*PED | Oct 10, 2026 | | | | |
| | 7759328 | Jan 29, 2023 | DP U-2001 | | | |
| | 7759328 | Jan 29, 2023 | DP U-2002 | | | |
| | 7759328 | Jan 29, 2023 | DP U-2122 | | | |
| | 7759328*PED | Jul 29, 2023 | | | | |
| | 7897646*PED | Mar 09, 2019 | | | | |
| | 7967011 | Aug 11, 2021 | DP | | | |
| | 7967011*PED | Feb 11, 2022 | | | | |
| | 8143239 | Jan 29, 2023 | DP U-2001 | | | |
| | 8143239 | Jan 29, 2023 | DP U-2002 | | | |
| | 8143239 | Jan 29, 2023 | DP U-2122 | | | |
| | 8143239*PED | Jul 29, 2023 | | | | |
| | 8387615 | Mar 26, 2027 | DP | | | |
| | 8387615*PED | Sep 26, 2027 | | | | |
| | 8461211*PED | Mar 09, 2019 | | | | |
| | 8528545 | Oct 16, 2028 | DP | | | |
| | 8528545*PED | Apr 16, 2029 | | | | |
| | 8575137 | Jan 29, 2023 | DP U-2001 | | | |
| | 8575137 | Jan 29, 2023 | DP U-2002 | | | |
| | 8575137 | Jan 29, 2023 | DP U-2122 | | | |
| | 8575137*PED | Jul 29, 2023 | | | | |
| | 8616196 | Apr 07, 2029 | DP | | | |
| | 8616196*PED | Oct 07, 2029 | | | | |
| | 8875699 | Nov 10, 2024 | DP | | | |
| | 8875699*PED | May 10, 2025 | | | | |
| <u>BUPIVACAINE - EXPAREL</u> | | | | | | |
| N 022496 001 | 9585838 | Dec 24, 2021 | DP | | I-771 | Apr 06, 2021 |
| <u>BUPIVACAINE - EXPAREL</u> | | | | | | |
| N 022496 002 | 9585838 | Dec 24, 2021 | DP | | I-771 | Apr 06, 2021 |
| <u>BUPRENORPHINE - SUBLOCADE</u> | | | | | | |
| N 209819 001 | 8921387 | Jan 06, 2032 | DP U-2173 | | NP | Nov 30, 2020 |
| | 8921387 | Jan 06, 2032 | DP U-2174 | | | |
| | 8975270 | Sep 05, 2031 | DP U-2175 | | | |
| | 8975270 | Sep 05, 2031 | DP U-2206 | | | |
| | 9272044 | Jun 06, 2031 | U-2176 | | | |
| | 9272044 | Jun 06, 2031 | U-2177 | | | |
| | 9272044 | Jun 06, 2031 | U-2178 | | | |
| | 9272044 | Jun 06, 2031 | U-2209 | | | |
| | 9498432 | Jun 06, 2031 | DP U-2179 | | | |
| | 9782402 | Jun 06, 2031 | DP U-2176 | | | |
| | 9782402 | Jun 06, 2031 | DP U-2180 | | | |
| | 9782402 | Jun 06, 2031 | DP U-2207 | | | |
| | 9782402 | Jun 06, 2031 | DP U-2208 | | | |
| | 9827241 | Jun 06, 2031 | DP U-2174 | | | |
| | 9827241 | Jun 06, 2031 | DP U-2181 | | | |
| | 9827241 | Jun 06, 2031 | DP U-2206 | | | |
| | 9827241 | Jun 06, 2031 | DP U-2210 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>BUPRENORPHINE - SUBLOCADE</u> | | | | | | |
| N 209819 001 | 9827241 | Jun 06, 2031 | DP U-2211 | | | |
| <u>BUPRENORPHINE - SUBLOCADE</u> | | | | | | |
| N 209819 002 | 8921387 | Jan 06, 2032 | DP U-2173 | | NP | Nov 30, 2020 |
| | 8921387 | Jan 06, 2032 | DP U-2174 | | | |
| | 8975270 | Sep 05, 2031 | DP U-2175 | | | |
| | 8975270 | Sep 05, 2031 | DP U-2206 | | | |
| | 9272044 | Jun 06, 2031 | U-2176 | | | |
| | 9272044 | Jun 06, 2031 | U-2177 | | | |
| | 9272044 | Jun 06, 2031 | U-2178 | | | |
| | 9272044 | Jun 06, 2031 | U-2209 | | | |
| | 9498432 | Jun 06, 2031 | DP U-2179 | | | |
| | 9782402 | Jun 06, 2031 | DP U-2176 | | | |
| | 9782402 | Jun 06, 2031 | DP U-2180 | | | |
| | 9782402 | Jun 06, 2031 | DP U-2207 | | | |
| | 9782402 | Jun 06, 2031 | DP U-2208 | | | |
| | 9827241 | Jun 06, 2031 | DP U-2174 | | | |
| | 9827241 | Jun 06, 2031 | DP U-2181 | | | |
| | 9827241 | Jun 06, 2031 | DP U-2206 | | | |
| | 9827241 | Jun 06, 2031 | DP U-2210 | | | |
| | 9827241 | Jun 06, 2031 | DP U-2211 | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE - PROBUPHINE</u> | | | | | | |
| N 204442 001 | 7736665 | Apr 25, 2024 | U-1878 | | NP | May 26, 2019 |
| <u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u> | | | | | | |
| N 207932 001 | 7579019 | Jan 22, 2020 | U-1769 | | | |
| | 8147866 | Jul 23, 2027 | DP U-1769 | | | |
| | 9655843 | Jul 23, 2027 | DP U-1556 | | | |
| | 9901539 | Dec 21, 2032 | U-1556 | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u> | | | | | | |
| N 207932 002 | 7579019 | Jan 22, 2020 | U-1769 | | | |
| | 8147866 | Jul 23, 2027 | DP U-1769 | | | |
| | 9655843 | Jul 23, 2027 | DP U-1556 | | | |
| | 9901539 | Dec 21, 2032 | U-1556 | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u> | | | | | | |
| N 207932 003 | 7579019 | Jan 22, 2020 | U-1769 | | | |
| | 8147866 | Jul 23, 2027 | DP U-1769 | | | |
| | 9655843 | Jul 23, 2027 | DP U-1556 | | | |
| | 9901539 | Dec 21, 2032 | U-1556 | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u> | | | | | | |
| N 207932 004 | 7579019 | Jan 22, 2020 | U-1769 | | | |
| | 8147866 | Jul 23, 2027 | DP U-1769 | | | |
| | 9655843 | Jul 23, 2027 | DP U-1556 | | | |
| | 9901539 | Dec 21, 2032 | U-1556 | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u> | | | | | | |
| N 207932 005 | 7579019 | Jan 22, 2020 | U-1769 | | | |
| | 8147866 | Jul 23, 2027 | DP U-1769 | | | |
| | 9655843 | Jul 23, 2027 | DP U-1556 | | | |
| | 9901539 | Dec 21, 2032 | U-1556 | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u> | | | | | | |
| N 207932 006 | 7579019 | Jan 22, 2020 | U-1769 | | | |
| | 8147866 | Jul 23, 2027 | DP U-1769 | | | |
| | 9655843 | Jul 23, 2027 | DP U-1556 | | | |
| | 9901539 | Dec 21, 2032 | U-1556 | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u> | | | | | | |
| N 207932 007 | 7579019 | Jan 22, 2020 | U-1769 | | | |
| | 8147866 | Jul 23, 2027 | DP U-1769 | | | |
| | 9655843 | Jul 23, 2027 | DP U-1556 | | | |
| | 9901539 | Dec 21, 2032 | U-1556 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u> | | | | | | |
| N 022410 001 | 8017150 | Feb 13, 2023 | DP | | | |
| | 8475832 | Mar 26, 2030 | DP U-1411 | | | |
| | 8603514 | Apr 03, 2024 | DP U-1464 | | | |
| | 9687454 | Aug 07, 2029 | DP U-1464 | | | |
| | 9855221 | Feb 14, 2022 | DP | | | |
| | 9931305 | Feb 14, 2022 | DP | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u> | | | | | | |
| N 022410 002 | 8017150 | Feb 13, 2023 | DP | | | |
| | 8475832 | Mar 26, 2030 | DP U-1411 | | | |
| | 8603514 | Apr 03, 2024 | DP U-1464 | | | |
| | 9687454 | Aug 07, 2029 | DP U-1464 | | | |
| | 9855221 | Feb 14, 2022 | DP | | | |
| | 9931305 | Feb 14, 2022 | DP | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u> | | | | | | |
| N 022410 003 | 8017150 | Feb 13, 2023 | DP | | | |
| | 8475832 | Mar 26, 2030 | DP U-1411 | | | |
| | 8603514 | Apr 03, 2024 | DP U-1464 | | | |
| | 9687454 | Aug 07, 2029 | DP U-1464 | | | |
| | 9855221 | Feb 14, 2022 | DP | | | |
| | 9931305 | Feb 14, 2022 | DP | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u> | | | | | | |
| N 022410 004 | 8017150 | Feb 13, 2023 | DP | | | |
| | 8475832 | Mar 26, 2030 | DP U-1411 | | | |
| | 8603514 | Apr 03, 2024 | DP U-1464 | | | |
| | 9687454 | Aug 07, 2029 | DP U-1464 | | | |
| | 9855221 | Feb 14, 2022 | DP | | | |
| | 9931305 | Feb 14, 2022 | DP | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u> | | | | | | |
| N 204242 001 | 8454996 | Sep 24, 2019 | U-1421 | | | |
| | 8470361 | May 22, 2030 | DP U-1425 | | | |
| | 8658198 | Dec 03, 2027 | DP U-1494 | | | |
| | 8940330 | Sep 18, 2032 | DP | | | |
| | 9259421 | Sep 18, 2032 | DP | | | |
| | 9439900 | Sep 18, 2032 | DP | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u> | | | | | | |
| N 204242 002 | 8454996 | Sep 24, 2019 | U-1421 | | | |
| | 8470361 | May 22, 2030 | DP U-1425 | | | |
| | 8658198 | Dec 03, 2027 | DP U-1494 | | | |
| | 8940330 | Sep 18, 2032 | DP | | | |
| | 9259421 | Sep 18, 2032 | DP | | | |
| | 9439900 | Sep 18, 2032 | DP | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u> | | | | | | |
| N 204242 003 | 8454996 | Sep 24, 2019 | U-1421 | | | |
| | 8470361 | May 22, 2030 | DP U-1425 | | | |
| | 8658198 | Dec 03, 2027 | DP U-1494 | | | |
| | 8940330 | Sep 18, 2032 | DP | | | |
| | 9259421 | Sep 18, 2032 | DP | | | |
| | 9439900 | Sep 18, 2032 | DP | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u> | | | | | | |
| N 204242 004 | 8454996 | Sep 24, 2019 | U-1421 | | | |
| | 8470361 | May 22, 2030 | DP U-1425 | | | |
| | 8658198 | Dec 03, 2027 | DP U-1494 | | | |
| | 8940330 | Sep 18, 2032 | DP | | | |
| | 9259421 | Sep 18, 2032 | DP | | | |
| | 9439900 | Sep 18, 2032 | DP | | | |
| <u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u> | | | | | | |
| N 204242 005 | 8454996 | Sep 24, 2019 | U-1421 | | | |
| | 8470361 | May 22, 2030 | DP U-1425 | | | |
| | 8658198 | Dec 03, 2027 | DP U-1494 | | | |
| | 8940330 | Sep 18, 2032 | DP | | | |
| | 9259421 | Sep 18, 2032 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u> | | | | | | |
| N 204242 | 005 | 9439900 | Sep 18, 2032 | DP | | |
| <u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u> | | | | | | |
| N 204242 | 006 | 8454996 | Sep 24, 2019 | | U-1421 | |
| | | 8470361 | May 22, 2030 | DP | U-1425 | |
| | | 8658198 | Dec 03, 2027 | DP | U-1494 | |
| | | 8940330 | Sep 18, 2032 | DP | | |
| | | 9259421 | Sep 18, 2032 | DP | | |
| | | 9439900 | Sep 18, 2032 | DP | | |
| | | | | | Y | |
| <u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - BUNAVAIL</u> | | | | | | |
| N 205637 | 001 | 7579019 | Jan 22, 2020 | | U-1521 | |
| | | 8147866 | Jul 23, 2027 | DP | U-1521 | |
| | | 8703177 | Aug 20, 2032 | DP | | |
| | | 9522188 | Apr 24, 2035 | DP | | |
| | | 9655843 | Jul 23, 2027 | DP | U-2017 | |
| <u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - BUNAVAIL</u> | | | | | | |
| N 205637 | 002 | 7579019 | Jan 22, 2020 | | U-1521 | |
| | | 8147866 | Jul 23, 2027 | DP | U-1521 | |
| | | 8703177 | Aug 20, 2032 | DP | | |
| | | 9522188 | Apr 24, 2035 | DP | | |
| | | 9655843 | Jul 23, 2027 | DP | U-2017 | |
| <u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - BUNAVAIL</u> | | | | | | |
| N 205637 | 003 | 7579019 | Jan 22, 2020 | | U-1521 | |
| | | 8147866 | Jul 23, 2027 | DP | U-1521 | |
| | | 8703177 | Aug 20, 2032 | DP | | |
| | | 9522188 | Apr 24, 2035 | DP | | |
| | | 9655843 | Jul 23, 2027 | DP | U-2017 | |
| <u>BUPROPION HYDROBROMIDE - APLENZIN</u> | | | | | | |
| N 022108 | 001 | 7241805 | Jun 27, 2026 | DP | | |
| | | 7569610 | Jun 27, 2026 | | U-997 | |
| | | 7572935 | Jun 27, 2026 | DP | | |
| | | 7585897 | Jun 27, 2026 | DP | | |
| | | 7645802 | Jun 27, 2026 | DP | | |
| | | 7649019 | Jun 27, 2026 | DP | | |
| | | 7662407 | Jun 27, 2026 | DP | | |
| | | 7671094 | Jun 27, 2026 | DP | | |
| <u>BUPROPION HYDROBROMIDE - APLENZIN</u> | | | | | | |
| N 022108 | 002 | 7241805 | Jun 27, 2026 | DP | | |
| | | 7569610 | Jun 27, 2026 | | U-997 | |
| | | 7572935 | Jun 27, 2026 | DP | | |
| | | 7585897 | Jun 27, 2026 | DP | | |
| | | 7645802 | Jun 27, 2026 | DP | | |
| | | 7649019 | Jun 27, 2026 | DP | | |
| | | 7662407 | Jun 27, 2026 | DP | | |
| | | 7671094 | Jun 27, 2026 | DP | | |
| <u>BUPROPION HYDROBROMIDE - APLENZIN</u> | | | | | | |
| N 022108 | 003 | 7241805 | Jun 27, 2026 | DP | | |
| | | 7569610 | Jun 27, 2026 | | U-997 | |
| | | 7572935 | Jun 27, 2026 | DP | | |
| | | 7585897 | Jun 27, 2026 | DP | | |
| | | 7645802 | Jun 27, 2026 | DP | | |
| | | 7649019 | Jun 27, 2026 | DP | | |
| | | 7662407 | Jun 27, 2026 | DP | | |
| | | 7671094 | Jun 27, 2026 | DP | | |
| <u>BUPROPION HYDROCHLORIDE - FORFIVO XL</u> | | | | | | |
| N 022497 | 001 | 7674479 | Jun 25, 2027 | DP | | |
| <u>BUPROPION HYDROCHLORIDE; NALTREXONE HYDROCHLORIDE - CONTRAVE</u> | | | | | | |
| N 200063 | 001 | 7375111 | Mar 26, 2025 | DP | | |
| | | 7462626 | Jul 20, 2024 | | U-1583 | |
| | | 8088786 | Feb 03, 2029 | DP | | |
| | | 8318788 | Nov 08, 2027 | | U-1584 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>BUPROPION HYDROCHLORIDE; NALTREXONE HYDROCHLORIDE - CONTRAVE</u> | | | | | | |
| N 200063 | 001 | 8722085 | Nov 08, 2027 | U-1585 | | |
| | | 8815889 | Jul 20, 2024 | U-1586 | | |
| | | 8916195 | Feb 02, 2030 | U-1639 | | |
| | | 9107837 | Jun 04, 2027 | U-1639 | | |
| | | 9125868 | Nov 08, 2027 | U-1585 | | |
| | | 9248123 | Jan 13, 2032 | U-1808 | | |
| <u>CABAZITAXEL - JEVTANA KIT</u> | | | | | | |
| N 201023 | 001 | 5847170 | Mar 26, 2021 | DS DP | M-201 | May 17, 2020 |
| | | 5847170*PED | Sep 26, 2021 | | M-209 | Sep 14, 2020 |
| | | 7241907 | Dec 10, 2025 | DS | PED | Nov 17, 2020 |
| | | 7241907*PED | Jun 10, 2026 | | | |
| | | 8927592 | Oct 27, 2030 | U-1630 | | |
| | | 8927592*PED | Apr 27, 2031 | | | |
| <u>CABOZANTINIB S-MALATE - COMETRIO</u> | | | | | | |
| N 203756 | 001 | 7579473 | Aug 14, 2026 | DS DP | ODE-33 | Nov 29, 2019 |
| | | 8877776 | Oct 08, 2030 | DS DP | U-1617 | |
| | | 9717720 | Feb 10, 2032 | DP | | |
| <u>CABOZANTINIB S-MALATE - COMETRIO</u> | | | | | | |
| N 203756 | 002 | 7579473 | Aug 14, 2026 | DS DP | ODE-33 | Nov 29, 2019 |
| | | 8877776 | Oct 08, 2030 | DS DP | U-1617 | |
| | | 9717720 | Feb 10, 2032 | DP | | |
| <u>CABOZANTINIB S-MALATE - CABOMETYX</u> | | | | | | |
| N 208692 | 001 | 10039757 | Jul 18, 2031 | | I-760 | Dec 19, 2020 |
| | | 7579473 | Aug 14, 2026 | DS DP | NP | Apr 25, 2019 |
| | | 8497284 | Sep 24, 2024 | | U-1220 | |
| | | 8877776 | Oct 08, 2030 | DS DP | | |
| | | 9724342 | Jul 09, 2033 | DP | | |
| <u>CABOZANTINIB S-MALATE - CABOMETYX</u> | | | | | | |
| N 208692 | 002 | 10039757 | Jul 18, 2031 | | I-760 | Dec 19, 2020 |
| | | 7579473 | Aug 14, 2026 | DS DP | NP | Apr 25, 2019 |
| | | 8497284 | Sep 24, 2024 | | U-1220 | |
| | | 8877776 | Oct 08, 2030 | DS DP | | |
| | | 9724342 | Jul 09, 2033 | DP | | |
| <u>CABOZANTINIB S-MALATE - CABOMETYX</u> | | | | | | |
| N 208692 | 003 | 10039757 | Jul 18, 2031 | | I-760 | Dec 19, 2020 |
| | | 7579473 | Aug 14, 2026 | DS DP | NP | Apr 25, 2019 |
| | | 8497284 | Sep 24, 2024 | | U-1220 | |
| | | 8877776 | Oct 08, 2030 | DS DP | | |
| | | 9724342 | Jul 09, 2033 | DP | | |
| <u>CALCIFEDIOL - RAYALDEE</u> | | | | | | |
| N 208010 | 001 | 6582727 | Aug 22, 2020 | DP | NP | Jun 17, 2019 |
| | | 8207149 | Apr 25, 2028 | | U-1871 | |
| | | 8361488 | Jul 19, 2028 | DP | | |
| | | 8426391 | Aug 27, 2028 | | U-1872 | |
| | | 8778373 | Apr 25, 2028 | | U-1873 | |
| | | 8906410 | Feb 02, 2027 | DP | | |
| | | 9408858 | Apr 25, 2028 | | U-1888 | |
| | | 9498486 | Apr 25, 2028 | | U-1920 | |
| | | 9861644 | Mar 14, 2034 | DP | | |
| | | 9925147 | Apr 25, 2028 | DP | U-2255 | |
| | | 9925147 | Apr 25, 2028 | DP | U-2256 | |
| | | 9925147 | Apr 25, 2028 | DP | U-2257 | |
| | | 9925147 | Apr 25, 2028 | DP | U-2258 | |
| | | 9925147 | Apr 25, 2028 | DP | U-2259 | |
| | | 9943530 | Feb 02, 2027 | | U-2274 | |
| <u>CALCIPOTRIENE - SORILUX</u> | | | | | | |
| N 022563 | 001 | 8263580 | Sep 27, 2028 | DP | U-1280 | |
| | | 8629128 | May 26, 2026 | DP | U-1280 | |
| | | 8629128 | May 26, 2026 | DP | U-1767 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>CALCITONIN SALMON RECOMBINANT - FORTICAL</u> | | | | | | |
| N 021406 | 001 | 6440392 | Feb 02, 2021 | DP U-227 | | |
| | | RE40812 | Feb 02, 2021 | DP | | |
| | | RE43580 | Feb 02, 2021 | DP U-227 | | |
| <u>CALCITRIOL - CALCIJEX</u> | | | | | | |
| N 018874 | 001 | 6051567 | Aug 02, 2019 | | | |
| | | 6265392 | Aug 02, 2019 | | | |
| | | 6274169 | Aug 02, 2019 | | | |
| <u>CALCITRIOL - CALCIJEX</u> | | | | | | |
| N 018874 | 002 | 6051567 | Aug 02, 2019 | | | |
| | | 6265392 | Aug 02, 2019 | | | |
| | | 6274169 | Aug 02, 2019 | | | |
| <u>CALCIUM ACETATE - PHOSLO</u> | | | | | | |
| N 021160 | 002 | 6576665 | Apr 03, 2021 | | | |
| <u>CALCIUM ACETATE - PHOSLO GELCAPS</u> | | | | | | |
| N 021160 | 003 | 6576665 | Apr 03, 2021 | | | |
| | | 6875445 | Jul 30, 2021 | DP | | |
| <u>CALCIUM ACETATE - PHOSLYRA</u> | | | | | | |
| N 022581 | 001 | 8591938 | Feb 23, 2030 | DP U-1469 | | |
| | | 8592480 | Jul 20, 2027 | U-1469 | | |
| | | 9089528 | Jul 20, 2027 | U-1469 | | |
| <u>CALCIUM CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE - PEPCID COMPLETE</u> | | | | | | |
| N 020958 | 001 | 6814978 | Aug 26, 2021 | DP | | |
| <u>CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; OXIGLUTATIONE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE - NAVSTEL</u> | | | | | | |
| N 022193 | 001 | 7084130 | Nov 29, 2021 | DP U-891 | | |
| <u>CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE - PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER</u> | | | | | | |
| N 207026 | 001 | | | | ODE-85 | Jan 13, 2022 |
| <u>CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE - PHOXILLUM B22K 4/0 IN PLASTIC CONTAINER</u> | | | | | | |
| N 207026 | 002 | | | | ODE-85 | Jan 13, 2022 |
| <u>CALCIUM GLUCONATE - CALCIUM GLUCONATE IN SODIUM CHLORIDE</u> | | | | | | |
| N 210906 | 001 | 10130646 | Jan 11, 2038 | DP | | |
| <u>CALCIUM GLUCONATE - CALCIUM GLUCONATE IN SODIUM CHLORIDE</u> | | | | | | |
| N 210906 | 002 | 10130646 | Jan 11, 2038 | DP | | |
| <u>CANAGLIFLOZIN - INVOKANA</u> | | | | | | |
| N 204042 | 001 | 7943582 | Feb 26, 2029 | DS DP U-2441 | I-733 | May 20, 2019 |
| | | 7943582 | Feb 26, 2029 | DS DP U-493 | I-788 | Oct 29, 2021 |
| | | 7943788 | Jul 14, 2027 | DS DP | M-197 | Feb 01, 2020 |
| | | 8222219 | Apr 11, 2025 | U-2441 | | |
| | | 8222219 | Apr 11, 2025 | U-493 | | |
| | | 8513202 | Dec 03, 2027 | DS DP U-2441 | | |
| | | 8513202 | Dec 03, 2027 | DS DP U-493 | | |
| <u>CANAGLIFLOZIN - INVOKANA</u> | | | | | | |
| N 204042 | 002 | 7943582 | Feb 26, 2029 | DS DP U-2441 | I-733 | May 20, 2019 |
| | | 7943582 | Feb 26, 2029 | DS DP U-493 | I-788 | Oct 29, 2021 |
| | | 7943788 | Jul 14, 2027 | DS DP | M-197 | Feb 01, 2020 |
| | | 8222219 | Apr 11, 2025 | U-2441 | | |
| | | 8222219 | Apr 11, 2025 | U-493 | | |
| | | 8513202 | Dec 03, 2027 | DS DP U-2441 | | |
| | | 8513202 | Dec 03, 2027 | DS DP U-493 | | |
| <u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u> | | | | | | |
| N 204353 | 001 | 7943582 | Feb 26, 2029 | DS DP U-2441 | I-735 | May 20, 2019 |
| | | 7943582 | Feb 26, 2029 | DS DP U-493 | I-788 | Oct 29, 2021 |
| | | 7943788 | Jul 14, 2027 | DS DP | M-197 | Feb 01, 2020 |
| | | 8222219 | Apr 11, 2025 | U-2441 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u> | | | | | | |
| N 204353 001 | 8222219 | Apr 11, 2025 | | U-493 | | |
| | 8513202 | Dec 03, 2027 | DS DP | U-2441 | | |
| | 8513202 | Dec 03, 2027 | DS DP | U-493 | | |
| | 8785403 | Jul 30, 2024 | | DP | | |
| <u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u> | | | | | | |
| N 204353 002 | 7943582 | Feb 26, 2029 | DS DP | U-2441 | I-735 | May 20, 2019 |
| | 7943582 | Feb 26, 2029 | DS DP | U-493 | I-788 | Oct 29, 2021 |
| | 7943788 | Jul 14, 2027 | DS DP | | M-197 | Feb 01, 2020 |
| | 8222219 | Apr 11, 2025 | | U-2441 | | |
| | 8222219 | Apr 11, 2025 | | U-493 | | |
| | 8513202 | Dec 03, 2027 | DS DP | U-2441 | | |
| | 8513202 | Dec 03, 2027 | DS DP | U-493 | | |
| | 8785403 | Jul 30, 2024 | | DP | | |
| <u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u> | | | | | | |
| N 204353 003 | 7943582 | Feb 26, 2029 | DS DP | U-2441 | I-735 | May 20, 2019 |
| | 7943582 | Feb 26, 2029 | DS DP | U-493 | I-788 | Oct 29, 2021 |
| | 7943788 | Jul 14, 2027 | DS DP | | M-197 | Feb 01, 2020 |
| | 8222219 | Apr 11, 2025 | | U-2441 | | |
| | 8222219 | Apr 11, 2025 | | U-493 | | |
| | 8513202 | Dec 03, 2027 | DS DP | U-2441 | | |
| | 8513202 | Dec 03, 2027 | DS DP | U-493 | | |
| | 8785403 | Jul 30, 2024 | | DP | | |
| <u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u> | | | | | | |
| N 204353 004 | 7943582 | Feb 26, 2029 | DS DP | U-2441 | I-735 | May 20, 2019 |
| | 7943582 | Feb 26, 2029 | DS DP | U-493 | I-788 | Oct 29, 2021 |
| | 7943788 | Jul 14, 2027 | DS DP | | M-197 | Feb 01, 2020 |
| | 8222219 | Apr 11, 2025 | | U-2441 | | |
| | 8222219 | Apr 11, 2025 | | U-493 | | |
| | 8513202 | Dec 03, 2027 | DS DP | U-2441 | | |
| | 8513202 | Dec 03, 2027 | DS DP | U-493 | | |
| | 8785403 | Jul 30, 2024 | | DP | | |
| <u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u> | | | | | | |
| N 205879 001 | 6723340 | Oct 25, 2021 | | DP | I-735 | May 20, 2019 |
| | 7943582 | Feb 26, 2029 | DS DP | U-2441 | I-788 | Oct 29, 2021 |
| | 7943582 | Feb 26, 2029 | DS DP | U-493 | | |
| | 7943788 | Jul 14, 2027 | DS DP | | | |
| | 8222219 | Apr 11, 2025 | | U-2441 | | |
| | 8222219 | Apr 11, 2025 | | U-493 | | |
| | 8513202 | Dec 03, 2027 | DS DP | U-2441 | | |
| | 8513202 | Dec 03, 2027 | DS DP | U-493 | | |
| | 8785403 | Jul 30, 2024 | | DP | | |
| <u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u> | | | | | | |
| N 205879 002 | 7943582 | Feb 26, 2029 | DS DP | U-2441 | I-735 | May 20, 2019 |
| | 7943582 | Feb 26, 2029 | DS DP | U-493 | I-788 | Oct 29, 2021 |
| | 7943788 | Jul 14, 2027 | DS DP | | | |
| | 8222219 | Apr 11, 2025 | | U-2441 | | |
| | 8222219 | Apr 11, 2025 | | U-493 | | |
| | 8513202 | Dec 03, 2027 | DS DP | U-2441 | | |
| | 8513202 | Dec 03, 2027 | DS DP | U-493 | | |
| | 8785403 | Jul 30, 2024 | | DP | | |
| <u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u> | | | | | | |
| N 205879 003 | 6723340 | Oct 25, 2021 | | DP | I-735 | May 20, 2019 |
| | 7943582 | Feb 26, 2029 | DS DP | U-2441 | I-788 | Oct 29, 2021 |
| | 7943582 | Feb 26, 2029 | DS DP | U-493 | | |
| | 7943788 | Jul 14, 2027 | DS DP | | | |
| | 8222219 | Apr 11, 2025 | | U-2441 | | |
| | 8222219 | Apr 11, 2025 | | U-493 | | |
| | 8513202 | Dec 03, 2027 | DS DP | U-2441 | | |
| | 8513202 | Dec 03, 2027 | DS DP | U-493 | | |
| | 8785403 | Jul 30, 2024 | | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</u> | | | | | | |
| N 205879 | 004 | 7943582 | Feb 26, 2029 | DS DP U-2441 | I-735 | May 20, 2019 |
| | | 7943582 | Feb 26, 2029 | DS DP U-493 | I-788 | Oct 29, 2021 |
| | | 7943788 | Jul 14, 2027 | DS DP | | |
| | | 8222219 | Apr 11, 2025 | | U-2441 | |
| | | 8222219 | Apr 11, 2025 | | U-493 | |
| | | 8513202 | Dec 03, 2027 | DS DP U-2441 | | |
| | | 8513202 | Dec 03, 2027 | DS DP U-493 | | |
| | | 8785403 | Jul 30, 2024 | DP | | |
| <u>CANGRELOR - KENGREAL</u> | | | | | | |
| N 204958 | 001 | 10039780 | Jul 10, 2035 | | U-2260 | NCE Jun 22, 2020 |
| | | 6130208 | Jun 29, 2023 | DP | U-1715 | |
| | | 8680052 | Mar 09, 2033 | | U-1715 | |
| | | 8759316 | May 13, 2029 | | U-1715 | |
| | | 9295687 | Jul 10, 2035 | DP | | |
| | | 9427448 | Nov 10, 2030 | | U-1926 | |
| | | 9439921 | Jul 10, 2035 | DP | | |
| | | 9700575 | Jul 10, 2035 | DP | | |
| | | 9925265 | May 13, 2029 | | U-2260 | |
| <u>CANNABIDIOL - EPIDIOLEX</u> | | | | | | |
| N 210365 | 001 | 10092525 | Jun 17, 2035 | | U-2427 | NCE Sep 28, 2023 |
| | | 10111840 | Jun 17, 2035 | | U-2442 | ODE-216 Sep 28, 2025 |
| | | 10111840 | Jun 17, 2035 | | U-2443 | |
| | | 10137095 | Jun 17, 2035 | | U-2454 | |
| | | 10137095 | Jun 17, 2035 | | U-2455 | |
| | | 9949937 | Jun 17, 2035 | | U-2421 | |
| | | 9956183 | Jun 17, 2035 | | U-2422 | |
| | | 9956183 | Jun 17, 2035 | | U-2423 | |
| | | 9956184 | Jun 17, 2035 | | U-2424 | |
| | | 9956185 | Jun 17, 2035 | | U-2425 | |
| | | 9956186 | Jun 17, 2035 | | U-2426 | |
| <u>CAPSAICIN - OUTENZA</u> | | | | | | |
| N 022395 | 001 | 6239180 | Jun 04, 2021 | DP | | |
| <u>CARBAMAZEPINE - EQUETRO</u> | | | | | | |
| N 021710 | 001 | 6977253 | May 19, 2024 | | U-693 | |
| <u>CARBAMAZEPINE - EQUETRO</u> | | | | | | |
| N 021710 | 002 | 6977253 | May 19, 2024 | | U-693 | |
| <u>CARBAMAZEPINE - EQUETRO</u> | | | | | | |
| N 021710 | 003 | 6977253 | May 19, 2024 | | U-693 | |
| <u>CARBAMAZEPINE - CARNEXIV</u> | | | | | | |
| N 206030 | 001 | 7635773 | Mar 13, 2029 | DP | | ODE-124 Oct 07, 2023 |
| | | 8410077 | Mar 13, 2029 | DP | | |
| | | 9493582 | Feb 27, 2033 | DP | | |
| | | 9629797 | Nov 10, 2028 | | U-2004 | |
| | | 9629797 | Nov 10, 2028 | | U-2005 | |
| | | 9629797 | Nov 10, 2028 | | U-2006 | |
| | | 9750822 | Mar 13, 2029 | DP | | |
| | | 9770407 | Nov 10, 2028 | DP | | |
| <u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 50</u> | | | | | | |
| N 021485 | 001 | 6500867 | Jun 29, 2020 | DP | U-219 | |
| | | 6797732 | Jun 29, 2020 | DP | | |
| <u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 100</u> | | | | | | |
| N 021485 | 002 | 6500867 | Jun 29, 2020 | DP | U-219 | |
| | | 6797732 | Jun 29, 2020 | DP | | |
| <u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 150</u> | | | | | | |
| N 021485 | 003 | 6500867 | Jun 29, 2020 | DP | U-219 | |
| | | 6797732 | Jun 29, 2020 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 200</u> | | | | | | |
| N 021485 | 004 | 6500867 | Jun 29, 2020 | DP U-219 | | |
| | | 6797732 | Jun 29, 2020 | DP | | |
| <u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 75</u> | | | | | | |
| N 021485 | 005 | 6500867 | Jun 29, 2020 | DP U-219 | | |
| | | 6797732 | Jun 29, 2020 | DP | | |
| <u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 125</u> | | | | | | |
| N 021485 | 006 | 6500867 | Jun 29, 2020 | DP U-219 | | |
| | | 6797732 | Jun 29, 2020 | DP | | |
| <u>CARBIDOPA; LEVODOPA - RYTARY</u> | | | | | | |
| N 203312 | 001 | 7094427 | May 29, 2022 | DP U-1645 | Y | |
| | | 8377474 | Dec 26, 2028 | DP U-1645 | | |
| | | 8377474 | Dec 26, 2028 | DP U-219 | | |
| | | 8454998 | Dec 26, 2028 | DP U-1645 | | |
| | | 8454998 | Dec 26, 2028 | DP U-1646 | | |
| | | 8454998 | Dec 26, 2028 | DP U-1647 | | |
| | | 8454998 | Dec 26, 2028 | DP U-1649 | | |
| | | 8454998 | Dec 26, 2028 | DP U-219 | | |
| | | 8557283 | Dec 26, 2028 | DP U-1645 | | |
| | | 8557283 | Dec 26, 2028 | DP U-219 | | |
| | | 9089607 | Dec 26, 2028 | DP U-1645 | | |
| | | 9089607 | Dec 26, 2028 | DP U-1720 | | |
| | | 9089608 | Dec 26, 2028 | DP | | |
| | | 9463246 | Dec 26, 2028 | DP U-219 | | |
| | | 9533046 | Dec 26, 2028 | DP U-219 | | |
| | | 9901640 | Dec 26, 2028 | DP U-219 | | |
| <u>CARBIDOPA; LEVODOPA - RYTARY</u> | | | | | | |
| N 203312 | 002 | 7094427 | May 29, 2022 | DP U-1645 | Y | |
| | | 8377474 | Dec 26, 2028 | DP U-1645 | | |
| | | 8377474 | Dec 26, 2028 | DP U-219 | | |
| | | 8454998 | Dec 26, 2028 | DP U-1645 | | |
| | | 8454998 | Dec 26, 2028 | DP U-1646 | | |
| | | 8454998 | Dec 26, 2028 | DP U-1647 | | |
| | | 8454998 | Dec 26, 2028 | DP U-1649 | | |
| | | 8454998 | Dec 26, 2028 | DP U-219 | | |
| | | 8557283 | Dec 26, 2028 | DP U-1645 | | |
| | | 8557283 | Dec 26, 2028 | DP U-219 | | |
| | | 9089607 | Dec 26, 2028 | DP U-1645 | | |
| | | 9089607 | Dec 26, 2028 | DP U-1720 | | |
| | | 9089608 | Dec 26, 2028 | DP | | |
| | | 9463246 | Dec 26, 2028 | DP U-219 | | |
| | | 9533046 | Dec 26, 2028 | DP U-219 | | |
| | | 9901640 | Dec 26, 2028 | DP U-219 | | |
| <u>CARBIDOPA; LEVODOPA - RYTARY</u> | | | | | | |
| N 203312 | 003 | 7094427 | May 29, 2022 | DP U-1645 | Y | |
| | | 8377474 | Dec 26, 2028 | DP U-1645 | | |
| | | 8377474 | Dec 26, 2028 | DP U-219 | | |
| | | 8454998 | Dec 26, 2028 | DP U-1645 | | |
| | | 8454998 | Dec 26, 2028 | DP U-1646 | | |
| | | 8454998 | Dec 26, 2028 | DP U-1647 | | |
| | | 8454998 | Dec 26, 2028 | DP U-1649 | | |
| | | 8454998 | Dec 26, 2028 | DP U-219 | | |
| | | 8557283 | Dec 26, 2028 | DP U-1645 | | |
| | | 8557283 | Dec 26, 2028 | DP U-219 | | |
| | | 9089607 | Dec 26, 2028 | DP U-1645 | | |
| | | 9089607 | Dec 26, 2028 | DP U-1720 | | |
| | | 9089608 | Dec 26, 2028 | DP | | |
| | | 9463246 | Dec 26, 2028 | DP U-219 | | |
| | | 9533046 | Dec 26, 2028 | DP U-219 | | |
| | | 9901640 | Dec 26, 2028 | DP U-219 | | |
| <u>CARBIDOPA; LEVODOPA - RYTARY</u> | | | | | | |
| N 203312 | 004 | 7094427 | May 29, 2022 | DP U-1645 | Y | |
| | | 8377474 | Dec 26, 2028 | DP U-1645 | | |
| | | 8377474 | Dec 26, 2028 | DP U-219 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>CARBIDOPA; LEVODOPA - RYTARY</u> | | | | | | |
| N 203312 004 | 8454998 | Dec 26, 2028 | DP U-1645 | | | |
| | 8454998 | Dec 26, 2028 | DP U-1646 | | | |
| | 8454998 | Dec 26, 2028 | DP U-1647 | | | |
| | 8454998 | Dec 26, 2028 | DP U-1649 | | | |
| | 8454998 | Dec 26, 2028 | DP U-219 | | | |
| | 8557283 | Dec 26, 2028 | DP U-1645 | | | |
| | 8557283 | Dec 26, 2028 | DP U-219 | | | |
| | 9089607 | Dec 26, 2028 | DP U-1645 | | | |
| | 9089607 | Dec 26, 2028 | DP U-1720 | | | |
| | 9089608 | Dec 26, 2028 | DP | | | |
| | 9463246 | Dec 26, 2028 | DP U-219 | | | |
| | 9533046 | Dec 26, 2028 | DP U-219 | | | |
| | 9901640 | Dec 26, 2028 | DP U-219 | | | |
| <u>CARBIDOPA; LEVODOPA - DUOPA</u> | | | | | | |
| N 203952 001 | | | | | ODE-84 | Jan 09, 2022 |
| <u>CARBINOXAMINE MALEATE - KARBINAL ER</u> | | | | | | |
| N 022556 001 | 8062667 | Mar 29, 2029 | DP | | | |
| | 9522191 | Jun 15, 2027 | DP | | | |
| <u>CARFILZOMIB - KYPROLIS</u> | | | | | | |
| N 202714 001 | 7232818 | Apr 14, 2025 | DS DP | | D-172 | Sep 28, 2021 |
| | 7417042 | Jul 20, 2026 | DS DP | | I-722 | Jan 21, 2019 |
| | 7491704 | Apr 14, 2025 | | U-1260 | I-723 | Jan 21, 2019 |
| | 7737112 | Dec 07, 2027 | DP | | ODE-27 | Jul 20, 2019 |
| | 8129346 | Apr 14, 2025 | | U-1260 | | |
| | 8207125 | Apr 14, 2025 | DS DP | | | |
| | 8207126 | Apr 14, 2025 | DP | | | |
| | 8207127 | Apr 14, 2025 | | U-1260 | | |
| | 8207297 | Apr 14, 2025 | DS DP | | | |
| | 9493582 | Feb 27, 2033 | DP | | | |
| | 9511109 | Oct 21, 2029 | | U-1924 | | |
| <u>CARFILZOMIB - KYPROLIS</u> | | | | | | |
| N 202714 002 | 7232818 | Apr 14, 2025 | DS DP | | D-172 | Sep 28, 2021 |
| | 7417042 | Jul 20, 2026 | DS DP | | I-722 | Jan 21, 2019 |
| | 7491704 | Apr 14, 2025 | | U-1260 | I-723 | Jan 21, 2019 |
| | 7737112 | Dec 07, 2027 | DP | | ODE-27 | Jul 20, 2019 |
| | 8129346 | Apr 14, 2025 | | U-1260 | | |
| | 8207125 | Apr 14, 2025 | DS DP | | | |
| | 8207126 | Apr 14, 2025 | DP | | | |
| | 8207127 | Apr 14, 2025 | | U-1260 | | |
| | 8207297 | Apr 14, 2025 | DS DP | | | |
| | 9493582 | Feb 27, 2033 | DP | | | |
| | 9511109 | Oct 21, 2029 | | U-1924 | | |
| <u>CARFILZOMIB - KYPROLIS</u> | | | | | | |
| N 202714 003 | 7232818 | Apr 14, 2025 | DS DP | | D-172 | Sep 28, 2021 |
| | 7417042 | Jul 20, 2026 | DS DP | | | |
| | 7491704 | Apr 14, 2025 | | U-2319 | | |
| | 7491704 | Apr 14, 2025 | | U-2320 | | |
| | 7737112 | Dec 07, 2027 | DP | | | |
| | 8129346 | Apr 14, 2025 | | U-2319 | | |
| | 8129346 | Apr 14, 2025 | | U-2320 | | |
| | 8207125 | Apr 14, 2025 | DS DP | | | |
| | 8207126 | Apr 14, 2025 | DP | | | |
| | 8207127 | Apr 14, 2025 | | U-2319 | | |
| | 8207127 | Apr 14, 2025 | | U-2320 | | |
| | 8207297 | Apr 14, 2025 | DS DP | | | |
| | 9493582 | Feb 27, 2033 | DP | | | |
| | 9511109 | Oct 21, 2029 | | U-1924 | | |
| <u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u> | | | | | | |
| N 204370 001 | 7737142 | Mar 27, 2027 | DS DP U-1750 | | M-213 | Nov 09, 2020 |
| | 7943621 | Dec 16, 2028 | DS DP | | NCE | Sep 17, 2020 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u> | | | | | | |
| N 204370 002 | 7737142 | Mar 27, 2027 | DS DP U-1750 | | M-213 | Nov 09, 2020 |
| | 7943621 | Dec 16, 2028 | DS DP | | NCE | Sep 17, 2020 |
| <u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u> | | | | | | |
| N 204370 003 | 7737142 | Mar 27, 2027 | DS DP U-1750 | | M-213 | Nov 09, 2020 |
| | 7943621 | Dec 16, 2028 | DS DP | | NCE | Sep 17, 2020 |
| <u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u> | | | | | | |
| N 204370 004 | 7737142 | Mar 27, 2027 | DS DP U-1750 | | M-213 | Nov 09, 2020 |
| | 7943621 | Dec 16, 2028 | DS DP | | NCE | Sep 17, 2020 |
| <u>CARVEDILOL PHOSPHATE - COREG CR</u> | | | | | | |
| N 022012 001 | 7268156 | Jun 27, 2023 | DS DP U-3 | | | |
| | 7268156 | Jun 27, 2023 | DS DP U-313 | | | |
| | 8101209 | Sep 11, 2025 | DP | | | |
| <u>CARVEDILOL PHOSPHATE - COREG CR</u> | | | | | | |
| N 022012 002 | 7268156 | Jun 27, 2023 | DS DP U-3 | | | |
| | 7268156 | Jun 27, 2023 | DS DP U-313 | | | |
| | 8101209 | Sep 11, 2025 | DP | | | |
| <u>CARVEDILOL PHOSPHATE - COREG CR</u> | | | | | | |
| N 022012 003 | 7268156 | Jun 27, 2023 | DS DP U-3 | | | |
| | 7268156 | Jun 27, 2023 | DS DP U-313 | | | |
| | 8101209 | Sep 11, 2025 | DP | | | |
| <u>CARVEDILOL PHOSPHATE - COREG CR</u> | | | | | | |
| N 022012 004 | 7268156 | Jun 27, 2023 | DS DP U-3 | | | |
| | 7268156 | Jun 27, 2023 | DS DP U-313 | | | |
| | 8101209 | Sep 11, 2025 | DP | | | |
| <u>CASPOFUNGIN ACETATE - CASPOFUNGIN ACETATE</u> | | | | | | |
| N 206110 001 | 9636407 | Dec 21, 2032 | DP | | | |
| <u>CASPOFUNGIN ACETATE - CASPOFUNGIN ACETATE</u> | | | | | | |
| N 206110 002 | 9636407 | Dec 21, 2032 | DP | | | |
| <u>CEFIXIME - SUPRAX</u> | | | | | | |
| N 202091 001 | 9233112 | Dec 14, 2028 | DP U-1676 | | | |
| <u>CEFTAROLINE FOSAMIL - TEFLARO</u> | | | | | | |
| N 200327 001 | 6417175 | Apr 11, 2022 | DS DP U-1676 | | NPP | May 27, 2019 |
| | 6906055 | Dec 15, 2021 | DS DP | | NPP | May 27, 2019 |
| | 7419973 | Dec 15, 2021 | DP | | | |
| | 8247400 | Feb 10, 2031 | DP U-282 | | | |
| | 9629861 | Sep 21, 2030 | DP | | | |
| <u>CEFTAROLINE FOSAMIL - TEFLARO</u> | | | | | | |
| N 200327 002 | 6417175 | Apr 11, 2022 | DS DP U-1676 | | NPP | May 27, 2019 |
| | 6906055 | Dec 15, 2021 | DS DP | | NPP | May 27, 2019 |
| | 7419973 | Dec 15, 2021 | DP | | | |
| | 8247400 | Feb 10, 2031 | DP U-282 | | | |
| | 9629861 | Sep 21, 2030 | DP | | | |
| <u>CEFTOLOZANE SULFATE; TAZOBACTAM SODIUM - ZERBAXA</u> | | | | | | |
| N 206829 001 | 10125149 | Aug 14, 2035 | DP | | NCE | Dec 19, 2019 |
| | 7129232 | Oct 21, 2024 | DS DP U-36 | | GAIN | Dec 19, 2024 |
| | 8476425 | Sep 27, 2032 | DS | | | |
| | 8685957 | Sep 27, 2032 | DS U-36 | | | |
| | 8906898 | May 28, 2034 | DS DP | | | |
| | 8968753 | Mar 14, 2034 | U-1672 | | | |
| | 8968753 | Mar 14, 2034 | U-1673 | | | |
| | 9320740 | Mar 14, 2034 | DP | | | |
| | 9872906 | Mar 14, 2034 | DP | | | |
| <u>CERITINIB - ZYKADIA</u> | | | | | | |
| N 205755 001 | 7153964 | Feb 26, 2021 | DS DP | | M-199 | May 26, 2020 |
| | 7893074 | Apr 25, 2026 | DS DP | | NCE | Apr 29, 2019 |
| | 7964592 | Jan 13, 2027 | DS DP | | ODE-145 | May 26, 2024 |
| | 8039479 | Jun 29, 2030 | DS DP | | ODE-66 | Apr 29, 2021 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>CERITINIB - ZYKADIA</u> | | | | | | |
| N 205755 | 001 | 8188276 | Jan 31, 2023 | DS DP | | |
| | | 8377921 | Nov 20, 2027 | | U-1179 | |
| | | 8399450 | Nov 20, 2027 | DS DP | | |
| | | 8703787 | Feb 02, 2032 | | U-1179 | |
| | | 8835430 | Jan 31, 2023 | DS DP | | |
| | | 9018204 | Jan 31, 2023 | DS DP | | |
| | | 9309229 | Jan 18, 2032 | DS DP | | |
| | | 9416112 | Jan 31, 2023 | DS DP | | |
| <u>CETIRIZINE HYDROCHLORIDE - ZERVIAE</u> | | | | | | |
| N 208694 | 001 | 8829005 | Mar 15, 2030 | | U-1680 | NDF May 30, 2020 |
| | | 8829005*PED | Sep 15, 2030 | | | PED Nov 30, 2020 |
| | | 9254286 | Jul 09, 2032 | DP | | |
| | | 9254286*PED | Jan 09, 2033 | | | |
| | | 9750684 | Mar 15, 2030 | DP | | |
| | | 9993471 | Mar 15, 2030 | | U-1680 | |
| <u>CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ZYRTEC-D 12 HOUR</u> | | | | | | |
| N 021150 | 002 | 6469009 | Jul 13, 2019 | DP | U-295 | |
| | | 7014867 | Jun 10, 2022 | DP | | |
| | | 7226614 | Jun 10, 2022 | | U-295 | |
| <u>CETRORELIX - CETROTIDE</u> | | | | | | |
| N 021197 | 001 | 6319192 | Apr 23, 2019 | | U-426 | |
| <u>CETRORELIX - CETROTIDE</u> | | | | | | |
| N 021197 | 002 | 6319192 | Apr 23, 2019 | | U-426 | |
| <u>CHLORHEXIDINE GLUCONATE - CHLORHEXIDINE GLUCONATE</u> | | | | | | |
| N 021669 | 001 | 7066916 | Feb 17, 2024 | | U-737 | |
| | | 7427574 | Apr 25, 2026 | DP | | |
| | | 7595021 | May 12, 2023 | DP | U-1022 | |
| | | 7717889 | Feb 27, 2025 | DP | U-1022 | |
| | | 7935093 | Oct 02, 2027 | DP | U-1022 | |
| <u>CHLORHEXIDINE GLUCONATE - READYPREP CHG</u> | | | | | | |
| N 207964 | 001 | | | | | NF Nov 20, 2021 |
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u> | | | | | | |
| N 020832 | 001 | 6536975 | Nov 10, 2020 | DP | | |
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u> | | | | | | |
| N 020832 | 002 | 6729786 | Mar 14, 2023 | DP | | |
| | | 6991394 | Jan 31, 2024 | DP | | |
| | | 7182536 | Dec 30, 2023 | DP | | |
| | | 7241065 | Mar 14, 2023 | DP | | |
| | | 7422388 | Apr 25, 2027 | DP | U-1397 | |
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u> | | | | | | |
| N 020832 | 004 | 6536975 | Nov 10, 2020 | DP | | |
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u> | | | | | | |
| N 020832 | 005 | 6536975 | Nov 10, 2020 | DP | | |
| | | 6729786 | Mar 14, 2023 | DP | | |
| | | 7241065 | Mar 14, 2023 | DP | | |
| | | 7422388 | Apr 25, 2027 | DP | U-1397 | |
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u> | | | | | | |
| N 020832 | 006 | 6991394 | Jan 31, 2024 | DP | | |
| | | 7182536 | Dec 30, 2023 | DP | | |
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u> | | | | | | |
| N 020832 | 007 | 6536975 | Nov 10, 2020 | DP | | |
| | | 6729786 | Mar 14, 2023 | DP | | |
| | | 7241065 | Mar 14, 2023 | DP | | |
| | | 7422388 | Apr 25, 2027 | DP | U-1397 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - PREVANTICS SWAB</u> | | | | | | |
| N 021524 | 001 | | | | M-221 | Feb 14, 2021 |
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - PREVANTICS SWABSTICK</u> | | | | | | |
| N 021524 | 002 | | | | M-221 | Feb 14, 2021 |
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - PREVANTICS MAXI SWABSTICK</u> | | | | | | |
| N 021524 | 003 | | | | M-221 | Feb 14, 2021 |
| <u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - SOLUPREP</u> | | | | | | |
| N 208288 | 001 | 8623935 | Jul 26, 2029 | DP U-1022 | NP | Aug 08, 2021 |
| <u>CHLOROPROCAINE HYDROCHLORIDE - CLOROTEKAL</u> | | | | | | |
| N 208791 | 001 | | | | NP | Sep 26, 2020 |
| <u>CHLORPHENIRAMINE MALEATE; CODEINE PHOSPHATE - TUXARIN ER</u> | | | | | | |
| N 206323 | 001 | 6248363 | Nov 23, 2019 | DP U-1716 | | |
| | | 6383471 | Apr 06, 2019 | DP U-1716 | | |
| | | 9066942 | Jan 03, 2032 | U-1716 | | |
| | | 9107921 | Jan 03, 2032 | DP | | |
| <u>CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE - ADVIL ALLERGY SINUS</u> | | | | | | |
| N 021441 | 001 | 7863287 | Feb 28, 2027 | DP | | |
| <u>CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX - TUZISTRA XR</u> | | | | | | |
| N 207768 | 001 | 8062667 | Mar 29, 2029 | DP | | |
| | | 8790700 | Mar 15, 2027 | DP | | |
| <u>CHOLIC ACID - CHOLBAM</u> | | | | | | |
| N 205750 | 001 | | | | NCE ODE-91 | Mar 17, 2020 Mar 17, 2022 |
| <u>CHOLIC ACID - CHOLBAM</u> | | | | | | |
| N 205750 | 002 | | | | NCE ODE-91 | Mar 17, 2020 Mar 17, 2022 |
| <u>CHOLINE FENOFIBRATE - TRILIPIX</u> | | | | | | |
| N 022224 | 001 | 7259186 | Jan 07, 2025 | DS | | |
| <u>CHOLINE FENOFIBRATE - TRILIPIX</u> | | | | | | |
| N 022224 | 002 | 7259186 | Jan 07, 2025 | DS | | |
| <u>CHORIOGONADOTROPIN ALFA - OVIDREL</u> | | | | | | |
| N 021149 | 002 | 6706681 | Mar 16, 2021 | DP | | |
| <u>CICLESONIDE - ALVESCO</u> | | | | | | |
| N 021658 | 002 | 8371292 | Feb 01, 2028 | U-1355 | | |
| <u>CICLESONIDE - ALVESCO</u> | | | | | | |
| N 021658 | 003 | 8371292 | Feb 01, 2028 | U-1355 | | |
| <u>CICLESONIDE - OMNARIS</u> | | | | | | |
| N 022004 | 001 | 6767901 | Oct 21, 2020 | DP | | |
| | | 6939559 | Apr 21, 2019 | DP | | |
| | | 7235247 | Apr 21, 2019 | DP | | |
| | | 8371292 | Feb 01, 2028 | U-1356 | | |
| | | 8383611 | Oct 20, 2020 | DP | | |
| <u>CICLESONIDE - ZETONNA</u> | | | | | | |
| N 202129 | 001 | 8371292 | Feb 01, 2028 | U-1357 | | |
| <u>CINACALCET HYDROCHLORIDE - SENSIPAR</u> | | | | | | |
| N 021688 | 001 | 7829595 | Sep 22, 2026 | DP U-1098 | M-200 | May 23, 2020 |
| | | 9375405 | Sep 22, 2026 | DP | ODE-78 | Nov 21, 2021 |
| <u>CINACALCET HYDROCHLORIDE - SENSIPAR</u> | | | | | | |
| N 021688 | 002 | 7829595 | Sep 22, 2026 | DP U-1098 | M-200 | May 23, 2020 |
| | | 9375405 | Sep 22, 2026 | DP | ODE-78 | Nov 21, 2021 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>CINACALCET HYDROCHLORIDE - SENSIPAR</u> | | | | | | |
| N 021688 | 003 | 7829595 | Sep 22, 2026 | DP U-1098 | M-200 | May 23, 2020 |
| | | 9375405 | Sep 22, 2026 | DP | ODE-78 | Nov 21, 2021 |
| <u>CIPROFLOXACIN - OTIPRIO</u> | | | | | | |
| N 207986 | 001 | 8318817 | Apr 27, 2030 | U-1792 | I-770 | Mar 02, 2021 |
| | | 9205048 | Apr 21, 2029 | U-1793 | | |
| | | 9220796 | Jul 01, 2035 | DP | | |
| | | 9233068 | Dec 11, 2029 | DP | | |
| | | 9603796 | Apr 21, 2029 | DS DP U-2252 | | |
| <u>CIPROFLOXACIN HYDROCHLORIDE - PROQUIN XR</u> | | | | | | |
| N 021744 | 001 | 6488962 | Jun 20, 2020 | DP | | |
| <u>CIPROFLOXACIN HYDROCHLORIDE; FLUOCINOLONE ACETONIDE - OTOVEL</u> | | | | | | |
| N 208251 | 001 | 8932610 | Mar 24, 2030 | DP U-1578 | NC | Apr 29, 2019 |
| <u>CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE - CIPRO XR</u> | | | | | | |
| N 021473 | 001 | 7709022 | Jun 23, 2021 | DP | | |
| | | 8187632 | Jun 23, 2021 | DP | | |
| | | 8187632*PED | Dec 23, 2021 | | | |
| <u>CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE - CIPRO XR</u> | | | | | | |
| N 021473 | 002 | 7709022 | Jun 23, 2021 | DP | | |
| | | 8187632 | Jun 23, 2021 | DP | | |
| | | 8187632*PED | Dec 23, 2021 | | | |
| <u>CIPROFLOXACIN; DEXAMETHASONE - CIPRODEX</u> | | | | | | |
| N 021537 | 001 | 6284804 | Aug 10, 2020 | | | |
| | | 6359016 | Aug 10, 2020 | | | |
| | | 8846650 | Jun 04, 2025 | DP U-1578 | | |
| | | 9149486 | Sep 13, 2022 | DP U-1578 | | |
| | | 9345714 | Sep 13, 2022 | DP U-1578 | | |
| | | 9402805 | Sep 13, 2022 | DP U-1578 | | |
| | | 9402805 | Sep 13, 2022 | DP U-1679 | | |
| <u>CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE - PREPOPIK</u> | | | | | | |
| N 202535 | 001 | 8450338 | Oct 10, 2028 | DP | NPP | Aug 15, 2021 |
| | | 8481083 | Oct 10, 2028 | DP | | |
| <u>CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE - CLENPIO</u> | | | | | | |
| N 209589 | 001 | 9827231 | Jun 23, 2034 | DP U-2162 | | |
| <u>CLEVIDIPINE - CLEVIPREX</u> | | | | | | |
| N 022156 | 001 | 10010537 | Oct 10, 2031 | DP | | |
| | | 5856346 | Jan 05, 2021 | DS DP U-893 | | |
| | | 8658676 | Oct 10, 2031 | DP | | |
| <u>CLEVIDIPINE - CLEVIPREX</u> | | | | | | |
| N 022156 | 002 | 10010537 | Oct 10, 2031 | DP | | |
| | | 5856346 | Jan 05, 2021 | DS DP U-893 | | |
| | | 8658676 | Oct 10, 2031 | DP | | |
| <u>CLEVIDIPINE - CLEVIPREX</u> | | | | | | |
| N 022156 | 003 | 10010537 | Oct 10, 2031 | DP | | |
| | | 5856346 | Jan 05, 2021 | DS DP U-893 | | |
| | | 8658676 | Oct 10, 2031 | DP | | |
| <u>CLINDAMYCIN PHOSPHATE - CLEOCIN</u> | | | | | | |
| N 050767 | 001 | 6495157 | Jul 20, 2020 | DP | | |
| <u>CLINDAMYCIN PHOSPHATE - CLINDAGEL</u> | | | | | | |
| N 050782 | 001 | 6387383 | Aug 03, 2020 | DP U-818 | | |
| <u>CLINDAMYCIN PHOSPHATE - CLINDESSE</u> | | | | | | |
| N 050793 | 001 | 6899890 | Apr 27, 2023 | DP U-137 | | |
| | | 9789057 | Dec 02, 2026 | DP U-137 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------|--------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>CLINDAMYCIN PHOSPHATE - EVOCLIN</u> | | | | | | |
| N 050801 | 001 7141237 | Jan 23, 2024 | DS DP | | | |
| | 7374747 | Aug 09, 2026 | DS DP U-921 | | | |
| <u>CLINDAMYCIN PHOSPHATE; TRETINOIN - ZIANA</u> | | | | | | |
| N 050802 | 001 6387383 | Aug 03, 2020 | DP U-916 | | | |
| <u>CLOBAZAM - SYMPAZAN</u> | | | | | | |
| N 210833 | 001 8603514 | Apr 03, 2024 | DP | | | |
| | 8765167 | Feb 20, 2024 | DP | | | |
| <u>CLOBAZAM - SYMPAZAN</u> | | | | | | |
| N 210833 | 002 8603514 | Apr 03, 2024 | DP | | | |
| | 8765167 | Feb 20, 2024 | DP | | | |
| <u>CLOBAZAM - SYMPAZAN</u> | | | | | | |
| N 210833 | 003 8603514 | Apr 03, 2024 | DP | | | |
| | 8765167 | Feb 20, 2024 | DP | | | |
| <u>CLOBETASOL PROPIONATE - CLOBEX</u> | | | | | | |
| N 021644 | 001 7316810 | Jun 17, 2019 | DP | | | |
| | 7700081 | Jan 03, 2022 | U-1044 | | | |
| | 8066975 | Jun 17, 2019 | DP | | | |
| | 8066976 | Jun 17, 2019 | DP | | | |
| <u>CLOBETASOL PROPIONATE - OLUX E</u> | | | | | | |
| N 022013 | 001 6730288 | Sep 08, 2019 | DP | | | |
| | 7029659 | Sep 08, 2019 | DP | | | |
| | 8460641 | Nov 05, 2028 | DP U-1410 | | | |
| | 8962000 | Aug 31, 2025 | DP U-1410 | | | |
| <u>CLOBETASOL PROPIONATE - IMPOYZ</u> | | | | | | |
| N 209483 | 001 10064875 | Aug 31, 2030 | DP U-1408 | | NP | Nov 28, 2020 |
| | 10064875 | Aug 31, 2030 | DP U-1858 | | | |
| | 10064875 | Aug 31, 2030 | DP U-193 | | | |
| | 10064875 | Aug 31, 2030 | DP U-742 | | | |
| | 10064875 | Aug 31, 2030 | DP U-88 | | | |
| | 9855334 | Mar 11, 2035 | DP | | | |
| | 9956231 | Aug 31, 2030 | DP U-1408 | | | |
| | 9956231 | Aug 31, 2030 | DP U-1761 | | | |
| | 9956231 | Aug 31, 2030 | DP U-1858 | | | |
| | 9956231 | Aug 31, 2030 | DP U-193 | | | |
| | 9956231 | Aug 31, 2030 | DP U-742 | | | |
| | 9956231 | Aug 31, 2030 | DP U-88 | | | |
| <u>CLOPIDOGREL BISULFATE - PLAVIX</u> | | | | | | |
| N 020839 | 001 6429210 | Jun 10, 2019 | DS DP | | | |
| | 6504030 | Jun 10, 2019 | DS | | | |
| <u>CLOPIDOGREL BISULFATE - PLAVIX</u> | | | | | | |
| N 020839 | 002 6429210 | Jun 10, 2019 | DS DP | | | |
| | 6504030 | Jun 10, 2019 | DS | | | |
| <u>COBICISTAT - TYBOST</u> | | | | | | |
| N 203094 | 001 10039718 | Oct 04, 2032 | DP | | | |
| | 8148374 | Sep 03, 2029 | DS DP U-1279 | | | |
| <u>COBICISTAT; DARUNAVIR ETHANOLATE - PREZCOBIX</u> | | | | | | |
| N 205395 | 001 10039718 | Oct 04, 2032 | DP | | | |
| | 7470506 | Jun 23, 2019 | U-1660 | | | |
| | 7470506*PED | Dec 23, 2019 | | | | |
| | 7700645 | Dec 26, 2026 | DS DP | | | |
| | 7700645*PED | Jun 26, 2027 | | | | |
| | 8148374 | Sep 03, 2029 | DS DP U-1279 | | | |
| | 8518987 | Feb 16, 2024 | DS DP | | | |
| | 8518987*PED | Aug 16, 2024 | | | | |
| | 8597876 | Jun 23, 2019 | U-1660 | | | |
| | 8597876*PED | Dec 23, 2019 | | | | |
| | 9889115 | Jun 23, 2019 | U-1660 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>COBICISTAT; DARUNAVIR ETHANOLATE; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - SYMTUZA</u> | | | | | | |
| N 210455 | 001 | 10039718 | Oct 04, 2032 | DP | NC | Jul 17, 2020 |
| | | 6642245 | Nov 04, 2020 | U-2352 | NCE | Nov 05, 2020 |
| | | 6703396 | Mar 09, 2021 | DS DP | | |
| | | 7390791 | May 07, 2022 | DS DP | | |
| | | 7470506 | Jun 23, 2019 | U-2352 | | |
| | | 7700645 | Dec 26, 2026 | DS DP | | |
| | | 7803788 | Feb 02, 2022 | U-2352 | | |
| | | 8148374 | Sep 03, 2029 | DS DP U-2353 | | |
| | | 8148374 | Sep 03, 2029 | DS DP U-2364 | | |
| | | 8148374 | Sep 03, 2029 | DS DP U-2365 | | |
| | | 8518987 | Feb 16, 2024 | DS DP | | |
| | | 8597876 | Jun 23, 2019 | U-2352 | | |
| | | 8754065 | Aug 15, 2032 | DS DP U-2352 | | |
| | | 9296769 | Aug 15, 2032 | DS DP U-2352 | | |
| | | 9889115 | Jun 23, 2019 | U-2352 | | |
| <u>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - GENVOYA</u> | | | | | | |
| N 207561 | 001 | 10039718 | Oct 04, 2032 | DP | D-173 | Dec 10, 2021 |
| | | 6642245 | Nov 04, 2020 | U-257 | NCE | Nov 05, 2020 |
| | | 6642245*PED | May 04, 2021 | | NPP | Sep 25, 2020 |
| | | 6703396 | Mar 09, 2021 | DS DP | | |
| | | 6703396*PED | Sep 09, 2021 | | | |
| | | 7176220 | Aug 27, 2026 | DS DP U-257 | | |
| | | 7390791 | May 07, 2022 | DS DP | | |
| | | 7635704 | Oct 26, 2026 | DS DP U-257 | | |
| | | 7803788 | Feb 02, 2022 | U-257 | | |
| | | 8148374 | Sep 03, 2029 | DS DP U-1279 | | |
| | | 8633219 | Apr 24, 2030 | DP U-257 | | |
| | | 8754065 | Aug 15, 2032 | DS DP U-257 | | |
| | | 8981103 | Oct 26, 2026 | DS DP | | |
| | | 9296769 | Aug 15, 2032 | DS DP U-257 | | |
| | | 9891239 | Sep 03, 2029 | DP U-257 | | |
| <u>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - STRIBILD</u> | | | | | | |
| N 203100 | 001 | 10039718 | Oct 04, 2032 | DP | NPP | Jan 27, 2020 |
| | | 6642245 | Nov 04, 2020 | U-257 | | |
| | | 6703396 | Mar 09, 2021 | DS DP | | |
| | | 7176220 | Aug 27, 2026 | DS DP U-257 | | |
| | | 7635704 | Oct 26, 2026 | DS DP U-257 | | |
| | | 8148374 | Sep 03, 2029 | DS DP U-1279 | | |
| | | 8592397 | Jan 13, 2024 | DP U-257 | | |
| | | 8633219 | Apr 24, 2030 | DP U-257 | | |
| | | 8716264 | Jan 13, 2024 | DP U-257 | | |
| | | 8981103 | Oct 26, 2026 | DS DP | | |
| | | 9457036 | Jan 13, 2024 | DP U-257 | | |
| | | 9744181 | Jan 13, 2024 | DP U-257 | | |
| | | 9891239 | Sep 03, 2029 | DP U-257 | | |
| <u>COBIMETINIB FUMARATE - COTELLIC</u> | | | | | | |
| N 206192 | 001 | 7803839 | Feb 01, 2027 | DS DP | NCE | Nov 10, 2020 |
| | | 8362002 | Oct 05, 2026 | U-1776 | ODE-101 | Nov 10, 2022 |
| <u>COCAINE HYDROCHLORIDE - GOPRELTO</u> | | | | | | |
| N 209963 | 001 | 10016407 | Feb 07, 2037 | U-2329 | NCE | Dec 14, 2022 |
| | | 10149843 | Feb 07, 2037 | U-2478 | | |
| | | 10149843 | Feb 07, 2037 | U-2479 | | |
| | | 9867815 | Feb 07, 2037 | U-2225 | | |
| | | 9867815 | Feb 07, 2037 | U-2226 | | |
| | | 9867815 | Feb 07, 2037 | U-2227 | | |
| <u>COLCHICINE - COLCRYS</u> | | | | | | |
| N 022352 | 001 | 7601758 | Feb 10, 2029 | U-1007 | | |
| | | 7619004 | Dec 03, 2028 | U-1020 | | |
| | | 7820681 | Feb 17, 2029 | U-1020 | | |
| | | 7906519 | Feb 17, 2029 | U-1116 | | |
| | | 7915269 | Feb 17, 2029 | U-1007 | | |
| | | 7935731 | Dec 03, 2028 | U-1116 | | |
| | | 7964647 | Oct 06, 2028 | U-1007 | | |
| | | 7964648 | Oct 06, 2028 | U-1161 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>COLCHICINE - COLCRYS</u> | | | | | | |
| N 022352 | 001 | 7981938 | Oct 06, 2028 | U-1166 | | |
| | | 8093296 | Oct 06, 2028 | U-1007 | | |
| | | 8093297 | Oct 06, 2028 | U-1161 | | |
| | | 8093298 | Oct 06, 2028 | U-1116 | | |
| | | 8097655 | Oct 06, 2028 | U-1020 | | |
| | | 8415395 | Oct 06, 2028 | U-1007 | | |
| | | 8415396 | Oct 06, 2028 | U-1007 | | |
| | | 8440721 | Feb 17, 2029 | U-1007 | | |
| | | 8440722 | Feb 17, 2029 | U-1020 | | |
| <u>COLCHICINE - MITIGARE</u> | | | | | | |
| N 204820 | 001 | 8927607 | Aug 22, 2033 | U-1020 | | |
| | | 9399036 | Aug 22, 2033 | U-1020 | | |
| | | 9555029 | Aug 22, 2033 | U-1020 | | |
| | | 9675613 | Aug 22, 2033 | U-1020 | | |
| | | 9789108 | Aug 22, 2033 | U-1020 | | |
| <u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u> | | | | | | |
| N 021176 | 001 | 7229613 | Apr 17, 2022 | U-851 | | |
| <u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u> | | | | | | |
| N 022362 | 001 | 7229613 | Apr 17, 2022 | U-493 | | |
| <u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u> | | | | | | |
| N 022362 | 002 | 7229613 | Apr 17, 2022 | U-493 | | |
| <u>CONIVAPTAN HYDROCHLORIDE - VAPRISOL</u> | | | | | | |
| N 021697 | 001 | 5723606 | Dec 15, 2019 | DS DP U-698 | | |
| | | 5723606 | Dec 15, 2019 | DS DP U-868 | | |
| <u>CONIVAPTAN HYDROCHLORIDE - VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER</u> | | | | | | |
| N 021697 | 002 | 5723606 | Dec 15, 2019 | DS DP U-698 | | |
| | | 5723606 | Dec 15, 2019 | DS DP U-868 | | |
| <u>COPANLISIB DIHYDROCHLORIDE - ALIOOPA</u> | | | | | | |
| N 209936 | 001 | 7511041 | May 13, 2024 | DS DP | NCE | Sep 14, 2022 |
| | | 9636344 | Mar 29, 2032 | U-2124 | ODE-155 | Sep 14, 2024 |
| | | RE46856 | Oct 22, 2029 | DS DP U-2124 | | |
| <u>CRISABOROLE - EUCRISA</u> | | | | | | |
| N 207695 | 001 | 8039451 | Jun 11, 2026 | DS DP | NCE | Dec 14, 2021 |
| | | 8168614 | Jan 20, 2030 | U-1932 | | |
| | | 8501712 | Feb 16, 2027 | U-1932 | | |
| | | 9682092 | Feb 16, 2027 | U-1932 | | |
| <u>CRIZOTINIB - XALKORI</u> | | | | | | |
| N 202570 | 001 | 7230098 | Aug 26, 2025 | DS | ODE-111 | Mar 11, 2023 |
| | | 7825137 | May 12, 2027 | U-1179 | | |
| | | 7858643 | Oct 08, 2029 | DS DP | | |
| | | 8217057 | Nov 06, 2029 | DS DP | | |
| | | 8785632 | Mar 01, 2025 | DS | | |
| <u>CRIZOTINIB - XALKORI</u> | | | | | | |
| N 202570 | 002 | 7230098 | Aug 26, 2025 | DS | ODE-111 | Mar 11, 2023 |
| | | 7825137 | May 12, 2027 | U-1179 | | |
| | | 7858643 | Oct 08, 2029 | DS DP | | |
| | | 8217057 | Nov 06, 2029 | DS DP | | |
| | | 8785632 | Mar 01, 2025 | DS | | |
| <u>CROFELEMER - MYTESI</u> | | | | | | |
| N 202292 | 001 | 8962680 | Oct 31, 2031 | U-1319 | | |
| | | 9585868 | Oct 31, 2031 | DS U-1319 | | |
| <u>CYANOCOBALAMIN - NASCOBAL</u> | | | | | | |
| N 021642 | 001 | 7229636 | Aug 01, 2024 | DP U-817 | | |
| | | 7404489 | Mar 12, 2024 | DP | | |
| | | 7879349 | Aug 01, 2024 | DP U-1152 | | |
| | | 8003353 | Aug 01, 2024 | U-817 | | |
| | | 8940714 | Feb 26, 2024 | U-1152 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>CYANOCOBALAMIN - NASCOBAL</u> | | | | | | |
| N 021642 | 001 | 9415007 | Jul 28, 2024 | U-1896 | | |
| <u>CYCLOBENZAPRINE HYDROCHLORIDE - AMRIX</u> | | | | | | |
| N 021777 | 001 | 7387793 | Feb 26, 2025 | DP | | |
| | | 7544372 | Nov 14, 2023 | U-979 | | |
| | | 7790199 | Nov 14, 2023 | DP | | |
| | | 7820203 | Nov 14, 2023 | DP | | |
| | | 7829121 | Nov 14, 2023 | U-1088 | | |
| | | 8877245 | Nov 14, 2023 | U-979 | | |
| | | 9375410 | Nov 14, 2023 | U-1088 | | |
| | | 9399025 | Nov 14, 2023 | DP U-979 | | |
| <u>CYCLOBENZAPRINE HYDROCHLORIDE - AMRIX</u> | | | | | | |
| N 021777 | 002 | 7387793 | Feb 26, 2025 | DP | | |
| | | 7544372 | Nov 14, 2023 | U-979 | | |
| | | 7790199 | Nov 14, 2023 | DP | | |
| | | 7820203 | Nov 14, 2023 | DP | | |
| | | 7829121 | Nov 14, 2023 | U-1088 | | |
| | | 8877245 | Nov 14, 2023 | U-979 | | |
| | | 9375410 | Nov 14, 2023 | U-1088 | | |
| | | 9399025 | Nov 14, 2023 | DP U-979 | | |
| <u>CYCLOSPORINE - RESTASIS</u> | | | | | | |
| N 050790 | 001 | 8629111 | Aug 27, 2024 | DP | | |
| | | 8633162 | Aug 27, 2024 | U-1479 | | |
| | | 8642556 | Aug 27, 2024 | DP | | |
| | | 8648048 | Aug 27, 2024 | U-1483 | | |
| | | 8685930 | Aug 27, 2024 | DP | | |
| | | 9248191 | Aug 27, 2024 | U-1479 | | |
| <u>CYCLOSPORINE - RESTASIS MULTIDOSE</u> | | | | | | |
| N 050790 | 002 | 8292129 | Feb 25, 2031 | DP | | |
| | | 8561859 | Apr 16, 2032 | DP | | |
| | | 8629111 | Aug 27, 2024 | DP | | |
| | | 8633162 | Aug 27, 2024 | U-1479 | | |
| | | 8642556 | Aug 27, 2024 | DP | | |
| | | 8648048 | Aug 27, 2024 | U-1483 | | |
| | | 8685930 | Aug 27, 2024 | DP | | |
| | | 9248191 | Aug 27, 2024 | U-1479 | | |
| | | 9669974 | May 11, 2034 | DP | | |
| | | 9676525 | Feb 07, 2034 | DP | | |
| <u>CYCLOSPORINE - CEQUA</u> | | | | | | |
| N 210913 | 001 | 8980839 | Aug 23, 2033 | DP U-1483 | | |
| | | 9937225 | Aug 23, 2033 | DP U-1483 | | |
| <u>CYSTEAMINE BITARTRATE - PROCYSBI</u> | | | | | | |
| N 203389 | 001 | 10143665 | Aug 16, 2036 | U-1399 | M-216 | Dec 22, 2020 |
| | | 8026284 | Sep 22, 2027 | U-1399 | ODE-162 | Dec 22, 2024 |
| | | 8026284*PED | Mar 22, 2028 | | ODE-45 | Apr 30, 2020 |
| | | 9173851 | Jun 17, 2034 | DP | ODE-97 | Aug 14, 2022 |
| | | 9173851*PED | Dec 17, 2034 | | PED | Oct 30, 2020 |
| | | 9192590 | Jan 26, 2027 | U-1399 | PED | Jun 22, 2021 |
| | | 9192590*PED | Jul 26, 2027 | | PED | Feb 14, 2023 |
| | | 9198882 | Jan 26, 2027 | U-1399 | | |
| | | 9198882*PED | Jul 26, 2027 | | | |
| | | 9233077 | Jun 17, 2034 | DP | | |
| | | 9233077*PED | Dec 17, 2034 | | | |
| | | 9925156 | Jan 26, 2027 | DS DP U-1399 | | |
| | | 9925157 | Jan 26, 2027 | DS DP U-1399 | | |
| | | 9925158 | Jan 26, 2027 | DS DP U-1399 | | |
| <u>CYSTEAMINE BITARTRATE - PROCYSBI</u> | | | | | | |
| N 203389 | 002 | 10143665 | Aug 16, 2036 | U-1399 | M-216 | Dec 22, 2020 |
| | | 8026284 | Sep 22, 2027 | U-1399 | ODE-162 | Dec 22, 2024 |
| | | 8026284*PED | Mar 22, 2028 | | ODE-45 | Apr 30, 2020 |
| | | 9173851 | Jun 17, 2034 | DP | ODE-97 | Aug 14, 2022 |
| | | 9173851*PED | Dec 17, 2034 | | PED | Oct 30, 2020 |
| | | 9192590 | Jan 26, 2027 | U-1399 | PED | Jun 22, 2021 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>CYSTEAMINE BITARTRATE - PROCYSBI</u> | | | | | | |
| N 203389 | 002 | 9192590*PED | Jul 26, 2027 | | PED | Feb 14, 2023 |
| | | 9198882 | Jan 26, 2027 | U-1399 | | |
| | | 9198882*PED | Jul 26, 2027 | | | |
| | | 9233077 | Jun 17, 2034 | DP | | |
| | | 9233077*PED | Dec 17, 2034 | | | |
| | | 9925156 | Jan 26, 2027 | DS DP U-1399 | | |
| | | 9925157 | Jan 26, 2027 | DS DP U-1399 | | |
| | | 9925158 | Jan 26, 2027 | DS DP U-1399 | | |
| <u>CYSTEAMINE HYDROCHLORIDE - CYSTARAN</u> | | | | | | |
| N 200740 | 001 | | | | ODE-31 | Oct 02, 2019 |
| <u>CYTARABINE; DAUNORUBICIN - VYXEOS</u> | | | | | | |
| N 209401 | 001 | 10028912 | Sep 29, 2034 | DP U-2341 | NP | Aug 03, 2020 |
| | | 10028912 | Sep 29, 2034 | DP U-2342 | | |
| | | 7850990 | Jan 23, 2027 | DP U-2090 | | |
| | | 8022279 | Sep 14, 2027 | DP U-2090 | | |
| | | 8092828 | Apr 01, 2029 | U-2090 | | |
| | | 8431806 | Apr 22, 2025 | DP U-2090 | | |
| | | 8518437 | Jun 07, 2026 | DP | | |
| | | 9271931 | Jan 23, 2027 | DP | | |
| <u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u> | | | | | | |
| N 022512 | 001 | 6087380 | Dec 28, 2021 | DS DP U-1931 | | |
| | | 7866474 | Aug 31, 2027 | DP | Y | |
| | | 7932273 | Sep 07, 2025 | DS DP | | |
| | | 9034822 | Jan 20, 2031 | U-1759 | | |
| | | 9925174 | Jun 14, 2023 | DP | | |
| <u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u> | | | | | | |
| N 022512 | 002 | 6087380 | Dec 28, 2021 | DS DP U-1931 | | |
| | | 7866474 | Aug 31, 2027 | DP | Y | |
| | | 7932273 | Sep 07, 2025 | DS DP | | |
| | | 9034822 | Jan 20, 2031 | U-1759 | | |
| | | 9925174 | Jun 14, 2023 | DP | | |
| <u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u> | | | | | | |
| N 022512 | 003 | 6087380 | Dec 28, 2021 | DS DP U-1931 | | |
| | | 7866474 | Aug 31, 2027 | DP | Y | |
| | | 7932273 | Sep 07, 2025 | DS DP | | |
| | | 9034822 | Jan 20, 2031 | U-1759 | | |
| | | 9925174 | Jun 14, 2023 | DP | | |
| <u>DABRAFENIB MESYLATE - TAFINLAR</u> | | | | | | |
| N 202806 | 001 | 7994185 | Jan 20, 2030 | DS DP U-1406 | I-745 | Jun 22, 2020 |
| | | 7994185 | Jan 20, 2030 | DS DP U-2031 | I-778 | Apr 30, 2021 |
| | | 7994185 | Jan 20, 2030 | DS DP U-2032 | I-781 | May 04, 2021 |
| | | 7994185 | Jan 20, 2030 | DS DP U-2296 | ODE-147 | Jun 22, 2024 |
| | | 8415345 | Jan 20, 2030 | DS DP U-1406 | ODE-182 | Apr 30, 2025 |
| | | 8415345 | Jan 20, 2030 | DS DP U-2031 | ODE-183 | May 04, 2025 |
| | | 8415345 | Jan 20, 2030 | DS DP U-2032 | ODE-47 | May 29, 2020 |
| | | 8415345 | Jan 20, 2030 | DS DP U-2296 | ODE-58 | Jan 09, 2021 |
| | | 8703781 | Oct 15, 2030 | DS DP U-1713 | | |
| | | 8703781 | Oct 15, 2030 | DS DP U-2032 | | |
| | | 8703781 | Oct 15, 2030 | DS DP U-2296 | | |
| | | 8703781 | Oct 15, 2030 | DS DP U-2297 | | |
| | | 8703781 | Oct 15, 2030 | DS DP U-2298 | | |
| | | 8835443 | Jun 10, 2025 | U-2026 | | |
| | | 8835443 | Jun 10, 2025 | U-2027 | | |
| | | 8835443 | Jun 10, 2025 | U-2296 | | |
| | | 8835443 | Jun 10, 2025 | U-2298 | | |
| | | 8952018 | Oct 15, 2030 | U-2027 | | |
| | | 9233956 | May 04, 2029 | U-1811 | | |
| | | 9233956 | May 04, 2029 | U-2031 | | |
| | | 9233956 | May 04, 2029 | U-2032 | | |
| | | 9233956 | May 04, 2029 | U-2296 | | |
| | | 9233956 | May 04, 2029 | U-2297 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DABRAFENIB MESYLATE - TAFINLAR</u> | | | | | | |
| N 202806 002 | 7994185 | Jan 20, 2030 | DS DP U-1406 | | I-745 | Jun 22, 2020 |
| | 7994185 | Jan 20, 2030 | DS DP U-2031 | | I-778 | Apr 30, 2021 |
| | 7994185 | Jan 20, 2030 | DS DP U-2032 | | I-781 | May 04, 2021 |
| | 7994185 | Jan 20, 2030 | DS DP U-2296 | | ODE-147 | Jun 22, 2024 |
| | 8415345 | Jan 20, 2030 | DS DP U-1406 | | ODE-182 | Apr 30, 2025 |
| | 8415345 | Jan 20, 2030 | DS DP U-2031 | | ODE-183 | May 04, 2025 |
| | 8415345 | Jan 20, 2030 | DS DP U-2032 | | ODE-47 | May 29, 2020 |
| | 8415345 | Jan 20, 2030 | DS DP U-2296 | | ODE-58 | Jan 09, 2021 |
| | 8703781 | Oct 15, 2030 | DS DP U-1713 | | | |
| | 8703781 | Oct 15, 2030 | DS DP U-2032 | | | |
| | 8703781 | Oct 15, 2030 | DS DP U-2296 | | | |
| | 8703781 | Oct 15, 2030 | DS DP U-2297 | | | |
| | 8703781 | Oct 15, 2030 | DS DP U-2298 | | | |
| | 8835443 | Jun 10, 2025 | U-2026 | | | |
| | 8835443 | Jun 10, 2025 | U-2027 | | | |
| | 8835443 | Jun 10, 2025 | U-2296 | | | |
| | 8835443 | Jun 10, 2025 | U-2298 | | | |
| | 8952018 | Oct 15, 2030 | U-2027 | | | |
| | 9233956 | May 04, 2029 | U-1811 | | | |
| | 9233956 | May 04, 2029 | U-2031 | | | |
| | 9233956 | May 04, 2029 | U-2032 | | | |
| | 9233956 | May 04, 2029 | U-2296 | | | |
| | 9233956 | May 04, 2029 | U-2297 | | | |
| <u>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</u> | | | | | | |
| N 206843 001 | 8329159 | Apr 13, 2028 | DS | | D-161 | Feb 05, 2019 |
| | 8629171 | Jun 13, 2031 | DS DP U-1724 | | D-162 | Feb 05, 2019 |
| | 8642025 | Aug 11, 2027 | DS DP U-1724 | | I-726 | Feb 05, 2019 |
| | 8642025 | Aug 11, 2027 | DS DP U-1725 | | I-727 | Feb 05, 2019 |
| | 8900566 | Aug 08, 2027 | U-1724 | | NCE | Jul 24, 2020 |
| | 8900566 | Aug 08, 2027 | U-1725 | | | |
| | 9421192 | Aug 08, 2027 | DS U-1724 | | | |
| | 9421192 | Aug 08, 2027 | DS U-1725 | | | |
| <u>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</u> | | | | | | |
| N 206843 002 | 8329159 | Apr 13, 2028 | DS | | D-161 | Feb 05, 2019 |
| | 8629171 | Jun 13, 2031 | DS DP U-1724 | | D-162 | Feb 05, 2019 |
| | 8642025 | Aug 11, 2027 | DS DP U-1724 | | I-726 | Feb 05, 2019 |
| | 8642025 | Aug 11, 2027 | DS DP U-1725 | | I-727 | Feb 05, 2019 |
| | 8900566 | Aug 08, 2027 | U-1724 | | NCE | Jul 24, 2020 |
| | 8900566 | Aug 08, 2027 | U-1725 | | | |
| | 9421192 | Aug 08, 2027 | DS U-1724 | | | |
| | 9421192 | Aug 08, 2027 | DS U-1725 | | | |
| <u>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</u> | | | | | | |
| N 206843 003 | 9421192 | Aug 08, 2027 | DS U-1724 | | | |
| | 9421192 | Aug 08, 2027 | DS U-1725 | | | |
| <u>DACOMITINIB - VIZIMPRO</u> | | | | | | |
| N 211288 001 | 7772243 | Aug 26, 2028 | DS DP | | NCE | Sep 27, 2023 |
| | 8623883 | May 05, 2025 | U-1403 | | ODE-206 | Sep 27, 2025 |
| | | | | | ODE-213 | Sep 27, 2025 |
| <u>DACOMITINIB - VIZIMPRO</u> | | | | | | |
| N 211288 002 | 7772243 | Aug 26, 2028 | DS DP | | NCE | Sep 27, 2023 |
| | 8623883 | May 05, 2025 | U-1403 | | ODE-206 | Sep 27, 2025 |
| | | | | | ODE-213 | Sep 27, 2025 |
| <u>DACOMITINIB - VIZIMPRO</u> | | | | | | |
| N 211288 003 | 7772243 | Aug 26, 2028 | DS DP | | NCE | Sep 27, 2023 |
| | 8623883 | May 05, 2025 | U-1403 | | ODE-206 | Sep 27, 2025 |
| | | | | | ODE-213 | Sep 27, 2025 |
| <u>DALBAVANCIN HYDROCHLORIDE - DALVANCE</u> | | | | | | |
| N 021883 001 | 6900175 | Dec 25, 2023 | U-1517 | | D-154 | Jan 20, 2019 |
| | 7115564 | Nov 14, 2023 | DP | | NCE | May 23, 2019 |
| | 7119061 | Nov 14, 2023 | DP | | GAIN | May 23, 2024 |
| | 8143212 | Nov 14, 2023 | U-1517 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DALFAMPRIDINE - DALFAMPRIDINE</u> | | | | | | |
| A 206863 | 001 | | | | PC | Mar 17, 2019 |
| <u>DALFAMPRIDINE - AMPYRA</u> | | | | | | |
| N 022250 | 001 | 8007826 | | U-1030 | | |
| | | 8354437 | | U-1030 | | |
| | | 8440703 | | U-1030 | | |
| | | 8663685 | | U-1030 | | |
| | | 9918973 | | U-1030 | | |
| <u>DANTROLENE SODIUM - RYANODEX</u> | | | | | | |
| N 205579 | 001 | 7758890 | | DP | ODE-69 | Jul 22, 2021 |
| | | 8110225 | | DP | | |
| | | 8604072 | | DP | | |
| | | 8685460 | | U-1546 | | |
| | | 9884044 | | DP U-1546 | | |
| <u>DAPAGLIFLOZIN - FARXIGA</u> | | | | | | |
| N 202293 | 001 | 6414126 | | DS DP U-2139 | M-212 | Oct 20, 2020 |
| | | 6414126 | | DS DP U-493 | NCE | Jan 08, 2019 |
| | | 6515117 | | DS DP U-2139 | | |
| | | 6515117 | | DS DP U-493 | | |
| | | 6936590 | | U-493 | | |
| | | 7456254 | | DP U-2139 | | |
| | | 7851502 | | DP | | |
| | | 7919598 | | DS | | |
| | | 8221786 | | DP | | |
| | | 8329648 | | U-2139 | | |
| | | 8329648 | | U-2212 | | |
| | | 8329648 | | U-2213 | | |
| | | 8361972 | | U-2139 | | |
| | | 8361972 | | U-493 | | |
| | | 8431685 | | DP U-2139 | | |
| | | 8461105 | | DP U-2139 | | |
| | | 8501698 | | DP U-493 | | |
| | | 8685934 | | U-1522 | | |
| | | 8716251 | | DP | | |
| | | 8721615 | | DP | Y | |
| | | 8906851 | | U-2139 | | |
| | | 9198925 | | U-2139 | | |
| | | 9198925 | | U-493 | | |
| | | 9238076 | | DP U-2139 | | |
| <u>DAPAGLIFLOZIN - FARXIGA</u> | | | | | | |
| N 202293 | 002 | 6414126 | | DS DP U-2139 | M-212 | Oct 20, 2020 |
| | | 6414126 | | DS DP U-493 | NCE | Jan 08, 2019 |
| | | 6515117 | | DS DP U-2139 | | |
| | | 6515117 | | DS DP U-493 | | |
| | | 6936590 | | U-493 | | |
| | | 7456254 | | DP U-2139 | | |
| | | 7851502 | | DP | | |
| | | 7919598 | | DS | | |
| | | 8221786 | | DP | | |
| | | 8329648 | | U-2139 | | |
| | | 8329648 | | U-2212 | | |
| | | 8329648 | | U-2213 | | |
| | | 8361972 | | U-2139 | | |
| | | 8361972 | | U-493 | | |
| | | 8431685 | | DP U-2139 | | |
| | | 8461105 | | DP U-2139 | | |
| | | 8501698 | | DP U-493 | | |
| | | 8685934 | | U-1522 | | |
| | | 8716251 | | DP | | |
| | | 8721615 | | DP | Y | |
| | | 8906851 | | U-2139 | | |
| | | 9198925 | | U-2139 | | |
| | | 9198925 | | U-493 | | |
| | | 9238076 | | DP U-2139 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u> | | | | | | |
| N 205649 001 | 6414126 | Oct 04, 2020 | DS DP U-493 | | NCE | Jan 08, 2019 |
| | 6515117 | Oct 04, 2020 | DS DP U-493 | | | |
| | 6936590 | Oct 04, 2020 | U-493 | | | |
| | 7919598 | Dec 16, 2029 | DS | | | |
| | 8501698 | Jun 20, 2027 | DP U-493 | | | |
| | 8685934 | May 26, 2030 | U-1522 | | | |
| | 9198925 | Oct 04, 2020 | U-493 | | | |
| | 9616028 | Nov 12, 2030 | DP | | | |
| <u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u> | | | | | | |
| N 205649 002 | 6414126 | Oct 04, 2020 | DS DP U-493 | | NCE | Jan 08, 2019 |
| | 6515117 | Oct 04, 2020 | DS DP U-493 | | | |
| | 6936590 | Oct 04, 2020 | U-493 | | | |
| | 7919598 | Dec 16, 2029 | DS | | | |
| | 8501698 | Jun 20, 2027 | DP U-493 | | | |
| | 8685934 | May 26, 2030 | U-1522 | | | |
| | 9198925 | Oct 04, 2020 | U-493 | | | |
| | 9616028 | Nov 12, 2030 | DP | | | |
| <u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u> | | | | | | |
| N 205649 003 | 6414126 | Oct 04, 2020 | DS DP U-493 | | NCE | Jan 08, 2019 |
| | 6515117 | Oct 04, 2020 | DS DP U-493 | | | |
| | 6936590 | Oct 04, 2020 | U-493 | | | |
| | 7919598 | Dec 16, 2029 | DS | | | |
| | 8501698 | Jun 20, 2027 | DP U-493 | | | |
| | 8685934 | May 26, 2030 | U-1522 | | | |
| | 9198925 | Oct 04, 2020 | U-493 | | | |
| | 9616028 | Nov 12, 2030 | DP | | | |
| <u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u> | | | | | | |
| N 205649 004 | 6414126 | Oct 04, 2020 | DS DP U-493 | | NCE | Jan 08, 2019 |
| | 6515117 | Oct 04, 2020 | DS DP U-493 | | | |
| | 6936590 | Oct 04, 2020 | U-493 | | | |
| | 7919598 | Dec 16, 2029 | DS | | | |
| | 8501698 | Jun 20, 2027 | DP U-493 | | | |
| | 8685934 | May 26, 2030 | U-1522 | | | |
| | 9198925 | Oct 04, 2020 | U-493 | | | |
| | 9616028 | Nov 12, 2030 | DP | | | |
| <u>DAPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - XIGDUO XR</u> | | | | | | |
| N 205649 005 | 6414126 | Oct 04, 2020 | DS DP U-493 | | NCE | Jan 08, 2019 |
| | 6515117 | Oct 04, 2020 | DS DP U-493 | | | |
| | 6936590 | Oct 04, 2020 | U-493 | | | |
| | 7919598 | Dec 16, 2029 | DS | | | |
| | 8501698 | Jun 20, 2027 | DP U-493 | | | |
| | 8685934 | May 26, 2030 | U-1522 | | | |
| | 9198925 | Oct 04, 2020 | U-493 | | | |
| | 9616028 | Nov 12, 2030 | DP | | | |
| <u>DAPAGLIFLOZIN; SAXAGLIPTIN HYDROCHLORIDE - QTERN</u> | | | | | | |
| N 209091 001 | 6414126 | Oct 04, 2020 | DS DP U-1976 | | M-175 | Apr 05, 2019 |
| | 6414126 | Oct 04, 2020 | DS DP U-1977 | | NC | Feb 27, 2020 |
| | 6515117 | Oct 04, 2020 | DS DP U-1976 | | NCE | Jan 08, 2019 |
| | 6515117 | Oct 04, 2020 | DS DP U-1977 | | | |
| | 6936590 | Oct 04, 2020 | U-1976 | | | |
| | 6936590 | Oct 04, 2020 | U-1977 | | | |
| | 7919598 | Dec 16, 2029 | DS | | | |
| | 8221786 | Mar 21, 2028 | DP | | | |
| | 8361972 | Mar 21, 2028 | U-1976 | | | |
| | 8361972 | Mar 21, 2028 | U-1977 | | | |
| | 8501698 | Jun 20, 2027 | DP U-1976 | | | |
| | 8501698 | Jun 20, 2027 | DP U-1977 | | | |
| | 8628799 | Jul 13, 2025 | DP | | | |
| | 8716251 | Mar 21, 2028 | DP | | | |
| | 9198925 | Oct 04, 2020 | U-1976 | | | |
| | 9198925 | Oct 04, 2020 | U-1977 | | | |
| | RE44186 | Jul 31, 2023 | DS DP U-1976 | | | |
| | RE44186 | Jul 31, 2023 | DS DP U-1977 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DAPSONE - ACZONE</u> | | | | | | |
| N 207154 001 | 9161926 | Nov 18, 2033 | DP | | NS | Feb 24, 2019 |
| | 9517219 | Nov 18, 2033 | U-1033 | | | |
| <u>DAPTOMYCIN - CUBICIN</u> | | | | | | |
| N 021572 002 | 8003673 | Sep 04, 2028 | U-1180 | | M-211 NPP | Sep 01, 2020 Mar 29, 2020 |
| <u>DAPTOMYCIN - CUBICIN RF</u> | | | | | | |
| N 021572 003 | 9138456 | Nov 23, 2030 | DP | | M-211 NPP | Sep 01, 2020 Mar 29, 2020 |
| <u>DARUNAVIR ETHANOLATE - PREZISTA</u> | | | | | | |
| N 021976 001 | 7470506 | Jun 23, 2019 | U-1209 | | | |
| | 7470506 | Jun 23, 2019 | U-1305 | | | |
| | 7470506 | Jun 23, 2019 | U-935 | | | |
| | 7700645 | Dec 26, 2026 | DS DP | | | |
| | 8518987 | Feb 16, 2024 | DS DP | | | |
| | 8518987*PED | Aug 16, 2024 | | | | |
| | 8597876 | Jun 23, 2019 | U-1305 | | | |
| | 8597876*PED | Dec 23, 2019 | | | | |
| <u>DARUNAVIR ETHANOLATE - PREZISTA</u> | | | | | | |
| N 021976 002 | 7470506 | Jun 23, 2019 | U-1209 | | | |
| | 7470506 | Jun 23, 2019 | U-1305 | | | |
| | 7470506 | Jun 23, 2019 | U-935 | | | |
| | 7700645 | Dec 26, 2026 | DS DP | | | |
| | 8518987 | Feb 16, 2024 | DS DP | | | |
| | 8518987*PED | Aug 16, 2024 | | | | |
| | 8597876 | Jun 23, 2019 | U-1305 | | | |
| | 8597876*PED | Dec 23, 2019 | | | | |
| | 9889115 | Jun 23, 2019 | U-1305 | | | |
| <u>DARUNAVIR ETHANOLATE - PREZISTA</u> | | | | | | |
| N 021976 003 | 7470506 | Jun 23, 2019 | U-1209 | | | |
| | 7470506 | Jun 23, 2019 | U-1305 | | | |
| | 7470506 | Jun 23, 2019 | U-935 | | | |
| | 7700645 | Dec 26, 2026 | DS DP | | | |
| | 8518987 | Feb 16, 2024 | DS DP | | | |
| | 8518987*PED | Aug 16, 2024 | | | | |
| | 8597876 | Jun 23, 2019 | U-1305 | | | |
| | 8597876*PED | Dec 23, 2019 | | | | |
| <u>DARUNAVIR ETHANOLATE - PREZISTA</u> | | | | | | |
| N 021976 004 | 7470506 | Jun 23, 2019 | U-1209 | | | |
| | 7470506 | Jun 23, 2019 | U-1305 | | | |
| | 7470506 | Jun 23, 2019 | U-935 | | | |
| | 7700645 | Dec 26, 2026 | DS DP | | | |
| | 8518987 | Feb 16, 2024 | DS DP | | | |
| | 8518987*PED | Aug 16, 2024 | | | | |
| | 8597876 | Jun 23, 2019 | U-1305 | | | |
| | 8597876*PED | Dec 23, 2019 | | | | |
| | 9889115 | Jun 23, 2019 | U-1305 | | | |
| <u>DARUNAVIR ETHANOLATE - PREZISTA</u> | | | | | | |
| N 021976 005 | 7470506 | Jun 23, 2019 | U-1209 | | | |
| | 7470506 | Jun 23, 2019 | U-1305 | | | |
| | 7470506 | Jun 23, 2019 | U-935 | | | |
| | 7700645 | Dec 26, 2026 | DS DP | | | |
| | 8518987 | Feb 16, 2024 | DS DP | | | |
| | 8518987*PED | Aug 16, 2024 | | | | |
| | 8597876 | Jun 23, 2019 | U-1305 | | | |
| | 8597876*PED | Dec 23, 2019 | | | | |
| | 9889115 | Jun 23, 2019 | U-1305 | | | |
| <u>DARUNAVIR ETHANOLATE - PREZISTA</u> | | | | | | |
| N 021976 006 | 7470506 | Jun 23, 2019 | U-1209 | | | |
| | 7470506 | Jun 23, 2019 | U-1305 | | | |
| | 7470506 | Jun 23, 2019 | U-935 | | | |
| | 7700645 | Dec 26, 2026 | DS DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DARUNAVIR ETHANOLATE - PREZISTA</u> | | | | | | |
| N 021976 | 006 | 8518987 | Feb 16, 2024 | DS DP | | |
| | | 8518987*PED | Aug 16, 2024 | | | |
| | | 8597876 | Jun 23, 2019 | | U-1305 | |
| | | 8597876*PED | Dec 23, 2019 | | | |
| | | 9889115 | Jun 23, 2019 | | U-1305 | |
| <u>DARUNAVIR ETHANOLATE - PREZISTA</u> | | | | | | |
| N 202895 | 001 | 7470506 | Jun 23, 2019 | | U-1209 | |
| | | 7470506 | Jun 23, 2019 | | U-1305 | |
| | | 7700645 | Dec 26, 2026 | DS DP | | |
| | | 8518987 | Feb 16, 2024 | DS DP | | |
| | | 8518987*PED | Aug 16, 2024 | | | |
| | | 8597876 | Jun 23, 2019 | | U-1305 | |
| | | 8597876*PED | Dec 23, 2019 | | | |
| | | 9889115 | Jun 23, 2019 | | U-1305 | |
| <u>DASABUVIR SODIUM ; OMBITASVIR; PARITAPREVIR; RITONAVIR - VIEKIRA PAK (COPACKAGED)</u> | | | | | | |
| N 206619 | 001 | 7148359 | Jul 19, 2019 | DP | | D-163 |
| | | 7364752 | Nov 10, 2020 | DP | | NCE |
| | | 8188104 | May 17, 2029 | DS DP | U-1636 | |
| | | 8268349 | Aug 25, 2024 | DP | | |
| | | 8399015 | Aug 25, 2024 | DP | | |
| | | 8420596 | Apr 10, 2031 | DS DP | | |
| | | 8466159 | Sep 04, 2032 | | U-1637 | |
| | | 8492386 | Sep 04, 2032 | | U-1840 | |
| | | 8501238 | Sep 17, 2028 | DS DP | U-1636 | |
| | | 8642538 | Sep 10, 2029 | DS DP | U-1638 | |
| | | 8680106 | Sep 04, 2032 | | U-1637 | |
| | | 8685984 | Sep 04, 2032 | | U-1840 | |
| | | 8686026 | Jun 09, 2031 | DP | | |
| | | 8691938 | Apr 13, 2032 | DS DP | | |
| | | 9006387 | Jun 10, 2030 | | U-1687 | |
| | | 9044480 | Apr 10, 2031 | | U-1638 | |
| | | 9139536 | Nov 09, 2028 | | U-1753 | |
| | | 9629841 | Oct 18, 2033 | DP | U-1753 | |
| <u>DASABUVIR SODIUM; OMBITASVIR; PARITAPREVIR; RITONAVIR - VIEKIRA XR</u> | | | | | | |
| N 208624 | 001 | 10105365 | Jan 02, 2035 | DP | U-1889 | NCE |
| | | 7148359 | Jul 19, 2019 | DP | | Dec 19, 2019 |
| | | 7364752 | Nov 10, 2020 | DP | | |
| | | 8188104 | May 17, 2029 | DS DP | U-1636 | |
| | | 8268349 | Aug 25, 2024 | DP | | |
| | | 8399015 | Aug 25, 2024 | DP | | |
| | | 8420596 | Apr 10, 2031 | DS DP | | |
| | | 8466159 | Sep 04, 2032 | | U-1637 | |
| | | 8492386 | Sep 04, 2032 | | U-1840 | |
| | | 8501238 | Sep 17, 2028 | DS DP | U-1636 | |
| | | 8642538 | Sep 10, 2029 | DS DP | U-1638 | |
| | | 8680106 | Sep 04, 2032 | | U-1637 | |
| | | 8685984 | Sep 04, 2032 | | U-1840 | |
| | | 8686026 | Jun 09, 2031 | DP | | |
| | | 8691938 | Apr 13, 2032 | DS DP | | |
| | | 9006387 | Jun 10, 2030 | | U-1687 | |
| | | 9044480 | Apr 10, 2031 | | U-1638 | |
| | | 9139536 | Nov 09, 2028 | | U-1753 | |
| | | 9333204 | Jan 02, 2035 | DP | U-1889 | |
| | | 9744170 | Jan 02, 2035 | DP | U-1889 | |
| <u>DASATINIB - SPRYCEL</u> | | | | | | |
| N 021986 | 001 | 6596746 | Jun 28, 2020 | DS DP | U-748 | I-791 |
| | | 6596746 | Jun 28, 2020 | DS DP | U-780 | NPP |
| | | 6596746*PED | Dec 28, 2020 | | | ODE-164 |
| | | 7125875 | Apr 13, 2020 | | U-779 | PED |
| | | 7125875 | Apr 13, 2020 | | U-780 | PED |
| | | 7125875*PED | Oct 13, 2020 | | | PED |
| | | 7153856 | Apr 28, 2020 | | U-780 | PED |
| | | 7153856*PED | Oct 28, 2020 | | | |
| | | 7491725 | Mar 28, 2026 | DS DP | | |
| | | 7491725*PED | Sep 28, 2026 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DASATINIB - SPRYCEL</u> | | | | | | |
| N 021986 001 | 8680103 | Feb 04, 2025 | DP | | | |
| | 8680103*PED | Aug 04, 2025 | | | | |
| <u>DASATINIB - SPRYCEL</u> | | | | | | |
| N 021986 002 | 6596746 | Jun 28, 2020 | DS DP U-748 | | I-791 | Dec 21, 2021 |
| | 6596746 | Jun 28, 2020 | DS DP U-780 | | NPP | Nov 09, 2020 |
| | 6596746*PED | Dec 28, 2020 | | | ODE-164 | Nov 09, 2024 |
| | 7125875 | Apr 13, 2020 | | U-779 | PED | May 09, 2021 |
| | 7125875 | Apr 13, 2020 | | U-780 | PED | Jun 21, 2022 |
| | 7125875*PED | Oct 13, 2020 | | | PED | May 09, 2025 |
| | 7153856 | Apr 28, 2020 | | U-780 | | |
| | 7153856*PED | Oct 28, 2020 | | | | |
| | 7491725 | Mar 28, 2026 | DS DP | | | |
| | 7491725*PED | Sep 28, 2026 | | | | |
| | 8680103 | Feb 04, 2025 | DP | | | |
| | 8680103*PED | Aug 04, 2025 | | | | |
| <u>DASATINIB - SPRYCEL</u> | | | | | | |
| N 021986 003 | 6596746 | Jun 28, 2020 | DS DP U-748 | | I-791 | Dec 21, 2021 |
| | 6596746 | Jun 28, 2020 | DS DP U-780 | | NPP | Nov 09, 2020 |
| | 6596746*PED | Dec 28, 2020 | | | ODE-164 | Nov 09, 2024 |
| | 7125875 | Apr 13, 2020 | | U-779 | PED | May 09, 2021 |
| | 7125875 | Apr 13, 2020 | | U-780 | PED | Jun 21, 2022 |
| | 7125875*PED | Oct 13, 2020 | | | PED | May 09, 2025 |
| | 7153856 | Apr 28, 2020 | | U-780 | | |
| | 7153856*PED | Oct 28, 2020 | | | | |
| | 7491725 | Mar 28, 2026 | DS DP | | | |
| | 7491725*PED | Sep 28, 2026 | | | | |
| | 8680103 | Feb 04, 2025 | DP | | | |
| | 8680103*PED | Aug 04, 2025 | | | | |
| <u>DASATINIB - SPRYCEL</u> | | | | | | |
| N 021986 004 | 6596746 | Jun 28, 2020 | DS DP U-748 | | I-791 | Dec 21, 2021 |
| | 6596746 | Jun 28, 2020 | DS DP U-780 | | NPP | Nov 09, 2020 |
| | 6596746*PED | Dec 28, 2020 | | | ODE-164 | Nov 09, 2024 |
| | 7125875 | Apr 13, 2020 | | U-779 | PED | May 09, 2021 |
| | 7125875 | Apr 13, 2020 | | U-780 | PED | Jun 21, 2022 |
| | 7125875*PED | Oct 13, 2020 | | | PED | May 09, 2025 |
| | 7153856 | Apr 28, 2020 | | U-780 | | |
| | 7153856*PED | Oct 28, 2020 | | | | |
| | 7491725 | Mar 28, 2026 | DS DP | | | |
| | 7491725*PED | Sep 28, 2026 | | | | |
| | 8680103 | Feb 04, 2025 | DP | | | |
| | 8680103*PED | Aug 04, 2025 | | | | |
| <u>DASATINIB - SPRYCEL</u> | | | | | | |
| N 021986 005 | 6596746 | Jun 28, 2020 | DS DP U-748 | | I-791 | Dec 21, 2021 |
| | 6596746 | Jun 28, 2020 | DS DP U-780 | | NPP | Nov 09, 2020 |
| | 6596746*PED | Dec 28, 2020 | | | ODE-164 | Nov 09, 2024 |
| | 7125875 | Apr 13, 2020 | | U-779 | PED | May 09, 2021 |
| | 7125875 | Apr 13, 2020 | | U-780 | PED | Jun 21, 2022 |
| | 7125875*PED | Oct 13, 2020 | | | PED | May 09, 2025 |
| | 7153856 | Apr 28, 2020 | | U-780 | | |
| | 7153856*PED | Oct 28, 2020 | | | | |
| | 7491725 | Mar 28, 2026 | DS DP | | | |
| | 7491725*PED | Sep 28, 2026 | | | | |
| | 8680103 | Feb 04, 2025 | DP | | | |
| | 8680103*PED | Aug 04, 2025 | | | | |
| <u>DASATINIB - SPRYCEL</u> | | | | | | |
| N 021986 006 | 6596746 | Jun 28, 2020 | DS DP U-748 | | I-791 | Dec 21, 2021 |
| | 6596746 | Jun 28, 2020 | DS DP U-780 | | NPP | Nov 09, 2020 |
| | 6596746*PED | Dec 28, 2020 | | | ODE-164 | Nov 09, 2024 |
| | 7125875 | Apr 13, 2020 | | U-779 | PED | May 09, 2021 |
| | 7125875 | Apr 13, 2020 | | U-780 | PED | Jun 21, 2022 |
| | 7125875*PED | Oct 13, 2020 | | | PED | May 09, 2025 |
| | 7153856 | Apr 28, 2020 | | U-780 | | |
| | 7153856*PED | Oct 28, 2020 | | | | |
| | 7491725 | Mar 28, 2026 | DS DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------|-----------|---------------------------------------|----------------------------------------------|------------------------------------|------------------------|-----------------------------------|
| <u>DASATINIB - SPRYCEL</u> | | | | | | |
| N 021986 | 006 | 7491725*PED 8680103 8680103*PED | Sep 28, 2026 Feb 04, 2025 Aug 04, 2025 | DP | | |
| <u>DEFERASIROX - EXJADE</u> | | | | | | |
| N 021882 | 001 | 6465504 | Apr 05, 2019 | DS DP | ODE-39 | Jan 23, 2020 |
| <u>DEFERASIROX - EXJADE</u> | | | | | | |
| N 021882 | 002 | 6465504 | Apr 05, 2019 | DS DP | ODE-39 | Jan 23, 2020 |
| <u>DEFERASIROX - EXJADE</u> | | | | | | |
| N 021882 | 003 | 6465504 | Apr 05, 2019 | DS DP | ODE-39 | Jan 23, 2020 |
| <u>DEFERASIROX - JADENU</u> | | | | | | |
| N 206910 | 001 | 6465504 9283209 | Apr 05, 2019 Nov 21, 2034 | DS DP DS DP | ODE-39 | Jan 23, 2020 |
| <u>DEFERASIROX - JADENU</u> | | | | | | |
| N 206910 | 002 | 6465504 9283209 | Apr 05, 2019 Nov 21, 2034 | DS DP DS DP | ODE-39 | Jan 23, 2020 |
| <u>DEFERASIROX - JADENU</u> | | | | | | |
| N 206910 | 003 | 6465504 9283209 | Apr 05, 2019 Nov 21, 2034 | DS DP DS DP | ODE-39 | Jan 23, 2020 |
| <u>DEFERASIROX - JADENU SPRINKLE</u> | | | | | | |
| N 207968 | 001 | 6465504 | Apr 05, 2019 | DS DP | | |
| <u>DEFERASIROX - JADENU SPRINKLE</u> | | | | | | |
| N 207968 | 002 | 6465504 | Apr 05, 2019 | DS DP | | |
| <u>DEFERASIROX - JADENU SPRINKLE</u> | | | | | | |
| N 207968 | 003 | 6465504 | Apr 05, 2019 | DS DP | | |
| <u>DEFERIPRONE - FERRIPROX</u> | | | | | | |
| N 021825 | 001 | 7049328 | Jun 28, 2021 | U-735 | | |
| <u>DEFERIPRONE - FERRIPROX</u> | | | | | | |
| N 208030 | 001 | 7049328 8703156 | Jun 28, 2021 Oct 29, 2029 | U-735 DP U-735 | | |
| <u>DEFIBROTIDE SODIUM - DEFITELIO</u> | | | | | | |
| N 208114 | 001 | | | | NCE ODE-112 | Mar 30, 2021 Mar 30, 2023 |
| <u>DEFLAZACORT - EMFLAZA</u> | | | | | | |
| N 208684 | 001 | | | | NCE ODE-130 | Feb 09, 2022 Feb 09, 2024 |
| <u>DEFLAZACORT - EMFLAZA</u> | | | | | | |
| N 208684 | 002 | | | | NCE ODE-130 | Feb 09, 2022 Feb 09, 2024 |
| <u>DEFLAZACORT - EMFLAZA</u> | | | | | | |
| N 208684 | 003 | | | | NCE ODE-130 | Feb 09, 2022 Feb 09, 2024 |
| <u>DEFLAZACORT - EMFLAZA</u> | | | | | | |
| N 208684 | 004 | | | | NCE ODE-130 | Feb 09, 2022 Feb 09, 2024 |
| <u>DEFLAZACORT - EMFLAZA</u> | | | | | | |
| N 208685 | 001 | | | | NCE ODE-130 | Feb 09, 2022 Feb 09, 2024 |
| <u>DEGARELIX ACETATE - FIRMAGON</u> | | | | | | |
| N 022201 | 001 | 5925730 9415085 9579359 | May 18, 2021 Apr 27, 2032 Feb 10, 2029 | DS DP U-943 U-1895 U-1978 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DEGARELIX ACETATE - FIRMAGON</u> | | | | | | |
| N 022201 | 002 | 5925730 | May 18, 2021 | DS DP U-943 | | |
| | | 9415085 | Apr 27, 2032 | U-1895 | | |
| | | 9579359 | Feb 10, 2029 | U-1978 | | |
| <u>DELAFLOXACIN MEGLUMINE - BAXDELA</u> | | | | | | |
| N 208610 | 001 | 7728143 | Nov 20, 2027 | DS | NCE | Jun 19, 2022 |
| | | 8252813 | Oct 02, 2026 | DP U-2028 | GAIN | Jun 19, 2027 |
| | | 8273892 | Aug 06, 2026 | DS | | |
| | | 8648093 | Oct 07, 2025 | DP U-2028 | | |
| | | 8871938 | Sep 23, 2029 | DS | | |
| | | 8969569 | Oct 07, 2025 | DP U-2028 | | |
| | | 9539250 | Oct 07, 2025 | DS DP U-2028 | | |
| | | RE46617 | Dec 28, 2029 | DS | | |
| <u>DELAFLOXACIN MEGLUMINE - BAXDELA</u> | | | | | | |
| N 208611 | 001 | 7635773 | Mar 13, 2029 | DP | NCE | Jun 19, 2022 |
| | | 7728143 | Nov 20, 2027 | DS | GAIN | Jun 19, 2027 |
| | | 8252813 | Oct 02, 2026 | DP U-2028 | | |
| | | 8273892 | Aug 06, 2026 | DS | | |
| | | 8410077 | Mar 13, 2029 | DP | | |
| | | 8648093 | Oct 07, 2025 | DP U-2028 | | |
| | | 8871938 | Sep 23, 2029 | DS | | |
| | | 9200088 | Mar 13, 2029 | DP | | |
| | | 9493582 | Feb 27, 2033 | DP | | |
| | | 9539250 | Oct 07, 2025 | DS DP U-2028 | | |
| | | 9750822 | Mar 13, 2029 | DP | | |
| | | RE46617 | Dec 28, 2029 | DS | | |
| <u>DELAVIRDINE MESYLATE - RESCRIPTOR</u> | | | | | | |
| N 020705 | 002 | 6177101 | Jun 07, 2019 | | | |
| <u>DEOXYCHOLIC ACID - KYBELLA</u> | | | | | | |
| N 206333 | 001 | 7622130 | Dec 10, 2027 | U-1690 | | |
| | | 7754230 | Dec 10, 2027 | U-1690 | | |
| | | 8101593 | Mar 02, 2030 | DP | | |
| | | 8242294 | May 16, 2028 | DS | | |
| | | 8298556 | Aug 03, 2025 | U-1690 | | |
| | | 8367649 | Mar 02, 2030 | DP | | |
| | | 8461140 | Feb 21, 2028 | DP | | |
| | | 8546367 | Feb 21, 2028 | DP U-1690 | | |
| | | 8653058 | Mar 02, 2030 | DP | | |
| | | 8846066 | Feb 08, 2025 | U-1690 | | |
| | | 8883770 | Feb 21, 2028 | DP | | |
| | | 9522155 | Feb 21, 2028 | DP U-1940 | | |
| | | 9636349 | Feb 21, 2028 | U-1940 | | |
| | | 9949986 | Feb 21, 2028 | U-1940 | | |
| <u>DESLORATADINE - CLARINEX</u> | | | | | | |
| N 021165 | 001 | 6100274 | Jul 07, 2019 | | | |
| | | 7405223 | Jul 07, 2019 | U-886 | | |
| <u>DESLORATADINE - CLARINEX</u> | | | | | | |
| N 021312 | 001 | 6100274 | Jul 07, 2019 | DP | | |
| | | 7618649 | Dec 19, 2020 | DP U-1017 | | |
| <u>DESLORATADINE - CLARINEX</u> | | | | | | |
| N 021312 | 002 | 6100274 | Jul 07, 2019 | DP | | |
| | | 7618649 | Dec 19, 2020 | DP U-1017 | | |
| <u>DESLORATADINE; PSEUDOEPHEDRINE SULFATE - CLARINEX-D 12 HOUR</u> | | | | | | |
| N 021313 | 001 | 6100274 | Jul 07, 2019 | DP | | |
| | | 6709676 | Feb 18, 2021 | DP U-707 | | |
| | | 7618649 | Dec 19, 2020 | DP U-1017 | | |
| | | 8187630 | Dec 19, 2020 | DP U-1017 | | |
| <u>DESLORATADINE; PSEUDOEPHEDRINE SULFATE - CLARINEX D 24 HOUR</u> | | | | | | |
| N 021605 | 001 | 6100274 | Jul 07, 2019 | DP | | |
| | | 6979463 | Mar 28, 2022 | DP | | |
| | | 7618649 | Dec 19, 2020 | DP U-1017 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DES Loratadine; Pseudoephedrine Sulfate - Clarinex D 24 Hour</u> | | | | | | |
| N 021605 001 | 7820199 | Mar 28, 2022 | DP | | | |
| <u>Desmopressin Acetate - NoCDURNA</u> | | | | | | |
| N 022517 001 | 7560429 | Feb 02, 2024 | DP U-2326 | | NP | Jun 21, 2021 |
| | 7947654 | Dec 29, 2023 | DP | | | |
| | 8802624 | Dec 29, 2023 | U-2326 | | | |
| | 9220747 | May 07, 2023 | U-2326 | | | |
| | 9504647 | May 07, 2023 | DP U-2326 | | | |
| | 9919025 | May 07, 2023 | U-2326 | | | |
| | 9974826 | Apr 13, 2030 | U-2326 | | | |
| <u>Desmopressin Acetate - NoCDURNA</u> | | | | | | |
| N 022517 002 | 10137167 | May 21, 2029 | U-2327 | | NP | Jun 21, 2021 |
| | 7560429 | Feb 02, 2024 | DP U-2326 | | | |
| | 7947654 | Dec 29, 2023 | DP | | | |
| | 8802624 | Dec 29, 2023 | U-2326 | | | |
| | 9220747 | May 07, 2023 | U-2326 | | | |
| | 9504647 | May 07, 2023 | DP U-2326 | | | |
| | 9919025 | May 07, 2023 | U-2326 | | | |
| | 9974826 | Apr 13, 2030 | U-2327 | | | |
| <u>Desmopressin Acetate - NoCTIVA</u> | | | | | | |
| N 201656 001 | 7405203 | May 06, 2023 | U-1980 | | NP | Mar 03, 2020 |
| | 7579321 | May 06, 2023 | U-1980 | | | |
| | 7799761 | Sep 26, 2024 | DP | | | |
| | 9539302 | Jun 15, 2030 | DP | | | |
| <u>Desmopressin Acetate - NoCTIVA</u> | | | | | | |
| N 201656 002 | 7405203 | May 06, 2023 | U-1980 | | NP | Mar 03, 2020 |
| | 7579321 | May 06, 2023 | U-1980 | | | |
| | 9539302 | Jun 15, 2030 | DP | | | |
| <u>Desonide - Desonate</u> | | | | | | |
| N 021844 001 | 6387383 | Aug 03, 2020 | DS DP U-783 | | | |
| <u>Desonide - Verdeso</u> | | | | | | |
| N 021978 001 | 6730288 | Sep 08, 2019 | DP | | | |
| | 7029659 | Sep 08, 2019 | DP | | | |
| | 8460641 | Nov 05, 2028 | DP U-1412 | | | |
| | 8962000 | Aug 31, 2025 | DP U-1412 | | | |
| | 9492384 | Aug 31, 2025 | DP U-1412 | | | |
| <u>Desoximetasone - Topicort</u> | | | | | | |
| N 204141 001 | 8277780 | Sep 01, 2028 | DP U-1408 | | | |
| | 8715624 | May 26, 2026 | DP U-1408 | | | |
| <u>Desvenlafaxine Succinate - Pristiq</u> | | | | | | |
| N 021992 001 | 6673838 | Mar 01, 2022 | DS U-1364 | | M-222 | Feb 06, 2021 |
| | 6673838 | Mar 01, 2022 | DS U-860 | | | |
| | 8269040 | Jul 05, 2027 | DS | | | |
| <u>Desvenlafaxine Succinate - Pristiq</u> | | | | | | |
| N 021992 002 | 6673838 | Mar 01, 2022 | DS U-1364 | | M-222 | Feb 06, 2021 |
| | 6673838 | Mar 01, 2022 | DS U-860 | | | |
| | 8269040 | Jul 05, 2027 | DS | | | |
| <u>Desvenlafaxine Succinate - Pristiq</u> | | | | | | |
| N 021992 003 | 6673838 | Mar 01, 2022 | DS U-1364 | | M-222 | Feb 06, 2021 |
| | 6673838 | Mar 01, 2022 | DS U-860 | | | |
| | 8269040 | Jul 05, 2027 | DS | | | |
| <u>Deutetrabenazine - Austedo</u> | | | | | | |
| N 208082 001 | 8524733 | Mar 27, 2031 | DS DP | | I-751 | Aug 30, 2020 |
| | 9233959 | Sep 18, 2033 | DP | | NCE | Apr 03, 2022 |
| | 9296739 | Sep 18, 2033 | DP | | ODE-134 | Apr 03, 2024 |
| | 9550780 | Sep 18, 2033 | DS DP U-1846 | | | |
| | 9550780 | Sep 18, 2033 | DS DP U-1995 | | | |
| | 9814708 | Sep 18, 2033 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DEUTETRABENAZINE - AUSTEDO</u> | | | | | | |
| N 208082 002 | 8524733 | Mar 27, 2031 | DS DP | | I-751 | Aug 30, 2020 |
| | 9233959 | Sep 18, 2033 | DP | | NCE | Apr 03, 2022 |
| | 9296739 | Sep 18, 2033 | DP | | ODE-134 | Apr 03, 2024 |
| | 9550780 | Sep 18, 2033 | DS DP U-1846 | | | |
| | 9550780 | Sep 18, 2033 | DS DP U-1995 | | | |
| | 9814708 | Sep 18, 2033 | DP | | | |
| <u>DEUTETRABENAZINE - AUSTEDO</u> | | | | | | |
| N 208082 003 | 8524733 | Mar 27, 2031 | DS DP | | I-751 | Aug 30, 2020 |
| | 9233959 | Sep 18, 2033 | DP | | NCE | Apr 03, 2022 |
| | 9296739 | Sep 18, 2033 | DP | | ODE-134 | Apr 03, 2024 |
| | 9550780 | Sep 18, 2033 | DS DP U-1846 | | | |
| | 9550780 | Sep 18, 2033 | DS DP U-1995 | | | |
| | 9814708 | Sep 18, 2033 | DP | | | |
| <u>DEXAMETHASONE - OZURDEX</u> | | | | | | |
| N 022315 001 | 10076526 | Jan 09, 2023 | DP | | | |
| | 6726918 | Oct 20, 2020 | DP U-1204 | | | |
| | 6726918 | Oct 20, 2020 | DP U-1205 | | | |
| | 6899717 | Nov 01, 2023 | U-1206 | | | |
| | 7033605 | Oct 20, 2020 | DP | | | |
| | 7767223 | Nov 28, 2021 | DP | | | |
| | 8034366 | Jan 09, 2023 | DP U-1204 | | | |
| | 8034366 | Jan 09, 2023 | DP U-1205 | | | |
| | 8034370 | Jan 09, 2023 | DP | | | |
| | 8043628 | Oct 20, 2020 | U-1205 | | | |
| | 8063031 | Oct 20, 2020 | DP | | | |
| | 8088407 | Oct 20, 2020 | U-1205 | | | |
| | 8506987 | Jan 09, 2023 | U-1204 | | | |
| | 8506987 | Jan 09, 2023 | U-1205 | | | |
| | 9012437 | Oct 20, 2020 | U-1205 | | | |
| | 9192511 | Jan 09, 2023 | DP | | | |
| | 9283178 | Oct 20, 2020 | U-1205 | | | |
| | 9592242 | Oct 20, 2020 | U-1989 | | | |
| | 9592242 | Oct 20, 2020 | U-1990 | | | |
| | 9775849 | Oct 20, 2020 | U-1989 | | | |
| | 9775849 | Oct 20, 2020 | U-1990 | | | |
| <u>DEXAMETHASONE - DEXYCU KIT</u> | | | | | | |
| N 208912 001 | 10022502 | Sep 12, 2020 | U-2340 | | NP | Feb 09, 2021 |
| | 10028965 | May 23, 2034 | U-2340 | | | |
| | 6960346 | Jul 03, 2023 | DP | | | |
| | 7560120 | Sep 05, 2022 | DP | | | |
| <u>DEXAMETHASONE; TOBRAMYCIN - TOBRADEX ST</u> | | | | | | |
| N 050818 001 | 7795316 | Aug 03, 2028 | DP U-1082 | | | |
| | 8101582 | Dec 19, 2027 | DP U-1082 | | | |
| | 8450287 | Dec 19, 2027 | DP | | | |
| <u>DEXLANSOPRAZOLE - DEXILANT</u> | | | | | | |
| N 022287 001 | 6462058 | Jun 15, 2020 | DS DP U-949 | | NPP | Jul 08, 2019 |
| | 6462058 | Jun 15, 2020 | DS DP U-950 | | NPP | Jul 08, 2019 |
| | 6462058 | Jun 15, 2020 | DS DP U-951 | | NPP | Jul 08, 2019 |
| | 6664276 | Jan 30, 2023 | DS DP U-1507 | | | |
| | 6664276 | Jan 30, 2023 | DS DP U-949 | | | |
| | 6664276 | Jan 30, 2023 | DS DP U-950 | | | |
| | 6664276 | Jan 30, 2023 | DS DP U-951 | | | |
| | 6664276*PED | Jul 30, 2023 | | | | |
| | 6939971 | Jun 15, 2020 | U-949 | | | |
| | 6939971 | Jun 15, 2020 | U-950 | | | |
| | 6939971 | Jun 15, 2020 | U-951 | | | |
| | 7285668 | Jun 15, 2020 | DS | | | |
| | 7790755 | Aug 02, 2026 | DP | | | |
| | 8105626 | Sep 27, 2026 | DP | | | |
| | 8173158 | Mar 17, 2030 | U-949 | | | |
| | 8173158 | Mar 17, 2030 | U-950 | | | |
| | 8173158 | Mar 17, 2030 | U-951 | | | |
| | 8461187 | Jan 17, 2026 | DP | | | |
| | 8461187*PED | Jul 17, 2026 | | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DEXLANSOPRAZOLE - DEXILANT</u> | | | | | | |
| N 022287 001 | 8722084 | Oct 15, 2023 | DP | | | |
| | 8722084*PED | Apr 15, 2024 | | | | |
| | 8784885 | Oct 15, 2023 | DP U-1552 | | | |
| | 8784885 | Oct 15, 2023 | DP U-1553 | | | |
| | 8784885 | Oct 15, 2023 | DP U-1554 | | | |
| | 8784885*PED | Apr 15, 2024 | | | | |
| | 8871273 | Jan 11, 2028 | DP | | | |
| | 9011926 | Feb 24, 2026 | DP | | | |
| | 9145389 | Jun 15, 2020 | DS DP | | | |
| | 9233103 | Mar 05, 2032 | | U-1805 | | |
| | 9238029 | Jan 17, 2026 | DP | | | |
| <u>DEXLANSOPRAZOLE - DEXILANT</u> | | | | | | |
| N 022287 002 | 6462058 | Jun 15, 2020 | DS DP U-949 | | NPP | Jul 08, 2019 |
| | 6462058 | Jun 15, 2020 | DS DP U-950 | | NPP | Jul 08, 2019 |
| | 6462058 | Jun 15, 2020 | DS DP U-951 | | NPP | Jul 08, 2019 |
| | 6664276 | Jan 30, 2023 | DS DP U-1507 | | | |
| | 6664276 | Jan 30, 2023 | DS DP U-949 | | | |
| | 6664276 | Jan 30, 2023 | DS DP U-950 | | | |
| | 6664276 | Jan 30, 2023 | DS DP U-951 | | | |
| | 6664276*PED | Jul 30, 2023 | | | | |
| | 6939971 | Jun 15, 2020 | | U-949 | | |
| | 6939971 | Jun 15, 2020 | | U-950 | | |
| | 6939971 | Jun 15, 2020 | | U-951 | | |
| | 7285668 | Jun 15, 2020 | DS | | | |
| | 7790755 | Aug 02, 2026 | DP | | | |
| | 8105626 | Sep 27, 2026 | DP | | | |
| | 8173158 | Mar 17, 2030 | | U-949 | | |
| | 8173158 | Mar 17, 2030 | | U-950 | | |
| | 8173158 | Mar 17, 2030 | | U-951 | | |
| | 8461187 | Jan 17, 2026 | DP | | | |
| | 8461187*PED | Jul 17, 2026 | | | | |
| | 8722084 | Oct 15, 2023 | DP | | | |
| | 8722084*PED | Apr 15, 2024 | | | | |
| | 8784885 | Oct 15, 2023 | DP U-1552 | | | |
| | 8784885 | Oct 15, 2023 | DP U-1553 | | | |
| | 8784885 | Oct 15, 2023 | DP U-1554 | | | |
| | 8784885*PED | Apr 15, 2024 | | | | |
| | 8871273 | Jan 11, 2028 | DP | | | |
| | 9011926 | Feb 24, 2026 | DP | | | |
| | 9145389 | Jun 15, 2020 | DS DP | | | |
| | 9233103 | Mar 05, 2032 | | U-1805 | | |
| | 9238029 | Jan 17, 2026 | DP | | | |
| <u>DEXLANSOPRAZOLE - DEXILANT SOLUTAB</u> | | | | | | |
| N 208056 001 | 6328994 | May 17, 2019 | DP | | | |
| | 6328994*PED | Nov 17, 2019 | | | | |
| | 6462058 | Jun 15, 2020 | DS DP U-950 | | | |
| | 6462058 | Jun 15, 2020 | DS DP U-951 | | | |
| | 6462058*PED | Dec 15, 2020 | | | | |
| | 6664276 | Jan 30, 2023 | DS DP U-950 | | | |
| | 6664276 | Jan 30, 2023 | DS DP U-951 | | | |
| | 6664276*PED | Jul 30, 2023 | | | | |
| | 6939971 | Jun 15, 2020 | | U-950 | | |
| | 6939971 | Jun 15, 2020 | | U-951 | | |
| | 6939971*PED | Dec 15, 2020 | | | | |
| | 7285668 | Jun 15, 2020 | DS | | | |
| | 7285668*PED | Dec 15, 2020 | | | | |
| | 7431942 | May 17, 2019 | DP | | | |
| | 7431942*PED | Nov 17, 2019 | | | | |
| | 7875292 | May 17, 2019 | DP | | | |
| | 7875292*PED | Nov 17, 2019 | | | | |
| | 8461187 | Jan 17, 2026 | DP | | | |
| | 8461187*PED | Jul 17, 2026 | | | | |
| | 8784885 | Oct 15, 2023 | DP | | | |
| | 8784885*PED | Apr 15, 2024 | | | | |
| | 8871273 | Jan 11, 2028 | DP | | | |
| | 8871273*PED | Jul 11, 2028 | | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DEXLANSOPRAZOLE - DEXILANT SOLUTAB</u> | | | | | | |
| N 208056 | 001 | 9011926 | Feb 24, 2026 | DP | | |
| | | 9145389 | Jun 15, 2020 | DS DP | | |
| | | 9238029 | Jan 17, 2026 | DP | | |
| | | 9241910 | Mar 10, 2029 | DP | | |
| <u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u> | | | | | | |
| N 021038 | 001 | 6716867 | Mar 31, 2019 | | U-1472 | |
| | | 6716867*PED | Oct 01, 2019 | | | |
| <u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u> | | | | | | |
| N 021038 | 002 | 10016396 | Jan 04, 2032 | DP | | |
| | | 6716867 | Mar 31, 2019 | | U-1472 | |
| | | 6716867*PED | Oct 01, 2019 | | | |
| | | 8242158 | Jan 04, 2032 | DP | | |
| | | 8242158*PED | Jul 04, 2032 | | | |
| | | 8338470 | Jan 04, 2032 | DP | | |
| | | 8338470*PED | Jul 04, 2032 | | | |
| | | 8455527 | Jan 04, 2032 | | U-421 | |
| | | 8455527*PED | Jul 04, 2032 | | | |
| | | 8648106 | Jan 04, 2032 | DP | | |
| | | 8648106*PED | Jul 04, 2032 | | | |
| | | 9320712 | Jan 04, 2032 | DP | | |
| | | 9320712*PED | Jul 04, 2032 | | | |
| | | 9616049 | Jan 04, 2032 | DP | | |
| | | 9616049*PED | Jul 04, 2032 | | | |
| <u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u> | | | | | | |
| N 021038 | 003 | 10016396 | Jan 04, 2032 | DP | | |
| | | 6716867 | Mar 31, 2019 | | U-1472 | |
| | | 6716867*PED | Oct 01, 2019 | | | |
| | | 8242158 | Jan 04, 2032 | DP | | |
| | | 8242158*PED | Jul 04, 2032 | | | |
| | | 8338470 | Jan 04, 2032 | DP | | |
| | | 8338470*PED | Jul 04, 2032 | | | |
| | | 8455527 | Jan 04, 2032 | | U-421 | |
| | | 8455527*PED | Jul 04, 2032 | | | |
| | | 8648106 | Jan 04, 2032 | DP | | |
| | | 8648106*PED | Jul 04, 2032 | | | |
| | | 9320712 | Jan 04, 2032 | DP | | |
| | | 9320712*PED | Jul 04, 2032 | | | |
| | | 9616049 | Jan 04, 2032 | DP | | |
| | | 9616049*PED | Jul 04, 2032 | | | |
| <u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u> | | | | | | |
| N 021038 | 004 | 6716867 | Mar 31, 2019 | | U-1472 | |
| | | 6716867*PED | Oct 01, 2019 | | | |
| | | 8242158 | Jan 04, 2032 | DP | | |
| | | 8242158*PED | Jul 04, 2032 | | | |
| | | 8338470 | Jan 04, 2032 | DP | | |
| | | 8338470*PED | Jul 04, 2032 | | | |
| | | 8455527 | Jan 04, 2032 | | U-421 | |
| | | 8455527*PED | Jul 04, 2032 | | | |
| | | 8648106 | Jan 04, 2032 | DP | | |
| | | 8648106*PED | Jul 04, 2032 | | | |
| | | 9320712 | Jan 04, 2032 | DP | | |
| | | 9320712*PED | Jul 04, 2032 | | | |
| | | 9616049 | Jan 04, 2032 | DP | | |
| | | 9616049*PED | Jul 04, 2032 | | | |
| <u>DEXMEDETOMIDINE HYDROCHLORIDE - DEXMEDETOMIDINE HYDROCHLORIDE</u> | | | | | | |
| N 206628 | 003 | 9649296 | Apr 20, 2036 | DP | | |
| | | 9717796 | Apr 20, 2036 | DP | | |
| <u>DEXMEDETOMIDINE HYDROCHLORIDE - DEXMEDETOMIDINE HYDROCHLORIDE</u> | | | | | | |
| N 206628 | 004 | 9649296 | Apr 20, 2036 | DP | | |
| | | 9717796 | Apr 20, 2036 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u> | | | | | | |
| N 021802 001 | 6228398 | Nov 01, 2019 | DP U-676 | | | |
| | 6730325 | Nov 01, 2019 | DP U-676 | | | |
| <u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u> | | | | | | |
| N 021802 002 | 6228398 | Nov 01, 2019 | DP U-676 | | | |
| | 6730325 | Nov 01, 2019 | DP U-676 | | | |
| <u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u> | | | | | | |
| N 021802 003 | 6228398 | Nov 01, 2019 | DP U-676 | | | |
| | 6730325 | Nov 01, 2019 | DP U-676 | | | |
| <u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u> | | | | | | |
| N 021802 004 | 6228398 | Nov 01, 2019 | DP U-676 | | | |
| | 6730325 | Nov 01, 2019 | DP U-676 | | | |
| <u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u> | | | | | | |
| N 021802 005 | 6228398 | Nov 01, 2019 | DP U-676 | | | |
| | 6730325 | Nov 01, 2019 | DP U-676 | | | |
| <u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u> | | | | | | |
| N 021802 006 | 6228398 | Nov 01, 2019 | DP U-676 | | | |
| | 6730325 | Nov 01, 2019 | DP U-676 | | | |
| <u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u> | | | | | | |
| N 021802 007 | 6228398 | Nov 01, 2019 | DP U-676 | | | |
| | 6730325 | Nov 01, 2019 | DP U-676 | | | |
| <u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u> | | | | | | |
| N 021802 008 | 6228398 | Nov 01, 2019 | DP U-676 | | | |
| | 6730325 | Nov 01, 2019 | DP U-676 | | | |
| <u>DEXRAZOXANE HYDROCHLORIDE - TOTECT</u> | | | | | | |
| N 022025 001 | 6727253 | Mar 13, 2020 | | U-829 | | |
| <u>DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN - MUCINEX DM</u> | | | | | | |
| N 021620 001 | 6372252 | Apr 28, 2020 | DP | | | |
| | 6955821 | Apr 28, 2020 | DP U-685 | | | |
| | 7838032 | Apr 28, 2020 | DP | | | |
| <u>DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN - MUCINEX DM</u> | | | | | | |
| N 021620 002 | 6372252 | Apr 28, 2020 | DP | | | |
| | 6955821 | Apr 28, 2020 | DP U-685 | | | |
| | 7838032 | Apr 28, 2020 | DP | | | |
| <u>DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE - NUEDEXTA</u> | | | | | | |
| N 021879 001 | 7659282 | Aug 13, 2026 | | U-1093 | | |
| | 8227484 | Jul 17, 2023 | | U-1093 | | |
| <u>DICHLORPHENAMIDE - KEVEYIS</u> | | | | | | |
| N 011366 002 | | | | | ODE-96 | Aug 07, 2022 |
| <u>DICLOFENAC - ZORVOLEX</u> | | | | | | |
| N 204592 001 | 8679544 | Apr 23, 2030 | DP | | | |
| | 8999387 | Apr 23, 2030 | | U-55 | | |
| | 9017721 | Apr 23, 2030 | DP | | | |
| | 9173854 | Apr 23, 2030 | DP | | | |
| | 9180095 | Apr 23, 2030 | | U-55 | | |
| | 9180096 | Apr 23, 2030 | DP | | | |
| | 9186328 | Apr 23, 2030 | | U-55 | | |
| <u>DICLOFENAC - ZORVOLEX</u> | | | | | | |
| N 204592 002 | 8679544 | Apr 23, 2030 | DP | | | |
| | 8999387 | Apr 23, 2030 | | U-55 | | |
| | 9017721 | Apr 23, 2030 | DP | | | |
| | 9173854 | Apr 23, 2030 | DP | | | |
| | 9180095 | Apr 23, 2030 | | U-55 | | |
| | 9180096 | Apr 23, 2030 | DP | | | |
| | 9186328 | Apr 23, 2030 | | U-55 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------------------------------------------------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DICLOFENAC EPOLAMINE - FLECTOR</u> | | | | | | |
| N 021234 | 001 5607690 | Apr 13, 2019 | DP | | | |
| <u>DICLOFENAC EPOLAMINE - LICART</u> | | | | | | |
| N 206976 | 001 | | | | NP | Dec 19, 2021 |
| <u>DICLOFENAC POTASSIUM - CAMBIA</u> | | | | | | |
| N 022165 | 001 7759394 | Jun 16, 2026 | DS DP U-436 | | | |
| | 8097651 | Jun 16, 2026 | DS DP U-436 | | | |
| | 8927604 | Jun 16, 2026 | U-436 | | | |
| | 9827197 | Jun 16, 2026 | DP | | | |
| <u>DICLOFENAC POTASSIUM - ZIPSOR</u> | | | | | | |
| N 022202 | 001 6287594 | Jan 15, 2019 | DP | | | |
| | 6365180 | Jul 15, 2019 | DP U-980 | | | |
| | 7662858 | Feb 24, 2029 | U-1035 | | | |
| | 7884095 | Feb 24, 2029 | U-1111 | | | |
| | 7939518 | Feb 24, 2029 | U-980 | | | |
| | 8110606 | Feb 24, 2029 | U-980 | | | |
| | 8623920 | Feb 24, 2029 | U-1482 | | | |
| | 9561200 | Feb 24, 2029 | U-1482 | | | |
| <u>DICLOFENAC SODIUM - PENNSAID</u> | | | | | | |
| N 020947 | 001 8217078 | Jul 10, 2029 | U-1248 | | | |
| | 8546450 | Aug 09, 2030 | U-1435 | | | |
| | 8546450 | Aug 09, 2030 | U-1436 | | | |
| | 8618164 | Jul 10, 2029 | U-1477 | | | |
| | 8741956 | Jul 10, 2029 | U-1435 | | | |
| <u>DICLOFENAC SODIUM - DYLOJECT</u> | | | | | | |
| N 022396 | 001 6407079 | Jun 18, 2019 | DP | | | |
| | 8946292 | Mar 22, 2027 | U-1659 | | | |
| <u>DICLOFENAC SODIUM - PENNSAID</u> | | | | | | |
| N 204623 | 001 8217078 | Jul 10, 2029 | U-1477 | | | |
| | 8252838 | Apr 21, 2028 | DP U-1489 | | | |
| | 8546450 | Aug 09, 2030 | U-1435 | | | |
| | 8546450 | Aug 09, 2030 | U-1436 | | | |
| | 8563613 | Oct 17, 2027 | DP U-1488 | | | |
| | 8618164 | Jul 10, 2029 | U-1477 | | | |
| | 8741956 | Jul 10, 2029 | U-1435 | | | |
| | 8871809 | Oct 17, 2027 | U-1614 | | | |
| | 9066913 | Oct 17, 2027 | DP U-1488 | | | |
| | 9101591 | Oct 17, 2027 | DP U-1488 | | | |
| | 9132110 | Oct 17, 2027 | U-1488 | | | |
| | 9168304 | Oct 17, 2027 | DP | | | |
| | 9168305 | Oct 17, 2027 | U-1488 | | | |
| | 9220784 | Oct 17, 2027 | U-1488 | | | |
| | 9339551 | Oct 17, 2027 | U-1488 | | | |
| | 9339552 | Oct 17, 2027 | DP U-1488 | | | |
| | 9370501 | Jul 10, 2029 | U-1614 | | | |
| | 9375412 | Jul 10, 2029 | U-1614 | | | |
| | 9415029 | Jul 10, 2029 | U-1614 | | | |
| | 9539335 | Oct 17, 2027 | U-1614 | | | |
| <u>DIENOGEST; DIENOGEST; DIENOGEST; ESTRADIOL VALERATE; ESTRADIOL VALERATE; ESTRADIOL VALERATE; ESTRADIOL VALERATE; ESTRADIOL VALERATE - NATAZIA</u> | | | | | | |
| N 022252 | 001 8071577 | May 13, 2026 | DP U-1 | | | |
| | 8153616 | Jan 30, 2028 | U-1240 | | | |
| <u>DIFLUPREDNATE - DUREZOL</u> | | | | | | |
| N 022212 | 001 6114319 | May 18, 2019 | DP | | ODE-26 | Jun 13, 2019 |
| | 6114319*PED | Nov 18, 2019 | | | PED | Dec 13, 2019 |
| <u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u> | | | | | | |
| N 021392 | 001 6923984 | Feb 25, 2021 | DP | | | |
| | 7108866 | Dec 17, 2019 | DP U-107 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u> | | | | | | |
| N 021392 002 | 6923984 | Feb 25, 2021 | DP | | | |
| | 7108866 | Dec 17, 2019 | DP U-107 | | | |
| <u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u> | | | | | | |
| N 021392 003 | 6923984 | Feb 25, 2021 | DP | | | |
| | 7108866 | Dec 17, 2019 | DP U-107 | | | |
| <u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u> | | | | | | |
| N 021392 004 | 6923984 | Feb 25, 2021 | DP | | | |
| | 7108866 | Dec 17, 2019 | DP U-107 | | | |
| <u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u> | | | | | | |
| N 021392 005 | 6923984 | Feb 25, 2021 | DP | | | |
| | 7108866 | Dec 17, 2019 | DP U-107 | | | |
| <u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u> | | | | | | |
| N 021392 006 | 6923984 | Feb 25, 2021 | DP | | | |
| | 7108866 | Dec 17, 2019 | DP U-107 | | | |
| <u>DIMETHYL FUMARATE - TECFIDERA</u> | | | | | | |
| N 204063 001 | 6509376 | Apr 01, 2019 | DP | | | |
| | 7320999 | Oct 20, 2019 | U-1384 | | | |
| | 7619001 | Jun 20, 2020 | U-1384 | | | |
| | 7803840 | Apr 01, 2019 | U-1385 | | | |
| | 8399514 | Feb 07, 2028 | U-1384 | | | |
| <u>DIMETHYL FUMARATE - TECFIDERA</u> | | | | | | |
| N 204063 002 | 6509376 | Apr 01, 2019 | DP | | | |
| | 7320999 | Oct 20, 2019 | U-1384 | | | |
| | 7619001 | Jun 20, 2020 | U-1384 | | | |
| | 7803840 | Apr 01, 2019 | U-1385 | | | |
| | 8399514 | Feb 07, 2028 | U-1384 | | | |
| <u>DIPHENHYDRAMINE CITRATE; IBUPROFEN - ADVIL PM</u> | | | | | | |
| N 021394 001 | 8263647 | May 30, 2022 | DP | | | |
| <u>DIPHENHYDRAMINE HYDROCHLORIDE; IBUPROFEN - ADVIL PM</u> | | | | | | |
| N 021393 001 | 8883849 | Jan 17, 2022 | U-1618 | | | |
| | 9155718 | Jan 17, 2022 | DP | | | |
| <u>DOCETAXEL - DOCETAXEL</u> | | | | | | |
| N 205934 001 | 8940786 | Sep 30, 2033 | DP U-1789 | | | |
| | 9308195 | Sep 30, 2033 | DP | | | |
| <u>DOCETAXEL - DOCETAXEL</u> | | | | | | |
| N 205934 002 | 8940786 | Sep 30, 2033 | DP U-1789 | | | |
| | 9308195 | Sep 30, 2033 | DP | | | |
| <u>DOCETAXEL - DOCETAXEL</u> | | | | | | |
| N 205934 003 | 8940786 | Sep 30, 2033 | DP U-1789 | | | |
| | 9308195 | Sep 30, 2033 | DP | | | |
| <u>DOLUTEGRAVIR SODIUM - TIVICAY</u> | | | | | | |
| N 204790 001 | 8129385 | Oct 05, 2027 | DS DP | | I-758 | Nov 21, 2020 |
| | 9242986 | Dec 08, 2029 | DS DP | | | |
| <u>DOLUTEGRAVIR SODIUM - TIVICAY</u> | | | | | | |
| N 204790 002 | 8129385 | Oct 05, 2027 | DS DP | | I-758 | Nov 21, 2020 |
| | 9242986 | Dec 08, 2029 | DS DP | | | |
| <u>DOLUTEGRAVIR SODIUM - TIVICAY</u> | | | | | | |
| N 204790 003 | 8129385 | Oct 05, 2027 | DS DP | | I-758 | Nov 21, 2020 |
| | 9242986 | Dec 08, 2029 | DS DP | | | |
| <u>DOLUTEGRAVIR SODIUM; RILPIVIRINE HYDROCHLORIDE - JULUCA</u> | | | | | | |
| N 210192 001 | 6838464 | Feb 26, 2021 | DS DP | | NC | Nov 21, 2020 |
| | 7067522 | Dec 20, 2019 | DS DP | | | |
| | 7125879 | Apr 21, 2025 | DS DP U-257 | | | |
| | 8080551 | Apr 11, 2023 | DS DP | | | |
| | 8101629 | Aug 09, 2022 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DOLUTEGRAVIR SODIUM; RILPIVIRINE HYDROCHLORIDE - JULUCA</u> | | | | | | |
| N 210192 | 001 | 8129385 | Oct 05, 2027 | DS DP | | |
| | | 9242986 | Dec 08, 2029 | DS DP | | |
| <u>DONEPEZIL HYDROCHLORIDE - ARICEPT</u> | | | | | | |
| N 022568 | 001 | 8481565 | Oct 04, 2026 | DP | | |
| <u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u> | | | | | | |
| N 206439 | 001 | 8039009 | Mar 24, 2029 | | U-1641 | |
| | | 8039009*PED | Sep 24, 2029 | | | |
| | | 8058291 | Dec 05, 2029 | | U-1641 | |
| | | 8168209 | Nov 22, 2025 | DP | | |
| | | 8168209*PED | May 22, 2026 | | | |
| | | 8173708 | Nov 22, 2025 | | U-1641 | |
| | | 8173708*PED | May 22, 2026 | | | |
| | | 8283379 | Nov 22, 2025 | | U-1641 | |
| | | 8283379*PED | May 22, 2026 | | | |
| | | 8293794 | Nov 22, 2025 | DP | | |
| | | 8329752 | Nov 22, 2025 | DP | | |
| | | 8329752*PED | May 22, 2026 | | | |
| | | 8338485 | Nov 22, 2025 | DP | | |
| | | 8338486 | Nov 22, 2025 | | U-1641 | |
| | | 8362085 | Nov 22, 2025 | | U-1641 | |
| | | 8362085*PED | May 22, 2026 | | | |
| | | 8580858 | Nov 22, 2025 | | U-1641 | |
| | | 8598233 | Nov 22, 2025 | DP | | |
| | | 8598233*PED | May 22, 2026 | | | |
| <u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u> | | | | | | |
| N 206439 | 002 | 8039009 | Mar 24, 2029 | | U-1641 | |
| | | 8039009*PED | Sep 24, 2029 | | | |
| | | 8058291 | Dec 05, 2029 | | U-1641 | |
| | | 8168209 | Nov 22, 2025 | DP | | |
| | | 8168209*PED | May 22, 2026 | | | |
| | | 8173708 | Nov 22, 2025 | | U-1641 | |
| | | 8173708*PED | May 22, 2026 | | | |
| | | 8283379 | Nov 22, 2025 | | U-1641 | |
| | | 8283379*PED | May 22, 2026 | | | |
| | | 8293794 | Nov 22, 2025 | DP | | |
| | | 8329752 | Nov 22, 2025 | DP | | |
| | | 8329752*PED | May 22, 2026 | | | |
| | | 8338485 | Nov 22, 2025 | DP | | |
| | | 8338486 | Nov 22, 2025 | | U-1641 | |
| | | 8362085 | Nov 22, 2025 | | U-1641 | |
| | | 8362085*PED | May 22, 2026 | | | |
| | | 8580858 | Nov 22, 2025 | | U-1641 | |
| | | 8598233 | Nov 22, 2025 | DP | | |
| | | 8598233*PED | May 22, 2026 | | | |
| <u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u> | | | | | | |
| N 206439 | 003 | 8039009 | Mar 24, 2029 | | U-1641 | |
| | | 8039009*PED | Sep 24, 2029 | | | |
| | | 8058291 | Dec 05, 2029 | | U-1641 | |
| | | 8168209 | Nov 22, 2025 | DP | | |
| | | 8168209*PED | May 22, 2026 | | | |
| | | 8173708 | Nov 22, 2025 | | U-1641 | |
| | | 8173708*PED | May 22, 2026 | | | |
| | | 8283379 | Nov 22, 2025 | | U-1641 | |
| | | 8283379*PED | May 22, 2026 | | | |
| | | 8293794 | Nov 22, 2025 | DP | | |
| | | 8329752 | Nov 22, 2025 | DP | | |
| | | 8329752*PED | May 22, 2026 | | | |
| | | 8338485 | Nov 22, 2025 | DP | | |
| | | 8338486 | Nov 22, 2025 | | U-1641 | |
| | | 8362085 | Nov 22, 2025 | | U-1641 | |
| | | 8362085*PED | May 22, 2026 | | | |
| | | 8580858 | Nov 22, 2025 | | U-1641 | |
| | | 8598233 | Nov 22, 2025 | DP | | |
| | | 8598233*PED | May 22, 2026 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</u> | | | | | | |
| N 206439 | 004 | 8039009 | Mar 24, 2029 | U-1641 | | |
| | | 8039009*PED | Sep 24, 2029 | | | |
| | | 8058291 | Dec 05, 2029 | U-1641 | | |
| | | 8168209 | Nov 22, 2025 | DP | | |
| | | 8168209*PED | May 22, 2026 | | | |
| | | 8173708 | Nov 22, 2025 | U-1641 | | |
| | | 8173708*PED | May 22, 2026 | | | |
| | | 8283379 | Nov 22, 2025 | U-1641 | | |
| | | 8283379*PED | May 22, 2026 | | | |
| | | 8293794 | Nov 22, 2025 | DP | | |
| | | 8329752 | Nov 22, 2025 | DP | | |
| | | 8329752*PED | May 22, 2026 | | | |
| | | 8338485 | Nov 22, 2025 | DP | | |
| | | 8338486 | Nov 22, 2025 | U-1641 | | |
| | | 8362085 | Nov 22, 2025 | U-1641 | | |
| | | 8362085*PED | May 22, 2026 | | | |
| | | 8580858 | Nov 22, 2025 | U-1641 | | |
| | | 8598233 | Nov 22, 2025 | DP | | |
| | | 8598233*PED | May 22, 2026 | | | |
| <u>DORAVIRINE - PIFELTRO</u> | | | | | | |
| N 210806 | 001 | 8486975 | Oct 07, 2031 | DS DP U-2394 | NCE | Aug 30, 2023 |
| <u>DORAVIRINE; LAMIVUDINE; TENOFOVIR DISOPROXIL FUMARATE - DELSTRIGO</u> | | | | | | |
| N 210807 | 001 | 8486975 | Oct 07, 2031 | DS DP U-2395 | NCE | Aug 30, 2023 |
| <u>DORIPENEM - DORIBAX</u> | | | | | | |
| N 022106 | 001 | 8247402 | Mar 30, 2021 | DS DP | | |
| <u>DORIPENEM - DORIBAX</u> | | | | | | |
| N 022106 | 002 | 8247402 | Mar 30, 2021 | DS DP | | |
| <u>DOXEPIIN HYDROCHLORIDE - SILENOR</u> | | | | | | |
| N 022036 | 001 | 6211229 | Feb 17, 2020 | U-620 | | |
| | | 7915307 | Aug 24, 2027 | U-620 | | |
| | | 8513299 | Sep 07, 2030 | U-620 | | |
| | | 9107898 | May 01, 2028 | U-620 | | |
| | | 9486437 | May 18, 2027 | U-620 | | |
| | | 9532971 | Jun 01, 2029 | DP | | |
| | | 9572814 | Jul 20, 2027 | U-620 | | |
| | | 9861607 | May 18, 2027 | U-620 | | |
| | | 9907780 | Apr 11, 2028 | DP | | |
| <u>DOXEPIIN HYDROCHLORIDE - SILENOR</u> | | | | | | |
| N 022036 | 002 | 6211229 | Feb 17, 2020 | U-620 | | |
| | | 7915307 | Aug 24, 2027 | U-620 | | |
| | | 8513299 | Sep 07, 2030 | U-620 | | |
| | | 9107898 | May 01, 2028 | U-620 | | |
| | | 9486437 | May 18, 2027 | U-620 | | |
| | | 9532971 | Jun 01, 2029 | DP | | |
| | | 9572814 | Jul 20, 2027 | U-620 | | |
| | | 9861607 | May 18, 2027 | U-620 | | |
| | | 9907780 | Apr 11, 2028 | DP | | |
| <u>DOXYCYCLINE - ORACEA</u> | | | | | | |
| N 050805 | 001 | 10058564 | Apr 05, 2022 | DS DP U-1063 | | |
| | | 7211267 | Apr 05, 2022 | U-925 | | |
| | | 7232572 | Apr 05, 2022 | U-925 | | |
| | | 7749532 | Dec 19, 2027 | DP U-1063 | | |
| | | 8206740 | Dec 24, 2025 | DP U-925 | | |
| | | 8394405 | Apr 07, 2024 | DP U-925 | | |
| | | 8394406 | Apr 07, 2024 | DP U-925 | | |
| | | 8470364 | Apr 07, 2024 | DP U-925 | | |
| | | 8603506 | Apr 05, 2022 | U-1063 | | |
| | | 8709478 | Apr 07, 2024 | U-1063 | | |
| | | 9241946 | Apr 05, 2022 | U-1063 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DOXYCYCLINE HYCLATE - DORYX</u> | | | | | | |
| N 050795 001 | 6958161 | Dec 15, 2022 | DP U-918 | | | |
| | 8715724 | Feb 03, 2028 | DP | | | |
| <u>DOXYCYCLINE HYCLATE - DORYX</u> | | | | | | |
| N 050795 002 | 6958161 | Dec 15, 2022 | DP U-918 | | | |
| | 8715724 | Feb 03, 2028 | DP | | | |
| <u>DOXYCYCLINE HYCLATE - DORYX</u> | | | | | | |
| N 050795 003 | 6958161 | Dec 15, 2022 | DP U-918 | | | |
| | 8715724 | Feb 03, 2028 | DP | | | |
| <u>DOXYCYCLINE HYCLATE - DORYX</u> | | | | | | |
| N 050795 004 | 6958161 | Dec 15, 2022 | DP U-918 | | | |
| | 8715724 | Feb 03, 2028 | DP | | | |
| <u>DOXYCYCLINE HYCLATE - DORYX</u> | | | | | | |
| N 050795 005 | 6958161 | Dec 15, 2022 | DP U-918 | | | |
| | 8715724 | Feb 03, 2028 | DP | | | |
| <u>DOXYCYCLINE HYCLATE - DORYX</u> | | | | | | |
| N 050795 006 | 6958161 | Dec 15, 2022 | DP U-918 | | | |
| | 8715724 | Feb 03, 2028 | DP | | | |
| <u>DOXYCYCLINE HYCLATE - DORYX MPC</u> | | | | | | |
| N 050795 007 | 6958161 | Dec 15, 2022 | DP U-918 | | | |
| | 8715724 | Feb 03, 2028 | DP | | | |
| | 9295652 | Oct 23, 2034 | DP U-918 | | | |
| | 9446057 | Dec 23, 2034 | DP U-918 | | | |
| | 9511031 | Oct 23, 2034 | DP | | | |
| <u>DOXYCYCLINE HYCLATE - DORYX MPC</u> | | | | | | |
| N 050795 008 | 6958161 | Dec 15, 2022 | DP U-918 | | | |
| | 8715724 | Feb 03, 2028 | DP | | | |
| | 9295652 | Oct 23, 2034 | DP U-918 | | | |
| | 9446057 | Dec 23, 2034 | DP U-918 | | | |
| | 9511031 | Oct 23, 2034 | DP | | | |
| <u>DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE - DICLEGIS</u> | | | | | | |
| N 021876 001 | 6340695 | Jun 21, 2021 | DP U-1382 | | | |
| | 7560122 | Jan 25, 2019 | DP | | | |
| <u>DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE - BONJESTA</u> | | | | | | |
| N 209661 001 | 7560122 | Jan 25, 2019 | DP | | | |
| | 9089489 | Feb 18, 2033 | DP U-1382 | | | |
| | 9375404 | Feb 18, 2033 | DP U-1382 | | | |
| | 9526703 | Feb 18, 2033 | DP U-1382 | | | |
| | 9937132 | Feb 18, 2033 | DP U-1382 | | | |
| <u>DRONABINOL - SYNDROS</u> | | | | | | |
| N 205525 001 | 8222292 | Aug 06, 2028 | DS DP | | | |
| | 9345771 | Aug 06, 2028 | DS DP | | | |
| <u>DRONEDARONE HYDROCHLORIDE - MULTAQ</u> | | | | | | |
| N 022425 001 | 8410167 | Apr 16, 2029 | U-1387 | | | |
| | 8410167 | Apr 16, 2029 | U-1388 | | | |
| | 8602215 | Jun 30, 2031 | U-1473 | | | |
| | 9107900 | Apr 16, 2029 | U-1726 | | | |
| | 9107900 | Apr 16, 2029 | U-1728 | | | |
| <u>DROSPIRENONE; ESTRADIOL - ANGELIO</u> | | | | | | |
| N 021355 001 | 8906890 | Oct 22, 2031 | DP | | | |
| <u>DROSPIRENONE; ETHINYL ESTRADIOL - YASMIN</u> | | | | | | |
| N 021098 001 | 6787531 | Aug 31, 2020 | DP | | | |
| <u>DROSPIRENONE; ETHINYL ESTRADIOL - YAZ</u> | | | | | | |
| N 021676 001 | 6787531 | Aug 31, 2020 | DP | | | |
| | 6958326 | Dec 20, 2021 | DP | | | |
| | 7163931 | Dec 20, 2021 | U-1 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM - BEYAZ</u> | | | | | | |
| N 022532 001 | 6441168 | Jul 30, 2022 | DS | | | |
| | 6958326 | Dec 20, 2021 | DP | | | |
| | 7163931 | Mar 03, 2022 | U-1 | | | |
| | 8617597 | Feb 08, 2030 | DP | | | |
| <u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM - SAFYRAL</u> | | | | | | |
| N 022574 001 | 6441168 | Apr 17, 2020 | DS | | | |
| | 6958326 | Dec 20, 2021 | DP | | | |
| | 7163931 | Mar 03, 2022 | U-1 | | | |
| | 8617597 | Feb 08, 2030 | DP | | | |
| <u>DROXIDOPA - NORTHERA</u> | | | | | | |
| N 203202 001 | | | | | NCE ODE-61 | Feb 18, 2019 Feb 18, 2021 |
| <u>DROXIDOPA - NORTHERA</u> | | | | | | |
| N 203202 002 | | | | | NCE ODE-61 | Feb 18, 2019 Feb 18, 2021 |
| <u>DROXIDOPA - NORTHERA</u> | | | | | | |
| N 203202 003 | | | | | NCE ODE-61 | Feb 18, 2019 Feb 18, 2021 |
| <u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u> | | | | | | |
| N 021427 001 | 6596756 | Sep 10, 2019 | U-882 | | | |
| <u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u> | | | | | | |
| N 021427 002 | 6596756 | Sep 10, 2019 | U-882 | | | |
| <u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u> | | | | | | |
| N 021427 004 | 6596756 | Sep 10, 2019 | U-882 | | | |
| <u>DUVELISIB - COPIKTRA</u> | | | | | | |
| N 211155 001 | 8193182 | Feb 13, 2030 | DS | | NCE | Sep 24, 2023 |
| | 9216982 | Jan 05, 2029 | U-2412 | | ODE-208 | Sep 24, 2025 |
| | 9216982 | Jan 05, 2029 | U-2413 | | ODE-209 | Sep 24, 2025 |
| | 9840505 | Jan 10, 2032 | U-2412 | | | |
| | 9840505 | Jan 10, 2032 | U-2413 | | | |
| | RE46621 | May 17, 2032 | DS DP | | | |
| <u>DUVELISIB - COPIKTRA</u> | | | | | | |
| N 211155 002 | 8193182 | Feb 13, 2030 | DS | | NCE | Sep 24, 2023 |
| | 9216982 | Jan 05, 2029 | U-2412 | | ODE-208 | Sep 24, 2025 |
| | 9216982 | Jan 05, 2029 | U-2413 | | ODE-209 | Sep 24, 2025 |
| | 9840505 | Jan 10, 2032 | U-2412 | | | |
| | 9840505 | Jan 10, 2032 | U-2413 | | | |
| | RE46621 | May 17, 2032 | DS DP | | | |
| <u>ECONAZOLE NITRATE - ECOZA</u> | | | | | | |
| N 205175 001 | 10071054 | Aug 08, 2031 | DP | | | |
| <u>EDARAVONE - RADICAVA</u> | | | | | | |
| N 209176 001 | 6933310 | Nov 13, 2020 | U-2013 | | NCE ODE-144 | May 05, 2022 May 05, 2024 |
| <u>EDARAVONE - RADICAVA</u> | | | | | | |
| N 209176 002 | 6933310 | Nov 13, 2020 | U-2013 | | | |
| <u>EDOXABAN TOSYLATE - SAVAYSA</u> | | | | | | |
| N 206316 001 | 7365205 | Jun 12, 2023 | DS | | NCE | Jan 08, 2020 |
| | 9149532 | Mar 28, 2028 | DP | | | |
| <u>EDOXABAN TOSYLATE - SAVAYSA</u> | | | | | | |
| N 206316 002 | 7365205 | Jun 12, 2023 | DS | | NCE | Jan 08, 2020 |
| | 9149532 | Mar 28, 2028 | DP | | | |
| <u>EDOXABAN TOSYLATE - SAVAYSA</u> | | | | | | |
| N 206316 003 | 7365205 | Jun 12, 2023 | DS | | NCE | Jan 08, 2020 |
| | 9149532 | Mar 28, 2028 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>EFAVIRENZ - SUSTIVA</u> | | | | | | |
| N 020972 | 001 | 6238695 | | DP | | |
| | | 6238695*PED | | | | |
| | | 6555133 | | U-248 | | |
| | | 6555133*PED | | | | |
| <u>EFAVIRENZ - SUSTIVA</u> | | | | | | |
| N 020972 | 002 | 6238695 | | DP | | |
| | | 6238695*PED | | | | |
| | | 6555133 | | U-248 | | |
| | | 6555133*PED | | | | |
| <u>EFAVIRENZ - SUSTIVA</u> | | | | | | |
| N 020972 | 003 | 6238695 | | DP | | |
| | | 6238695*PED | | | | |
| | | 6555133 | | U-248 | | |
| | | 6555133*PED | | | | |
| <u>EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - ATRIPLA</u> | | | | | | |
| N 021937 | 001 | 6642245 | | U-1170 | | |
| | | 6642245 | | U-750 | | |
| | | 6703396 | DS DP | | | |
| | | 8592397 | | DP U-1170 | | |
| | | 8592397 | | DP U-750 | | |
| | | 8598185 | | DP | | |
| | | 8716264 | | DP U-257 | | |
| | | 9018192 | | U-1170 | | |
| | | 9018192 | | U-750 | | |
| | | 9457036 | | DP U-257 | | |
| | | 9545414 | | DP U-1170 | | |
| | | 9545414 | | DP U-750 | | |
| | | 9744181 | | DP U-257 | | |
| <u>EFINACONAZOLE - JUBLIA</u> | | | | | | |
| N 203567 | 001 | 10105444 | | DP | NCE | Jun 06, 2019 |
| | | 7214506 | | U-281 | | |
| | | 8039494 | | U-281 | | |
| | | 8486978 | | DP | | |
| | | 9302009 | | DP | | |
| | | 9566272 | | U-1969 | | |
| | | 9662394 | | DP | | |
| | | 9861698 | | DP | | |
| | | 9877955 | | U-1969 | | |
| <u>ELAGOLIX SODIUM - ORILISSA</u> | | | | | | |
| N 210450 | 001 | 6872728 | | DS DP | NCE | Jul 23, 2023 |
| | | 7056927 | | DS DP | | |
| | | 7176211 | | U-2360 | | |
| | | 7179815 | | U-2360 | | |
| | | 7419983 | | DS DP U-2360 | | |
| | | 7462625 | | DS DP U-2360 | | |
| <u>ELAGOLIX SODIUM - ORILISSA</u> | | | | | | |
| N 210450 | 002 | 6872728 | | DS DP | NCE | Jul 23, 2023 |
| | | 7056927 | | DS DP | | |
| | | 7176211 | | U-2360 | | |
| | | 7179815 | | U-2360 | | |
| | | 7419983 | | DS DP U-2360 | | |
| | | 7462625 | | DS DP U-2360 | | |
| <u>ELBASVIR; GRAZOPREVIR - ZEPATIER</u> | | | | | | |
| N 208261 | 001 | 7973040 | | DS DP U-1813 | NCE | Jan 28, 2021 |
| | | 8871759 | | DS DP U-1813 | | |
| <u>ELIGLUSTAT TARTRATE - CERDELGA</u> | | | | | | |
| N 205494 | 001 | 6916802 | | DS U-1571 | NCE | Aug 19, 2019 |
| | | 7196205 | | DS | ODE-73 | Aug 19, 2021 |
| | | 7253185 | | DP | | |
| | | 7615573 | | U-1571 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ELTROMBOPAG OLAMINE - PROMACTA</u> | | | | | | |
| N 022291 001 | 6280959*PED | Apr 30, 2019 | | | ODE-210 | Nov 16, 2025 |
| | 7160870 | Nov 20, 2022 | DS DP U-1306 | | ODE-75 | Aug 26, 2021 |
| | 7160870 | Nov 20, 2022 | DS DP U-1575 | | PED | Feb 26, 2022 |
| | 7160870 | Nov 20, 2022 | DS DP U-1714 | | | |
| | 7160870 | Nov 20, 2022 | DS DP U-2451 | | | |
| | 7160870 | Nov 20, 2022 | DS DP U-930 | | | |
| | 7160870*PED | May 20, 2023 | | | | |
| | 7332481 | May 24, 2021 | U-1306 | | | |
| | 7332481 | May 24, 2021 | U-1575 | | | |
| | 7332481 | May 24, 2021 | U-1714 | | | |
| | 7332481 | May 24, 2021 | U-2451 | | | |
| | 7332481 | May 24, 2021 | U-930 | | | |
| | 7332481*PED | Nov 24, 2021 | | | | |
| | 7452874 | May 24, 2021 | DS DP U-1714 | | | |
| | 7452874*PED | Nov 24, 2021 | | | | |
| | 7473686 | May 24, 2021 | DS DP U-1306 | | | |
| | 7473686 | May 24, 2021 | DS DP U-1575 | | | |
| | 7473686 | May 24, 2021 | DS DP U-1714 | | | |
| | 7473686 | May 24, 2021 | DS DP U-2451 | | | |
| | 7473686 | May 24, 2021 | DS DP U-930 | | | |
| | 7473686*PED | Nov 24, 2021 | | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-1306 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-1575 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-1714 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-2451 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-2452 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-930 | | | |
| | 7547719*PED | Jan 13, 2026 | | | | |
| | 7790704 | May 24, 2021 | U-1306 | | | |
| | 7790704 | May 24, 2021 | U-1575 | | | |
| | 7790704 | May 24, 2021 | U-1714 | | | |
| | 7790704 | May 24, 2021 | U-2451 | | | |
| | 7790704 | May 24, 2021 | U-930 | | | |
| | 7790704*PED | Nov 24, 2021 | | | | |
| | 7795293 | May 21, 2023 | U-1306 | | | |
| | 7795293 | May 21, 2023 | U-1575 | | | |
| | 7795293 | May 21, 2023 | U-1714 | | | |
| | 7795293 | May 21, 2023 | U-2451 | | | |
| | 7795293 | May 21, 2023 | U-930 | | | |
| | 7795293*PED | Nov 21, 2023 | | | | |
| | 8052993 | Aug 01, 2027 | DP U-1306 | | | |
| | 8052993 | Aug 01, 2027 | DP U-1575 | | | |
| | 8052993 | Aug 01, 2027 | DP U-1714 | | | |
| | 8052993 | Aug 01, 2027 | DP U-2451 | | | |
| | 8052993 | Aug 01, 2027 | DP U-930 | | | |
| | 8052993*PED | Feb 01, 2028 | | | | |
| | 8052994 | Aug 01, 2027 | DP U-1714 | | | |
| | 8052994*PED | Feb 01, 2028 | | | | |
| | 8062665 | Aug 01, 2027 | DP U-1714 | | | |
| | 8062665*PED | Feb 01, 2028 | | | | |
| | 8071129 | Aug 01, 2027 | DP U-1714 | | | |
| | 8071129*PED | Feb 01, 2028 | | | | |
| | 8828430 | Aug 01, 2027 | DP U-1306 | | | |
| | 8828430 | Aug 01, 2027 | DP U-1619 | | | |
| | 8828430 | Aug 01, 2027 | DP U-1714 | | | |
| | 8828430 | Aug 01, 2027 | DP U-2451 | | | |
| | 8828430*PED | Feb 01, 2028 | | | | |
| <u>ELTROMBOPAG OLAMINE - PROMACTA</u> | | | | | | |
| N 022291 002 | 6280959*PED | Apr 30, 2019 | | | ODE-210 | Nov 16, 2025 |
| | 7160870 | Nov 20, 2022 | DS DP U-1306 | | ODE-75 | Aug 26, 2021 |
| | 7160870 | Nov 20, 2022 | DS DP U-1575 | | PED | Feb 26, 2022 |
| | 7160870 | Nov 20, 2022 | DS DP U-1714 | | | |
| | 7160870 | Nov 20, 2022 | DS DP U-2451 | | | |
| | 7160870 | Nov 20, 2022 | DS DP U-930 | | | |
| | 7160870*PED | May 20, 2023 | | | | |
| | 7332481 | May 24, 2021 | U-1306 | | | |
| | 7332481 | May 24, 2021 | U-1575 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ELTROMBOPAG OLAMINE - PROMACTA</u> | | | | | | |
| N 022291 002 | 7332481 | May 24, 2021 | U-1714 | | | |
| | 7332481 | May 24, 2021 | U-2451 | | | |
| | 7332481 | May 24, 2021 | U-930 | | | |
| | 7332481*PED | Nov 24, 2021 | | | | |
| | 7452874 | May 24, 2021 | DS DP U-1714 | | | |
| | 7452874*PED | Nov 24, 2021 | | | | |
| | 7473686 | May 24, 2021 | DS DP U-1306 | | | |
| | 7473686 | May 24, 2021 | DS DP U-1575 | | | |
| | 7473686 | May 24, 2021 | DS DP U-1714 | | | |
| | 7473686 | May 24, 2021 | DS DP U-2451 | | | |
| | 7473686 | May 24, 2021 | DS DP U-930 | | | |
| | 7473686*PED | Nov 24, 2021 | | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-1306 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-1575 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-1714 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-2451 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-2452 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-930 | | | |
| | 7547719*PED | Jan 13, 2026 | | | | |
| | 7790704 | May 24, 2021 | U-1306 | | | |
| | 7790704 | May 24, 2021 | U-1575 | | | |
| | 7790704 | May 24, 2021 | U-1714 | | | |
| | 7790704 | May 24, 2021 | U-2451 | | | |
| | 7790704 | May 24, 2021 | U-930 | | | |
| | 7790704*PED | Nov 24, 2021 | | | | |
| | 7795293 | May 21, 2023 | U-1306 | | | |
| | 7795293 | May 21, 2023 | U-1575 | | | |
| | 7795293 | May 21, 2023 | U-1714 | | | |
| | 7795293 | May 21, 2023 | U-2451 | | | |
| | 7795293 | May 21, 2023 | U-930 | | | |
| | 7795293*PED | Nov 21, 2023 | | | | |
| | 8052993 | Aug 01, 2027 | DP U-1714 | | | |
| | 8052993*PED | Feb 01, 2028 | | | | |
| | 8052994 | Aug 01, 2027 | DP U-1306 | | | |
| | 8052994 | Aug 01, 2027 | DP U-1575 | | | |
| | 8052994 | Aug 01, 2027 | DP U-1714 | | | |
| | 8052994 | Aug 01, 2027 | DP U-2451 | | | |
| | 8052994 | Aug 01, 2027 | DP U-930 | | | |
| | 8052994*PED | Feb 01, 2028 | | | | |
| | 8062665 | Aug 01, 2027 | DP U-1714 | | | |
| | 8062665*PED | Feb 01, 2028 | | | | |
| | 8071129 | Aug 01, 2027 | DP U-1714 | | | |
| | 8071129*PED | Feb 01, 2028 | | | | |
| | 8828430 | Aug 01, 2027 | DP U-1306 | | | |
| | 8828430 | Aug 01, 2027 | DP U-1619 | | | |
| | 8828430 | Aug 01, 2027 | DP U-1714 | | | |
| | 8828430 | Aug 01, 2027 | DP U-2451 | | | |
| | 8828430*PED | Feb 01, 2028 | | | | |
| <u>ELTROMBOPAG OLAMINE - PROMACTA</u> | | | | | | |
| N 022291 003 | 6280959*PED | Apr 30, 2019 | | | ODE-210 | Nov 16, 2025 |
| | 7160870 | Nov 20, 2022 | DS DP U-1306 | | ODE-75 | Aug 26, 2021 |
| | 7160870 | Nov 20, 2022 | DS DP U-1575 | | PED | Feb 26, 2022 |
| | 7160870 | Nov 20, 2022 | DS DP U-1714 | | | |
| | 7160870 | Nov 20, 2022 | DS DP U-2451 | | | |
| | 7160870 | Nov 20, 2022 | DS DP U-930 | | | |
| | 7160870*PED | May 20, 2023 | | | | |
| | 7332481 | May 24, 2021 | U-1306 | | | |
| | 7332481 | May 24, 2021 | U-1575 | | | |
| | 7332481 | May 24, 2021 | U-1714 | | | |
| | 7332481 | May 24, 2021 | U-2451 | | | |
| | 7332481 | May 24, 2021 | U-930 | | | |
| | 7332481*PED | Nov 24, 2021 | | | | |
| | 7452874 | May 24, 2021 | DS DP U-1714 | | | |
| | 7452874*PED | Nov 24, 2021 | | | | |
| | 7473686 | May 24, 2021 | DS DP U-1306 | | | |
| | 7473686 | May 24, 2021 | DS DP U-1575 | | | |
| | 7473686 | May 24, 2021 | DS DP U-1714 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ELTROMBOPAG OLAMINE - PROMACTA</u> | | | | | | |
| N 022291 003 | 7473686 | May 24, 2021 | DS DP U-2451 | | | |
| | 7473686 | May 24, 2021 | DS DP U-930 | | | |
| | 7473686*PED | Nov 24, 2021 | | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-1306 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-1575 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-1714 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-2451 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-2452 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-930 | | | |
| | 7547719*PED | Jan 13, 2026 | | | | |
| | 7790704 | May 24, 2021 | U-1306 | | | |
| | 7790704 | May 24, 2021 | U-1575 | | | |
| | 7790704 | May 24, 2021 | U-1714 | | | |
| | 7790704 | May 24, 2021 | U-2451 | | | |
| | 7790704 | May 24, 2021 | U-930 | | | |
| | 7790704*PED | Nov 24, 2021 | | | | |
| | 7795293 | May 21, 2023 | U-1306 | | | |
| | 7795293 | May 21, 2023 | U-1575 | | | |
| | 7795293 | May 21, 2023 | U-1714 | | | |
| | 7795293 | May 21, 2023 | U-2451 | | | |
| | 7795293 | May 21, 2023 | U-930 | | | |
| | 7795293*PED | Nov 21, 2023 | | | | |
| | 8052993 | Aug 01, 2027 | DP U-1714 | | | |
| | 8052993*PED | Feb 01, 2028 | | | | |
| | 8052994 | Aug 01, 2027 | DP U-1714 | | | |
| | 8052994*PED | Feb 01, 2028 | | | | |
| | 8062665 | Aug 01, 2027 | DP U-1306 | | | |
| | 8062665 | Aug 01, 2027 | DP U-1575 | | | |
| | 8062665 | Aug 01, 2027 | DP U-1714 | | | |
| | 8062665 | Aug 01, 2027 | DP U-2451 | | | |
| | 8062665 | Aug 01, 2027 | DP U-930 | | | |
| | 8062665*PED | Feb 01, 2028 | | | | |
| | 8071129 | Aug 01, 2027 | DP U-1714 | | | |
| | 8071129*PED | Feb 01, 2028 | | | | |
| | 8828430 | Aug 01, 2027 | DP U-1306 | | | |
| | 8828430 | Aug 01, 2027 | DP U-1619 | | | |
| | 8828430 | Aug 01, 2027 | DP U-1714 | | | |
| | 8828430 | Aug 01, 2027 | DP U-2451 | | | |
| | 8828430*PED | Feb 01, 2028 | | | | |
| <u>ELTROMBOPAG OLAMINE - PROMACTA</u> | | | | | | |
| N 022291 004 | 6280959*PED | Apr 30, 2019 | | | ODE-210 | Nov 16, 2025 |
| | 7160870 | Nov 20, 2022 | DS DP U-1306 | | ODE-75 | Aug 26, 2021 |
| | 7160870 | Nov 20, 2022 | DS DP U-1575 | | PED | Feb 26, 2022 |
| | 7160870 | Nov 20, 2022 | DS DP U-1714 | | | |
| | 7160870 | Nov 20, 2022 | DS DP U-2451 | | | |
| | 7160870 | Nov 20, 2022 | DS DP U-930 | | | |
| | 7160870*PED | May 20, 2023 | | | | |
| | 7332481 | May 24, 2021 | U-1306 | | | |
| | 7332481 | May 24, 2021 | U-1575 | | | |
| | 7332481 | May 24, 2021 | U-1714 | | | |
| | 7332481 | May 24, 2021 | U-2451 | | | |
| | 7332481 | May 24, 2021 | U-930 | | | |
| | 7332481*PED | Nov 24, 2021 | | | | |
| | 7452874 | May 24, 2021 | DS DP U-1714 | | | |
| | 7452874*PED | Nov 24, 2021 | | | | |
| | 7473686 | May 24, 2021 | DS DP U-1306 | | | |
| | 7473686 | May 24, 2021 | DS DP U-1575 | | | |
| | 7473686 | May 24, 2021 | DS DP U-1714 | | | |
| | 7473686 | May 24, 2021 | DS DP U-2451 | | | |
| | 7473686 | May 24, 2021 | DS DP U-930 | | | |
| | 7473686*PED | Nov 24, 2021 | | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-1306 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-1575 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-1714 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-2451 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-2452 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-930 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ELTROMBOPAG OLAMINE - PROMACTA</u> | | | | | | |
| N 022291 004 | 7547719*PED | Jan 13, 2026 | | | | |
| | 7790704 | May 24, 2021 | U-1306 | | | |
| | 7790704 | May 24, 2021 | U-1575 | | | |
| | 7790704 | May 24, 2021 | U-1714 | | | |
| | 7790704 | May 24, 2021 | U-2451 | | | |
| | 7790704 | May 24, 2021 | U-930 | | | |
| | 7790704*PED | Nov 24, 2021 | | | | |
| | 7795293 | May 21, 2023 | U-1306 | | | |
| | 7795293 | May 21, 2023 | U-1575 | | | |
| | 7795293 | May 21, 2023 | U-1714 | | | |
| | 7795293 | May 21, 2023 | U-2451 | | | |
| | 7795293 | May 21, 2023 | U-930 | | | |
| | 7795293*PED | Nov 21, 2023 | | | | |
| | 8052993 | Aug 01, 2027 | DP U-1714 | | | |
| | 8052993*PED | Feb 01, 2028 | | | | |
| | 8052994 | Aug 01, 2027 | DP U-1714 | | | |
| | 8052994*PED | Feb 01, 2028 | | | | |
| | 8062665 | Aug 01, 2027 | DP U-1714 | | | |
| | 8062665*PED | Feb 01, 2028 | | | | |
| | 8071129 | Aug 01, 2027 | DP U-1306 | | | |
| | 8071129 | Aug 01, 2027 | DP U-1575 | | | |
| | 8071129 | Aug 01, 2027 | DP U-1714 | | | |
| | 8071129 | Aug 01, 2027 | DP U-2451 | | | |
| | 8071129 | Aug 01, 2027 | DP U-930 | | | |
| | 8071129*PED | Feb 01, 2028 | | | | |
| | 8828430 | Aug 01, 2027 | DP U-1306 | | | |
| | 8828430 | Aug 01, 2027 | DP U-1619 | | | |
| | 8828430 | Aug 01, 2027 | DP U-1714 | | | |
| | 8828430 | Aug 01, 2027 | DP U-2451 | | | |
| | 8828430*PED | Feb 01, 2028 | | | | |
| <u>ELTROMBOPAG OLAMINE - PROMACTA</u> | | | | | | |
| N 022291 005 | 6280959*PED | Apr 30, 2019 | | | ODE-210 | Nov 16, 2025 |
| | 7160870 | Nov 20, 2022 | DS DP U-1306 | | ODE-75 | Aug 26, 2021 |
| | 7160870 | Nov 20, 2022 | DS DP U-1575 | | PED | Feb 26, 2022 |
| | 7160870 | Nov 20, 2022 | DS DP U-1714 | | | |
| | 7160870 | Nov 20, 2022 | DS DP U-930 | | | |
| | 7160870*PED | May 20, 2023 | | | | |
| | 7332481 | May 24, 2021 | U-1306 | | | |
| | 7332481 | May 24, 2021 | U-1575 | | | |
| | 7332481 | May 24, 2021 | U-1714 | | | |
| | 7332481 | May 24, 2021 | U-930 | | | |
| | 7332481*PED | Nov 24, 2021 | | | | |
| | 7452874 | May 24, 2021 | DS DP U-1714 | | | |
| | 7452874*PED | Nov 24, 2021 | | | | |
| | 7473686 | May 24, 2021 | DS DP U-1306 | | | |
| | 7473686 | May 24, 2021 | DS DP U-1575 | | | |
| | 7473686 | May 24, 2021 | DS DP U-1714 | | | |
| | 7473686 | May 24, 2021 | DS DP U-930 | | | |
| | 7473686*PED | Nov 24, 2021 | | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-1306 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-1575 | | | |
| | 7547719 | Jul 13, 2025 | DS DP U-930 | | | |
| | 7547719*PED | Jan 13, 2026 | | | | |
| | 7790704 | May 24, 2021 | U-1306 | | | |
| | 7790704 | May 24, 2021 | U-1575 | | | |
| | 7790704 | May 24, 2021 | U-930 | | | |
| | 7790704*PED | Nov 24, 2021 | | | | |
| | 7795293 | May 21, 2023 | U-1306 | | | |
| | 7795293 | May 21, 2023 | U-1575 | | | |
| | 7795293 | May 21, 2023 | U-930 | | | |
| | 7795293*PED | Nov 21, 2023 | | | | |
| | 8052995 | Aug 01, 2027 | DP U-1306 | | | |
| | 8052995 | Aug 01, 2027 | DP U-1575 | | | |
| | 8052995*PED | Feb 01, 2028 | | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ELTROMBOPAG OLAMINE - PROMACTA KIT</u> | | | | | | |
| N 207027 | 001 | 6280959*PED | Apr 30, 2019 | | ODE-74 | Aug 26, 2021 |
| | | 7160870 | Nov 20, 2022 | DS DP U-1736 | PED | Feb 26, 2022 |
| | | 7160870*PED | May 20, 2023 | | | |
| | | 7332481 | May 24, 2021 | U-1736 | | |
| | | 7332481*PED | Nov 24, 2021 | | | |
| | | 7452874 | May 24, 2021 | DS DP | | |
| | | 7452874*PED | Nov 24, 2021 | | | |
| | | 7473686 | May 24, 2021 | DS DP U-1736 | | |
| | | 7473686*PED | Nov 24, 2021 | | | |
| | | 7547719 | Jul 13, 2025 | DS DP U-1736 | | |
| | | 7547719*PED | Jan 13, 2026 | | | |
| | | 7790704 | May 24, 2021 | U-1736 | | |
| | | 7790704*PED | Nov 24, 2021 | | | |
| | | 7795293 | May 24, 2023 | U-1736 | | |
| | | 7795293*PED | Nov 24, 2023 | | | |
| <u>ELUXADOLINE - VIBERZI</u> | | | | | | |
| N 206940 | 001 | 7741356 | Mar 25, 2028 | DS DP | NCE | May 27, 2020 |
| | | 7786158 | Mar 14, 2025 | DS | | |
| | | 8344011 | Mar 14, 2025 | U-1709 | | |
| | | 8609709 | Mar 14, 2025 | DS | | |
| | | 8691860 | Jul 07, 2028 | DS U-1709 | | |
| | | 8772325 | Mar 14, 2025 | U-1709 | | |
| | | 9115091 | Jul 07, 2028 | DS DP U-1738 | | |
| | | 9205076 | Mar 14, 2025 | U-1709 | | |
| | | 9364489 | Jul 07, 2028 | U-1709 | | |
| | | 9675587 | Mar 14, 2033 | DP | | |
| | | 9700542 | Mar 14, 2025 | DP | | |
| | | 9789125 | Jul 07, 2028 | DP U-1709 | | |
| | | 9789125 | Jul 07, 2028 | DP U-2152 | | |
| <u>ELUXADOLINE - VIBERZI</u> | | | | | | |
| N 206940 | 002 | 7741356 | Mar 25, 2028 | DS DP | NCE | May 27, 2020 |
| | | 7786158 | Mar 14, 2025 | DS | | |
| | | 8344011 | Mar 14, 2025 | U-1709 | | |
| | | 8609709 | Mar 14, 2025 | DS | | |
| | | 8691860 | Jul 07, 2028 | DS U-1709 | | |
| | | 8772325 | Mar 14, 2025 | U-1709 | | |
| | | 9115091 | Jul 07, 2028 | DS DP U-1738 | | |
| | | 9205076 | Mar 14, 2025 | U-1709 | | |
| | | 9364489 | Jul 07, 2028 | U-1709 | | |
| | | 9675587 | Mar 14, 2033 | DP | | |
| | | 9700542 | Mar 14, 2025 | DP | | |
| | | 9789125 | Jul 07, 2028 | DP U-1709 | | |
| | | 9789125 | Jul 07, 2028 | DP U-2152 | | |
| <u>ELVITEGRAVIR - VITEKTA</u> | | | | | | |
| N 203093 | 001 | 7176220 | Aug 27, 2026 | DS DP U-257 | | |
| | | 7635704 | Oct 26, 2026 | DS DP U-257 | | |
| | | 8981103 | Oct 26, 2026 | DS DP | | |
| <u>ELVITEGRAVIR - VITEKTA</u> | | | | | | |
| N 203093 | 002 | 7176220 | Aug 27, 2026 | DS DP U-257 | | |
| | | 7635704 | Oct 26, 2026 | DS DP U-257 | | |
| | | 8981103 | Oct 26, 2026 | DS DP | | |
| <u>EMPAGLIFLOZIN - JARDIANCE</u> | | | | | | |
| N 204629 | 001 | 7579449 | Nov 05, 2025 | DS | I-739 | Dec 02, 2019 |
| | | 7713938 | Apr 15, 2027 | DS DP | M-174 | Mar 18, 2019 |
| | | 8551957 | Oct 14, 2029 | U-1651 | NCE | Aug 01, 2019 |
| | | 9949997 | May 17, 2034 | U-2292 | | |
| | | 9949998 | Jun 11, 2034 | U-2290 | | |
| <u>EMPAGLIFLOZIN - JARDIANCE</u> | | | | | | |
| N 204629 | 002 | 7579449 | Nov 05, 2025 | DS | I-739 | Dec 02, 2019 |
| | | 7713938 | Apr 15, 2027 | DS DP | M-174 | Mar 18, 2019 |
| | | 8551957 | Oct 14, 2029 | U-1651 | NCE | Aug 01, 2019 |
| | | 9949997 | May 17, 2034 | U-2292 | | |
| | | 9949998 | Jun 11, 2034 | U-2290 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>EMPAGLIFLOZIN - JARDIANCE</u> | | | | | | |
| N 204629 002 | 7579449 | Nov 05, 2025 | DS | | I-739 | Dec 02, 2019 |
| | 7713938 | Apr 15, 2027 | DS DP | | M-174 | Mar 18, 2019 |
| | 8551957 | Oct 14, 2029 | | U-1651 | NCE | Aug 01, 2019 |
| | 9949997 | May 17, 2034 | | U-2292 | | |
| | 9949998 | Jun 11, 2034 | | U-2290 | | |
| <u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u> | | | | | | |
| N 206073 001 | 6890898 | Feb 02, 2019 | | U-1652 | I-739 | Dec 02, 2019 |
| | 7078381 | Feb 02, 2019 | | U-1651 | NCE | Aug 01, 2019 |
| | 7407955 | May 02, 2025 | DS DP | | | |
| | 7459428 | Feb 02, 2019 | | U-1651 | | |
| | 7579449 | Nov 05, 2025 | DS | | | |
| | 7713938 | Apr 15, 2027 | DS DP | | | |
| | 8119648 | Aug 12, 2023 | | U-1651 | | |
| | 8178541 | Aug 12, 2023 | DP | U-1653 | | |
| | 8178541 | Aug 12, 2023 | DP | U-1654 | | |
| | 8551957 | Oct 14, 2029 | DP | U-1651 | | |
| | 8673927 | May 04, 2027 | | U-1652 | | |
| | 8883805 | Nov 26, 2025 | DP | | | |
| | 9173859 | May 04, 2027 | DP | U-1772 | | |
| | 9949998 | Jun 11, 2034 | | U-2290 | | |
| <u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u> | | | | | | |
| N 206073 002 | 6890898 | Feb 02, 2019 | | U-1652 | I-739 | Dec 02, 2019 |
| | 7078381 | Feb 02, 2019 | | U-1651 | NCE | Aug 01, 2019 |
| | 7407955 | May 02, 2025 | DS DP | | | |
| | 7459428 | Feb 02, 2019 | | U-1651 | | |
| | 7579449 | Nov 05, 2025 | DS | | | |
| | 7713938 | Apr 15, 2027 | DS DP | | | |
| | 8119648 | Aug 12, 2023 | | U-1651 | | |
| | 8178541 | Aug 12, 2023 | DP | U-1653 | | |
| | 8178541 | Aug 12, 2023 | DP | U-1654 | | |
| | 8551957 | Oct 14, 2029 | DP | U-1651 | | |
| | 8673927 | May 04, 2027 | | U-1652 | | |
| | 8883805 | Nov 26, 2025 | DP | | | |
| | 9173859 | May 04, 2027 | DP | U-1772 | | |
| | 9949998 | Jun 11, 2034 | | U-2290 | | |
| <u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u> | | | | | | |
| N 206111 001 | 7579449 | Nov 05, 2025 | DS | | I-739 | Dec 02, 2019 |
| | 7713938 | Apr 15, 2027 | DS DP | | M-174 | Mar 18, 2019 |
| | | | | | NCE | Aug 01, 2019 |
| <u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u> | | | | | | |
| N 206111 002 | 7579449 | Nov 05, 2025 | DS | | I-739 | Dec 02, 2019 |
| | 7713938 | Apr 15, 2027 | DS DP | | M-174 | Mar 18, 2019 |
| | | | | | NCE | Aug 01, 2019 |
| <u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u> | | | | | | |
| N 206111 003 | 7579449 | Nov 05, 2025 | DS | | I-739 | Dec 02, 2019 |
| | 7713938 | Apr 15, 2027 | DS DP | | M-174 | Mar 18, 2019 |
| | | | | | NCE | Aug 01, 2019 |
| <u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u> | | | | | | |
| N 206111 004 | 7579449 | Nov 05, 2025 | DS | | I-739 | Dec 02, 2019 |
| | 7713938 | Apr 15, 2027 | DS DP | | M-174 | Mar 18, 2019 |
| | | | | | NCE | Aug 01, 2019 |
| <u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u> | | | | | | |
| N 208658 001 | 6488962 | Jun 20, 2020 | | DP | I-739 | Dec 02, 2019 |
| | 7579449 | Nov 05, 2025 | DS | | NCE | Aug 01, 2019 |
| | 7713938 | Apr 15, 2027 | DS DP | | | |
| | 9949998 | Jun 11, 2034 | | U-2290 | | |
| <u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u> | | | | | | |
| N 208658 002 | 6488962 | Jun 20, 2020 | | DP | I-739 | Dec 02, 2019 |
| | 7579449 | Nov 05, 2025 | DS | | NCE | Aug 01, 2019 |
| | 7713938 | Apr 15, 2027 | DS DP | | | |
| | 9949998 | Jun 11, 2034 | | U-2290 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u> | | | | | | |
| N 208658 002 | 6488962 | Jun 20, 2020 | DP | | I-739 | Dec 02, 2019 |
| | 7579449 | Nov 05, 2025 | DS | | NCE | Aug 01, 2019 |
| | 7713938 | Apr 15, 2027 | DS DP | | | |
| | 9949998 | Jun 11, 2034 | | U-2290 | | |
| <u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u> | | | | | | |
| N 208658 003 | 6488962 | Jun 20, 2020 | DP | | I-739 | Dec 02, 2019 |
| | 7579449 | Nov 05, 2025 | DS | | NCE | Aug 01, 2019 |
| | 7713938 | Apr 15, 2027 | DS DP | | | |
| | 9949998 | Jun 11, 2034 | | U-2290 | | |
| <u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u> | | | | | | |
| N 208658 004 | 6488962 | Jun 20, 2020 | DP | | I-739 | Dec 02, 2019 |
| | 7579449 | Nov 05, 2025 | DS | | NCE | Aug 01, 2019 |
| | 7713938 | Apr 15, 2027 | DS DP | | | |
| | 9949998 | Jun 11, 2034 | | U-2290 | | |
| <u>EMTRICITABINE - EMTRIVA</u> | | | | | | |
| N 021500 001 | 6642245 | Nov 04, 2020 | | U-257 | | |
| | 6642245 | Nov 04, 2020 | | U-541 | | |
| | 6703396 | Mar 09, 2021 | DS DP | | | |
| <u>EMTRICITABINE - EMTRIVA</u> | | | | | | |
| N 021896 001 | 6642245 | Nov 04, 2020 | | U-257 | | |
| | 6703396 | Mar 09, 2021 | DS DP | | | |
| <u>EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR ALAFENAMIDE FUMARATE - ODEFSEY</u> | | | | | | |
| N 208351 001 | 6642245 | Nov 04, 2020 | | U-257 | M-206 | Aug 21, 2020 |
| | 6642245*PED | May 04, 2021 | | | M-207 | Aug 21, 2020 |
| | 6703396 | Mar 09, 2021 | DS DP | | NCE | Nov 05, 2020 |
| | 6703396*PED | Sep 09, 2021 | | | | |
| | 6838464 | Feb 26, 2021 | DS DP | | | |
| | 7067522 | Dec 20, 2019 | DS DP | | | |
| | 7125879 | Apr 21, 2025 | DS DP | U-257 | | |
| | 7390791 | May 07, 2022 | DS DP | | | |
| | 7803788 | Feb 02, 2022 | | U-257 | | |
| | 8080551 | Apr 11, 2023 | DS DP | | | |
| | 8101629 | Aug 09, 2022 | DP | | | |
| | 8754065 | Aug 15, 2032 | DS DP | U-257 | | |
| | 9296769 | Aug 15, 2032 | DS DP | U-257 | | |
| <u>EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR DISOPROXIL FUMARATE - COMPLERA</u> | | | | | | |
| N 202123 001 | 6642245 | Nov 04, 2020 | | U-257 | | |
| | 6703396 | Mar 09, 2021 | DS DP | | | |
| | 6838464 | Feb 26, 2021 | DS DP | | | |
| | 7067522 | Dec 20, 2019 | DS DP | | | |
| | 7125879 | Apr 21, 2025 | DS DP | U-257 | | |
| | 8080551 | Apr 11, 2023 | DS DP | | | |
| | 8101629 | Aug 09, 2022 | DP | | | |
| | 8592397 | Jan 13, 2024 | DP | U-257 | | |
| | 8716264 | Jan 13, 2024 | DP | U-257 | | |
| | 8841310 | Dec 09, 2025 | DP | U-257 | | |
| | 9457036 | Jan 13, 2024 | DP | U-257 | | |
| | 9744181 | Jan 13, 2024 | DP | U-257 | | |
| <u>EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - DESCOVY</u> | | | | | | |
| N 208215 001 | 6642245 | Nov 04, 2020 | | U-257 | NCE | Nov 05, 2020 |
| | 6642245*PED | May 04, 2021 | | | NPP | Sep 25, 2020 |
| | 6703396 | Mar 09, 2021 | DS DP | | | |
| | 6703396*PED | Sep 09, 2021 | | | | |
| | 7390791 | May 07, 2022 | DS DP | | | |
| | 7803788 | Feb 02, 2022 | | U-257 | | |
| | 8754065 | Aug 15, 2032 | DS DP | U-257 | | |
| | 9296769 | Aug 15, 2032 | DS DP | U-257 | | |
| <u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u> | | | | | | |
| N 021752 001 | 6642245 | Nov 04, 2020 | | U-1170 | | |
| | 6642245 | Nov 04, 2020 | | U-248 | | |
| | 6642245 | Nov 04, 2020 | | U-541 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u> | | | | | | |
| N 021752 001 | 6703396 | Mar 09, 2021 | DS DP | | | |
| | 8592397 | Jan 13, 2024 | DP U-1170 | | | |
| | 8592397 | Jan 13, 2024 | DP U-248 | | | |
| | 8592397 | Jan 13, 2024 | DP U-541 | | | |
| | 8716264 | Jan 13, 2024 | DP U-257 | | | |
| | 9457036 | Jan 13, 2024 | DP U-257 | | | |
| | 9744181 | Jan 13, 2024 | DP U-257 | | | |
| <u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u> | | | | | | |
| N 021752 002 | 6642245 | Nov 04, 2020 | | | U-1170 | |
| | 6642245 | Nov 04, 2020 | | | U-248 | |
| | 6642245 | Nov 04, 2020 | | | U-541 | |
| | 6642245*PED | May 04, 2021 | | | | |
| | 6703396 | Mar 09, 2021 | DS DP | | | |
| | 6703396*PED | Sep 09, 2021 | | | | |
| <u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u> | | | | | | |
| N 021752 003 | 6642245 | Nov 04, 2020 | | | U-1170 | |
| | 6642245 | Nov 04, 2020 | | | U-248 | |
| | 6642245 | Nov 04, 2020 | | | U-541 | |
| | 6642245*PED | May 04, 2021 | | | | |
| | 6703396 | Mar 09, 2021 | DS DP | | | |
| | 6703396*PED | Sep 09, 2021 | | | | |
| <u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u> | | | | | | |
| N 021752 004 | 6642245 | Nov 04, 2020 | | | U-1170 | |
| | 6642245 | Nov 04, 2020 | | | U-248 | |
| | 6642245 | Nov 04, 2020 | | | U-541 | |
| | 6642245*PED | May 04, 2021 | | | | |
| | 6703396 | Mar 09, 2021 | DS DP | | | |
| | 6703396*PED | Sep 09, 2021 | | | | |
| <u>ENALAPRIL MALEATE - EPANED KIT</u> | | | | | | |
| N 204308 001 | 8568747 | Nov 06, 2032 | DP | | | |
| | 8778366 | Nov 06, 2032 | | | U-1723 | |
| | 8778366 | Nov 06, 2032 | | | U-185 | |
| | 8778366 | Nov 06, 2032 | | | U-1892 | |
| | 8778366 | Nov 06, 2032 | | | U-3 | |
| | 8778366 | Nov 06, 2032 | | | U-71 | |
| | 9855214 | Nov 06, 2032 | DP | | | |
| | 9968553 | Nov 06, 2032 | | | U-1723 | |
| | 9968553 | Nov 06, 2032 | | | U-185 | |
| | 9968553 | Nov 06, 2032 | | | U-1892 | |
| | 9968553 | Nov 06, 2032 | | | U-3 | |
| | 9968553 | Nov 06, 2032 | | | U-71 | |
| <u>ENALAPRIL MALEATE - EPANED</u> | | | | | | |
| N 208686 001 | 10039745 | Mar 25, 2036 | DP | | | |
| | 10154987 | Mar 25, 2036 | | | U-1723 | |
| | 10154987 | Mar 25, 2036 | | | U-185 | |
| | 10154987 | Mar 25, 2036 | | | U-1892 | |
| | 10154987 | Mar 25, 2036 | | | U-3 | |
| | 10154987 | Mar 25, 2036 | | | U-71 | |
| | 9669008 | Mar 25, 2036 | DP | | | |
| | 9808442 | Mar 25, 2036 | | | U-1723 | |
| | 9808442 | Mar 25, 2036 | | | U-185 | |
| | 9808442 | Mar 25, 2036 | | | U-1892 | |
| | 9808442 | Mar 25, 2036 | | | U-3 | |
| | 9808442 | Mar 25, 2036 | | | U-71 | |
| <u>ENASIDENIB MESYLATE - IDHIFA</u> | | | | | | |
| N 209606 001 | 10093654 | Aug 01, 2034 | DS DP | | U-2087 | |
| | 9512107 | Jan 07, 2033 | DS DP | | U-2087 | |
| | 9732062 | Sep 16, 2034 | DS | | | |
| | 9738625 | Aug 01, 2034 | DS | | | |
| <u>ENASIDENIB MESYLATE - IDHIFA</u> | | | | | | |
| N 209606 002 | 10093654 | Aug 01, 2034 | DS DP | | U-2087 | |
| | 9512107 | Jan 07, 2033 | DS DP | | U-2087 | |
| | | | | | | NCE Aug 01, 2022 |
| | | | | | | ODE-151 Aug 01, 2024 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ENASIDENIB MESYLATE - IDHIFA</u> | | | | | | |
| N 209606 | 002 | 9732062 | Sep 16, 2034 | DS | | |
| | | 9738625 | Aug 01, 2034 | DS | | |
| <u>ENCORAFENIB - BRAFTOVI</u> | | | | | | |
| N 210496 | 001 | 10005761 | Aug 27, 2030 | | U-2335 | NCE |
| | | 8501758 | Mar 04, 2031 | DS DP | | Jun 27, 2023 |
| | | 8541575 | Feb 26, 2030 | DS DP | U-2335 | Jun 27, 2025 |
| | | 8946250 | Jul 23, 2029 | DS DP | | |
| | | 9314464 | Jul 04, 2031 | | U-2336 | |
| | | 9387208 | Nov 21, 2032 | DP | | |
| | | 9593099 | Aug 27, 2030 | DS | | |
| | | 9593100 | Aug 27, 2030 | DP | | |
| | | 9763941 | Nov 21, 2032 | | U-2335 | |
| | | 9850229 | Aug 27, 2030 | | U-2337 | |
| | | 9850230 | Aug 27, 2030 | | U-2334 | |
| <u>ENCORAFENIB - BRAFTOVI</u> | | | | | | |
| N 210496 | 002 | 10005761 | Aug 27, 2030 | | U-2335 | NCE |
| | | 8501758 | Mar 04, 2031 | DS DP | | Jun 27, 2023 |
| | | 8541575 | Feb 26, 2030 | DS DP | U-2335 | Jun 27, 2025 |
| | | 8946250 | Jul 23, 2029 | DS DP | | |
| | | 9314464 | Jul 04, 2031 | | U-2336 | |
| | | 9387208 | Nov 21, 2032 | DP | | |
| | | 9593099 | Aug 27, 2030 | DS | | |
| | | 9593100 | Aug 27, 2030 | DP | | |
| | | 9763941 | Nov 21, 2032 | | U-2335 | |
| | | 9850229 | Aug 27, 2030 | | U-2337 | |
| | | 9850230 | Aug 27, 2030 | | U-2334 | |
| <u>ENZALUTAMIDE - XTANDI</u> | | | | | | |
| N 203415 | 001 | 7709517 | Aug 13, 2027 | DS DP | | I-786 |
| | | 8183274 | Aug 24, 2026 | | U-1281 | Jul 13, 2021 |
| | | 8183274 | Aug 24, 2026 | | U-1588 | |
| | | 8183274 | Aug 24, 2026 | | U-2345 | |
| | | 9126941 | May 15, 2026 | | U-1588 | |
| | | 9126941 | May 15, 2026 | | U-2345 | |
| <u>EPINEPHRINE - EPIPEN</u> | | | | | | |
| N 019430 | 001 | 7449012 | Sep 11, 2025 | DP | | |
| | | 7794432 | Sep 11, 2025 | DP | | |
| | | 8048035 | Sep 11, 2025 | DP | | |
| | | 8870827 | Sep 11, 2025 | DP | | |
| | | 9586010 | Sep 11, 2025 | DP | | |
| <u>EPINEPHRINE - EPIPEN JR.</u> | | | | | | |
| N 019430 | 002 | 7449012 | Sep 11, 2025 | DP | | |
| | | 7794432 | Sep 11, 2025 | DP | | |
| | | 8048035 | Sep 11, 2025 | DP | | |
| | | 8870827 | Sep 11, 2025 | DP | | |
| | | 9586010 | Sep 11, 2025 | DP | | |
| <u>EPINEPHRINE - TWINJECT 0.3</u> | | | | | | |
| N 020800 | 001 | 7297136 | Jan 18, 2025 | DP | | |
| | | 7621891 | Feb 04, 2025 | DP | | |
| <u>EPINEPHRINE - TWINJECT 0.15</u> | | | | | | |
| N 020800 | 002 | 7297136 | Jan 18, 2025 | DP | | |
| | | 7621891 | Feb 04, 2025 | DP | | |
| <u>EPINEPHRINE - ADRENACLICK</u> | | | | | | |
| N 020800 | 003 | 7905352 | Apr 12, 2027 | DP | | |
| <u>EPINEPHRINE - ADRENACLICK</u> | | | | | | |
| N 020800 | 004 | 7905352 | Apr 12, 2027 | DP | | |
| <u>EPINEPHRINE - AUVI-Q</u> | | | | | | |
| N 201739 | 001 | 7731686 | Jun 01, 2026 | DP | | |
| | | 7731690 | Jan 15, 2025 | DP | | |
| | | 7749194 | Oct 30, 2028 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>EPINEPHRINE - AUVI-Q</u> | | | | | | |
| N 201739 001 | 7918823 | Nov 23, 2024 | DP | | | |
| | 7947017 | Mar 12, 2028 | DP | | | |
| | 8016788 | Mar 21, 2025 | DP | | | |
| | 8021344 | Nov 02, 2029 | DP | | | |
| | 8206360 | Feb 27, 2027 | DP | | | |
| | 8226610 | Apr 10, 2029 | DP | | | |
| | 8231573 | Nov 25, 2028 | DP | | | |
| | 8313466 | Nov 23, 2024 | DP | | | |
| | 8361029 | Nov 23, 2024 | DP | | | |
| | 8425462 | Nov 23, 2024 | DP | | | |
| | 8608698 | Nov 23, 2024 | DP | | | |
| | 8920377 | Nov 23, 2024 | DP | | | |
| | 8926594 | Mar 31, 2026 | DP | | | |
| | 9056170 | Nov 23, 2024 | DP | | | |
| | 9149579 | Jul 19, 2025 | | U-1758 | | |
| | 9238108 | Feb 20, 2027 | DP | | | |
| | 9259539 | Feb 01, 2026 | DP | | | |
| | 9278182 | Feb 01, 2026 | DP | | | |
| | 9724471 | May 23, 2027 | DP | U-2092 | | |
| | 9737669 | Nov 23, 2024 | DP | | | |
| <u>EPINEPHRINE - AUVI-Q</u> | | | | | | |
| N 201739 002 | 7731686 | Jun 01, 2026 | DP | | | |
| | 7731690 | Jan 15, 2025 | DP | | | |
| | 7749194 | Oct 30, 2028 | DP | | | |
| | 7918823 | Nov 23, 2024 | DP | | | |
| | 7947017 | Mar 12, 2028 | DP | | | |
| | 8016788 | Mar 21, 2025 | DP | | | |
| | 8021344 | Nov 02, 2029 | DP | | | |
| | 8206360 | Feb 27, 2027 | DP | | | |
| | 8226610 | Apr 10, 2029 | DP | | | |
| | 8231573 | Nov 25, 2028 | DP | | | |
| | 8313466 | Nov 23, 2024 | DP | | | |
| | 8361029 | Nov 23, 2024 | DP | | | |
| | 8425462 | Nov 23, 2024 | DP | | | |
| | 8608698 | Nov 23, 2024 | DP | | | |
| | 8920377 | Nov 23, 2024 | DP | | | |
| | 8926594 | Mar 31, 2026 | DP | | | |
| | 9056170 | Nov 23, 2024 | DP | | | |
| | 9149579 | Jul 19, 2025 | | U-1758 | | |
| | 9238108 | Feb 20, 2027 | DP | | | |
| | 9259539 | Feb 01, 2026 | DP | | | |
| | 9278182 | Feb 01, 2026 | DP | | | |
| | 9724471 | May 23, 2027 | DP | U-2092 | | |
| | 9737669 | Nov 23, 2024 | DP | | | |
| <u>EPINEPHRINE - AUVI-Q</u> | | | | | | |
| N 201739 003 | 7731686 | Jun 01, 2026 | DP | | | |
| | 7731690 | Jan 15, 2025 | DP | | | |
| | 7749194 | Oct 30, 2028 | DP | | | |
| | 7918823 | Nov 23, 2024 | DP | | | |
| | 7947017 | Mar 12, 2028 | DP | | | |
| | 8016788 | Mar 21, 2025 | DP | | | |
| | 8021344 | Nov 02, 2029 | DP | | | |
| | 8206360 | Feb 27, 2027 | DP | | | |
| | 8226610 | Apr 10, 2029 | DP | | | |
| | 8231573 | Nov 25, 2028 | DP | | | |
| | 8313466 | Nov 23, 2024 | DP | | | |
| | 8361029 | Nov 23, 2024 | DP | | | |
| | 8425462 | Nov 23, 2024 | DP | | | |
| | 8608698 | Nov 23, 2024 | DP | | | |
| | 8920377 | Nov 23, 2024 | DP | | | |
| | 8926594 | Mar 31, 2026 | DP | | | |
| | 9056170 | Nov 23, 2024 | DP | | | |
| | 9149579 | Jul 19, 2025 | | U-1758 | | |
| | 9238108 | Feb 20, 2027 | DP | | | |
| | 9259539 | Feb 01, 2026 | DP | | | |
| | 9278182 | Feb 01, 2026 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>EPINEPHRINE - AUVI-Q</u> | | | | | | |
| N 201739 | 003 | 9724471 | May 23, 2027 | DP U-2092 | | |
| | | 9737669 | Nov 23, 2024 | DP | | |
| | | 9833573 | Nov 23, 2024 | U-2172 | | |
| <u>EPINEPHRINE - ADRENALIN</u> | | | | | | |
| N 204200 | 001 | 9119876 | Mar 13, 2035 | DP | | |
| | | 9295657 | Mar 13, 2035 | U-1829 | | |
| <u>EPINEPHRINE - ADRENALIN</u> | | | | | | |
| N 204640 | 001 | 10130592 | Mar 13, 2035 | DP | | |
| | | 9119876 | Mar 13, 2035 | DP | | |
| | | 9295657 | Mar 13, 2035 | U-1829 | | |
| <u>EPINEPHRINE - EPINEPHRINE</u> | | | | | | |
| N 205029 | 001 | 10004700 | Aug 14, 2034 | DP U-2325 | | |
| | | 10039728 | Aug 14, 2034 | U-1828 | | |
| | | 9283197 | Aug 15, 2034 | DP U-1828 | | |
| | | 9283197 | Aug 15, 2034 | DP U-1829 | | |
| | | 9283197 | Aug 15, 2034 | DP U-1830 | | |
| <u>EPINEPHRINE - PRIMATENE MIST</u> | | | | | | |
| N 205920 | 001 | 8367734 | Jan 26, 2026 | DP | NP | Nov 07, 2021 |
| <u>EPINEPHRINE; LIDOCAINE HYDROCHLORIDE - LIDOSITE TOPICAL SYSTEM KIT</u> | | | | | | |
| N 021504 | 001 | 6629968 | Jun 30, 2020 | DS DP | | |
| | | 6635045 | Jun 29, 2021 | DS DP | | |
| <u>EPLERENONE - INSPRA</u> | | | | | | |
| N 021437 | 001 | 6410054 | Dec 08, 2019 | U-3 | | |
| | | 6410054 | Dec 08, 2019 | U-537 | | |
| | | 6410524 | Nov 05, 2019 | U-467 | | |
| | | 6495165 | Dec 08, 2019 | U-3 | | |
| | | 6495165 | Dec 08, 2019 | U-537 | | |
| | | 6534093 | Dec 08, 2019 | U-3 | | |
| | | 6534093 | Dec 08, 2019 | U-537 | | |
| | | 6558707 | Dec 08, 2019 | DP U-537 | | |
| | | 6747020 | Nov 05, 2019 | U-587 | | |
| | | 7157101 | Dec 08, 2019 | DP U-664 | | |
| <u>EPLERENONE - INSPRA</u> | | | | | | |
| N 021437 | 002 | 6410054 | Dec 08, 2019 | U-3 | | |
| | | 6410054 | Dec 08, 2019 | U-537 | | |
| | | 6410524 | Nov 05, 2019 | U-467 | | |
| | | 6495165 | Dec 08, 2019 | U-3 | | |
| | | 6495165 | Dec 08, 2019 | U-537 | | |
| | | 6534093 | Dec 08, 2019 | U-3 | | |
| | | 6534093 | Dec 08, 2019 | U-537 | | |
| | | 6558707 | Dec 08, 2019 | DP U-537 | | |
| | | 6747020 | Nov 05, 2019 | U-587 | | |
| | | 7157101 | Dec 08, 2019 | DP U-664 | | |
| <u>EPLERENONE - INSPRA</u> | | | | | | |
| N 021437 | 003 | 6410054 | Dec 08, 2019 | U-3 | | |
| | | 6410054 | Dec 08, 2019 | U-537 | | |
| | | 6410524 | Nov 05, 2019 | U-467 | | |
| | | 6495165 | Dec 08, 2019 | U-3 | | |
| | | 6495165 | Dec 08, 2019 | U-537 | | |
| | | 6534093 | Dec 08, 2019 | U-3 | | |
| | | 6534093 | Dec 08, 2019 | U-537 | | |
| | | 6558707 | Dec 08, 2019 | DP U-537 | | |
| | | 6747020 | Nov 05, 2019 | U-587 | | |
| | | 7157101 | Dec 08, 2019 | DP U-664 | | |
| <u>EPOPROSTENOL SODIUM - VELETRI</u> | | | | | | |
| N 022260 | 001 | 8318802 | Mar 15, 2027 | DP | | |
| | | 8598227 | Feb 02, 2027 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>EPROSTENOL SODIUM - VELETRI</u> | | | | | | |
| N 022260 002 | 8318802 | Mar 15, 2027 | DP | | | |
| | 8598227 | Feb 02, 2027 | | | | |
| <u>ERAVACYCLINE DIHYDROCHLORIDE - XERAVA</u> | | | | | | |
| N 211109 001 | | | | | NCE GAIN | Aug 27, 2023 Aug 27, 2028 |
| <u>ERIBULIN MESYLATE - HALAVEN</u> | | | | | | |
| N 201532 001 | 6214865 | Jul 20, 2023 | DS | | I-721 | Jan 28, 2019 |
| | 6469182 | Jun 16, 2019 | | U-1096 | ODE-107 | Jan 28, 2023 |
| | 6469182 | Jun 16, 2019 | | U-1812 | | |
| | 7470720 | Jun 16, 2019 | DP | | | |
| | 8097648 | Jan 22, 2021 | | U-1096 | | |
| | RE46965 | Jan 08, 2027 | DP | | | |
| <u>ERLOTINIB HYDROCHLORIDE - TARCEVA</u> | | | | | | |
| N 021743 001 | 5747498*PED | May 08, 2019 | | | D-164 | May 20, 2019 |
| | 6900221 | Nov 09, 2020 | DS DP | U-1046 | M-181 | Jun 01, 2019 |
| | 6900221 | Nov 09, 2020 | DS DP | U-1403 | M-190 | Oct 18, 2019 |
| | 6900221 | Nov 09, 2020 | DS DP | U-659 | | |
| | 6900221 | Nov 09, 2020 | DS DP | U-875 | | |
| | 6900221*PED | May 09, 2021 | | | | |
| | 7087613 | Nov 09, 2020 | | U-1045 | | |
| | 7087613 | Nov 09, 2020 | | U-1403 | | |
| | 7087613 | Nov 09, 2020 | | U-659 | | |
| | 7087613*PED | May 09, 2021 | | | | |
| | RE41065*PED | May 08, 2019 | | | | |
| <u>ERLOTINIB HYDROCHLORIDE - TARCEVA</u> | | | | | | |
| N 021743 002 | 5747498*PED | May 08, 2019 | | | D-164 | May 20, 2019 |
| | 6900221 | Nov 09, 2020 | DS DP | U-1046 | M-181 | Jun 01, 2019 |
| | 6900221 | Nov 09, 2020 | DS DP | U-1403 | M-190 | Oct 18, 2019 |
| | 6900221 | Nov 09, 2020 | DS DP | U-659 | | |
| | 6900221 | Nov 09, 2020 | DS DP | U-875 | | |
| | 6900221*PED | May 09, 2021 | | | | |
| | 7087613 | Nov 09, 2020 | | U-1045 | | |
| | 7087613 | Nov 09, 2020 | | U-1403 | | |
| | 7087613 | Nov 09, 2020 | | U-659 | | |
| | 7087613*PED | May 09, 2021 | | | | |
| | RE41065*PED | May 08, 2019 | | | | |
| <u>ERLOTINIB HYDROCHLORIDE - TARCEVA</u> | | | | | | |
| N 021743 003 | 5747498*PED | May 08, 2019 | | | D-164 | May 20, 2019 |
| | 6900221 | Nov 09, 2020 | DS DP | U-1046 | M-181 | Jun 01, 2019 |
| | 6900221 | Nov 09, 2020 | DS DP | U-1403 | M-190 | Oct 18, 2019 |
| | 6900221 | Nov 09, 2020 | DS DP | U-659 | | |
| | 6900221 | Nov 09, 2020 | DS DP | U-875 | | |
| | 6900221*PED | May 09, 2021 | | | | |
| | 7087613 | Nov 09, 2020 | | U-1045 | | |
| | 7087613 | Nov 09, 2020 | | U-1403 | | |
| | 7087613 | Nov 09, 2020 | | U-659 | | |
| | 7087613*PED | May 09, 2021 | | | | |
| | RE41065*PED | May 08, 2019 | | | | |
| <u>ERTUGLIFLOZIN - STEGLATRO</u> | | | | | | |
| N 209803 001 | 8080580 | Jul 13, 2030 | DS DP | U-2214 | NCE | Dec 19, 2022 |
| <u>ERTUGLIFLOZIN - STEGLATRO</u> | | | | | | |
| N 209803 002 | 8080580 | Jul 13, 2030 | DS DP | U-2214 | NCE | Dec 19, 2022 |
| <u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUROMET</u> | | | | | | |
| N 209806 001 | 8080580 | Jul 13, 2030 | DS DP | U-2214 | NCE | Dec 19, 2022 |
| | 9308204 | Oct 21, 2030 | DP | | | |
| | 9439902 | Oct 21, 2030 | | U-2214 | | |
| <u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUROMET</u> | | | | | | |
| N 209806 002 | 8080580 | Jul 13, 2030 | DS DP | U-2214 | NCE | Dec 19, 2022 |
| | 9308204 | Oct 21, 2030 | DP | | | |
| | 9439902 | Oct 21, 2030 | | U-2214 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUROMET</u> | | | | | | |
| N 209806 002 | 8080580 | Jul 13, 2030 | DS DP U-2214 | | NCE | Dec 19, 2022 |
| | 9308204 | Oct 21, 2030 | DP | | | |
| | 9439902 | Oct 21, 2030 | U-2214 | | | |
| <u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUROMET</u> | | | | | | |
| N 209806 003 | 8080580 | Jul 13, 2030 | DS DP U-2214 | | NCE | Dec 19, 2022 |
| | 9308204 | Oct 21, 2030 | DP | | | |
| | 9439902 | Oct 21, 2030 | U-2214 | | | |
| <u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUROMET</u> | | | | | | |
| N 209806 004 | 8080580 | Jul 13, 2030 | DS DP U-2214 | | NCE | Dec 19, 2022 |
| | 9308204 | Oct 21, 2030 | DP | | | |
| | 9439902 | Oct 21, 2030 | U-2214 | | | |
| <u>ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE - STEGLUJAN</u> | | | | | | |
| N 209805 001 | 6699871 | Jul 26, 2022 | DS DP U-2214 | | NCE | Dec 19, 2022 |
| | 6890898 | Feb 02, 2019 | U-2215 | | | |
| | 7078381 | Feb 02, 2019 | U-2216 | | | |
| | 7326708 | Nov 24, 2026 | DS DP U-2214 | | | |
| | 7459428 | Feb 02, 2019 | U-2215 | | | |
| | 8080580 | Jul 13, 2030 | DS DP U-2214 | | | |
| | 9308204 | Oct 21, 2030 | DP | | | |
| | 9439901 | Oct 21, 2030 | U-2214 | | | |
| <u>ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE - STEGLUJAN</u> | | | | | | |
| N 209805 002 | 6699871 | Jul 26, 2022 | DS DP U-2214 | | NCE | Dec 19, 2022 |
| | 6890898 | Feb 02, 2019 | U-2215 | | | |
| | 7078381 | Feb 02, 2019 | U-2216 | | | |
| | 7326708 | Nov 24, 2026 | DS DP U-2214 | | | |
| | 7459428 | Feb 02, 2019 | U-2215 | | | |
| | 8080580 | Jul 13, 2030 | DS DP U-2214 | | | |
| | 9308204 | Oct 21, 2030 | DP | | | |
| | 9439901 | Oct 21, 2030 | U-2214 | | | |
| <u>ESCITALOPRAM OXALATE - LEXAPRO</u> | | | | | | |
| N 021323 001 | 6916941 | Aug 12, 2022 | DS DP | | | |
| | 7420069 | Aug 12, 2022 | DP | | | |
| <u>ESCITALOPRAM OXALATE - LEXAPRO</u> | | | | | | |
| N 021323 002 | 6916941 | Aug 12, 2022 | DS DP | | | |
| | 7420069 | Aug 12, 2022 | DP | | | |
| <u>ESCITALOPRAM OXALATE - LEXAPRO</u> | | | | | | |
| N 021323 003 | 6916941 | Aug 12, 2022 | DS DP | | | |
| | 7420069 | Aug 12, 2022 | DP | | | |
| <u>ESLICARBAZEPINE ACETATE - APTIOM</u> | | | | | | |
| N 022416 001 | 5753646 | Jun 27, 2021 | DS DP U-2041 | | | |
| | 8372431 | Apr 17, 2030 | DP | | | |
| | 9206135 | Apr 21, 2026 | DS | | | |
| | 9566244 | Oct 23, 2028 | DP | | | |
| | 9643929 | Apr 21, 2026 | DP | | | |
| | 9750747 | Aug 24, 2032 | U-2041 | | | |
| | 9750747 | Aug 24, 2032 | U-2121 | | | |
| | 9763954 | Sep 13, 2028 | U-2123 | | | |
| <u>ESLICARBAZEPINE ACETATE - APTIOM</u> | | | | | | |
| N 022416 002 | 5753646 | Jun 27, 2021 | DS DP U-2041 | | | |
| | 8372431 | Apr 17, 2030 | DP | | | |
| | 9206135 | Apr 21, 2026 | DS | | | |
| | 9566244 | Oct 23, 2028 | DP | | | |
| | 9643929 | Apr 21, 2026 | DP | | | |
| | 9750747 | Aug 24, 2032 | U-2041 | | | |
| | 9750747 | Aug 24, 2032 | U-2121 | | | |
| | 9763954 | Sep 13, 2028 | U-2123 | | | |
| <u>ESLICARBAZEPINE ACETATE - APTIOM</u> | | | | | | |
| N 022416 003 | 5753646 | Jun 27, 2021 | DS DP U-2041 | | | |
| | 8372431 | Apr 17, 2030 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ESLICARBAZEPINE ACETATE - APTIOM</u> | | | | | | |
| N 022416 003 | 9206135 | Apr 21, 2026 | DS | | | |
| | 9566244 | Oct 23, 2028 | DP | | | |
| | 9643929 | Apr 21, 2026 | DP | | | |
| | 9750747 | Aug 24, 2032 | | U-2041 | | |
| | 9750747 | Aug 24, 2032 | | U-2121 | | |
| | 9763954 | Sep 13, 2028 | | U-2123 | | |
| <u>ESLICARBAZEPINE ACETATE - APTIOM</u> | | | | | | |
| N 022416 004 | 5753646 | Jun 27, 2021 | DS DP | U-2041 | | |
| | 8372431 | Apr 17, 2030 | DP | | | |
| | 9206135 | Apr 21, 2026 | DS | | | |
| | 9566244 | Oct 23, 2028 | DP | | | |
| | 9643929 | Apr 21, 2026 | DP | | | |
| | 9750747 | Aug 24, 2032 | | U-2041 | | |
| | 9750747 | Aug 24, 2032 | | U-2121 | | |
| | 9763954 | Sep 13, 2028 | | U-2123 | | |
| <u>ESMOLOL HYDROCHLORIDE - BREVIBLOC IN PLASTIC CONTAINER</u> | | | | | | |
| N 019386 004 | 6310094 | Jan 12, 2021 | | | | |
| | 6528540 | Jan 12, 2021 | | | | |
| <u>ESMOLOL HYDROCHLORIDE - BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER</u> | | | | | | |
| N 019386 005 | 6310094 | Jan 12, 2021 | | | | |
| | 6528540 | Jan 12, 2021 | | | | |
| <u>ESMOLOL HYDROCHLORIDE - BREVIBLOC</u> | | | | | | |
| N 019386 006 | 6310094 | Jan 12, 2021 | | | | |
| | 6528540 | Jan 12, 2021 | | | | |
| <u>ESMOLOL HYDROCHLORIDE - BREVIBLOC</u> | | | | | | |
| N 019386 007 | 6310094 | Jan 12, 2021 | | | | |
| | 6528540 | Jan 12, 2021 | | | | |
| <u>ESMOLOL HYDROCHLORIDE - ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER</u> | | | | | | |
| N 205703 001 | 6310094 | Jan 12, 2021 | DP | | | |
| | 6528540 | Jan 12, 2021 | DP | | | |
| | 8829054 | Mar 15, 2033 | DP | | | |
| | 8835505 | Mar 15, 2033 | DP | | | |
| <u>ESMOLOL HYDROCHLORIDE - ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER</u> | | | | | | |
| N 205703 002 | 6310094 | Jan 12, 2021 | DP | | | |
| | 6528540 | Jan 12, 2021 | DP | | | |
| | 8829054 | Mar 15, 2033 | DP | | | |
| | 8835505 | Mar 15, 2033 | DP | | | |
| <u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u> | | | | | | |
| N 021153 001 | 6428810 | Nov 03, 2019 | DP | U-469 | | |
| | 6428810 | Nov 03, 2019 | DP | U-729 | | |
| | 6428810 | Nov 03, 2019 | DP | U-770 | | |
| <u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u> | | | | | | |
| N 021153 002 | 6428810 | Nov 03, 2019 | DP | U-469 | | |
| | 6428810 | Nov 03, 2019 | DP | U-729 | | |
| | 6428810 | Nov 03, 2019 | DP | U-770 | | |
| <u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u> | | | | | | |
| N 021957 001 | 6428810 | Nov 03, 2019 | DP | U-1207 | | |
| | 6428810 | Nov 03, 2019 | DP | U-729 | | |
| | 6428810 | Nov 03, 2019 | DP | U-773 | | |
| <u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u> | | | | | | |
| N 021957 002 | 6428810 | Nov 03, 2019 | DP | U-1207 | | |
| | 6428810 | Nov 03, 2019 | DP | U-729 | | |
| | 6428810 | Nov 03, 2019 | DP | U-773 | | |
| <u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u> | | | | | | |
| N 021957 003 | 6428810 | Nov 03, 2019 | DP | U-1207 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u> | | | | | | |
| N 021957 | 004 6428810 | Nov 03, 2019 | DP U-1207 | | | |
| <u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u> | | | | | | |
| N 022101 | 001 6428810 | Nov 03, 2019 | DP U-858 | | | |
| <u>ESOMEPRAZOLE MAGNESIUM - NEXIUM 24HR</u> | | | | | | |
| N 204655 | 001 6428810 | Nov 03, 2019 | DP U-1509 | | | |
| | 6428810 | Nov 03, 2019 | DP U-1874 | | | |
| | 6428810*PED | May 03, 2020 | | | | |
| <u>ESOMEPRAZOLE MAGNESIUM - NEXIUM 24HR</u> | | | | | | |
| N 207920 | 001 6428810 | Nov 03, 2019 | DP U-1785 | | | |
| | 6428810*PED | May 03, 2020 | | | | |
| <u>ESOMEPRAZOLE MAGNESIUM; NAPROXEN - VIMOVO</u> | | | | | | |
| N 022511 | 001 6926907 | Feb 28, 2023 | DP U-1052 | | NPP | Jul 06, 2020 |
| | 8557285 | May 31, 2022 | DP | | | |
| | 8852636 | May 31, 2022 | DP U-1052 | | | |
| | 8858996 | May 31, 2022 | DP U-1052 | | | |
| | 8945621 | Oct 17, 2031 | U-1661 | | | |
| | 9161920 | May 31, 2022 | U-1760 | | | |
| | 9198888 | May 31, 2022 | U-1781 | | | |
| | 9220698 | Mar 10, 2031 | U-1781 | | | |
| | 9345695 | May 31, 2022 | DP | | | |
| | 9393208 | Sep 03, 2029 | U-1781 | | | |
| | 9707181 | May 31, 2022 | DP | | | |
| <u>ESOMEPRAZOLE MAGNESIUM; NAPROXEN - VIMOVO</u> | | | | | | |
| N 022511 | 002 6926907 | Feb 28, 2023 | DP U-1052 | | NPP | Jul 06, 2020 |
| | 8557285 | May 31, 2022 | DP | | | |
| | 8852636 | May 31, 2022 | DP U-1052 | | | |
| | 8858996 | May 31, 2022 | DP U-1052 | | | |
| | 8945621 | Oct 17, 2031 | U-1661 | | | |
| | 9161920 | May 31, 2022 | U-1760 | | | |
| | 9198888 | May 31, 2022 | U-1781 | | | |
| | 9345695 | May 31, 2022 | DP | | | |
| | 9393208 | Sep 03, 2029 | U-1781 | | | |
| | 9707181 | May 31, 2022 | DP | | | |
| <u>ESTRADIOL - VAGIFEM</u> | | | | | | |
| N 020908 | 002 7018992 | Sep 17, 2022 | U-1023 | | | |
| <u>ESTRADIOL - ELESTRIN</u> | | | | | | |
| N 021813 | 001 7198801 | Jun 25, 2022 | DP | | | |
| | 7470433 | Aug 03, 2021 | DP | | | |
| <u>ESTRADIOL - EVAMIST</u> | | | | | | |
| N 022014 | 001 6978945 | Jul 31, 2022 | DP | | | |
| <u>ESTRADIOL - MINIVELLE</u> | | | | | | |
| N 203752 | 001 6841716 | Apr 27, 2020 | DP | | | |
| | 8231906 | Jul 04, 2030 | DS DP | | | |
| | 9730900 | Jul 10, 2028 | | | U-2086 | |
| | 9833419 | Jul 10, 2028 | DP | | | |
| <u>ESTRADIOL - MINIVELLE</u> | | | | | | |
| N 203752 | 002 6841716 | Apr 27, 2020 | DP | | | |
| | 8231906 | Jul 04, 2030 | DS DP | | | |
| | 9730900 | Jul 10, 2028 | | | U-2086 | |
| | 9833419 | Jul 10, 2028 | DP | | | |
| <u>ESTRADIOL - MINIVELLE</u> | | | | | | |
| N 203752 | 003 6841716 | Apr 27, 2020 | DP | | | |
| | 8231906 | Jul 04, 2030 | DS DP | | | |
| | 9730900 | Jul 10, 2028 | | | U-2086 | |
| | 9833419 | Jul 10, 2028 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ESTRADIOL - MINIVELLE</u> | | | | | | |
| N 203752 | 004 | 6841716 | Apr 27, 2020 | DP | | |
| | | 8231906 | Jul 04, 2030 | DS DP | | |
| | | 9730900 | Jul 10, 2028 | | U-2086 | |
| | | 9833419 | Jul 10, 2028 | DP | | |
| <u>ESTRADIOL - MINIVELLE</u> | | | | | | |
| N 203752 | 005 | 6841716 | Apr 27, 2020 | DP | | |
| | | 8231906 | Jul 04, 2030 | DS DP | | |
| | | 9724310 | Jul 10, 2028 | DS DP | | |
| | | 9730900 | Jul 10, 2028 | DP | U-2086 | |
| | | 9833419 | Jul 10, 2028 | DP | | |
| <u>ESTRADIOL - IMVEXXY</u> | | | | | | |
| N 208564 | 001 | 9180091 | Dec 20, 2033 | DP U-2316 | NP | May 29, 2021 |
| | | 9180091 | Dec 20, 2033 | DP U-2317 | | |
| | | 9289382 | Nov 21, 2032 | DP | | |
| <u>ESTRADIOL - IMVEXXY</u> | | | | | | |
| N 208564 | 002 | 9180091 | Dec 20, 2033 | DP U-2316 | NP | May 29, 2021 |
| | | 9180091 | Dec 20, 2033 | DP U-2317 | | |
| | | 9289382 | Nov 21, 2032 | DP | | |
| <u>ESTRADIOL ACETATE - FEMTRACE</u> | | | | | | |
| N 021633 | 001 | 6962908 | Dec 21, 2021 | DP | | |
| | | 7572779 | Oct 02, 2025 | | U-904 | |
| | | 7799771 | Dec 21, 2021 | DP | | |
| <u>ESTRADIOL ACETATE - FEMTRACE</u> | | | | | | |
| N 021633 | 002 | 6962908 | Dec 21, 2021 | DP | | |
| | | 7572779 | Oct 02, 2025 | | U-904 | |
| | | 7799771 | Dec 21, 2021 | DP | | |
| <u>ESTRADIOL ACETATE - FEMTRACE</u> | | | | | | |
| N 021633 | 003 | 6962908 | Dec 21, 2021 | DP | | |
| | | 7572779 | Oct 02, 2025 | | U-904 | |
| | | 7799771 | Dec 21, 2021 | DP | | |
| <u>ESTRADIOL; ESTRADIOL; NORGESTIMATE - PREFEST</u> | | | | | | |
| N 021040 | 001 | 6747019 | Mar 20, 2020 | | U-311 | |
| | | 7320970 | Mar 30, 2020 | DP | U-844 | |
| <u>ESTRADIOL; PROGESTERONE - BIJUVA</u> | | | | | | |
| N 210132 | 001 | 10052386 | Nov 21, 2032 | DP | | NP Oct 28, 2021 |
| | | 8633178 | Nov 21, 2032 | DP | | |
| | | 8846648 | Nov 21, 2032 | | U-2439 | |
| | | 8846649 | Nov 21, 2032 | DP | U-2439 | |
| | | 8987237 | Nov 21, 2032 | DP | | |
| | | 8993548 | Nov 21, 2032 | DP | | |
| | | 8993549 | Nov 21, 2032 | DP | | |
| | | 9006222 | Nov 21, 2032 | DP | U-2439 | |
| | | 9114145 | Nov 21, 2032 | | U-2439 | |
| | | 9114146 | Nov 21, 2032 | DP | U-2439 | |
| | | 9301920 | Nov 21, 2032 | DP | U-2439 | |
| <u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVA</u> | | | | | | |
| N 021443 | 001 | 6660726 | Mar 08, 2021 | DS DP | U-904 | |
| | | 6660726 | Mar 08, 2021 | DS DP | U-905 | |
| | | 6855703 | Feb 12, 2021 | DS DP | U-904 | |
| | | 6855703 | Feb 12, 2021 | DS DP | U-905 | |
| <u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVA</u> | | | | | | |
| N 021443 | 002 | 6660726 | Mar 08, 2021 | DS DP | U-904 | |
| | | 6660726 | Mar 08, 2021 | DS DP | U-905 | |
| | | 6855703 | Feb 12, 2021 | DS DP | U-904 | |
| | | 6855703 | Feb 12, 2021 | DS DP | U-905 | |
| <u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVA</u> | | | | | | |
| N 021443 | 003 | 6660726 | Mar 08, 2021 | DS DP | U-904 | |
| | | 6660726 | Mar 08, 2021 | DS DP | U-905 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUWIA</u> | | | | | | |
| N 021443 | 003 | 6855703 | Feb 12, 2021 | DS DP U-904 | | |
| | | 6855703 | Feb 12, 2021 | DS DP U-905 | | |
| <u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUWIA</u> | | | | | | |
| N 021443 | 004 | 6660726 | Mar 08, 2021 | DS DP U-904 | | |
| | | 6660726 | Mar 08, 2021 | DS DP U-905 | | |
| | | 6855703 | Feb 12, 2021 | DS DP U-904 | | |
| | | 6855703 | Feb 12, 2021 | DS DP U-905 | | |
| <u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUWIA</u> | | | | | | |
| N 021443 | 005 | 6660726 | Mar 08, 2021 | DS DP U-904 | | |
| | | 6660726 | Mar 08, 2021 | DS DP U-905 | | |
| | | 6855703 | Feb 12, 2021 | DS DP U-904 | | |
| | | 6855703 | Feb 12, 2021 | DS DP U-905 | | |
| <u>ETELCALCETIDE - PARSABIV</u> | | | | | | |
| N 208325 | 001 | 8377880 | Jul 29, 2030 | DS DP | NCE | Feb 07, 2022 |
| | | 8999932 | Jul 29, 2030 | DS DP U-2014 | | |
| | | 9278995 | Jul 29, 2030 | DS | | |
| | | 9701712 | Jul 29, 2030 | DS DP U-2014 | | |
| | | 9820938 | Jun 27, 2034 | DP | | |
| <u>ETELCALCETIDE - PARSABIV</u> | | | | | | |
| N 208325 | 002 | 8377880 | Jul 29, 2030 | DS DP | NCE | Feb 07, 2022 |
| | | 8999932 | Jul 29, 2030 | DS DP U-2014 | | |
| | | 9278995 | Jul 29, 2030 | DS | | |
| | | 9701712 | Jul 29, 2030 | DS DP U-2014 | | |
| | | 9820938 | Jun 27, 2034 | DP | | |
| <u>ETELCALCETIDE - PARSABIV</u> | | | | | | |
| N 208325 | 003 | 8377880 | Jul 29, 2030 | DS DP | NCE | Feb 07, 2022 |
| | | 8999932 | Jul 29, 2030 | DS DP U-2014 | | |
| | | 9278995 | Jul 29, 2030 | DS | | |
| | | 9701712 | Jul 29, 2030 | DS DP U-2014 | | |
| | | 9820938 | Jun 27, 2034 | DP | | |
| <u>ETEPLIRSEN - EXONDYS 51</u> | | | | | | |
| N 206488 | 001 | 8486907 | Jun 28, 2025 | U-1904 | Y | NCE |
| | | 9018368 | Jun 28, 2025 | DS DP | | ODE-122 |
| | | 9243245 | Oct 27, 2028 | DS U-2097 | | Sep 19, 2021 |
| | | 9243245 | Oct 27, 2028 | DS U-2098 | | Sep 19, 2023 |
| | | 9416361 | May 04, 2021 | DS | | |
| | | 9506058 | Mar 14, 2034 | U-1918 | | |
| | | 9506058 | Mar 14, 2034 | U-1919 | | |
| <u>ETEPLIRSEN - EXONDYS 51</u> | | | | | | |
| N 206488 | 002 | 8486907 | Jun 28, 2025 | U-1904 | Y | NCE |
| | | 9018368 | Jun 28, 2025 | DS DP | | ODE-122 |
| | | 9243245 | Oct 27, 2028 | DS U-2097 | | Sep 19, 2021 |
| | | 9243245 | Oct 27, 2028 | DS U-2098 | | Sep 19, 2023 |
| | | 9416361 | May 04, 2021 | DS | | |
| | | 9506058 | Mar 14, 2034 | U-1918 | | |
| | | 9506058 | Mar 14, 2034 | U-1919 | | |
| <u>ETHINYL ESTRADIOL; ETHINYL ESTRADIOL; ETHINYL ESTRADIOL; NORGESTIMATE; NORGESTIMATE; NORGESTIMATE - ORTHO TRI-CYCLEN LO</u> | | | | | | |
| N 021241 | 001 | 6214815 | Jun 09, 2019 | U-112 | | |
| <u>ETHINYL ESTRADIOL; ETHINYL ESTRADIOL; LEVONORGESTREL - SEASONIQUE</u> | | | | | | |
| N 021840 | 001 | 7320969 | Jan 30, 2024 | U-828 | | |
| | | 7615545 | Jun 15, 2023 | U-1 | | |
| | | 7855190 | Dec 05, 2028 | U-1 | | |
| | | 7858605 | Jun 23, 2023 | DP | | |
| <u>ETHINYL ESTRADIOL; LEVONORGESTREL - PREVEN EMERGENCY CONTRACEPTIVE KIT</u> | | | | | | |
| N 020946 | 001 | 6156742 | Dec 05, 2020 | U-374 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ETHINYL ESTRADIOL; LEVONORGESTREL - LOSEASONIQUE</u> | | | | | | |
| N 022262 | 001 | 7615545 | Jun 15, 2023 | U-1 | | |
| | | 7855190 | Dec 05, 2028 | U-1 | | |
| | | 7858605 | Jun 23, 2023 | DP | | |
| <u>ETHINYL ESTRADIOL; LEVONORGESTREL - QUARTETTE</u> | | | | | | |
| N 204061 | 001 | 8415332 | Mar 11, 2029 | DP | | |
| | | 8450299 | Oct 07, 2025 | U-1 | | |
| <u>ETHINYL ESTRADIOL; LEVONORGESTREL - BALCOLTRA</u> | | | | | | |
| N 208612 | 001 | 6716814 | Aug 16, 2021 | DS DP | | |
| <u>ETHINYL ESTRADIOL; NORETHINDRONE - FEMCON FE</u> | | | | | | |
| N 021490 | 001 | 6667050 | Apr 06, 2019 | DP U-1 | | |
| <u>ETHINYL ESTRADIOL; NORETHINDRONE - NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u> | | | | | | |
| N 022573 | 001 | 6667050 | Apr 06, 2019 | DP U-828 | | |
| <u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LO LOESTRIN FE</u> | | | | | | |
| N 022501 | 001 | 7704984 | Feb 02, 2029 | U-1090 | | |
| <u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - MINASTRIN 24 FE</u> | | | | | | |
| N 203667 | 001 | 6667050 | Apr 06, 2019 | DP U-1 | | |
| <u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - TAYTULLA</u> | | | | | | |
| N 204426 | 001 | 6652880 | Mar 29, 2020 | DP | | |
| <u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LO MINASTRIN FE</u> | | | | | | |
| N 204654 | 001 | 6667050 | Apr 06, 2019 | DP U-1 | | |
| | | 7704984 | Feb 02, 2029 | U-1 | | |
| <u>ETHINYL ESTRADIOL; SEGESTERONE ACETATE - ANNOVERA</u> | | | | | | |
| N 209627 | 001 | | | | NCE | Aug 10, 2023 |
| <u>ETHIODIZED OIL - LIPIODOL</u> | | | | | | |
| N 009190 | 001 | | | | ODE-64 | Apr 04, 2021 |
| <u>ETONOGESTREL - IMPLANON</u> | | | | | | |
| N 021529 | 001 | 9757552 | Jul 28, 2030 | DP U-1 | | |
| <u>ETONOGESTREL - NEXPLANON</u> | | | | | | |
| N 021529 | 002 | 8722037 | Sep 28, 2027 | DP | | |
| | | 8888745 | Aug 28, 2026 | DP | | |
| | | 9757552 | Jul 28, 2030 | DP U-1 | | |
| <u>ETRAVIRINE - INTELENCE</u> | | | | | | |
| N 022187 | 001 | 6878717 | Nov 05, 2019 | U-1016 | NPP | Jul 16, 2021 |
| | | 6878717 | Nov 05, 2019 | U-1237 | PED | Jan 16, 2022 |
| | | 6878717 | Nov 05, 2019 | U-2354 | | |
| | | 6878717 | Nov 05, 2019 | U-256 | | |
| | | 6878717*PED | May 05, 2020 | | | |
| | | 7037917 | Dec 13, 2020 | DS DP U-1016 | | |
| | | 7037917 | Dec 13, 2020 | DS DP U-1237 | | |
| | | 7037917 | Dec 13, 2020 | DS DP U-2354 | | |
| | | 7037917 | Dec 13, 2020 | DS DP U-256 | | |
| | | 7037917*PED | Jun 13, 2021 | | | |
| | | 7887845 | Mar 25, 2019 | DP | | |
| | | 7887845*PED | Sep 25, 2019 | | | |
| | | 8003789 | Nov 01, 2019 | DS DP | | |
| | | 8003789*PED | May 01, 2020 | | | |
| <u>ETRAVIRINE - INTELENCE</u> | | | | | | |
| N 022187 | 002 | 6878717 | Nov 05, 2019 | U-1016 | NPP | Jul 16, 2021 |
| | | 6878717 | Nov 05, 2019 | U-1237 | PED | Jan 16, 2022 |
| | | 6878717 | Nov 05, 2019 | U-2354 | | |
| | | 6878717 | Nov 05, 2019 | U-256 | | |
| | | 6878717*PED | May 05, 2020 | | | |
| | | 7037917 | Dec 13, 2020 | DS DP U-1016 | | |
| | | 7037917 | Dec 13, 2020 | DS DP U-1237 | | |
| | | 7037917 | Dec 13, 2020 | DS DP U-2354 | | |
| | | 7037917 | Dec 13, 2020 | DS DP U-256 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ETRAVIRINE - INTELENCE</u> | | | | | | |
| N 022187 | 002 | 7037917*PED | Jun 13, 2021 | | | |
| | | 7887845 | Mar 25, 2019 | DP | | |
| | | 7887845*PED | Sep 25, 2019 | | | |
| | | 8003789 | Nov 01, 2019 | DS DP | | |
| | | 8003789*PED | May 01, 2020 | | | |
| <u>ETRAVIRINE - INTELENCE</u> | | | | | | |
| N 022187 | 003 | 6878717 | Nov 05, 2019 | U-1016 | NPP | Jul 16, 2021 |
| | | 6878717 | Nov 05, 2019 | U-1237 | PED | Jan 16, 2022 |
| | | 6878717 | Nov 05, 2019 | U-2354 | | |
| | | 6878717 | Nov 05, 2019 | U-256 | | |
| | | 6878717*PED | May 05, 2020 | | | |
| | | 7037917 | Dec 13, 2020 | DS DP U-1237 | | |
| | | 7037917 | Dec 13, 2020 | DS DP U-2354 | | |
| | | 7037917*PED | Jun 13, 2021 | | | |
| | | 7887845 | Mar 25, 2019 | DP | | |
| | | 7887845*PED | Sep 25, 2019 | | | |
| | | 8003789 | Nov 01, 2019 | DS DP | | |
| | | 8003789*PED | May 01, 2020 | | | |
| <u>EVEROLIMUS - ZORTRESS</u> | | | | | | |
| N 021560 | 001 | 5665772 | Sep 09, 2019 | DS DP U-1049 | | |
| | | 5665772 | Sep 09, 2019 | DS DP U-1365 | | |
| | | 5665772*PED | Mar 09, 2020 | | | |
| <u>EVEROLIMUS - ZORTRESS</u> | | | | | | |
| N 021560 | 002 | 5665772 | Sep 09, 2019 | DS DP U-1049 | | |
| | | 5665772 | Sep 09, 2019 | DS DP U-1365 | | |
| | | 5665772*PED | Mar 09, 2020 | | | |
| <u>EVEROLIMUS - ZORTRESS</u> | | | | | | |
| N 021560 | 003 | 5665772 | Sep 09, 2019 | DS DP U-1049 | | |
| | | 5665772 | Sep 09, 2019 | DS DP U-1365 | | |
| | | 5665772*PED | Mar 09, 2020 | | | |
| <u>EVEROLIMUS - AFINITOR</u> | | | | | | |
| N 022334 | 001 | 5665772 | Sep 09, 2019 | DS DP | I-724 | Feb 26, 2019 |
| | | 7297703 | Dec 06, 2019 | DP | ODE-108 | Feb 26, 2023 |
| | | 7741338 | Dec 06, 2019 | DP | ODE-24 | Apr 26, 2019 |
| | | 8410131 | Nov 01, 2025 | U-1368 | | |
| | | 8410131*PED | May 01, 2026 | | | |
| | | 8436010 | Feb 22, 2022 | U-1396 | | |
| | | 8436010*PED | Aug 22, 2022 | | | |
| | | 8778962 | Feb 18, 2022 | U-1541 | | |
| | | 8778962*PED | Aug 18, 2022 | | | |
| | | 9006224 | Jul 01, 2028 | U-1681 | | |
| <u>EVEROLIMUS - AFINITOR</u> | | | | | | |
| N 022334 | 002 | 5665772 | Sep 09, 2019 | DS DP | I-724 | Feb 26, 2019 |
| | | 7297703 | Dec 06, 2019 | DP | ODE-108 | Feb 26, 2023 |
| | | 7741338 | Dec 06, 2019 | DP | ODE-24 | Apr 26, 2019 |
| | | 8410131 | Nov 01, 2025 | U-1368 | | |
| | | 8410131*PED | May 01, 2026 | | | |
| | | 8436010 | Feb 22, 2022 | U-1396 | | |
| | | 8436010*PED | Aug 22, 2022 | | | |
| | | 8778962 | Feb 18, 2022 | U-1541 | | |
| | | 8778962*PED | Aug 18, 2022 | | | |
| | | 9006224 | Jul 01, 2028 | U-1681 | | |
| <u>EVEROLIMUS - AFINITOR</u> | | | | | | |
| N 022334 | 003 | 5665772 | Sep 09, 2019 | DS DP | I-724 | Feb 26, 2019 |
| | | 7297703 | Dec 06, 2019 | DP | ODE-108 | Feb 26, 2023 |
| | | 7741338 | Dec 06, 2019 | DP | ODE-24 | Apr 26, 2019 |
| | | 8410131 | Nov 01, 2025 | U-1368 | | |
| | | 8410131*PED | May 01, 2026 | | | |
| | | 8436010 | Feb 22, 2022 | U-1396 | | |
| | | 8436010*PED | Aug 22, 2022 | | | |
| | | 8778962 | Feb 18, 2022 | U-1541 | | |
| | | 8778962*PED | Aug 18, 2022 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>EVEROLIMUS - AFINITOR</u> | | | | | | |
| N 022334 | 003 | 9006224 | | | | |
| | | Jul 01, 2028 | | U-1681 | | |
| <u>EVEROLIMUS - AFINITOR</u> | | | | | | |
| N 022334 | 004 | 5665772 | Sep 09, 2019 | DS DP | I-724 | Feb 26, 2019 |
| | | 7297703 | Dec 06, 2019 | DP | ODE-108 | Feb 26, 2023 |
| | | 7741338 | Dec 06, 2019 | DP | ODE-24 | Apr 26, 2019 |
| | | 8410131 | Nov 01, 2025 | | | |
| | | 8410131*PED | May 01, 2026 | | | |
| | | 8436010 | Feb 22, 2022 | | U-1368 | |
| | | 8436010*PED | Aug 22, 2022 | | U-1396 | |
| | | 8778962 | Feb 18, 2022 | | U-1541 | |
| | | 8778962*PED | Aug 18, 2022 | | | |
| | | 9006224 | Jul 01, 2028 | | U-1681 | |
| <u>EVEROLIMUS - AFINITOR DISPERZ</u> | | | | | | |
| N 203985 | 001 | 5665772 | Sep 09, 2019 | DS DP | I-773 | Apr 10, 2021 |
| | | 7297703 | Dec 06, 2019 | DP | ODE-169 | Apr 10, 2025 |
| | | 8617598 | Sep 27, 2022 | DP | | |
| | | 8617598*PED | Mar 27, 2023 | | | |
| | | 8778962 | Feb 18, 2022 | | U-1541 | |
| | | 8778962 | Feb 18, 2022 | | U-2280 | |
| | | 8778962*PED | Aug 18, 2022 | | | |
| <u>EVEROLIMUS - AFINITOR DISPERZ</u> | | | | | | |
| N 203985 | 002 | 5665772 | Sep 09, 2019 | DS DP | I-773 | Apr 10, 2021 |
| | | 7297703 | Dec 06, 2019 | DP | ODE-169 | Apr 10, 2025 |
| | | 8617598 | Sep 27, 2022 | DP | | |
| | | 8617598*PED | Mar 27, 2023 | | | |
| | | 8778962 | Feb 18, 2022 | | U-1541 | |
| | | 8778962 | Feb 18, 2022 | | U-2280 | |
| | | 8778962*PED | Aug 18, 2022 | | | |
| <u>EVEROLIMUS - AFINITOR DISPERZ</u> | | | | | | |
| N 203985 | 003 | 5665772 | Sep 09, 2019 | DS DP | I-773 | Apr 10, 2021 |
| | | 7297703 | Dec 06, 2019 | DP | ODE-169 | Apr 10, 2025 |
| | | 8617598 | Sep 27, 2022 | DP | | |
| | | 8617598*PED | Mar 27, 2023 | | | |
| | | 8778962 | Feb 18, 2022 | | U-1541 | |
| | | 8778962 | Feb 18, 2022 | | U-2280 | |
| | | 8778962*PED | Aug 18, 2022 | | | |
| <u>EXENATIDE - BYDUREON BCISE</u> | | | | | | |
| N 209210 | 001 | 6479065 | Aug 10, 2020 | DP | NP | Oct 20, 2020 |
| | | 6667061 | May 25, 2020 | DP | | |
| | | 6824822 | Oct 09, 2022 | DP | | |
| | | 6872700 | Jan 14, 2020 | | U-654 | |
| | | 7223440 | Aug 31, 2021 | DP | | |
| | | 7456254 | Jun 30, 2025 | DP | U-1223 | |
| | | 7563871 | Apr 15, 2024 | DP | | |
| | | 7612176 | Apr 13, 2025 | DP | U-1223 | |
| | | 8329648 | Aug 18, 2026 | | U-1313 | |
| | | 8329648 | Aug 18, 2026 | | U-2154 | |
| | | 8329648 | Aug 18, 2026 | | U-2155 | |
| | | 8329648 | Aug 18, 2026 | | U-2156 | |
| | | 8431685 | Apr 13, 2025 | DP | U-412 | |
| | | 8461105 | Apr 13, 2025 | DP | U-412 | |
| | | 8895033 | Oct 04, 2030 | DP | U-1313 | |
| | | 8895033 | Oct 04, 2030 | DP | U-2157 | |
| | | 8895033 | Oct 04, 2030 | DP | U-2158 | |
| | | 8906851 | Aug 18, 2026 | | U-1313 | |
| | | 9238076 | Apr 15, 2024 | DP | U-412 | |
| | | 9884092 | Aug 18, 2026 | | U-1313 | |
| | | 9884092 | Aug 18, 2026 | | U-2154 | |
| | | 9884092 | Aug 18, 2026 | | U-2155 | |
| | | 9884092 | Aug 18, 2026 | | U-2156 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>EXENATIDE SYNTHETIC - BYETTA</u> | | | | | | |
| N 021773 001 | 6872700 | Jan 14, 2020 | U-654 | | | |
| | 6902744 | Jan 14, 2020 | DP | | | |
| <u>EXENATIDE SYNTHETIC - BYETTA</u> | | | | | | |
| N 021773 002 | 6872700 | Jan 14, 2020 | U-654 | | | |
| | 6902744 | Jan 14, 2020 | DP | | | |
| <u>EXENATIDE SYNTHETIC - BYDUREON</u> | | | | | | |
| N 022200 001 | 6414126 | Oct 04, 2020 | DS DP U-2139 | | M-212 | Oct 20, 2020 |
| | 6414126 | Oct 04, 2020 | DS DP U-493 | | M-224 | Apr 02, 2021 |
| | 6479065 | Aug 10, 2020 | DP | | | |
| | 6495164 | May 25, 2020 | DP | | | |
| | 6515117 | Oct 04, 2020 | DS DP U-2139 | | | |
| | 6515117 | Oct 04, 2020 | DS DP U-493 | | | |
| | 6667061 | May 25, 2020 | DP | | | |
| | 6824822 | Oct 09, 2022 | DP | | | |
| | 6872700 | Jan 14, 2020 | U-2288 | | | |
| | 6872700 | Jan 14, 2020 | U-654 | | | |
| | 6936590 | Oct 04, 2020 | U-493 | | | |
| | 7223440 | Aug 31, 2021 | DP | | | |
| | 7456254 | Jun 30, 2025 | DP U-1223 | | | |
| | 7563871 | Apr 15, 2024 | DP | | | |
| | 7612176 | Apr 13, 2025 | DP U-1223 | | | |
| | 7851502 | Aug 19, 2028 | DP | | | |
| | 7919598 | Dec 16, 2029 | DS | | | |
| | 8221786 | Mar 21, 2028 | DP | | | |
| | 8329648 | Aug 18, 2026 | U-1313 | | | |
| | 8361972 | Mar 21, 2028 | U-2139 | | | |
| | 8361972 | Mar 21, 2028 | U-493 | | | |
| | 8431685 | Apr 13, 2025 | DP U-412 | | | |
| | 8461105 | Apr 13, 2025 | DP U-412 | | | |
| | 8501698 | Jun 20, 2027 | DP U-493 | | | |
| | 8685934 | May 26, 2030 | U-1522 | | | |
| | 8716251 | Mar 21, 2028 | DP | | | |
| | 8906851 | Aug 18, 2026 | U-1313 | | | |
| | 9198925 | Oct 04, 2020 | U-2139 | | | |
| | 9198925 | Oct 04, 2020 | U-493 | | | |
| | 9238076 | Apr 15, 2024 | DP U-412 | | | |
| | 9884092 | Aug 18, 2026 | U-1313 | | | |
| | 9884092 | Aug 18, 2026 | U-2154 | | | |
| | 9884092 | Aug 18, 2026 | U-2155 | | | |
| | 9884092 | Aug 18, 2026 | U-2156 | | | |
| <u>EXENATIDE SYNTHETIC - BYDUREON PEN</u> | | | | | | |
| N 022200 002 | 6414126 | Oct 04, 2020 | DS DP U-2139 | | M-224 | Apr 02, 2021 |
| | 6414126 | Oct 04, 2020 | DS DP U-493 | | | |
| | 6479065 | Aug 10, 2020 | DP | | | |
| | 6495164 | May 25, 2020 | DP | | | |
| | 6515117 | Oct 04, 2020 | DS DP U-2139 | | | |
| | 6515117 | Oct 04, 2020 | DS DP U-493 | | | |
| | 6667061 | May 25, 2020 | DP | | | |
| | 6824822 | Oct 09, 2022 | DP | | | |
| | 6872700 | Jan 14, 2020 | U-2288 | | | |
| | 6872700 | Jan 14, 2020 | U-654 | | | |
| | 6936590 | Oct 04, 2020 | U-493 | | | |
| | 7223440 | Aug 31, 2021 | DP | | | |
| | 7456254 | Jun 30, 2025 | DP U-1223 | | | |
| | 7563871 | Apr 15, 2024 | DP | | | |
| | 7612176 | Apr 13, 2025 | DP U-1223 | | | |
| | 7851502 | Aug 19, 2028 | DP | | | |
| | 7919598 | Dec 16, 2029 | DS | | | |
| | 8216180 | Jan 12, 2028 | DP | | | |
| | 8221786 | Mar 21, 2028 | DP | | | |
| | 8329648 | Aug 18, 2026 | U-1313 | | | |
| | 8361972 | Mar 21, 2028 | U-2139 | | | |
| | 8361972 | Mar 21, 2028 | U-493 | | | |
| | 8431685 | Apr 13, 2025 | DP U-412 | | | |
| | 8439864 | Mar 25, 2028 | DP | | | |
| | 8461105 | Apr 13, 2025 | DP U-412 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>EXENATIDE SYNTHETIC - BYDUREON PEN</u> | | | | | | |
| N 022200 002 | 8501698 | Jun 20, 2027 | DP U-493 | | | |
| | 8685934 | May 26, 2030 | U-1522 | | | |
| | 8690837 | May 19, 2029 | DP | | | |
| | 8716251 | Mar 21, 2028 | DP | | | |
| | 8721615 | Jan 18, 2030 | DP | | | |
| | 8758292 | Nov 12, 2027 | DP | | | |
| | 8827963 | Feb 04, 2029 | DP | | | |
| | 8906851 | Aug 18, 2026 | U-1313 | | | |
| | 8998876 | Jan 07, 2030 | DP | | | |
| | 9198925 | Oct 04, 2020 | U-2139 | | | |
| | 9198925 | Oct 04, 2020 | U-493 | | | |
| | 9238076 | Apr 15, 2024 | DP U-412 | | | |
| | 9320853 | Mar 25, 2028 | DP | | | |
| | 9884092 | Aug 18, 2026 | U-1313 | | | |
| | 9884092 | Aug 18, 2026 | U-2154 | | | |
| | 9884092 | Aug 18, 2026 | U-2155 | | | |
| | 9884092 | Aug 18, 2026 | U-2156 | | | |
| <u>EZETIMIBE - ZETIA</u> | | | | | | |
| N 021445 001 | 7030106 | Jan 25, 2022 | DP | | | |
| | 7612058 | Oct 30, 2025 | U-1027 | | | |
| | 7612058 | Oct 30, 2025 | U-1173 | | | |
| | 7612058*PED | Apr 30, 2026 | | | | |
| <u>FAMOTIDINE - PEPCID AC</u> | | | | | | |
| N 020801 002 | 6814978 | Aug 26, 2021 | DP | | | |
| <u>FAMOTIDINE; IBUPROFEN - DUEXIS</u> | | | | | | |
| N 022519 001 | 8067033 | Jul 18, 2026 | DP | | | |
| | 8067451 | Jul 18, 2026 | DP U-1196 | | | |
| | 8309127 | Jul 18, 2026 | DP | | | |
| | 8318202 | Jul 18, 2026 | DP | | | |
| | 8449910 | Jul 18, 2026 | DP | | | |
| | 8501228 | Jul 18, 2026 | U-1196 | | | |
| <u>FEBUXOSTAT - ULORIC</u> | | | | | | |
| N 021856 001 | 5614520 | Mar 25, 2019 | DS DP U-954 | | M-205 | Aug 15, 2020 |
| | 6225474 | Jun 18, 2019 | DS | | | |
| | 7361676 | Mar 08, 2024 | DP | | | |
| | 8372872 | Sep 08, 2031 | U-1346 | | | |
| | 9107912 | Sep 08, 2031 | U-1346 | | | |
| <u>FEBUXOSTAT - ULORIC</u> | | | | | | |
| N 021856 002 | 5614520 | Mar 25, 2019 | DS DP U-954 | | M-205 | Aug 15, 2020 |
| | 6225474 | Jun 18, 2019 | DS | | | |
| | 7361676 | Mar 08, 2024 | DP | | | |
| | 8372872 | Sep 08, 2031 | U-1346 | | | |
| | 9107912 | Sep 08, 2031 | U-1346 | | | |
| <u>FENOFIBRATE - TRIGLIDE</u> | | | | | | |
| N 021350 001 | 6696084 | Sep 11, 2021 | DS DP U-680 | | | |
| <u>FENOFIBRATE - TRIGLIDE</u> | | | | | | |
| N 021350 002 | 6696084 | Sep 11, 2021 | DS DP U-680 | | | |
| <u>FENOFIBRATE - TRICOR</u> | | | | | | |
| N 021656 001 | 6375986 | Sep 21, 2020 | DP U-615 | | | |
| | 7276249 | Feb 21, 2023 | DP | | | |
| | 7320802 | Feb 21, 2023 | U-847 | | | |
| <u>FENOFIBRATE - TRICOR</u> | | | | | | |
| N 021656 002 | 6375986 | Sep 21, 2020 | DP U-615 | | | |
| | 7276249 | Feb 21, 2023 | DP | | | |
| | 7320802 | Feb 21, 2023 | U-847 | | | |
| <u>FENOFIBRATE - ANTARA (MICRONIZED)</u> | | | | | | |
| N 021695 001 | 7101574 | Aug 20, 2020 | DS DP | | | |
| | 7863331 | Aug 08, 2020 | U-1106 | | | |
| | 7863331 | Aug 08, 2020 | U-1107 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>FENOFIBRATE - ANTARA (MICRONIZED)</u> | | | | | | |
| N 021695 | 001 | 7101574 | Aug 20, 2020 | DS DP | | |
| | | 7863331 | Aug 08, 2020 | | U-1106 | |
| | | 7863331 | Aug 08, 2020 | | U-1107 | |
| <u>FENOFIBRATE - ANTARA (MICRONIZED)</u> | | | | | | |
| N 021695 | 003 | 7101574 | Aug 20, 2020 | DS DP | | |
| | | 7863331 | Aug 08, 2020 | | U-1106 | |
| | | 7863331 | Aug 08, 2020 | | U-1107 | |
| <u>FENOFIBRATE - ANTARA (MICRONIZED)</u> | | | | | | |
| N 021695 | 004 | 8026281 | Apr 22, 2025 | | U-1447 | |
| | | 8026281 | Apr 22, 2025 | | U-1448 | |
| <u>FENOFIBRATE - ANTARA (MICRONIZED)</u> | | | | | | |
| N 021695 | 005 | 8026281 | Apr 22, 2025 | | U-1447 | |
| | | 8026281 | Apr 22, 2025 | | U-1448 | |
| | | 9314447 | May 31, 2033 | DP | U-1447 | |
| | | 9314447 | May 31, 2033 | DP | U-1448 | |
| <u>FENOFIBRATE - FENOGLIDE</u> | | | | | | |
| N 022118 | 001 | 7658944 | Dec 09, 2024 | DP | | |
| | | 8124125 | Oct 01, 2024 | DP | U-1234 | |
| | | 8481078 | Oct 01, 2024 | DP | U-1416 | |
| | | 9173847 | Oct 01, 2024 | DP | | |
| <u>FENOFIBRATE - FENOGLIDE</u> | | | | | | |
| N 022118 | 002 | 7658944 | Dec 09, 2024 | DP | | |
| | | 8124125 | Oct 01, 2024 | DP | U-1234 | |
| | | 8481078 | Oct 01, 2024 | DP | U-1416 | |
| | | 9173847 | Oct 01, 2024 | DP | | |
| <u>FENOFIBRIC ACID - FIBRICOR</u> | | | | | | |
| N 022418 | 001 | 7569612 | Aug 20, 2027 | | U-1000 | |
| | | 7741373 | Aug 20, 2027 | | U-1059 | |
| | | 7741374 | Aug 20, 2027 | | U-1060 | |
| | | 7741374 | Aug 20, 2027 | | U-1061 | |
| | | 7915247 | Aug 20, 2027 | | U-1000 | |
| | | 7915247 | Aug 20, 2027 | | U-1059 | |
| | | 7915247 | Aug 20, 2027 | | U-1061 | |
| <u>FENOFIBRIC ACID - FIBRICOR</u> | | | | | | |
| N 022418 | 002 | 7569612 | Aug 20, 2027 | | U-1000 | |
| | | 7741373 | Aug 20, 2027 | | U-1059 | |
| | | 7741374 | Aug 20, 2027 | | U-1060 | |
| | | 7741374 | Aug 20, 2027 | | U-1061 | |
| | | 7915247 | Aug 20, 2027 | | U-1000 | |
| | | 7915247 | Aug 20, 2027 | | U-1059 | |
| | | 7915247 | Aug 20, 2027 | | U-1061 | |
| <u>FENTANYL - SUBSYS</u> | | | | | | |
| N 202788 | 001 | 10016403 | Jan 25, 2027 | DP | | |
| | | 8486972 | Apr 27, 2030 | DP | | |
| | | 8486973 | Apr 27, 2030 | | U-55 | |
| | | 8835459 | Jan 25, 2027 | DP | | |
| | | 8835460 | Jan 25, 2027 | DP | U-55 | |
| | | 9241935 | Jan 25, 2027 | DP | | |
| | | 9289387 | Jan 25, 2027 | DP | U-55 | |
| | | 9642797 | Jan 25, 2027 | DP | U-55 | |
| | | 9642844 | Jan 25, 2027 | DP | | |
| <u>FENTANYL - SUBSYS</u> | | | | | | |
| N 202788 | 002 | 10016403 | Jan 25, 2027 | DP | | |
| | | 8486972 | Apr 27, 2030 | DP | | |
| | | 8486973 | Apr 27, 2030 | | U-55 | |
| | | 8835460 | Jan 25, 2027 | DP | U-55 | |
| | | 9241935 | Jan 25, 2027 | DP | | |
| | | 9289387 | Jan 25, 2027 | DP | U-55 | |
| | | 9642797 | Jan 25, 2027 | DP | U-55 | |
| | | 9642844 | Jan 25, 2027 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>FENTANYL - SUBSYS</u> | | | | | | |
| N 202788 002 | 10016403 | Jan 25, 2027 | DP | | | |
| | 8486972 | Apr 27, 2030 | DP | | | |
| | 8486973 | Apr 27, 2030 | | U-55 | | |
| | 8835460 | Jan 25, 2027 | DP | U-55 | | |
| | 9241935 | Jan 25, 2027 | DP | | | |
| | 9289387 | Jan 25, 2027 | DP | U-55 | | |
| | 9642797 | Jan 25, 2027 | DP | U-55 | | |
| | 9642844 | Jan 25, 2027 | DP | | | |
| <u>FENTANYL - SUBSYS</u> | | | | | | |
| N 202788 003 | 10016403 | Jan 25, 2027 | DP | | | |
| | 8486972 | Apr 27, 2030 | DP | | | |
| | 8486973 | Apr 27, 2030 | | U-55 | | |
| | 8835459 | Jan 25, 2027 | DP | | | |
| | 8835460 | Jan 25, 2027 | DP | U-55 | | |
| | 9241935 | Jan 25, 2027 | DP | | | |
| | 9289387 | Jan 25, 2027 | DP | U-55 | | |
| | 9642797 | Jan 25, 2027 | DP | U-55 | | |
| | 9642844 | Jan 25, 2027 | DP | | | |
| <u>FENTANYL - SUBSYS</u> | | | | | | |
| N 202788 004 | 10016403 | Jan 25, 2027 | DP | | | |
| | 8486972 | Apr 27, 2030 | DP | | | |
| | 8486973 | Apr 27, 2030 | | U-55 | | |
| | 8835459 | Jan 25, 2027 | DP | | | |
| | 8835460 | Jan 25, 2027 | DP | U-55 | | |
| | 9241935 | Jan 25, 2027 | DP | | | |
| | 9289387 | Jan 25, 2027 | DP | U-55 | | |
| | 9642797 | Jan 25, 2027 | DP | U-55 | | |
| | 9642844 | Jan 25, 2027 | DP | | | |
| <u>FENTANYL - SUBSYS</u> | | | | | | |
| N 202788 005 | 10016403 | Jan 25, 2027 | DP | | | |
| | 8486972 | Apr 27, 2030 | DP | | | |
| | 8486973 | Apr 27, 2030 | | U-55 | | |
| | 8835460 | Jan 25, 2027 | DP | U-55 | | |
| | 9241935 | Jan 25, 2027 | DP | | | |
| | 9289387 | Jan 25, 2027 | DP | U-55 | | |
| | 9642797 | Jan 25, 2027 | DP | U-55 | | |
| | 9642844 | Jan 25, 2027 | DP | | | |
| <u>FENTANYL - SUBSYS</u> | | | | | | |
| N 202788 006 | 10016403 | Jan 25, 2027 | DP | | | |
| | 8486972 | Apr 27, 2030 | DP | | | |
| | 8486973 | Apr 27, 2030 | | U-55 | | |
| | 8835459 | Jan 25, 2027 | DP | | | |
| | 8835460 | Jan 25, 2027 | DP | U-55 | | |
| | 9241935 | Jan 25, 2027 | DP | | | |
| | 9289387 | Jan 25, 2027 | DP | U-55 | | |
| | 9642797 | Jan 25, 2027 | DP | U-55 | | |
| | 9642844 | Jan 25, 2027 | DP | | | |
| <u>FENTANYL - SUBSYS</u> | | | | | | |
| N 202788 007 | 10016403 | Jan 25, 2027 | DP | | | |
| | 8486972 | Apr 27, 2030 | DP | | | |
| | 8486973 | Apr 27, 2030 | | U-55 | | |
| | 8835459 | Jan 25, 2027 | DP | | | |
| | 8835460 | Jan 25, 2027 | DP | U-55 | | |
| | 9241935 | Jan 25, 2027 | DP | | | |
| | 9289387 | Jan 25, 2027 | DP | U-55 | | |
| | 9642797 | Jan 25, 2027 | DP | U-55 | | |
| | 9642844 | Jan 25, 2027 | DP | | | |
| <u>FENTANYL CITRATE - FENTORA</u> | | | | | | |
| N 021947 001 | 6200604 | Mar 26, 2019 | | U-767 | | |
| | 6974590 | Mar 26, 2019 | | U-767 | | |
| | 7862832 | Jun 15, 2028 | DP | | | |
| | 7862833 | Jun 15, 2028 | DP | | | |
| | 8092832 | Dec 30, 2024 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>FENTANYL CITRATE - FENTORA</u> | | | | | | |
| N 021947 001 | 8728441 | Mar 26, 2019 | | | | |
| | 8753611 | Mar 26, 2019 | | | | |
| | 8765100 | Mar 26, 2019 | DP | | | |
| <u>FENTANYL CITRATE - FENTORA</u> | | | | | | |
| N 021947 002 | 6200604 | Mar 26, 2019 | | | | |
| | 6974590 | Mar 26, 2019 | | | | |
| | 7862832 | Jun 15, 2028 | DP | | | |
| | 7862833 | Jun 15, 2028 | DP | | | |
| | 8092832 | Dec 30, 2024 | DP | | | |
| | 8119158 | Dec 30, 2024 | DP | | | |
| | 8728441 | Mar 26, 2019 | | | | |
| | 8753611 | Mar 26, 2019 | | | | |
| | 8765100 | Mar 26, 2019 | DP | | | |
| <u>FENTANYL CITRATE - FENTORA</u> | | | | | | |
| N 021947 003 | 6200604 | Mar 26, 2019 | | | | |
| | 6974590 | Mar 26, 2019 | | | | |
| | 7862832 | Jun 15, 2028 | DP | | | |
| | 7862833 | Jun 15, 2028 | DP | | | |
| | 8092832 | Dec 30, 2024 | DP | | | |
| | 8119158 | Dec 30, 2024 | DP | | | |
| | 8728441 | Mar 26, 2019 | | | | |
| | 8753611 | Mar 26, 2019 | | | | |
| | 8765100 | Mar 26, 2019 | DP | | | |
| <u>FENTANYL CITRATE - FENTORA</u> | | | | | | |
| N 021947 004 | 6200604 | Mar 26, 2019 | | | | |
| | 6974590 | Mar 26, 2019 | | | | |
| | 7862832 | Jun 15, 2028 | DP | | | |
| | 7862833 | Jun 15, 2028 | DP | | | |
| | 8092832 | Dec 30, 2024 | DP | | | |
| | 8119158 | Dec 30, 2024 | DP | | | |
| | 8728441 | Mar 26, 2019 | | | | |
| | 8753611 | Mar 26, 2019 | | | | |
| | 8765100 | Mar 26, 2019 | DP | | | |
| <u>FENTANYL CITRATE - FENTORA</u> | | | | | | |
| N 021947 005 | 6200604 | Mar 26, 2019 | | | | |
| | 6974590 | Mar 26, 2019 | | | | |
| | 7862832 | Jun 15, 2028 | DP | | | |
| | 7862833 | Jun 15, 2028 | DP | | | |
| | 8092832 | Dec 30, 2024 | DP | | | |
| | 8119158 | Dec 30, 2024 | DP | | | |
| | 8728441 | Mar 26, 2019 | | | | |
| | 8753611 | Mar 26, 2019 | | | | |
| | 8765100 | Mar 26, 2019 | DP | | | |
| <u>FENTANYL CITRATE - FENTORA</u> | | | | | | |
| N 021947 006 | 6200604 | Mar 26, 2019 | | | | |
| | 6974590 | Mar 26, 2019 | | | | |
| <u>FENTANYL CITRATE - ONSOLIS</u> | | | | | | |
| N 022266 001 | 7579019 | Jan 22, 2020 | | | | |
| | 9597288 | Jul 23, 2027 | DP | | | |
| <u>FENTANYL CITRATE - ONSOLIS</u> | | | | | | |
| N 022266 002 | 7579019 | Jan 22, 2020 | | | | |
| | 9597288 | Jul 23, 2027 | DP | | | |
| <u>FENTANYL CITRATE - ONSOLIS</u> | | | | | | |
| N 022266 003 | 7579019 | Jan 22, 2020 | | | | |
| | 9597288 | Jul 23, 2027 | DP | | | |
| <u>FENTANYL CITRATE - ONSOLIS</u> | | | | | | |
| N 022266 004 | 7579019 | Jan 22, 2020 | | | | |
| | 9597288 | Jul 23, 2027 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>FENTANYL CITRATE - ONSOLIS</u> | | | | | | |
| N 022266 | 005 7579019 | Jan 22, 2020 | | | | |
| | 9597288 | Jul 23, 2027 | DP U-767 | | | |
| <u>FENTANYL CITRATE - ABSTRAL</u> | | | | | | |
| N 022510 | 001 6759059 | Sep 24, 2019 | | | | |
| | 6761910 | Sep 24, 2019 | DP U-767 | | | |
| | 7910132 | Sep 24, 2019 | DP U-767 | | | |
| <u>FENTANYL CITRATE - ABSTRAL</u> | | | | | | |
| N 022510 | 002 6759059 | Sep 24, 2019 | | | | |
| | 6761910 | Sep 24, 2019 | DP U-767 | | | |
| | 7910132 | Sep 24, 2019 | DP U-767 | | | |
| <u>FENTANYL CITRATE - ABSTRAL</u> | | | | | | |
| N 022510 | 003 6759059 | Sep 24, 2019 | | | | |
| | 6761910 | Sep 24, 2019 | DP U-767 | | | |
| | 7910132 | Sep 24, 2019 | DP U-767 | | | |
| <u>FENTANYL CITRATE - ABSTRAL</u> | | | | | | |
| N 022510 | 004 6759059 | Sep 24, 2019 | | | | |
| | 6761910 | Sep 24, 2019 | DP U-767 | | | |
| | 7910132 | Sep 24, 2019 | DP U-767 | | | |
| <u>FENTANYL CITRATE - ABSTRAL</u> | | | | | | |
| N 022510 | 005 6759059 | Sep 24, 2019 | | | | |
| | 6761910 | Sep 24, 2019 | DP U-767 | | | |
| | 7910132 | Sep 24, 2019 | DP U-767 | | | |
| <u>FENTANYL CITRATE - ABSTRAL</u> | | | | | | |
| N 022510 | 006 6759059 | Sep 24, 2019 | | | | |
| | 6761910 | Sep 24, 2019 | DP U-767 | | | |
| | 7910132 | Sep 24, 2019 | DP U-767 | | | |
| <u>FENTANYL CITRATE - LAZANDA</u> | | | | | | |
| N 022569 | 001 8216604 | Oct 03, 2024 | | | | |
| | 8889176 | Jan 16, 2024 | | | | |
| | 9078814 | Jan 08, 2024 | DP | | | |
| | 9731869 | Jan 26, 2032 | DP | | | |
| | 9814705 | Jan 08, 2024 | DP | | | |
| <u>FENTANYL CITRATE - LAZANDA</u> | | | | | | |
| N 022569 | 002 8216604 | Oct 03, 2024 | | | | |
| | 8889176 | Jan 16, 2024 | | | | |
| | 9078814 | Jan 08, 2024 | DP | | | |
| | 9731869 | Jan 26, 2032 | DP | | | |
| | 9814705 | Jan 08, 2024 | DP | | | |
| <u>FENTANYL CITRATE - LAZANDA</u> | | | | | | |
| N 022569 | 003 9731869 | Jan 26, 2032 | | | | |
| | 9814705 | Jan 08, 2024 | DP | | | |
| <u>FENTANYL HYDROCHLORIDE - IONSYS</u> | | | | | | |
| N 021338 | 001 6181963 | Nov 02, 2019 | | | | |
| | 6195582 | Jan 28, 2019 | DP U-736 | | | |
| | 6881208 | Apr 19, 2022 | | | | |
| | 6975902 | Apr 01, 2024 | DP | | | |
| | 8301238 | Sep 30, 2031 | DP | | | |
| | 8428708 | May 21, 2032 | | | | |
| | 8428709 | Jun 11, 2032 | DP U-736 | | | |
| | 8781571 | Mar 31, 2032 | DP U-736 | | | |
| | 9095706 | Feb 03, 2033 | DP | | | |
| | 9364656 | Sep 30, 2031 | | | | |
| | 9731121 | Oct 17, 2031 | DP | | | |
| <u>FERRIC CARBOXYMALTOSE - INJECTAFER</u> | | | | | | |
| N 203565 | 001 7612109 | Feb 05, 2024 | DS DP | | | |
| | 7754702 | Feb 13, 2027 | DP U-1432 | | | |
| | 8895612 | Jan 08, 2027 | DP U-1620 | | | |
| | 9376505 | Oct 20, 2023 | DS DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>FERRIC CITRATE - AURYXIA</u> | | | | | | |
| N 205874 001 | 5753706 | Feb 03, 2019 | DP U-1577 | | | |
| | 7767851 | Feb 18, 2024 | DS DP | | | |
| | 8093423 | Apr 21, 2026 | U-1577 | | | |
| | 8299298 | Feb 18, 2024 | DP | | | |
| | 8338642 | Feb 18, 2024 | DS DP U-1577 | | | |
| | 8609896 | Feb 18, 2024 | DP | | | |
| | 8754257 | Feb 18, 2024 | DP | | | |
| | 8754258 | Feb 18, 2024 | DP | | | |
| | 8846976 | Feb 18, 2024 | U-1577 | | | |
| | 8901349 | Feb 18, 2024 | U-1577 | | | |
| | 9050316 | Feb 18, 2024 | U-1577 | | | |
| | 9328133 | Feb 18, 2024 | DS DP U-1577 | | | |
| | 9387191 | Jul 21, 2030 | DP | | | |
| | 9757416 | Feb 18, 2024 | DS DP U-1577 | | | |
| <u>FERRIC PYROPHOSPHATE CITRATE - TRIFERIC</u> | | | | | | |
| N 206317 001 | 7816404 | Apr 17, 2029 | DP U-1656 | | | |
| <u>FERUMOXYTOL - FERAHEME</u> | | | | | | |
| N 022180 001 | 6599498 | Jun 30, 2023 | DS DP | | I-767 | Feb 02, 2021 |
| | 7553479 | Mar 08, 2020 | DS DP | | | |
| | 7871597 | Mar 08, 2020 | DS DP | | | |
| | 8501158 | Mar 08, 2020 | U-1422 | | | |
| | 8591864 | Mar 08, 2020 | DP | | | |
| | 8926947 | Mar 08, 2020 | DS DP | | | |
| <u>FESOTERODINE FUMARATE - TOVIAZ</u> | | | | | | |
| N 022030 001 | 6858650 | Jul 03, 2022 | DS U-913 | | | |
| | 7384980 | May 11, 2019 | DS DP U-913 | | | |
| | 7807715 | Jun 07, 2027 | DP U-913 | | | |
| | 7855230 | May 11, 2019 | U-913 | | | |
| | 7985772 | May 11, 2019 | DS DP U-913 | | | |
| | 8088398 | Jun 07, 2027 | DP U-913 | | | |
| | 8338478 | May 11, 2019 | DS DP U-913 | | | |
| | 8501723 | Jun 07, 2027 | DP | | | |
| <u>FESOTERODINE FUMARATE - TOVIAZ</u> | | | | | | |
| N 022030 002 | 6858650 | Jul 03, 2022 | DS U-913 | | | |
| | 7384980 | May 11, 2019 | DS DP U-913 | | | |
| | 7807715 | Jun 07, 2027 | DP U-913 | | | |
| | 7855230 | May 11, 2019 | U-913 | | | |
| | 7985772 | May 11, 2019 | DS DP U-913 | | | |
| | 8088398 | Jun 07, 2027 | DP U-913 | | | |
| | 8338478 | May 11, 2019 | DS DP U-913 | | | |
| | 8501723 | Jun 07, 2027 | DP | | | |
| <u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA ALLERGY</u> | | | | | | |
| N 021909 002 | 6723348 | Nov 26, 2021 | DP U-1466 | | | |
| <u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA HIVES</u> | | | | | | |
| N 021909 003 | 6723348 | Nov 26, 2021 | DP | | | |
| <u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA ALLERGY</u> | | | | | | |
| N 201373 001 | 8933097 | Aug 16, 2032 | DP | | | |
| <u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA HIVES</u> | | | | | | |
| N 201373 002 | 8933097 | Aug 16, 2032 | DP | | | |
| <u>FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION</u> | | | | | | |
| N 021704 002 | 6613357 | Dec 25, 2020 | DP U-1159 | | | |
| <u>FIDAXOMICIN - DIFICID</u> | | | | | | |
| N 201699 001 | 7378508 | Jul 31, 2027 | DS DP | | | |
| | 7863249 | Jul 31, 2027 | DS DP | | | |
| | 7906489 | Mar 04, 2027 | U-319 | | | |
| | 8586551 | Jul 15, 2023 | DS DP | | | |
| | 8859510 | Jul 31, 2027 | U-319 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>FINAFLOXACIN - XTORO</u> | | | | | | |
| N 206307 001 | 8536167 | Aug 08, 2031 | | U-1679 | NCE | Dec 17, 2019 |
| | 9119859 | Jul 02, 2030 | | U-1679 | PED | Jun 17, 2020 |
| | 9504691 | Nov 21, 2033 | DP | U-1679 | | |
| <u>FINGOLIMOD HYDROCHLORIDE - GILENYA</u> | | | | | | |
| N 022527 001 | 5604229 | Feb 18, 2019 | DS | U-1086 | NPP | May 11, 2021 |
| | 5604229*PED | Aug 18, 2019 | | | PED | Nov 11, 2021 |
| | 8324283 | Mar 29, 2026 | DP | | | |
| | 8324283*PED | Sep 29, 2026 | | | | |
| | 9187405 | Jun 25, 2027 | | U-1086 | | |
| | 9187405*PED | Dec 25, 2027 | | | | |
| <u>FINGOLIMOD HYDROCHLORIDE - GILENYA</u> | | | | | | |
| N 022527 002 | 5604229 | Feb 18, 2019 | DS | U-1086 | NS | May 11, 2021 |
| | 5604229*PED | Aug 18, 2019 | | | PED | Nov 11, 2021 |
| | 9592208 | Mar 30, 2032 | DP | U-2315 | | |
| | 9592208*PED | Sep 30, 2032 | | | | |
| <u>FISH OIL TRIGLYCERIDES - OMEGAVEN</u> | | | | | | |
| N 210589 001 | 9566260 | Nov 12, 2023 | DP | U-2366 | NCE | Jul 27, 2023 |
| | 9629821 | Nov 12, 2023 | DP | U-2367 | ODE-202 | Jul 27, 2025 |
| <u>FISH OIL TRIGLYCERIDES - OMEGAVEN</u> | | | | | | |
| N 210589 002 | 9566260 | Nov 12, 2023 | DP | U-2366 | NCE | Jul 27, 2023 |
| | 9629821 | Nov 12, 2023 | DP | U-2367 | ODE-202 | Jul 27, 2025 |
| <u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u> | | | | | | |
| N 207648 001 | | | | | NCE | Jul 13, 2021 |
| <u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u> | | | | | | |
| N 207648 002 | | | | | NCE | Jul 13, 2021 |
| <u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u> | | | | | | |
| N 207648 003 | | | | | NCE | Jul 13, 2021 |
| <u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u> | | | | | | |
| N 207648 004 | | | | | NCE | Jul 13, 2021 |
| <u>FLIBANSERIN - ADDYI</u> | | | | | | |
| N 022526 001 | 7151103 | May 09, 2023 | | U-1734 | NCE | Aug 18, 2020 |
| | 7420057 | Aug 01, 2022 | DS DP | | | |
| | 8227471 | May 09, 2023 | | U-1734 | | |
| | 9468639 | Oct 16, 2022 | | U-1734 | | |
| <u>FLORBETABEN F-18 - NEURACEQ</u> | | | | | | |
| N 204677 001 | 7807135 | Mar 18, 2029 | DS DP | U-1497 | NCE | Mar 21, 2019 |
| <u>FLORBETAPIR F-18 - AMYVID</u> | | | | | | |
| N 202008 001 | 7687052 | Apr 30, 2027 | DS DP | | | |
| | 8506929 | Apr 30, 2027 | DS DP | U-1423 | | |
| <u>FLORBETAPIR F-18 - AMYVID</u> | | | | | | |
| N 202008 002 | 7687052 | Apr 30, 2027 | DS DP | | | |
| | 8506929 | Apr 30, 2027 | DS DP | U-1423 | | |
| <u>FLORBETAPIR F-18 - AMYVID</u> | | | | | | |
| N 202008 003 | 7687052 | Apr 30, 2027 | DS DP | | | |
| | 8506929 | Apr 30, 2027 | DS DP | U-1423 | | |
| <u>FLUCICLOVINE F-18 - AXUMIN</u> | | | | | | |
| N 208054 001 | 10010632 | Nov 28, 2026 | DP | | NCE | May 27, 2021 |
| | 10124079 | Dec 30, 2035 | | U-2450 | | |
| | 5808146 | Nov 09, 2020 | DS | | | |
| | 9387266 | Nov 28, 2026 | | U-1879 | | |
| <u>FLUDARABINE PHOSPHATE - OFORTA</u> | | | | | | |
| N 022273 001 | 7148207 | Dec 20, 2022 | DP | U-944 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>FLUOCINOLONE ACETONIDE - RETISERT</u> | | | | | | |
| N 021737 | 001 | 6217895 | Mar 22, 2019 | DP U-708 | | |
| | | 6548078 | Mar 22, 2019 | DP U-708 | | |
| <u>FLUOCINOLONE ACETONIDE - ILUVIEN</u> | | | | | | |
| N 201923 | 001 | 6217895 | Mar 22, 2019 | DP U-1597 | | |
| | | 6375972 | Apr 26, 2020 | DP U-1597 | | |
| | | 6548078 | Mar 22, 2019 | DP U-1597 | | |
| | | 8252307 | Jun 27, 2019 | DP | | |
| | | 8871241 | Aug 12, 2027 | DP | | |
| <u>FLUOCINOLONE ACETONIDE - YUTIQ</u> | | | | | | |
| N 210331 | 001 | 6217895 | Mar 22, 2019 | DP U-708 | NP | Nov 12, 2021 |
| | | 6375972 | Apr 26, 2020 | DP U-708 | | |
| | | 6548078 | Mar 22, 2019 | DP U-708 | | |
| | | 8252307 | Jun 27, 2019 | DP | | |
| | | 8574613 | Apr 26, 2020 | DP | | |
| | | 8574659 | Apr 26, 2020 | DP | | |
| | | 8871241 | Aug 12, 2027 | DP | | |
| | | 9192579 | Apr 26, 2020 | DP | | |
| | | 9849085 | Apr 26, 2020 | U-708 | | |
| <u>FLUOCINOLONE ACETONIDE; HYDROQUINONE; TRETINOIN - TRI-LUMA</u> | | | | | | |
| N 021112 | 001 | 7915243 | Sep 08, 2023 | DP | | |
| | | 7939516 | Sep 08, 2023 | DP | | |
| | | 8247395 | Oct 25, 2022 | DP | | |
| | | 8653053 | Oct 25, 2022 | DP | | |
| <u>FLUOCINONIDE - VANOS</u> | | | | | | |
| N 021758 | 001 | 6765001 | Dec 21, 2021 | DP | | |
| | | 7220424 | Jan 07, 2023 | U-861 | | |
| | | 7794738 | Sep 11, 2022 | U-1084 | | |
| | | 8232264 | Mar 09, 2023 | DP | | |
| <u>FLUOROURACIL - CARAC</u> | | | | | | |
| N 020985 | 001 | 6670335 | Jun 02, 2021 | DP U-68 | | |
| <u>FLUOROURACIL - TOLAK</u> | | | | | | |
| N 022259 | 001 | 7169401 | Jul 18, 2023 | DP | | |
| <u>FLUTEMETAMOL F-18 - VIZAMYL</u> | | | | | | |
| N 203137 | 001 | 7270800 | Sep 03, 2025 | DS DP U-336 | | |
| | | 7351401 | Jan 24, 2023 | DS DP U-336 | | |
| | | 8236282 | May 21, 2024 | DS DP | | |
| | | 8691185 | Jan 24, 2023 | U-336 | | |
| | | 8916131 | Sep 16, 2028 | DP | | |
| <u>FLUTEMETAMOL F-18 - VIZAMYL</u> | | | | | | |
| N 203137 | 002 | 7270800 | Sep 03, 2025 | DS DP U-336 | | |
| | | 7351401 | Jan 24, 2023 | DS DP U-336 | | |
| | | 8236282 | May 21, 2024 | DS DP | | |
| | | 8691185 | Jan 24, 2023 | U-336 | | |
| | | 8916131 | Sep 16, 2028 | DP | | |
| <u>FLUTICASONE FUROATE - FLONASE SENSIMIST ALLERGY RELIEF</u> | | | | | | |
| N 022051 | 002 | 6858596 | Aug 03, 2021 | DP U-1890 | | |
| | | 7101866 | Aug 03, 2021 | DS DP U-1890 | | |
| | | 7541350 | Aug 03, 2021 | DP U-1890 | | |
| | | 8062264 | Apr 05, 2026 | DP | | |
| | | 8147461 | Oct 15, 2028 | DP | | |
| | | 8347879 | Jul 15, 2028 | DP | | |
| | | 8752543 | Apr 05, 2026 | DP | | |
| | | 9320862 | Nov 06, 2024 | DP | | |
| <u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u> | | | | | | |
| N 205625 | 001 | 7101866 | Aug 03, 2021 | DS DP U-1559 | NPP | May 17, 2021 |
| | | 7629335 | Aug 03, 2021 | DP | | |
| | | 8113199 | Oct 23, 2027 | DP | | |
| | | 8161968 | Feb 05, 2028 | DP | | |
| | | 8201556 | Feb 05, 2029 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------------------------------|--------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u> | | | | | | |
| N 205625 001 | 8534281 | Mar 08, 2030 | DP | | | |
| | 8746242 | Oct 11, 2030 | DP | | | |
| | 9333310 | Oct 02, 2027 | DP | | | |
| <u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u> | | | | | | |
| N 205625 002 | 7101866 | Aug 03, 2021 | DS DP U-1559 | | | |
| | 7629335 | Aug 03, 2021 | DP | | | |
| | 8113199 | Oct 23, 2027 | DP | | | |
| | 8161968 | Feb 05, 2028 | DP | | | |
| | 8201556 | Feb 05, 2029 | DP | | | |
| | 8534281 | Mar 08, 2030 | DP | | | |
| | 8746242 | Oct 11, 2030 | DP | | | |
| | 9333310 | Oct 02, 2027 | DP | | | |
| <u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u> | | | | | | |
| N 205625 003 | 7101866 | Aug 03, 2021 | DS DP U-2349 | | NS | May 17, 2021 |
| | 7629335 | Aug 03, 2021 | DP | | | |
| | 8113199 | Oct 23, 2027 | DP | | | |
| | 8161968 | Feb 05, 2028 | DP | | | |
| | 8201556 | Feb 05, 2029 | DP | | | |
| | 8534281 | Mar 08, 2030 | DP | | | |
| | 8746242 | Oct 11, 2030 | DP | | | |
| | 9333310 | Oct 02, 2027 | DP | | | |
| <u>FLUTICASONE FUROATE; UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - TRELEGY ELLIPTA</u> | | | | | | |
| N 209482 001 | 6537983 | Aug 03, 2021 | DP U-2125 | | I-775 | Apr 24, 2021 |
| | 6759398 | Aug 03, 2021 | DP U-2125 | | | |
| | 6878698 | Aug 03, 2021 | U-2134 | | | |
| | 7101866 | Aug 03, 2021 | DS DP U-2126 | | | |
| | 7439393 | May 21, 2025 | DS DP U-2127 | | | |
| | 7488827 | Dec 18, 2027 | DS DP | | | |
| | 7498440 | Apr 27, 2025 | DS DP | | | |
| | 7629335 | Aug 03, 2021 | DP | | | |
| | 7776895 | Sep 11, 2022 | DP | | | |
| | 8113199 | Oct 23, 2027 | DP | | | |
| | 8161968 | Feb 05, 2028 | DP | | | |
| | 8183257 | Jul 27, 2025 | U-2128 | | | |
| | 8309572 | Apr 27, 2025 | U-2129 | | | |
| | 8511304 | Jun 14, 2027 | DP | | | |
| | 8534281 | Mar 08, 2030 | DP | | | |
| | 8746242 | Oct 11, 2030 | DP | | | |
| | 9333310 | Oct 02, 2027 | DP | | | |
| | 9750726 | Nov 29, 2030 | DP | | | |
| RE44874 | Mar 23, 2023 | DS DP U-2127 | | | | |
| <u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u> | | | | | | |
| N 204275 001 | 6537983 | Aug 03, 2021 | DP U-1401 | | M-202 | May 15, 2020 |
| | 6537983 | Aug 03, 2021 | DP U-1691 | | | |
| | 6759398 | Aug 03, 2021 | DP U-1401 | | | |
| | 6759398 | Aug 03, 2021 | DP U-1691 | | | |
| | 6878698 | Aug 03, 2021 | U-1401 | | | |
| | 7101866 | Aug 03, 2021 | DS DP U-1401 | | | |
| | 7101866 | Aug 03, 2021 | DS DP U-1691 | | | |
| | 7439393 | May 21, 2025 | DS DP U-1401 | | | |
| | 7439393 | May 21, 2025 | DS DP U-1691 | | | |
| | 7439393 | May 21, 2025 | DS DP U-2099 | | | |
| | 7439393 | May 21, 2025 | DS DP U-2100 | | | |
| | 7629335 | Aug 03, 2021 | DP | | | |
| | 7776895 | Sep 11, 2022 | DP | | | |
| | 8113199 | Oct 23, 2027 | DP | | | |
| | 8161968 | Feb 05, 2028 | DP | | | |
| | 8511304 | Jun 14, 2027 | DP U-1424 | | | |
| | 8511304 | Jun 14, 2027 | DP U-1691 | | | |
| | 8534281 | Mar 08, 2030 | DP | | | |
| | 8746242 | Oct 11, 2030 | DP | | | |
| | 9333310 | Oct 02, 2027 | DP | | | |
| | RE44874 | Mar 23, 2023 | DS DP U-1548 | | | |
| RE44874 | Mar 23, 2023 | DS DP U-1691 | | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u> | | | | | | |
| N 204275 | 002 | 6537983 | Aug 03, 2021 | DP U-1691 | M-202 | May 15, 2020 |
| | | 6759398 | Aug 03, 2021 | DP U-1691 | | |
| | | 7101866 | Aug 03, 2021 | DS DP U-1691 | | |
| | | 7439393 | May 21, 2025 | DS DP U-1691 | | |
| | | 7439393 | May 21, 2025 | DS DP U-2099 | | |
| | | 7439393 | May 21, 2025 | DS DP U-2100 | | |
| | | 7629335 | Aug 03, 2021 | DP | | |
| | | 7776895 | Sep 11, 2022 | DP | | |
| | | 8113199 | Oct 23, 2027 | DP | | |
| | | 8161968 | Feb 05, 2028 | DP | | |
| | | 8511304 | Jun 14, 2027 | DP U-1691 | | |
| | | 8534281 | Mar 08, 2030 | DP | | |
| | | 8746242 | Oct 11, 2030 | DP | | |
| | | 9333310 | Oct 02, 2027 | DP | | |
| | | RE44874 | Mar 23, 2023 | DS DP U-1691 | | |
| <u>FLUTICASONE PROPIONATE - CUTIVATE</u> | | | | | | |
| N 021152 | 001 | 7300669 | Oct 20, 2019 | DP U-835 | | |
| <u>FLUTICASONE PROPIONATE - FLOVENT HFA</u> | | | | | | |
| N 021433 | 001 | 7500444 | Feb 26, 2026 | DP | | |
| | | 7500444*PED | Aug 26, 2026 | | | |
| | | 7832351 | Jun 19, 2023 | DP | | |
| | | 9861771 | Oct 11, 2020 | DP | | |
| <u>FLUTICASONE PROPIONATE - FLOVENT HFA</u> | | | | | | |
| N 021433 | 002 | 6743413 | Jun 01, 2021 | U-581 | Y | |
| | | 7500444 | Feb 26, 2026 | DP | | |
| | | 7500444*PED | Aug 26, 2026 | | | |
| | | 7832351 | Jun 19, 2023 | DP | | |
| | | 9861771 | Oct 11, 2020 | DP | | |
| <u>FLUTICASONE PROPIONATE - FLOVENT HFA</u> | | | | | | |
| N 021433 | 003 | 7500444 | Feb 26, 2026 | DP | | |
| | | 7500444*PED | Aug 26, 2026 | | | |
| | | 7832351 | Jun 19, 2023 | DP | | |
| | | 9861771 | Oct 11, 2020 | DP | | |
| <u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u> | | | | | | |
| N 208798 | 001 | 10022510 | May 18, 2031 | DP | NP | Jan 27, 2020 |
| | | 10124131 | May 18, 2031 | DP | | |
| | | 6701917 | Jun 23, 2021 | DP | | |
| | | 6718972 | Jun 23, 2021 | DP | | |
| | | 6748947 | Jun 23, 2021 | DP | | |
| | | 6871646 | Jun 23, 2021 | DP | | |
| | | 7540282 | May 06, 2023 | DP | | |
| | | 8006690 | Jun 23, 2021 | DP | | |
| | | 8651103 | Mar 26, 2028 | DP | | |
| | | 8714149 | Feb 25, 2032 | DP | | |
| | | 8978966 | Jan 13, 2032 | DP | | |
| | | 9216260 | Jun 28, 2031 | DP | | |
| | | 9463288 | May 19, 2025 | DP | | |
| | | 9616024 | Sep 01, 2024 | DP | | |
| | | 9731087 | May 18, 2031 | DP | | |
| <u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u> | | | | | | |
| N 208798 | 002 | 10022510 | May 18, 2031 | DP | NP | Jan 27, 2020 |
| | | 10124131 | May 18, 2031 | DP | | |
| | | 6701917 | Jun 23, 2021 | DP | | |
| | | 6718972 | Jun 23, 2021 | DP | | |
| | | 6748947 | Jun 23, 2021 | DP | | |
| | | 6871646 | Jun 23, 2021 | DP | | |
| | | 7540282 | May 06, 2023 | DP | | |
| | | 8006690 | Jun 23, 2021 | DP | | |
| | | 8651103 | Mar 26, 2028 | DP | | |
| | | 8714149 | Feb 25, 2032 | DP | | |
| | | 8978966 | Jan 13, 2032 | DP | | |
| | | 9216260 | Jun 28, 2031 | DP | | |
| | | 9463288 | May 19, 2025 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u> | | | | | | |
| N 208798 | 002 | 9616024 | Sep 01, 2024 | DP | | |
| | | 9731087 | May 18, 2031 | DP | | |
| <u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u> | | | | | | |
| N 208798 | 003 | 10022510 | May 18, 2031 | DP | NP | Jan 27, 2020 |
| | | 10124131 | May 18, 2031 | DP | | |
| | | 6701917 | Jun 23, 2021 | DP | | |
| | | 6718972 | Jun 23, 2021 | DP | | |
| | | 6748947 | Jun 23, 2021 | DP | | |
| | | 6871646 | Jun 23, 2021 | DP | | |
| | | 7540282 | May 06, 2023 | DP | | |
| | | 8006690 | Jun 23, 2021 | DP | | |
| | | 8651103 | Mar 26, 2028 | DP | | |
| | | 8714149 | Feb 25, 2032 | DP | | |
| | | 8978966 | Jan 13, 2032 | DP | | |
| | | 9216260 | Jun 28, 2031 | DP | | |
| | | 9463288 | May 19, 2025 | DP | | |
| | | 9616024 | Sep 01, 2024 | DP | | |
| | | 9731087 | May 18, 2031 | DP | | |
| <u>FLUTICASONE PROPIONATE - XHANCE</u> | | | | | | |
| N 209022 | 001 | 10076614 | Oct 20, 2034 | DP | NP | Sep 18, 2020 |
| | | 10076615 | Jul 30, 2029 | U-2133 | | |
| | | 10124132 | Mar 06, 2027 | DP U-2133 | | |
| | | 6715485 | Mar 03, 2020 | DP | | |
| | | 7975690 | Dec 29, 2025 | U-2133 | | |
| | | 8327844 | Oct 08, 2023 | U-2133 | | |
| | | 8522778 | May 11, 2022 | DP | | |
| | | 8550073 | Oct 22, 2029 | DP | | |
| | | 8555878 | Mar 20, 2020 | DP | | |
| | | 8978647 | Aug 06, 2030 | DP | | |
| | | 9072857 | Apr 10, 2021 | DP | | |
| | | 9468727 | Jul 30, 2020 | DP | | |
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 100/50</u> | | | | | | |
| N 021077 | 001 | | | | M-214 | Dec 20, 2020 |
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 250/50</u> | | | | | | |
| N 021077 | 002 | | | | M-214 | Dec 20, 2020 |
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 500/50</u> | | | | | | |
| N 021077 | 003 | | | | M-214 | Dec 20, 2020 |
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u> | | | | | | |
| N 021254 | 001 | 7500444 | Feb 26, 2026 | DP | | |
| | | 7500444*PED | Aug 26, 2026 | | | |
| | | 7832351 | Jun 19, 2023 | DP | | |
| | | 9861771 | Oct 11, 2020 | DP | | |
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u> | | | | | | |
| N 021254 | 002 | 7500444 | Feb 26, 2026 | DP | | |
| | | 7500444*PED | Aug 26, 2026 | | | |
| | | 7832351 | Jun 19, 2023 | DP | | |
| | | 9861771 | Oct 11, 2020 | DP | | |
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u> | | | | | | |
| N 021254 | 003 | 7500444 | Feb 26, 2026 | DP | | |
| | | 7500444*PED | Aug 26, 2026 | | | |
| | | 7832351 | Jun 19, 2023 | DP | | |
| | | 9861771 | Oct 11, 2020 | DP | | |
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u> | | | | | | |
| N 208799 | 001 | 10022510 | May 18, 2031 | DP | NP | Jan 27, 2020 |
| | | 10124131 | May 18, 2031 | DP | | |
| | | 6701917 | Jun 23, 2021 | DP | | |
| | | 6718972 | Jun 23, 2021 | DP | | |
| | | 6748947 | Jun 23, 2021 | DP | | |
| | | 6871646 | Jun 23, 2021 | DP | | |
| | | 7540282 | May 06, 2023 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------------|-----------|------------------------|--------------|-------------------------|---------------------|-----------------------------|
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u> | | | | | | |
| N 208799 001 | 8006690 | Jun 23, 2021 | DP | | | |
| | 8651103 | Mar 26, 2028 | DP | | | |
| | 8714149 | Feb 25, 2032 | DP | | | |
| | 8978966 | Jan 13, 2032 | DP | | | |
| | 9066957 | Oct 06, 2034 | DP U-645 | | | |
| | 9216260 | Jun 28, 2031 | DP | | | |
| | 9415008 | Oct 06, 2034 | DP U-645 | | | |
| | 9463288 | May 19, 2025 | DP | | | |
| | 9616024 | Sep 01, 2024 | DP | | | |
| | 9731087 | May 18, 2031 | DP | | | |
| | 9987229 | Sep 01, 2024 | DP | | | |
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u> | | | | | | |
| N 208799 002 | 10022510 | May 18, 2031 | DP | | NP | Jan 27, 2020 |
| | 10124131 | May 18, 2031 | DP | | | |
| | 6701917 | Jun 23, 2021 | DP | | | |
| | 6718972 | Jun 23, 2021 | DP | | | |
| | 6748947 | Jun 23, 2021 | DP | | | |
| | 6871646 | Jun 23, 2021 | DP | | | |
| | 7540282 | May 06, 2023 | DP | | | |
| | 8006690 | Jun 23, 2021 | DP | | | |
| | 8651103 | Mar 26, 2028 | DP | | | |
| | 8714149 | Feb 25, 2032 | DP | | | |
| | 8978966 | Jan 13, 2032 | DP | | | |
| | 9066957 | Oct 06, 2034 | DP U-645 | | | |
| | 9216260 | Jun 28, 2031 | DP | | | |
| | 9463288 | May 19, 2025 | DP | | | |
| | 9616024 | Sep 01, 2024 | DP | | | |
| | 9731087 | May 18, 2031 | DP | | | |
| | 9987229 | Sep 01, 2024 | DP | | | |
| <u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u> | | | | | | |
| N 208799 003 | 10022510 | May 18, 2031 | DP | | NP | Jan 27, 2020 |
| | 10124131 | May 18, 2031 | DP | | | |
| | 6701917 | Jun 23, 2021 | DP | | | |
| | 6718972 | Jun 23, 2021 | DP | | | |
| | 6748947 | Jun 23, 2021 | DP | | | |
| | 6871646 | Jun 23, 2021 | DP | | | |
| | 7540282 | May 06, 2023 | DP | | | |
| | 8006690 | Jun 23, 2021 | DP | | | |
| | 8651103 | Mar 26, 2028 | DP | | | |
| | 8714149 | Feb 25, 2032 | DP | | | |
| | 8978966 | Jan 13, 2032 | DP | | | |
| | 9066957 | Oct 06, 2034 | DP U-645 | | | |
| | 9216260 | Jun 28, 2031 | DP | | | |
| | 9463288 | May 19, 2025 | DP | | | |
| | 9616024 | Sep 01, 2024 | DP | | | |
| | 9731087 | May 18, 2031 | DP | | | |
| | 9987229 | Sep 01, 2024 | DP | | | |
| <u>FLUVASTATIN SODIUM - LESCOL XL</u> | | | | | | |
| N 021192 001 | 6242003 | Apr 13, 2020 | | | | |
| <u>FLUVOXAMINE MALEATE - LUVOX CR</u> | | | | | | |
| N 022033 001 | 7465462 | May 10, 2020 | DP U-929 | | | |
| <u>FLUVOXAMINE MALEATE - LUVOX CR</u> | | | | | | |
| N 022033 002 | 7465462 | May 10, 2020 | DP U-929 | | | |
| <u>FOLLITROPIN ALFA/BETA - GONAL-F</u> | | | | | | |
| N 020378 004 | 7563763 | Aug 23, 2019 | DP | | | |
| <u>FOLLITROPIN ALFA/BETA - GONAL-F</u> | | | | | | |
| N 020378 005 | 7563763 | Aug 23, 2019 | DP | | | |
| <u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u> | | | | | | |
| N 021211 001 | 7446090 | Aug 23, 2019 | DP | | | |
| | 7563763 | Aug 23, 2019 | | | U-1183 | |
| | 7563763 | Aug 23, 2019 | | | U-1367 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u> | | | | | | |
| N 021211 | 001 | 7563763 | Aug 23, 2019 | U-993 | | |
| <u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u> | | | | | | |
| N 021211 | 002 | 7446090 | Aug 23, 2019 | DP | | |
| | | 7563763 | Aug 23, 2019 | U-1183 | | |
| | | 7563763 | Aug 23, 2019 | U-1367 | | |
| | | 7563763 | Aug 23, 2019 | U-993 | | |
| <u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u> | | | | | | |
| N 021211 | 003 | 7446090 | Aug 23, 2019 | DP | | |
| | | 7563763 | Aug 23, 2019 | U-1183 | | |
| | | 7563763 | Aug 23, 2019 | U-1367 | | |
| | | 7563763 | Aug 23, 2019 | U-993 | | |
| <u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u> | | | | | | |
| N 021211 | 004 | 7446090 | Aug 23, 2019 | DP | | |
| | | 7563763 | Aug 23, 2019 | U-1183 | | |
| | | 7563763 | Aug 23, 2019 | U-1367 | | |
| | | 7563763 | Aug 23, 2019 | U-993 | | |
| <u>FOLLITROPIN ALFA/BETA - GONAL-F RFF REDI-JECT</u> | | | | | | |
| N 021684 | 001 | 7446090 | Aug 23, 2019 | DP | | |
| | | 7741268 | Apr 02, 2024 | DP | | |
| <u>FOLLITROPIN ALFA/BETA - GONAL-F RFF REDI-JECT</u> | | | | | | |
| N 021684 | 002 | 7446090 | Aug 23, 2019 | DP | | |
| | | 7741268 | Apr 02, 2024 | DP | | |
| <u>FOLLITROPIN ALFA/BETA - GONAL-F RFF REDI-JECT</u> | | | | | | |
| N 021684 | 003 | 7446090 | Aug 23, 2019 | DP | | |
| | | 7741268 | Apr 02, 2024 | DP | | |
| <u>FOMEPIZOLE - ANTIZOL</u> | | | | | | |
| N 020696 | 001 | 7553863 | Jun 30, 2027 | DS DP | | |
| <u>FORMOTEROL FUMARATE - FORADIL</u> | | | | | | |
| N 020831 | 001 | 6488027 | Mar 08, 2019 | | | |
| | | 6887459 | Nov 28, 2020 | U-762 | | |
| <u>FORMOTEROL FUMARATE - PERFOROMIST</u> | | | | | | |
| N 022007 | 001 | 6667344 | Jun 22, 2021 | DP | | |
| | | 6814953 | Jun 22, 2021 | DP U-813 | | |
| | | 7348362 | Jun 22, 2021 | DP | | |
| | | 7462645 | Jun 22, 2021 | DP U-813 | | |
| | | 8623922 | Jun 22, 2021 | DP | | |
| | | 9730890 | Jun 22, 2021 | DP | | |
| <u>FORMOTEROL FUMARATE; GLYCOPYRROLATE - BEVESPI AEROSPHERE</u> | | | | | | |
| N 208294 | 001 | 8324266 | May 28, 2030 | U-1841 | NP | Apr 25, 2019 |
| | | 8703806 | May 28, 2030 | U-1841 | | |
| | | 8808713 | May 28, 2030 | DP U-1841 | | |
| | | 8815258 | Mar 17, 2031 | U-1841 | | |
| | | 9415009 | May 28, 2030 | U-1841 | | |
| | | 9463161 | May 28, 2030 | DP U-1841 | | |
| <u>FORMOTEROL FUMARATE; MOMETASONE FUROATE - DULERA</u> | | | | | | |
| N 022518 | 001 | 7067502 | May 21, 2020 | DP U-1068 | M-214 | Dec 20, 2020 |
| | | 7566705 | May 21, 2020 | DP U-1068 | | |
| <u>FORMOTEROL FUMARATE; MOMETASONE FUROATE - DULERA</u> | | | | | | |
| N 022518 | 002 | 7067502 | May 21, 2020 | DP U-1068 | M-214 | Dec 20, 2020 |
| | | 7566705 | May 21, 2020 | DP U-1068 | | |
| <u>FOSAMPRENAVIR CALCIUM - LEXIVA</u> | | | | | | |
| N 021548 | 001 | 6514953 | Jul 15, 2019 | DS DP U-257 | | |
| <u>FOSAPREPITANT DIMEGLUMINE - EMEND</u> | | | | | | |
| N 022023 | 001 | 5691336 | Mar 04, 2019 | DS DP | NPP | Apr 03, 2021 |
| | | 5691336*PED | Sep 04, 2019 | | PED | Oct 03, 2021 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>FOSAPREPITANT DIMEGLUMINE - EMEND</u> | | | | | | |
| N 022023 | 002 | 5691336 | Mar 04, 2019 | DS DP U-2265 | D-155 | Feb 01, 2019 |
| | | 5691336*PED | Sep 04, 2019 | | NPP | Apr 03, 2021 |
| | | | | | PED | Oct 03, 2021 |
| <u>FOSNETUPITANT CHLORIDE HYDROCHLORIDE; PALONOSETRON HYDROCHLORIDE - AKYNZEO</u> | | | | | | |
| N 210493 | 001 | 8426450 | May 23, 2032 | DS DP | NCE | Apr 19, 2023 |
| | | 8895586 | May 23, 2032 | U-2301 | | |
| | | 9186357 | Nov 18, 2030 | U-2301 | | |
| | | 9403772 | May 23, 2032 | DS U-2301 | | |
| | | 9908907 | May 23, 2032 | DS DP | | |
| <u>FOSPROPOFOL DISODIUM - LUSEDRA</u> | | | | | | |
| N 022244 | 001 | 6204257 | Jul 01, 2022 | DS DP U-945 | | |
| <u>FOSTAMATINIB DISODIUM - TAVALISSE</u> | | | | | | |
| N 209299 | 001 | 7449458 | Sep 04, 2026 | DS | NCE | Apr 17, 2023 |
| | | 7538108 | Mar 28, 2026 | DS U-2294 | ODE-174 | Apr 17, 2025 |
| | | 7989448 | Jun 12, 2026 | DS U-2294 | | |
| | | 8163902 | Jun 17, 2026 | DS U-2294 | | |
| | | 8211889 | Jan 19, 2026 | DS | | |
| | | 8263122 | Nov 24, 2030 | DP | | |
| | | 8445485 | Jun 17, 2026 | DP | | |
| | | 8652492 | Nov 06, 2028 | DP | | |
| | | 8771648 | Jul 27, 2032 | DP | | |
| | | 8912170 | Jun 17, 2026 | U-2294 | | |
| | | 8951504 | Jul 27, 2032 | U-2294 | | |
| | | 9266912 | Jan 19, 2026 | U-2294 | | |
| | | 9283238 | Jun 17, 2026 | U-2294 | | |
| | | 9737554 | Jan 19, 2026 | DP | | |
| <u>FOSTAMATINIB DISODIUM - TAVALISSE</u> | | | | | | |
| N 209299 | 002 | 7449458 | Sep 04, 2026 | DS | NCE | Apr 17, 2023 |
| | | 7538108 | Mar 28, 2026 | DS U-2294 | ODE-174 | Apr 17, 2025 |
| | | 7989448 | Jun 12, 2026 | DS U-2294 | | |
| | | 8163902 | Jun 17, 2026 | DS U-2294 | | |
| | | 8211889 | Jan 19, 2026 | DS | | |
| | | 8263122 | Nov 24, 2030 | DP | | |
| | | 8445485 | Jun 17, 2026 | DP | | |
| | | 8652492 | Nov 06, 2028 | DP | | |
| | | 8771648 | Jul 27, 2032 | DP | | |
| | | 8912170 | Jun 17, 2026 | U-2294 | | |
| | | 8951504 | Jul 27, 2032 | U-2294 | | |
| | | 9266912 | Jan 19, 2026 | U-2294 | | |
| | | 9283238 | Jun 17, 2026 | U-2294 | | |
| | | 9737554 | Jan 19, 2026 | DP | | |
| <u>FULVESTRANT - FASLODEX</u> | | | | | | |
| N 021344 | 001 | 6774122 | Jan 09, 2021 | U-1826 | I-725 | Feb 19, 2019 |
| | | 6774122 | Jan 09, 2021 | U-2108 | I-749 | Aug 25, 2020 |
| | | 6774122 | Jan 09, 2021 | U-2163 | | |
| | | 6774122 | Jan 09, 2021 | U-596 | | |
| | | 6774122*PED | Jul 09, 2021 | | | |
| | | 7456160 | Jan 09, 2021 | U-1826 | | |
| | | 7456160 | Jan 09, 2021 | U-2108 | | |
| | | 7456160 | Jan 09, 2021 | U-2163 | | |
| | | 7456160 | Jan 09, 2021 | U-596 | | |
| | | 7456160*PED | Jul 09, 2021 | | | |
| | | 8329680 | Jan 09, 2021 | U-1826 | | |
| | | 8329680 | Jan 09, 2021 | U-2108 | | |
| | | 8329680 | Jan 09, 2021 | U-2163 | | |
| | | 8329680 | Jan 09, 2021 | U-596 | | |
| | | 8329680*PED | Jul 09, 2021 | | | |
| | | 8466139 | Jan 09, 2021 | U-1826 | | |
| | | 8466139 | Jan 09, 2021 | U-2108 | | |
| | | 8466139 | Jan 09, 2021 | U-2163 | | |
| | | 8466139 | Jan 09, 2021 | U-596 | | |
| | | 8466139*PED | Jul 09, 2021 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>GABAPENTIN - NEURONTIN</u> | | | | | | |
| N 021129 | 001 7256216 | May 28, 2022 | DP | | | |
| <u>GABAPENTIN - GRALISE</u> | | | | | | |
| N 022544 | 001 6488962 | Jun 20, 2020 | DP | | | |
| | 6723340 | Oct 25, 2021 | DP | | | |
| | 7438927 | Feb 26, 2024 | U-1114 | | | |
| | 7731989 | Oct 25, 2022 | DP | | | |
| | 8192756 | Oct 25, 2022 | DP U-1114 | | | |
| | 8252332 | Oct 25, 2022 | DP U-1114 | | | |
| | 8333992 | Oct 25, 2022 | DP U-1114 | | | |
| <u>GABAPENTIN - GRALISE</u> | | | | | | |
| N 022544 | 002 6488962 | Jun 20, 2020 | DP | | | |
| | 6723340 | Oct 25, 2021 | DP | | | |
| | 7438927 | Feb 26, 2024 | U-1114 | | | |
| | 7731989 | Oct 25, 2022 | DP | | | |
| | 8192756 | Oct 25, 2022 | DP U-1114 | | | |
| | 8252332 | Oct 25, 2022 | DP U-1114 | | | |
| | 8333992 | Oct 25, 2022 | DP U-1114 | | | |
| <u>GABAPENTIN ENACARBIL - HORIZANT</u> | | | | | | |
| N 022399 | 001 6818787 | Nov 06, 2022 | DS DP | | ODE-25 | Jun 06, 2019 |
| | 8026279 | Nov 10, 2026 | DS DP | | | |
| | 8048917 | Nov 06, 2022 | DS DP U-1247 | | | |
| | 8114909 | Apr 11, 2026 | U-1231 | | | |
| | 8686034 | Jan 24, 2025 | U-1231 | | | |
| | 8686034 | Jan 24, 2025 | U-1247 | | | |
| | 8795725 | Jun 10, 2029 | DP U-1231 | | | |
| | 8795725 | Jun 10, 2029 | DP U-1247 | | | |
| <u>GABAPENTIN ENACARBIL - HORIZANT</u> | | | | | | |
| N 022399 | 002 6818787 | Nov 06, 2022 | DS DP | | ODE-25 | Jun 06, 2019 |
| | 8026279 | Nov 10, 2026 | DS DP | | | |
| | 8048917 | Nov 06, 2022 | DS DP U-1247 | | | |
| | 8114909 | Apr 11, 2026 | U-1231 | | | |
| | 8686034 | Jan 24, 2025 | U-1231 | | | |
| | 8686034 | Jan 24, 2025 | U-1247 | | | |
| | 8795725 | Jun 10, 2029 | DP U-1231 | | | |
| | 8795725 | Jun 10, 2029 | DP U-1247 | | | |
| <u>GADOBUTROL - GADAVIST</u> | | | | | | |
| N 201277 | 002 | | | | I-731 | Apr 27, 2019 |
| <u>GADOBUTROL - GADAVIST</u> | | | | | | |
| N 201277 | 006 5980864 | Nov 09, 2021 | DS DP U-1119 | | | |
| <u>GADOFOSVESET TRISODIUM - ABLAVAR</u> | | | | | | |
| N 021711 | 001 6676929 | May 04, 2020 | DP | | | |
| <u>GADOFOSVESET TRISODIUM - ABLAVAR</u> | | | | | | |
| N 021711 | 002 6676929 | May 04, 2020 | DP | | | |
| <u>GADOTERATE MEGLUMINE - DOTAREM</u> | | | | | | |
| N 204781 | 001 | | | | NPP | Aug 25, 2020 |
| <u>GADOTERATE MEGLUMINE - DOTAREM</u> | | | | | | |
| N 204781 | 002 | | | | NPP | Aug 25, 2020 |
| <u>GADOTERATE MEGLUMINE - DOTAREM</u> | | | | | | |
| N 204781 | 003 | | | | NPP | Aug 25, 2020 |
| <u>GADOTERATE MEGLUMINE - DOTAREM</u> | | | | | | |
| N 204781 | 004 | | | | NPP | Aug 25, 2020 |
| <u>GADOTERATE MEGLUMINE - DOTAREM</u> | | | | | | |
| N 204781 | 005 | | | | NPP | Aug 25, 2020 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>GADOXETATE DISODIUM - EOVI</u> | | | | | | |
| N 022090 | 001 6039931 | Nov 13, 2021 | U-1239 | | | |
| <u>GADOXETATE DISODIUM - EOVI</u> | | | | | | |
| N 022090 | 002 6039931 | Nov 13, 2021 | U-1239 | | | |
| <u>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</u> | | | | | | |
| N 021615 | 001 7160559 | Dec 20, 2019 | DP | | | |
| <u>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</u> | | | | | | |
| N 021615 | 002 7160559 | Dec 20, 2019 | DP | | | |
| <u>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</u> | | | | | | |
| N 021615 | 003 7160559 | Dec 20, 2019 | DP | | | |
| <u>GALLIUM DOTATATE GA-68 - NETSPOT</u> | | | | | | |
| N 208547 | 001 9375498 | Aug 10, 2032 | DP | | NCE ODE-120 | Jun 01, 2021 Jun 01, 2023 |
| <u>GANCICLOVIR - GANCICLOVIR</u> | | | | | | |
| N 209347 | 001 9486530 | Sep 02, 2034 | DP | | | |
| <u>GATIFLOXACIN - ZYMAR</u> | | | | | | |
| N 021493 | 001 6333045 | Aug 20, 2019 | DP | Y | | |
| | 6333045*PED | Feb 20, 2020 | | | | |
| <u>GEFITINIB - IRESSA</u> | | | | | | |
| N 206995 | 001 | | | | ODE-95 | Jul 13, 2022 |
| <u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u> | | | | | | |
| N 208313 | 001 9241948 | Jul 01, 2033 | DP | | | |
| <u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u> | | | | | | |
| N 208313 | 002 9241948 | Jul 01, 2033 | DP | | | |
| <u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u> | | | | | | |
| N 208313 | 003 9241948 | Jul 01, 2033 | DP | | | |
| <u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u> | | | | | | |
| N 208313 | 004 9241948 | Jul 01, 2033 | DP | | | |
| <u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u> | | | | | | |
| N 208313 | 005 9241948 | Jul 01, 2033 | DP | | | |
| <u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u> | | | | | | |
| N 208313 | 006 9241948 | Jul 01, 2033 | DP | | | |
| <u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u> | | | | | | |
| N 208313 | 007 9241948 | Jul 01, 2033 | DP | | | |
| <u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u> | | | | | | |
| N 208313 | 008 9241948 | Jul 01, 2033 | DP | | | |
| <u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u> | | | | | | |
| N 208313 | 009 9241948 | Jul 01, 2033 | DP | | | |
| <u>GEMCITABINE HYDROCHLORIDE - INFUGEM</u> | | | | | | |
| N 208313 | 010 9241948 | Jul 01, 2033 | DP | | | |
| <u>GEMIFLOXACIN MESYLATE - FACTIVE</u> | | | | | | |
| N 021158 | 001 6262071 | Sep 21, 2019 | U-513 | | | |
| | 6331550 | Sep 21, 2019 | U-511 | | | |
| | 6340689 | Sep 14, 2019 | U-512 | | | |
| | 6455540 | Sep 21, 2019 | U-511 | | | |
| | 6803376 | Sep 21, 2019 | DS DP U-608 | | | |
| | 6803376 | Sep 21, 2019 | DS DP U-609 | | | |
| <u>GILTERITINIB FUMARATE - XOSPATA</u> | | | | | | |
| N 211349 | 001 8969336 | Jan 27, 2031 | DS DP | | ODE-222 | Nov 28, 2025 |
| | 9487491 | Jul 28, 2030 | U-2456 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>GLASDEGIB - DAURISMO</u> | | | | | | |
| N 210656 001 | 8148401 | Jan 30, 2031 | DS DP | | NCE | Nov 21, 2023 |
| | 8431597 | Jun 29, 2028 | DP | | ODE-224 | Nov 21, 2025 |
| <u>GLASDEGIB - DAURISMO</u> | | | | | | |
| N 210656 002 | 8148401 | Jan 30, 2031 | DS DP | | NCE | Nov 21, 2023 |
| | 8431597 | Jun 29, 2028 | DP | | ODE-224 | Nov 21, 2025 |
| <u>GLATIRAMER ACETATE - COPAXONE</u> | | | | | | |
| N 020622 003 | 8232250 | Aug 19, 2030 | | U-441 | | |
| | 8399413 | Aug 19, 2030 | | U-441 | | |
| | 8969302 | Aug 19, 2030 | | U-441 | | |
| | 9155776 | Aug 19, 2030 | | U-441 | | |
| | 9402874 | Aug 19, 2030 | | U-441 | | |
| <u>GLECAPREVIR; PIBRENTASVIR - MAVYRET</u> | | | | | | |
| N 209394 001 | 10028937 | Jun 10, 2030 | | U-2141 | M-230 | Aug 06, 2021 |
| | 10039754 | Jun 10, 2030 | | U-2141 | NCE | Aug 03, 2022 |
| | 8648037 | Jan 19, 2032 | DS DP | U-2141 | | |
| | 8937150 | May 18, 2032 | DS DP | | | |
| | 9321807 | Jun 05, 2035 | DS | | | |
| | 9586978 | Jun 10, 2030 | | U-2141 | | |
| <u>GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT</u> | | | | | | |
| N 021925 001 | 7700128 | Jan 30, 2027 | DP | | | |
| | 8071130 | Jun 08, 2028 | DP | | | |
| <u>GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT</u> | | | | | | |
| N 021925 002 | 7700128 | Jan 30, 2027 | DP | | | |
| | 8071130 | Jun 08, 2028 | DP | | | |
| <u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u> | | | | | | |
| N 021700 001 | 7358366 | Apr 19, 2020 | DS | | Y | |
| | 7358366*PED | Oct 19, 2020 | | | | |
| <u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u> | | | | | | |
| N 021700 002 | 7358366 | Apr 19, 2020 | DS | | Y | |
| | 7358366*PED | Oct 19, 2020 | | | | |
| <u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u> | | | | | | |
| N 021700 003 | 7358366 | Apr 19, 2020 | DS | | Y | |
| | 7358366*PED | Oct 19, 2020 | | | | |
| <u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u> | | | | | | |
| N 021700 004 | 7358366 | Apr 19, 2020 | DS | | Y | |
| | 7358366*PED | Oct 19, 2020 | | | | |
| <u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u> | | | | | | |
| N 021700 005 | 7358366 | Apr 19, 2020 | DS | | Y | |
| | 7358366*PED | Oct 19, 2020 | | | | |
| <u>GLIPIZIDE - GLUCOTROL XL</u> | | | | | | |
| N 020329 001 | RE44459 | Mar 26, 2019 | | U-1431 | | |
| <u>GLIPIZIDE - GLUCOTROL XL</u> | | | | | | |
| N 020329 002 | RE44459 | Mar 26, 2019 | | U-1431 | | |
| <u>GLIPIZIDE - GLUCOTROL XL</u> | | | | | | |
| N 020329 003 | RE44459 | Mar 26, 2019 | | U-1431 | | |
| <u>GLYBURIDE; METFORMIN HYDROCHLORIDE - GLUCOVANCE</u> | | | | | | |
| N 021178 001 | 6303146 | Jul 14, 2019 | | U-412 | | |
| <u>GLYBURIDE; METFORMIN HYDROCHLORIDE - GLUCOVANCE</u> | | | | | | |
| N 021178 002 | 6303146 | Jul 14, 2019 | | U-412 | | |
| <u>GLYBURIDE; METFORMIN HYDROCHLORIDE - GLUCOVANCE</u> | | | | | | |
| N 021178 003 | 6303146 | Jul 14, 2019 | | U-412 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>GLYCEROL PHENYLBUTYRATE - RAVICTI</u> | | | | | | |
| N 203284 001 | 10045958 | Sep 22, 2030 | U-1816 | | NPP | Apr 28, 2020 |
| | 10045959 | Sep 22, 2030 | U-1816 | | ODE-157 | Apr 28, 2024 |
| | 8404215 | Mar 09, 2032 | U-1383 | | ODE-42 | Feb 01, 2020 |
| | 8642012 | Sep 22, 2030 | U-1383 | | | |
| | 9095559 | Mar 09, 2032 | U-1383 | | | |
| | 9254278 | Mar 09, 2032 | U-1816 | | | |
| | 9326966 | Mar 09, 2032 | U-1816 | | | |
| | 9561197 | Sep 22, 2030 | U-1383 | | | |
| | 9962359 | Sep 22, 2030 | U-1816 | | | |
| | 9999608 | Sep 22, 2030 | U-1816 | | | |
| <u>GLYCOPYRROLATE - CUVPOSA</u> | | | | | | |
| N 022571 001 | 7638552 | Aug 20, 2023 | U-1076 | | | |
| | 7816396 | Aug 20, 2023 | U-1076 | | | |
| <u>GLYCOPYRROLATE - SEEBRI</u> | | | | | | |
| N 207923 001 | 7229607 | Apr 09, 2021 | U-1773 | | | |
| | 7736670 | Jun 27, 2021 | DP | | | |
| | 8029768 | Apr 09, 2021 | U-1773 | | | |
| | 8048451 | Jun 27, 2021 | DP | | | |
| | 8182838 | Oct 20, 2028 | DP | | | |
| | 8303991 | Jun 27, 2021 | DP | | | |
| | 8435567 | Jun 27, 2021 | DP | | | |
| | 8479730 | Oct 11, 2028 | DP | | | |
| | 8580306 | Jun 27, 2021 | DP | | | |
| | 8956661 | Jun 27, 2021 | DP | | | |
| | 9931304 | Jun 27, 2021 | DP | | | |
| | 9962338 | Jun 27, 2021 | DP | | | |
| <u>GLYCOPYRROLATE - LONHALA MAGNAIR KIT</u> | | | | | | |
| N 208437 001 | 6962151 | Oct 27, 2020 | DP | | NP | Dec 05, 2020 |
| | 7316067 | Sep 06, 2022 | DP | | | |
| | 7458372 | Nov 18, 2024 | DP | | | |
| | 7931212 | Nov 25, 2025 | DP | | | |
| | 8511581 | Nov 08, 2023 | DP | | | |
| | 9168556 | Sep 01, 2032 | DP | | | |
| | 9265900 | Dec 07, 2028 | DP | | | |
| | 9604018 | May 16, 2033 | DP | | | |
| | 9789270 | Oct 30, 2030 | DP | | | |
| <u>GLYCOPYRROLATE ; INDACATEROL MALEATE - UTIBRON</u> | | | | | | |
| N 207930 001 | 6878721 | Feb 25, 2025 | DS DP U-1773 | | | |
| | 7229607 | Apr 09, 2021 | U-1773 | | | |
| | 7736670 | Jun 27, 2021 | DP | | | |
| | 7820694 | Jun 02, 2020 | DP U-1773 | | | |
| | 8029768 | Apr 09, 2021 | U-1773 | | | |
| | 8048451 | Jun 27, 2021 | DP | | | |
| | 8067437 | Jun 02, 2020 | U-1773 | | | |
| | 8182838 | Oct 20, 2028 | DP | | | |
| | 8283362 | Jun 02, 2020 | DP U-1773 | | | |
| | 8303991 | Jun 27, 2021 | DP | | | |
| | 8435567 | Jun 27, 2021 | DP | | | |
| | 8479730 | Oct 11, 2028 | DP | | | |
| | 8580306 | Jun 27, 2021 | DP | | | |
| | 8658673 | Jun 02, 2020 | DP U-1773 | | | |
| | 8796307 | Jun 02, 2020 | DP | | | |
| | 8956661 | Jun 27, 2021 | DP | | | |
| | 9931304 | Jun 27, 2021 | DP | | | |
| | 9962338 | Jun 27, 2021 | DP | | | |
| <u>GLYCOPYRROLATE ; INDACATEROL MALEATE - UTIBRON</u> | | | | | | |
| N 210361 001 | 10004717 | Feb 28, 2033 | DP U-2398 | | | |
| | 10052267 | Oct 17, 2028 | DP U-2398 | | | |
| | 6433003 | Apr 10, 2020 | U-2398 | | | |
| | 8618160 | Dec 10, 2029 | DP U-2398 | | | |
| | 8859610 | Feb 28, 2033 | DP U-2398 | | | |
| | 9006462 | Feb 28, 2033 | DP | | | |
| | 9259414 | Feb 28, 2033 | U-2398 | | | |
| | 9744105 | Jul 18, 2030 | DP U-2398 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>GLYCOPYRRONIUM TOSYLATE - OBREXZA</u> | | | | | | |
| N 210361 | 001 | 10004717 | Feb 28, 2033 | DP U-2398 | | |
| | | 10052267 | Oct 17, 2028 | DP U-2398 | | |
| | | 6433003 | Apr 10, 2020 | U-2398 | | |
| | | 8618160 | Dec 10, 2029 | DP U-2398 | | |
| | | 8859610 | Feb 28, 2033 | DP U-2398 | | |
| | | 9006462 | Feb 28, 2033 | DP | | |
| | | 9259414 | Feb 28, 2033 | U-2398 | | |
| | | 9744105 | Jul 18, 2030 | DP U-2398 | | |
| <u>GOSERELIN ACETATE - ZOLADEX</u> | | | | | | |
| N 019726 | 001 | 7118552 | Apr 13, 2022 | DP | | |
| | | 7220247 | Apr 09, 2022 | DP | | |
| | | 7500964 | Feb 26, 2021 | DP | | |
| <u>GOSERELIN ACETATE - ZOLADEX</u> | | | | | | |
| N 020578 | 001 | 7118552 | Apr 13, 2022 | DP | | |
| | | 7220247 | Apr 09, 2022 | DP | | |
| | | 7500964 | Feb 26, 2021 | DP | | |
| <u>GRANISETRON - SANCUSO</u> | | | | | | |
| N 022198 | 001 | 7608282 | Jan 22, 2025 | DP U-1011 | | |
| <u>GRANISETRON - SUSTOL</u> | | | | | | |
| N 022445 | 001 | 6613355 | Jun 28, 2021 | DP | NDF | Aug 09, 2019 |
| | | 6790458 | May 11, 2021 | DP | | |
| | | 8252304 | Sep 28, 2024 | DP | | |
| | | 8252305 | Sep 28, 2024 | U-1891 | | |
| | | 8715710 | Sep 28, 2024 | DP | | |
| | | 9913910 | Sep 28, 2024 | U-2253 | | |
| <u>GUAIFENESIN - MUCINEX</u> | | | | | | |
| N 021282 | 001 | 6372252 | Apr 28, 2020 | U-489 | | |
| | | 6955821 | Apr 28, 2020 | DP U-489 | | |
| | | 7838032 | Apr 28, 2020 | DP | | |
| <u>GUAIFENESIN - MUCINEX</u> | | | | | | |
| N 021282 | 002 | 6372252 | Apr 28, 2020 | U-489 | | |
| | | 6955821 | Apr 28, 2020 | DP U-489 | | |
| | | 7838032 | Apr 28, 2020 | DP | | |
| <u>GUAIFENESIN; HYDROCODONE BITARTRATE - OBREDON</u> | | | | | | |
| N 205474 | 001 | 10105324 | Nov 13, 2035 | DS DP U-2023 | | |
| | | 9549907 | Nov 13, 2035 | DS DP U-2023 | | |
| | | 9808431 | Nov 13, 2035 | DS DP U-2023 | | |
| <u>GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE - MUCINEX D</u> | | | | | | |
| N 021585 | 001 | 6372252 | Apr 28, 2020 | DP | | |
| | | 6955821 | Apr 28, 2020 | DP U-686 | | |
| | | 7838032 | Apr 28, 2020 | DP | | |
| <u>GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE - MUCINEX D</u> | | | | | | |
| N 021585 | 002 | 6372252 | Apr 28, 2020 | DP | | |
| | | 6955821 | Apr 28, 2020 | DP U-686 | | |
| | | 7838032 | Apr 28, 2020 | DP | | |
| <u>GUANFACINE HYDROCHLORIDE - INTUNIV</u> | | | | | | |
| N 022037 | 001 | 6287599 | Dec 20, 2020 | DP | | |
| | | 6287599*PED | Jun 20, 2021 | | | |
| | | 6811794 | Jul 04, 2022 | DP U-494 | | |
| | | 6811794*PED | Jan 04, 2023 | | | |
| <u>GUANFACINE HYDROCHLORIDE - INTUNIV</u> | | | | | | |
| N 022037 | 002 | 6287599 | Dec 20, 2020 | DP | | |
| | | 6287599*PED | Jun 20, 2021 | | | |
| | | 6811794 | Jul 04, 2022 | DP U-494 | | |
| | | 6811794*PED | Jan 04, 2023 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>GUANFACINE HYDROCHLORIDE - INTUNIV</u> | | | | | | |
| N 022037 | 003 | 6287599 | Dec 20, 2020 | DP | | |
| | | 6287599*PED | Jun 20, 2021 | | | |
| | | 6811794 | Jul 04, 2022 | DP U-494 | | |
| | | 6811794*PED | Jan 04, 2023 | | | |
| <u>GUANFACINE HYDROCHLORIDE - INTUNIV</u> | | | | | | |
| N 022037 | 004 | 6287599 | Dec 20, 2020 | DP | | |
| | | 6287599*PED | Jun 20, 2021 | | | |
| | | 6811794 | Jul 04, 2022 | DP U-494 | | |
| | | 6811794*PED | Jan 04, 2023 | | | |
| <u>HALOBETASOL PROPIONATE - ULTRAVATE</u> | | | | | | |
| N 208183 | 001 | 8962028 | Jun 19, 2033 | DP U-1775 | | |
| <u>HALOBETASOL PROPIONATE - BRYHALI</u> | | | | | | |
| N 209355 | 001 | | | | NP | Nov 06, 2021 |
| <u>HALOBETASOL PROPIONATE - HALOBETASOL PROPIONATE</u> | | | | | | |
| N 210566 | 001 | | | | NDF | May 24, 2021 |
| <u>HEXAMINOLEVULINATE HYDROCHLORIDE - CYSVIEW KIT</u> | | | | | | |
| N 022555 | 001 | 7348361 | Nov 06, 2020 | DP U-1087 | M-220 | Feb 15, 2021 |
| | | 7348361 | Nov 06, 2020 | DP U-2250 | | |
| <u>HISTRELIN ACETATE - SUPPRELIN LA</u> | | | | | | |
| N 022058 | 001 | 8062652 | Jun 16, 2026 | U-1197 | | |
| <u>HYALURONIDASE RECOMBINANT HUMAN - HYLENEX RECOMBINANT</u> | | | | | | |
| N 021859 | 001 | 7767429 | Sep 23, 2027 | DS DP | | |
| <u>HYDRALAZINE HYDROCHLORIDE; ISOSORBIDE DINITRATE - BIDIL</u> | | | | | | |
| N 020727 | 001 | 6465463 | Sep 08, 2020 | U-71 | | |
| | | 6784177 | Sep 08, 2020 | U-71 | | |
| <u>HYDROCHLOROTHIAZIDE; TELMISARTAN - MICARDIS HCT</u> | | | | | | |
| N 021162 | 001 | 6358986 | Jan 10, 2020 | | | |
| <u>HYDROCHLOROTHIAZIDE; TELMISARTAN - MICARDIS HCT</u> | | | | | | |
| N 021162 | 002 | 6358986 | Jan 10, 2020 | | | |
| <u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u> | | | | | | |
| N 202880 | 001 | 10028946 | Jul 25, 2033 | U-1810 | | |
| | | 10092559 | Sep 12, 2034 | U-55 | | |
| | | 6228398 | Nov 01, 2019 | DP | | |
| | | 6902742 | Nov 01, 2019 | DP | | |
| | | 9132096 | Sep 12, 2034 | DP | | |
| | | 9265760 | Jul 25, 2033 | U-1810 | | |
| | | 9326982 | Jul 25, 2033 | U-1810 | | |
| | | 9333201 | Jul 25, 2033 | U-1810 | | |
| | | 9339499 | Jul 25, 2033 | U-1810 | | |
| | | 9421200 | Jul 25, 2033 | U-1810 | | |
| | | 9433619 | Jul 25, 2033 | U-1810 | | |
| | | 9452163 | Sep 12, 2034 | U-55 | | |
| | | 9486451 | Sep 12, 2034 | U-55 | | |
| | | 9610286 | Jul 25, 2033 | U-1810 | | |
| | | 9713611 | Sep 12, 2034 | DP U-55 | | |
| <u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u> | | | | | | |
| N 202880 | 002 | 10028946 | Jul 25, 2033 | U-1810 | | |
| | | 10092559 | Sep 12, 2034 | U-55 | | |
| | | 6228398 | Nov 01, 2019 | DP | | |
| | | 6902742 | Nov 01, 2019 | DP | | |
| | | 9132096 | Sep 12, 2034 | DP | | |
| | | 9265760 | Jul 25, 2033 | U-1810 | | |
| | | 9326982 | Jul 25, 2033 | U-1810 | | |
| | | 9333201 | Jul 25, 2033 | U-1810 | | |
| | | 9339499 | Jul 25, 2033 | U-1810 | | |
| | | 9421200 | Jul 25, 2033 | U-1810 | | |
| | | 9433619 | Jul 25, 2033 | U-1810 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u> | | | | | | |
| N 202880 | 002 | 9452163 | Sep 12, 2034 | | U-55 | |
| | | 9486451 | Sep 12, 2034 | | U-55 | |
| | | 9610286 | Jul 25, 2033 | | U-1810 | |
| | | 9713611 | Sep 12, 2034 | DP | U-55 | |
| <u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u> | | | | | | |
| N 202880 | 003 | 10028946 | Jul 25, 2033 | | U-1810 | |
| | | 10092559 | Sep 12, 2034 | | U-55 | |
| | | 6228398 | Nov 01, 2019 | DP | | |
| | | 6902742 | Nov 01, 2019 | DP | | |
| | | 9132096 | Sep 12, 2034 | DP | | |
| | | 9265760 | Jul 25, 2033 | | U-1810 | |
| | | 9326982 | Jul 25, 2033 | | U-1810 | |
| | | 9333201 | Jul 25, 2033 | | U-1810 | |
| | | 9339499 | Jul 25, 2033 | | U-1810 | |
| | | 9421200 | Jul 25, 2033 | | U-1810 | |
| | | 9433619 | Jul 25, 2033 | | U-1810 | |
| | | 9452163 | Sep 12, 2034 | | U-55 | |
| | | 9486451 | Sep 12, 2034 | | U-55 | |
| | | 9610286 | Jul 25, 2033 | | U-1810 | |
| | | 9713611 | Sep 12, 2034 | DP | U-55 | |
| <u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u> | | | | | | |
| N 202880 | 004 | 10028946 | Jul 25, 2033 | | U-1810 | |
| | | 10092559 | Sep 12, 2034 | | U-55 | |
| | | 6228398 | Nov 01, 2019 | DP | | |
| | | 6902742 | Nov 01, 2019 | DP | | |
| | | 9132096 | Sep 12, 2034 | DP | | |
| | | 9265760 | Jul 25, 2033 | | U-1810 | |
| | | 9326982 | Jul 25, 2033 | | U-1810 | |
| | | 9333201 | Jul 25, 2033 | | U-1810 | |
| | | 9339499 | Jul 25, 2033 | | U-1810 | |
| | | 9421200 | Jul 25, 2033 | | U-1810 | |
| | | 9433619 | Jul 25, 2033 | | U-1810 | |
| | | 9452163 | Sep 12, 2034 | | U-55 | |
| | | 9486451 | Sep 12, 2034 | | U-55 | |
| | | 9610286 | Jul 25, 2033 | | U-1810 | |
| | | 9713611 | Sep 12, 2034 | DP | U-55 | |
| <u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u> | | | | | | |
| N 202880 | 005 | 10028946 | Jul 25, 2033 | | U-1810 | |
| | | 10092559 | Sep 12, 2034 | | U-55 | |
| | | 6228398 | Nov 01, 2019 | DP | | |
| | | 6902742 | Nov 01, 2019 | DP | | |
| | | 9132096 | Sep 12, 2034 | DP | | |
| | | 9265760 | Jul 25, 2033 | | U-1810 | |
| | | 9326982 | Jul 25, 2033 | | U-1810 | |
| | | 9333201 | Jul 25, 2033 | | U-1810 | |
| | | 9339499 | Jul 25, 2033 | | U-1810 | |
| | | 9421200 | Jul 25, 2033 | | U-1810 | |
| | | 9433619 | Jul 25, 2033 | | U-1810 | |
| | | 9452163 | Sep 12, 2034 | | U-55 | |
| | | 9486451 | Sep 12, 2034 | | U-55 | |
| | | 9610286 | Jul 25, 2033 | | U-1810 | |
| | | 9713611 | Sep 12, 2034 | DP | U-55 | |
| <u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u> | | | | | | |
| N 202880 | 006 | 10028946 | Jul 25, 2033 | | U-1810 | |
| | | 10092559 | Sep 12, 2034 | | U-55 | |
| | | 6228398 | Nov 01, 2019 | DP | | |
| | | 6902742 | Nov 01, 2019 | DP | | |
| | | 9132096 | Sep 12, 2034 | DP | | |
| | | 9265760 | Jul 25, 2033 | | U-1810 | |
| | | 9326982 | Jul 25, 2033 | | U-1810 | |
| | | 9333201 | Jul 25, 2033 | | U-1810 | |
| | | 9339499 | Jul 25, 2033 | | U-1810 | |
| | | 9421200 | Jul 25, 2033 | | U-1810 | |
| | | 9433619 | Jul 25, 2033 | | U-1810 | |
| | | 9452163 | Sep 12, 2034 | | U-55 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u> | | | | | | |
| N 202880 | 006 | 9486451 | Sep 12, 2034 | | U-55 | |
| | | 9610286 | Jul 25, 2033 | | U-1810 | |
| | | 9713611 | Sep 12, 2034 | DP | U-55 | |
| <u>HYDROCODONE BITARTRATE - HYSINGLA</u> | | | | | | |
| N 206627 | 001 | 10130591 | Nov 20, 2023 | DP | U-1819 | |
| | | 6733783 | Oct 30, 2021 | DP | U-1556 | |
| | | 8309060 | Nov 20, 2023 | DP | U-1556 | |
| | | 8361499 | Oct 30, 2021 | DP | | |
| | | 8529948 | Aug 06, 2022 | DP | | |
| | | 8551520 | Oct 30, 2021 | DP | | |
| | | 8647667 | Oct 30, 2021 | DP | | |
| | | 8808740 | Dec 21, 2031 | DP | U-1556 | |
| | | 9023401 | Oct 30, 2020 | DP | | |
| | | 9056052 | Oct 30, 2020 | DP | | |
| | | 9060940 | Oct 30, 2020 | | U-1556 | |
| | | 9084816 | Aug 24, 2027 | DP | | |
| | | 9095614 | Aug 24, 2027 | | U-1556 | |
| | | 9095615 | Aug 24, 2027 | DP | | |
| | | 9198863 | Oct 30, 2020 | DP | | |
| | | 9205056 | Oct 30, 2020 | DP | | |
| | | 9289391 | Oct 30, 2020 | DP | | |
| | | 9486412 | Aug 24, 2027 | DP | | |
| | | 9486413 | Aug 24, 2027 | DP | | |
| | | 9492389 | Aug 24, 2027 | DP | | |
| | | 9492390 | Aug 24, 2027 | | U-1556 | |
| | | 9492391 | Aug 24, 2027 | | U-1556 | |
| | | 9517236 | Oct 30, 2020 | DP | | |
| | | 9545380 | Aug 24, 2027 | | U-1556 | |
| | | 9572779 | Dec 21, 2031 | DP | | |
| | | 9572804 | Oct 30, 2020 | DP | | |
| | | 9669023 | Oct 30, 2020 | DP | | |
| | | 9669024 | Oct 30, 2020 | DP | | |
| | | 9675610 | Jun 16, 2023 | DP | | |
| | | 9675611 | Oct 30, 2020 | | U-1556 | |
| | | 9682077 | Oct 30, 2020 | | U-1556 | |
| | | 9750703 | Dec 21, 2031 | DP | | |
| | | 9763933 | Aug 24, 2027 | DP | | |
| | | 9770416 | Aug 24, 2027 | DP | | |
| | | 9775809 | Aug 24, 2027 | DP | | |
| | | 9861584 | Dec 21, 2031 | DP | | |
| | | 9872837 | Dec 21, 2031 | DP | | |
| <u>HYDROCODONE BITARTRATE - HYSINGLA</u> | | | | | | |
| N 206627 | 002 | 10130591 | Nov 20, 2023 | DP | U-1819 | |
| | | 6733783 | Oct 30, 2021 | DP | U-1556 | |
| | | 8309060 | Nov 20, 2023 | DP | U-1556 | |
| | | 8361499 | Oct 30, 2021 | DP | | |
| | | 8529948 | Aug 06, 2022 | DP | | |
| | | 8551520 | Oct 30, 2021 | DP | | |
| | | 8647667 | Oct 30, 2021 | DP | | |
| | | 8808740 | Dec 21, 2031 | DP | U-1556 | |
| | | 9023401 | Oct 30, 2020 | DP | | |
| | | 9056052 | Oct 30, 2020 | DP | | |
| | | 9060940 | Oct 30, 2020 | | U-1556 | |
| | | 9084816 | Aug 24, 2027 | DP | | |
| | | 9095614 | Aug 24, 2027 | | U-1556 | |
| | | 9095615 | Aug 24, 2027 | DP | | |
| | | 9198863 | Oct 30, 2020 | DP | | |
| | | 9205056 | Oct 30, 2020 | DP | | |
| | | 9289391 | Oct 30, 2020 | DP | | |
| | | 9486412 | Aug 24, 2027 | DP | | |
| | | 9486413 | Aug 24, 2027 | DP | | |
| | | 9492389 | Aug 24, 2027 | DP | | |
| | | 9492390 | Aug 24, 2027 | | U-1556 | |
| | | 9492391 | Aug 24, 2027 | | U-1556 | |
| | | 9517236 | Oct 30, 2020 | DP | | |
| | | 9545380 | Aug 24, 2027 | | U-1556 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>HYDROCODONE BITARTRATE - HYSINGLA</u> | | | | | | |
| N 206627 002 | 9572779 | Dec 21, 2031 | DP | | | |
| | 9572804 | Oct 30, 2020 | DP | | | |
| | 9669023 | Oct 30, 2020 | DP | | | |
| | 9669024 | Oct 30, 2020 | DP | | | |
| | 9675610 | Jun 16, 2023 | DP | | | |
| | 9675611 | Oct 30, 2020 | | U-1556 | | |
| | 9682077 | Oct 30, 2020 | | U-1556 | | |
| | 9750703 | Dec 21, 2031 | DP | | | |
| | 9763933 | Aug 24, 2027 | DP | | | |
| | 9770416 | Aug 24, 2027 | DP | | | |
| | 9775809 | Aug 24, 2027 | DP | | | |
| | 9861584 | Dec 21, 2031 | DP | | | |
| | 9872837 | Dec 21, 2031 | DP | | | |
| <u>HYDROCODONE BITARTRATE - HYSINGLA</u> | | | | | | |
| N 206627 003 | 10130591 | Nov 20, 2023 | DP | U-1819 | | |
| | 6733783 | Oct 30, 2021 | DP | U-1556 | | |
| | 8309060 | Nov 20, 2023 | DP | U-1556 | | |
| | 8361499 | Oct 30, 2021 | DP | | | |
| | 8529948 | Aug 06, 2022 | DP | | | |
| | 8551520 | Oct 30, 2021 | DP | | | |
| | 8647667 | Oct 30, 2021 | DP | | | |
| | 8808740 | Dec 21, 2031 | DP | U-1556 | | |
| | 9023401 | Oct 30, 2020 | DP | | | |
| | 9056052 | Oct 30, 2020 | DP | | | |
| | 9060940 | Oct 30, 2020 | | U-1556 | | |
| | 9084816 | Aug 24, 2027 | DP | | | |
| | 9095614 | Aug 24, 2027 | | U-1556 | | |
| | 9095615 | Aug 24, 2027 | DP | | | |
| | 9198863 | Oct 30, 2020 | DP | | | |
| | 9205056 | Oct 30, 2020 | DP | | | |
| | 9289391 | Oct 30, 2020 | DP | | | |
| | 9486412 | Aug 24, 2027 | DP | | | |
| | 9486413 | Aug 24, 2027 | DP | | | |
| | 9492389 | Aug 24, 2027 | DP | | | |
| | 9492390 | Aug 24, 2027 | | U-1556 | | |
| | 9492391 | Aug 24, 2027 | | U-1556 | | |
| | 9517236 | Oct 30, 2020 | DP | | | |
| | 9545380 | Aug 24, 2027 | | U-1556 | | |
| | 9572779 | Dec 21, 2031 | DP | | | |
| | 9572804 | Oct 30, 2020 | DP | | | |
| | 9669023 | Oct 30, 2020 | DP | | | |
| | 9669024 | Oct 30, 2020 | DP | | | |
| | 9675610 | Jun 16, 2023 | DP | | | |
| | 9675611 | Oct 30, 2020 | | U-1556 | | |
| | 9682077 | Oct 30, 2020 | | U-1556 | | |
| | 9750703 | Dec 21, 2031 | DP | | | |
| | 9763933 | Aug 24, 2027 | DP | | | |
| | 9770416 | Aug 24, 2027 | DP | | | |
| | 9775809 | Aug 24, 2027 | DP | | | |
| | 9861584 | Dec 21, 2031 | DP | | | |
| | 9872837 | Dec 21, 2031 | DP | | | |
| <u>HYDROCODONE BITARTRATE - HYSINGLA</u> | | | | | | |
| N 206627 004 | 10130591 | Nov 20, 2023 | DP | U-1819 | | |
| | 6733783 | Oct 30, 2021 | DP | U-1556 | | |
| | 8309060 | Nov 20, 2023 | DP | U-1556 | | |
| | 8361499 | Oct 30, 2021 | DP | | | |
| | 8529948 | Aug 06, 2022 | DP | | | |
| | 8551520 | Oct 30, 2021 | DP | | | |
| | 8647667 | Oct 30, 2021 | DP | | | |
| | 8808740 | Dec 21, 2031 | DP | U-1556 | | |
| | 9023401 | Oct 30, 2020 | DP | | | |
| | 9056052 | Oct 30, 2020 | DP | | | |
| | 9060940 | Oct 30, 2020 | | U-1556 | | |
| | 9084816 | Aug 24, 2027 | DP | | | |
| | 9095614 | Aug 24, 2027 | | U-1556 | | |
| | 9095615 | Aug 24, 2027 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>HYDROCODONE BITARTRATE - HYSINGLA</u> | | | | | | |
| N 206627 004 | 9198863 | Oct 30, 2020 | DP | | | |
| | 9205056 | Oct 30, 2020 | DP | | | |
| | 9289391 | Oct 30, 2020 | DP | | | |
| | 9486412 | Aug 24, 2027 | DP | | | |
| | 9486413 | Aug 24, 2027 | DP | | | |
| | 9492389 | Aug 24, 2027 | DP | | | |
| | 9492390 | Aug 24, 2027 | | U-1556 | | |
| | 9492391 | Aug 24, 2027 | | U-1556 | | |
| | 9517236 | Oct 30, 2020 | DP | | | |
| | 9545380 | Aug 24, 2027 | | U-1556 | | |
| | 9572779 | Dec 21, 2031 | DP | | | |
| | 9572804 | Oct 30, 2020 | DP | | | |
| | 9669023 | Oct 30, 2020 | DP | | | |
| | 9669024 | Oct 30, 2020 | DP | | | |
| | 9675610 | Jun 16, 2023 | DP | | | |
| | 9675611 | Oct 30, 2020 | | U-1556 | | |
| | 9682077 | Oct 30, 2020 | | U-1556 | | |
| | 9750703 | Dec 21, 2031 | DP | | | |
| | 9763933 | Aug 24, 2027 | DP | | | |
| | 9770416 | Aug 24, 2027 | DP | | | |
| | 9775809 | Aug 24, 2027 | DP | | | |
| | 9861584 | Dec 21, 2031 | DP | | | |
| | 9872837 | Dec 21, 2031 | DP | | | |
| <u>HYDROCODONE BITARTRATE - HYSINGLA</u> | | | | | | |
| N 206627 005 | 10130591 | Nov 20, 2023 | DP | U-1819 | | |
| | 6733783 | Oct 30, 2021 | DP | U-1556 | | |
| | 8309060 | Nov 20, 2023 | DP | U-1556 | | |
| | 8361499 | Oct 30, 2021 | DP | | | |
| | 8529948 | Aug 06, 2022 | DP | | | |
| | 8551520 | Oct 30, 2021 | DP | | | |
| | 8647667 | Oct 30, 2021 | DP | | | |
| | 8808740 | Dec 21, 2031 | DP | U-1556 | | |
| | 9056052 | Oct 30, 2020 | DP | | | |
| | 9060940 | Oct 30, 2020 | | U-1556 | | |
| | 9084816 | Aug 24, 2027 | DP | | | |
| | 9095614 | Aug 24, 2027 | | U-1556 | | |
| | 9095615 | Aug 24, 2027 | DP | | | |
| | 9198863 | Oct 30, 2020 | DP | | | |
| | 9205056 | Oct 30, 2020 | DP | | | |
| | 9289391 | Oct 30, 2020 | DP | | | |
| | 9486412 | Aug 24, 2027 | DP | | | |
| | 9486413 | Aug 24, 2027 | DP | | | |
| | 9492389 | Aug 24, 2027 | DP | | | |
| | 9492390 | Aug 24, 2027 | | U-1556 | | |
| | 9492391 | Aug 24, 2027 | | U-1556 | | |
| | 9517236 | Oct 30, 2020 | DP | | | |
| | 9545380 | Aug 24, 2027 | | U-1556 | | |
| | 9572779 | Dec 21, 2031 | DP | | | |
| | 9572804 | Oct 30, 2020 | DP | | | |
| | 9669023 | Oct 30, 2020 | DP | | | |
| | 9669024 | Oct 30, 2020 | DP | | | |
| | 9675610 | Jun 16, 2023 | DP | | | |
| | 9675611 | Oct 30, 2020 | | U-1556 | | |
| | 9682077 | Oct 30, 2020 | | U-1556 | | |
| | 9750703 | Dec 21, 2031 | DP | | | |
| | 9763933 | Aug 24, 2027 | DP | | | |
| | 9770416 | Aug 24, 2027 | DP | | | |
| | 9775809 | Aug 24, 2027 | DP | | | |
| | 9861584 | Dec 21, 2031 | DP | | | |
| | 9872837 | Dec 21, 2031 | DP | | | |
| <u>HYDROCODONE BITARTRATE - HYSINGLA</u> | | | | | | |
| N 206627 006 | 10130591 | Nov 20, 2023 | DP | U-1819 | | |
| | 6733783 | Oct 30, 2021 | DP | U-1556 | | |
| | 8309060 | Nov 20, 2023 | DP | U-1556 | | |
| | 8361499 | Oct 30, 2021 | DP | | | |
| | 8529948 | Aug 06, 2022 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>HYDROCODONE BITARTRATE - HYSINGLA</u> | | | | | | |
| N 206627 006 | 8551520 | Oct 30, 2021 | DP | | | |
| | 8647667 | Oct 30, 2021 | DP | | | |
| | 8808740 | Dec 21, 2031 | DP | U-1556 | | |
| | 9056052 | Oct 30, 2020 | DP | | | |
| | 9060940 | Oct 30, 2020 | | U-1556 | | |
| | 9084816 | Aug 24, 2027 | DP | | | |
| | 9095614 | Aug 24, 2027 | | U-1556 | | |
| | 9095615 | Aug 24, 2027 | DP | | | |
| | 9198863 | Oct 30, 2020 | DP | | | |
| | 9205056 | Oct 30, 2020 | DP | | | |
| | 9289391 | Oct 30, 2020 | DP | | | |
| | 9486412 | Aug 24, 2027 | DP | | | |
| | 9486413 | Aug 24, 2027 | DP | | | |
| | 9492389 | Aug 24, 2027 | DP | | | |
| | 9492390 | Aug 24, 2027 | | U-1556 | | |
| | 9492391 | Aug 24, 2027 | | U-1556 | | |
| | 9517236 | Oct 30, 2020 | DP | | | |
| | 9545380 | Aug 24, 2027 | | U-1556 | | |
| | 9572779 | Dec 21, 2031 | DP | | | |
| | 9572804 | Oct 30, 2020 | DP | | | |
| | 9669023 | Oct 30, 2020 | DP | | | |
| | 9669024 | Oct 30, 2020 | DP | | | |
| | 9675610 | Jun 16, 2023 | DP | | | |
| | 9675611 | Oct 30, 2020 | | U-1556 | | |
| | 9682077 | Oct 30, 2020 | | U-1556 | | |
| | 9750703 | Dec 21, 2031 | DP | | | |
| | 9763933 | Aug 24, 2027 | DP | | | |
| | 9770416 | Aug 24, 2027 | DP | | | |
| | 9775809 | Aug 24, 2027 | DP | | | |
| | 9861584 | Dec 21, 2031 | DP | | | |
| | 9872837 | Dec 21, 2031 | DP | | | |
| <u>HYDROCODONE BITARTRATE - HYSINGLA</u> | | | | | | |
| N 206627 007 | 10130591 | Nov 20, 2023 | DP | U-1819 | | |
| | 6733783 | Oct 30, 2021 | DP | U-1556 | | |
| | 8309060 | Nov 20, 2023 | DP | U-1556 | | |
| | 8361499 | Oct 30, 2021 | DP | | | |
| | 8529948 | Aug 06, 2022 | DP | | | |
| | 8551520 | Oct 30, 2021 | DP | | | |
| | 8647667 | Oct 30, 2021 | DP | | | |
| | 8808740 | Dec 21, 2031 | DP | U-1556 | | |
| | 9056052 | Oct 30, 2020 | DP | | | |
| | 9060940 | Oct 30, 2020 | | U-1556 | | |
| | 9084816 | Aug 24, 2027 | DP | | | |
| | 9095614 | Aug 24, 2027 | | U-1556 | | |
| | 9095615 | Aug 24, 2027 | DP | | | |
| | 9198863 | Oct 30, 2020 | DP | | | |
| | 9205056 | Oct 30, 2020 | DP | | | |
| | 9289391 | Oct 30, 2020 | DP | | | |
| | 9486412 | Aug 24, 2027 | DP | | | |
| | 9486413 | Aug 24, 2027 | DP | | | |
| | 9492389 | Aug 24, 2027 | DP | | | |
| | 9492390 | Aug 24, 2027 | | U-1556 | | |
| | 9492391 | Aug 24, 2027 | | U-1556 | | |
| | 9517236 | Oct 30, 2020 | DP | | | |
| | 9545380 | Aug 24, 2027 | | U-1556 | | |
| | 9572779 | Dec 21, 2031 | DP | | | |
| | 9572804 | Oct 30, 2020 | DP | | | |
| | 9669023 | Oct 30, 2020 | DP | | | |
| | 9669024 | Oct 30, 2020 | DP | | | |
| | 9675610 | Jun 16, 2023 | DP | | | |
| | 9675611 | Oct 30, 2020 | | U-1556 | | |
| | 9682077 | Oct 30, 2020 | | U-1556 | | |
| | 9750703 | Dec 21, 2031 | DP | | | |
| | 9763933 | Aug 24, 2027 | DP | | | |
| | 9770416 | Aug 24, 2027 | DP | | | |
| | 9775809 | Aug 24, 2027 | DP | | | |
| | 9861584 | Dec 21, 2031 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>HYDROCODONE BITARTRATE - HYSINGLA</u> | | | | | | |
| N 206627 | 007 | 9872837 | Dec 21, 2031 | DP | | |
| <u>HYDROCODONE BITARTRATE - VANTRELA ER</u> | | | | | | |
| N 207975 | 001 | 8445018 | Jul 31, 2029 | DP | | |
| | | 9216176 | Sep 13, 2027 | DP | | |
| | | 9572803 | Sep 13, 2027 | DP | | |
| <u>HYDROCODONE BITARTRATE - VANTRELA ER</u> | | | | | | |
| N 207975 | 002 | 8445018 | Jul 31, 2029 | DP | | |
| | | 9216176 | Sep 13, 2027 | DP | | |
| | | 9572803 | Sep 13, 2027 | DP | | |
| <u>HYDROCODONE BITARTRATE - VANTRELA ER</u> | | | | | | |
| N 207975 | 003 | 8445018 | Jul 31, 2029 | DP | | |
| | | 9216176 | Sep 13, 2027 | DP | | |
| | | 9572803 | Sep 13, 2027 | DP | | |
| <u>HYDROCODONE BITARTRATE - VANTRELA ER</u> | | | | | | |
| N 207975 | 004 | 8445018 | Jul 31, 2029 | DP | | |
| | | 9216176 | Sep 13, 2027 | DP | | |
| | | 9572803 | Sep 13, 2027 | DP | | |
| <u>HYDROCODONE BITARTRATE - VANTRELA ER</u> | | | | | | |
| N 207975 | 005 | 8445018 | Jul 31, 2029 | DP | | |
| | | 9216176 | Sep 13, 2027 | DP | | |
| | | 9572803 | Sep 13, 2027 | DP | | |
| <u>HYDROCORTISONE BUTYRATE - LOCOID</u> | | | | | | |
| N 022076 | 001 | 7378405 | Dec 19, 2026 | DP | | |
| | | 7981877 | Jan 23, 2025 | DP | | |
| <u>HYDROGEN PEROXIDE - ESKATA</u> | | | | | | |
| N 209305 | 001 | 10098910 | Apr 21, 2035 | DP | U-2205 | |
| | | 7381427 | Jun 08, 2022 | | U-2205 | |
| | | 9675639 | Jul 04, 2035 | DP | U-2205 | |
| | | 9980983 | Apr 21, 2035 | | U-2205 | |
| <u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID-HP</u> | | | | | | |
| N 019034 | 001 | 6589960 | Nov 09, 2020 | DP | | |
| | | 9248229 | Mar 12, 2034 | DP | | |
| | | 9731082 | Apr 23, 2032 | DP | | |
| <u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID-HP</u> | | | | | | |
| N 019034 | 002 | 6589960 | Nov 09, 2020 | DP | | |
| | | 9248229 | Mar 12, 2034 | DP | | |
| | | 9731082 | Apr 23, 2032 | DP | | |
| <u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u> | | | | | | |
| N 019034 | 003 | 6589960 | Nov 09, 2020 | DS DP | | |
| | | 9248229 | Mar 12, 2034 | DP | | |
| | | 9731082 | Apr 23, 2032 | DP | | |
| <u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u> | | | | | | |
| N 019034 | 004 | 6589960 | Nov 09, 2020 | DS DP | | |
| | | 9248229 | Mar 12, 2034 | DP | | |
| | | 9731082 | Apr 23, 2032 | DP | | |
| <u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u> | | | | | | |
| N 019034 | 005 | 6589960 | Nov 09, 2020 | DS DP | | |
| | | 9248229 | Mar 12, 2034 | DP | | |
| | | 9731082 | Apr 23, 2032 | DP | | |
| <u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u> | | | | | | |
| N 019891 | 001 | 6589960 | Nov 09, 2020 | DS DP | | |
| <u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u> | | | | | | |
| N 019892 | 001 | 6589960 | Nov 09, 2020 | DS DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u> | | | | | | |
| N 019892 002 | 6589960 | Nov 09, 2020 | DS DP | | | |
| <u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u> | | | | | | |
| N 019892 003 | 6589960 | Nov 09, 2020 | DS DP | | | |
| <u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u> | | | | | | |
| N 021044 001 | 6589960 | Nov 09, 2020 | DP | | | |
| <u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u> | | | | | | |
| N 021044 002 | 6589960 | Nov 09, 2020 | DP | | | |
| <u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u> | | | | | | |
| N 021044 003 | 6589960 | Nov 09, 2020 | DP | | | |
| <u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u> | | | | | | |
| N 021044 004 | 6589960 | Nov 09, 2020 | DP | | | |
| <u>HYDROXYPROGESTERONE CAPROATE - MAKENA (AUTOINJECTOR)</u> | | | | | | |
| N 021945 004 | 8021335 | Oct 04, 2026 | DP | | | |
| | 8562564 | Jan 24, 2026 | DP | | | |
| | 9180259 | Jan 24, 2026 | DP | | | |
| | 9533102 | Jan 24, 2026 | DP | | | |
| | 9629959 | Jan 24, 2026 | DP | | | |
| | 9789257 | Feb 11, 2034 | DP | | | |
| | 9844558 | May 02, 2036 | | U-2236 | | |
| | RE44846 | Aug 10, 2019 | DP | | | |
| <u>HYDROXYUREA - SIKLOS</u> | | | | | | |
| N 208843 001 | | | | | NP ODE-177 | Dec 21, 2020 Dec 21, 2024 |
| <u>HYDROXYUREA - SIKLOS</u> | | | | | | |
| N 208843 002 | | | | | NP ODE-177 | Dec 21, 2020 Dec 21, 2024 |
| <u>IBANDRONATE SODIUM - BONIVA</u> | | | | | | |
| N 021455 001 | 6294196 | Oct 07, 2019 | DP | | | |
| <u>IBANDRONATE SODIUM - BONIVA</u> | | | | | | |
| N 021455 002 | 6294196 | Oct 07, 2019 | DP | | | |
| | 7192938 | May 06, 2023 | | U-798 | | |
| | 7410957 | May 06, 2023 | | U-887 | | |
| | 7718634 | May 06, 2023 | | U-642 | | |
| <u>IBRUTINIB - IMBRUVICA</u> | | | | | | |
| N 205552 001 | 10004746 | Jun 03, 2031 | | U-1684 | D-165 | May 06, 2019 |
| | 10004746 | Jun 03, 2031 | | U-1946 | I-729 | Mar 04, 2019 |
| | 10004746 | Jun 03, 2031 | | U-2241 | I-736 | May 06, 2019 |
| | 10004746 | Jun 03, 2031 | | U-2242 | I-737 | May 06, 2019 |
| | 10016435 | Jun 03, 2031 | | U-1650 | I-741 | Jan 18, 2020 |
| | 10106548 | Jun 03, 2033 | DS DP | | I-753 | Aug 02, 2020 |
| | 10125140 | Jun 03, 2033 | DS DP | | ODE-109 | Mar 04, 2023 |
| | 7514444 | Dec 28, 2026 | DS DP | | ODE-117 | May 06, 2023 |
| | 8008309 | Dec 28, 2026 | DS DP | | ODE-128 | Jan 18, 2024 |
| | 8476284 | Dec 28, 2026 | | U-1456 | ODE-152 | Aug 02, 2024 |
| | 8476284 | Dec 28, 2026 | | U-1650 | ODE-55 | Nov 13, 2020 |
| | 8476284 | Dec 28, 2026 | | U-1946 | ODE-60 | Feb 12, 2021 |
| | 8476284 | Dec 28, 2026 | | U-1947 | ODE-72 | Jul 28, 2021 |
| | 8497277 | Dec 28, 2026 | | U-1456 | ODE-86 | Jan 29, 2022 |
| | 8497277 | Dec 28, 2026 | | U-1491 | | |
| | 8497277 | Dec 28, 2026 | | U-1650 | | |
| | 8497277 | Dec 28, 2026 | | U-1946 | | |
| | 8497277 | Dec 28, 2026 | | U-1947 | | |
| | 8563563 | Apr 26, 2027 | | U-1491 | | |
| | 8563563 | Apr 26, 2027 | | U-1650 | | |
| | 8563563 | Apr 26, 2027 | | U-1946 | | |
| | 8563563 | Apr 26, 2027 | | U-2219 | | |
| | 8697711 | Dec 28, 2026 | DS DP | | | |
| | 8703780 | Dec 28, 2026 | | U-1491 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>IBRUTINIB - IMBRUVICA</u> | | | | | | |
| N 205552 001 | 8735403 | Dec 28, 2026 | DS DP | | | |
| | 8754090 | Jun 03, 2031 | | U-1456 | | |
| | 8754091 | Dec 28, 2026 | DP | | | |
| | 8952015 | Dec 28, 2026 | | U-1456 | | |
| | 8952015 | Dec 28, 2026 | | U-1491 | | |
| | 8952015 | Dec 28, 2026 | | U-1650 | | |
| | 8952015 | Dec 28, 2026 | | U-1946 | | |
| | 8952015 | Dec 28, 2026 | | U-1947 | | |
| | 8957079 | Dec 28, 2026 | DS DP | | | |
| | 8999999 | Jun 03, 2031 | | U-1683 | | |
| | 8999999 | Jun 03, 2031 | | U-1684 | | |
| | 9125889 | Jun 03, 2031 | | U-1745 | | |
| | 9181257 | Dec 28, 2026 | DS DP | | | |
| | 9296753 | Oct 30, 2033 | DS DP | | | |
| | 9540382 | Aug 18, 2033 | | U-1456 | | |
| | 9540382 | Aug 18, 2033 | | U-1650 | | |
| | 9540382 | Aug 18, 2033 | | U-1684 | | |
| | 9540382 | Aug 18, 2033 | | U-1946 | | |
| | 9540382 | Aug 18, 2033 | | U-1947 | | |
| | 9713617 | Jun 03, 2033 | DP | | | |
| | 9725455 | Jun 03, 2033 | DS | | | |
| | 9795604 | Oct 24, 2034 | | U-2150 | | |
| | 9801881 | Jun 03, 2031 | | U-1491 | | |
| | 9801883 | Jun 03, 2031 | | U-2159 | | |
| | 9814721 | Jun 03, 2031 | | U-1947 | | |
| <u>IBRUTINIB - IMBRUVICA</u> | | | | | | |
| N 205552 002 | 10004746 | Jun 03, 2031 | | U-1684 | | |
| | 10004746 | Jun 03, 2031 | | U-1946 | | |
| | 10004746 | Jun 03, 2031 | | U-2241 | | |
| | 10004746 | Jun 03, 2031 | | U-2242 | | |
| | 10016435 | Jun 03, 2031 | | U-1650 | | |
| | 10106548 | Jun 03, 2033 | DS DP | | | |
| | 10125140 | Jun 03, 2033 | DS DP | | | |
| | 7514444 | Dec 28, 2026 | DS DP | | | |
| | 8008309 | Dec 28, 2026 | DS DP | | | |
| | 8476284 | Dec 28, 2026 | | U-1456 | | |
| | 8476284 | Dec 28, 2026 | | U-1650 | | |
| | 8476284 | Dec 28, 2026 | | U-1946 | | |
| | 8476284 | Dec 28, 2026 | | U-1947 | | |
| | 8497277 | Dec 28, 2026 | | U-1456 | | |
| | 8497277 | Dec 28, 2026 | | U-1491 | | |
| | 8497277 | Dec 28, 2026 | | U-1650 | | |
| | 8497277 | Dec 28, 2026 | | U-1946 | | |
| | 8497277 | Dec 28, 2026 | | U-1947 | | |
| | 8563563 | Apr 26, 2027 | | U-1491 | | |
| | 8563563 | Apr 26, 2027 | | U-1650 | | |
| | 8563563 | Apr 26, 2027 | | U-1946 | | |
| | 8563563 | Apr 26, 2027 | | U-2219 | | |
| | 8697711 | Dec 28, 2026 | DS DP | | | |
| | 8703780 | Dec 28, 2026 | | U-1491 | | |
| | 8735403 | Dec 28, 2026 | DS DP | | | |
| | 8754090 | Jun 03, 2031 | | U-1456 | | |
| | 8754091 | Dec 28, 2026 | DP | | | |
| | 8952015 | Dec 28, 2026 | | U-1456 | | |
| | 8952015 | Dec 28, 2026 | | U-1491 | | |
| | 8952015 | Dec 28, 2026 | | U-1650 | | |
| | 8952015 | Dec 28, 2026 | | U-1946 | | |
| | 8952015 | Dec 28, 2026 | | U-1947 | | |
| | 8957079 | Dec 28, 2026 | DS DP | | | |
| | 8999999 | Jun 03, 2031 | | U-1491 | | |
| | 8999999 | Jun 03, 2031 | | U-1946 | | |
| | 8999999 | Jun 03, 2031 | | U-2228 | | |
| | 9125889 | Jun 03, 2031 | | U-1650 | | |
| | 9181257 | Dec 28, 2026 | DS | | | |
| | 9296753 | Oct 30, 2033 | DS | | | |
| | 9540382 | Aug 18, 2033 | | U-1456 | | |
| | 9540382 | Aug 18, 2033 | | U-1491 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>IBRUTINIB - IMBRUVICA</u> | | | | | | |
| N 205552 | 002 | 9540382 | Aug 18, 2033 | | U-1650 | |
| | | 9540382 | Aug 18, 2033 | | U-1946 | |
| | | 9540382 | Aug 18, 2033 | | U-1947 | |
| | | 9713617 | Jun 03, 2033 | DP | | |
| | | 9725455 | Jun 03, 2033 | DS | | |
| | | 9795604 | Oct 24, 2034 | | U-2150 | |
| | | 9801881 | Jun 03, 2031 | | U-1491 | |
| | | 9801883 | Jun 03, 2031 | | U-2159 | |
| | | 9814721 | Jun 03, 2031 | | U-1947 | |
| <u>IBRUTINIB - IMBRUVICA</u> | | | | | | |
| N 210563 | 001 | 10004746 | Jun 03, 2031 | | U-1684 | |
| | | 10004746 | Jun 03, 2031 | | U-1946 | |
| | | 10004746 | Jun 03, 2031 | | U-2241 | |
| | | 10004746 | Jun 03, 2031 | | U-2242 | |
| | | 10010507 | Mar 03, 2036 | DP | | |
| | | 10016435 | Jun 03, 2031 | | U-1650 | |
| | | 10106548 | Jun 03, 2033 | DS DP | | |
| | | 10125140 | Jun 03, 2033 | DS DP | | |
| | | 7514444 | Dec 28, 2026 | DS DP | | |
| | | 8008309 | Dec 28, 2026 | DS DP | | |
| | | 8476284 | Dec 28, 2026 | | U-1456 | |
| | | 8476284 | Dec 28, 2026 | | U-1650 | |
| | | 8476284 | Dec 28, 2026 | | U-1946 | |
| | | 8476284 | Dec 28, 2026 | | U-1947 | |
| | | 8476284 | Dec 28, 2026 | | U-2241 | |
| | | 8497277 | Dec 28, 2026 | | U-1456 | |
| | | 8497277 | Dec 28, 2026 | | U-1491 | |
| | | 8497277 | Dec 28, 2026 | | U-1650 | |
| | | 8497277 | Dec 28, 2026 | | U-1946 | |
| | | 8497277 | Dec 28, 2026 | | U-1947 | |
| | | 8497277 | Dec 28, 2026 | | U-2241 | |
| | | 8497277 | Dec 28, 2026 | | U-2242 | |
| | | 8563563 | Apr 26, 2027 | | U-1491 | |
| | | 8563563 | Apr 26, 2027 | | U-1650 | |
| | | 8563563 | Apr 26, 2027 | | U-1946 | |
| | | 8563563 | Apr 26, 2027 | | U-2241 | |
| | | 8563563 | Apr 26, 2027 | | U-2242 | |
| | | 8697711 | Dec 28, 2026 | DS DP | | |
| | | 8703780 | Dec 28, 2026 | | U-1491 | |
| | | 8703780 | Dec 28, 2026 | | U-2242 | |
| | | 8735403 | Dec 28, 2026 | DS DP | | |
| | | 8754090 | Jun 03, 2031 | | U-1456 | |
| | | 8754091 | Dec 28, 2026 | DP | | |
| | | 8952015 | Dec 28, 2026 | | U-1456 | |
| | | 8952015 | Dec 28, 2026 | | U-1491 | |
| | | 8952015 | Dec 28, 2026 | | U-1650 | |
| | | 8952015 | Dec 28, 2026 | | U-1946 | |
| | | 8952015 | Dec 28, 2026 | | U-1947 | |
| | | 8952015 | Dec 28, 2026 | | U-2241 | |
| | | 8952015 | Dec 28, 2026 | | U-2242 | |
| | | 8957079 | Dec 28, 2026 | DS DP | | |
| | | 8999999 | Jun 03, 2031 | | U-1491 | |
| | | 8999999 | Jun 03, 2031 | | U-1946 | |
| | | 8999999 | Jun 03, 2031 | | U-2241 | |
| | | 8999999 | Jun 03, 2031 | | U-2242 | |
| | | 9125889 | Jun 03, 2031 | | U-1650 | |
| | | 9181257 | Dec 28, 2026 | DS | | |
| | | 9296753 | Oct 30, 2033 | DS | | |
| | | 9655857 | Mar 03, 2036 | DP | | |
| | | 9725455 | Jun 03, 2033 | DS | | |
| | | 9795604 | Oct 24, 2034 | | U-2150 | |
| | | 9801881 | Jun 03, 2031 | | U-1491 | |
| | | 9801881 | Jun 03, 2031 | | U-2242 | |
| | | 9801883 | Jun 03, 2031 | | U-2159 | |
| | | 9801883 | Jun 03, 2031 | | U-2243 | |
| | | 9814721 | Jun 03, 2031 | | U-1947 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>IBRUTINIB - IMBRUVICA</u> | | | | | | |
| N 210563 002 | 10004746 | Jun 03, 2031 | | | | U-1684 |
| | 10004746 | Jun 03, 2031 | | | | U-1946 |
| | 10004746 | Jun 03, 2031 | | | | U-2241 |
| | 10004746 | Jun 03, 2031 | | | | U-2242 |
| | 10010507 | Mar 03, 2036 | | DP | | |
| | 10016435 | Jun 03, 2031 | | | | U-1650 |
| | 10106548 | Jun 03, 2033 | DS DP | | | |
| | 10125140 | Jun 03, 2033 | DS DP | | | |
| | 7514444 | Dec 28, 2026 | DS DP | | | |
| | 8008309 | Dec 28, 2026 | DS DP | | | |
| | 8476284 | Dec 28, 2026 | | | | U-1456 |
| | 8476284 | Dec 28, 2026 | | | | U-1650 |
| | 8476284 | Dec 28, 2026 | | | | U-1946 |
| | 8476284 | Dec 28, 2026 | | | | U-1947 |
| | 8476284 | Dec 28, 2026 | | | | U-2241 |
| | 8497277 | Dec 28, 2026 | | | | U-1456 |
| | 8497277 | Dec 28, 2026 | | | | U-1491 |
| | 8497277 | Dec 28, 2026 | | | | U-1650 |
| | 8497277 | Dec 28, 2026 | | | | U-1946 |
| | 8497277 | Dec 28, 2026 | | | | U-1947 |
| | 8497277 | Dec 28, 2026 | | | | U-2241 |
| | 8497277 | Dec 28, 2026 | | | | U-2242 |
| | 8563563 | Apr 26, 2027 | | | | U-1491 |
| | 8563563 | Apr 26, 2027 | | | | U-1650 |
| | 8563563 | Apr 26, 2027 | | | | U-1946 |
| | 8563563 | Apr 26, 2027 | | | | U-2241 |
| | 8563563 | Apr 26, 2027 | | | | U-2242 |
| | 8697711 | Dec 28, 2026 | DS DP | | | |
| | 8703780 | Dec 28, 2026 | | | | U-1491 |
| | 8703780 | Dec 28, 2026 | | | | U-2242 |
| | 8735403 | Dec 28, 2026 | DS DP | | | |
| | 8754090 | Jun 03, 2031 | | | | U-1456 |
| | 8754091 | Dec 28, 2026 | | DP | | |
| | 8952015 | Dec 28, 2026 | | | | U-1456 |
| | 8952015 | Dec 28, 2026 | | | | U-1491 |
| | 8952015 | Dec 28, 2026 | | | | U-1650 |
| | 8952015 | Dec 28, 2026 | | | | U-1946 |
| | 8952015 | Dec 28, 2026 | | | | U-1947 |
| | 8952015 | Dec 28, 2026 | | | | U-2241 |
| | 8952015 | Dec 28, 2026 | | | | U-2242 |
| | 8957079 | Dec 28, 2026 | DS DP | | | |
| | 8999999 | Jun 03, 2031 | | | | U-1491 |
| | 8999999 | Jun 03, 2031 | | | | U-1946 |
| | 8999999 | Jun 03, 2031 | | | | U-2241 |
| | 8999999 | Jun 03, 2031 | | | | U-2242 |
| | 9125889 | Jun 03, 2031 | | | | U-1650 |
| | 9181257 | Dec 28, 2026 | DS | | | |
| | 9296753 | Oct 30, 2033 | DS | | | |
| | 9655857 | Mar 03, 2036 | | DP | | |
| | 9725455 | Jun 03, 2033 | DS | | | |
| | 9795604 | Oct 24, 2034 | | | | U-2150 |
| | 9801881 | Jun 03, 2031 | | | | U-1491 |
| | 9801881 | Jun 03, 2031 | | | | U-2242 |
| | 9801883 | Jun 03, 2031 | | | | U-2159 |
| | 9801883 | Jun 03, 2031 | | | | U-2243 |
| | 9814721 | Jun 03, 2031 | | | | U-1947 |
| <u>IBRUTINIB - IMBRUVICA</u> | | | | | | |
| N 210563 003 | 10004746 | Jun 03, 2031 | | | | U-1684 |
| | 10004746 | Jun 03, 2031 | | | | U-1946 |
| | 10004746 | Jun 03, 2031 | | | | U-2241 |
| | 10004746 | Jun 03, 2031 | | | | U-2242 |
| | 10010507 | Mar 03, 2036 | | DP | | |
| | 10016435 | Jun 03, 2031 | | | | U-1650 |
| | 10106548 | Jun 03, 2033 | DS DP | | | |
| | 10125140 | Jun 03, 2033 | DS DP | | | |
| | 7514444 | Dec 28, 2026 | DS DP | | | |
| | 8008309 | Dec 28, 2026 | DS DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>IBRUTINIB - IMBRUVICA</u> | | | | | | |
| N 210563 | 003 | 8476284 | | | | |
| | | 8476284 | | | U-1456 | |
| | | 8476284 | | | U-1650 | |
| | | 8476284 | | | U-1946 | |
| | | 8476284 | | | U-1947 | |
| | | 8476284 | | | U-2241 | |
| | | 8497277 | | | U-1456 | |
| | | 8497277 | | | U-1491 | |
| | | 8497277 | | | U-1650 | |
| | | 8497277 | | | U-1946 | |
| | | 8497277 | | | U-1947 | |
| | | 8497277 | | | U-2241 | |
| | | 8497277 | | | U-2242 | |
| | | 8563563 | | | U-1491 | |
| | | 8563563 | | | U-1650 | |
| | | 8563563 | | | U-1946 | |
| | | 8563563 | | | U-2241 | |
| | | 8563563 | | | U-2242 | |
| | | 8697711 | | | | |
| | | 8703780 | | DS DP | | |
| | | 8703780 | | | U-1491 | |
| | | 8703780 | | | U-2242 | |
| | | 8735403 | | DS DP | | |
| | | 8754090 | | | U-1456 | |
| | | 8754091 | | DP | | |
| | | 8952015 | | | U-1456 | |
| | | 8952015 | | | U-1491 | |
| | | 8952015 | | | U-1650 | |
| | | 8952015 | | | U-1946 | |
| | | 8952015 | | | U-1947 | |
| | | 8952015 | | | U-2241 | |
| | | 8952015 | | | U-2242 | |
| | | 8957079 | | DS DP | | |
| | | 8999999 | | | U-1491 | |
| | | 8999999 | | | U-1946 | |
| | | 8999999 | | | U-2241 | |
| | | 8999999 | | | U-2242 | |
| | | 9125889 | | | U-1650 | |
| | | 9181257 | | DS | | |
| | | 9296753 | | DS | | |
| | | 9655857 | | DP | | |
| | | 9725455 | | DS | | |
| | | 9795604 | | | U-2150 | |
| | | 9801881 | | | U-1491 | |
| | | 9801881 | | | U-2242 | |
| | | 9801883 | | | U-2159 | |
| | | 9801883 | | | U-2243 | |
| | | 9814721 | | | U-1947 | |
| <u>IBRUTINIB - IMBRUVICA</u> | | | | | | |
| N 210563 | 004 | 10004746 | | | | |
| | | 10004746 | | | U-1684 | |
| | | 10004746 | | | U-1946 | |
| | | 10004746 | | | U-2241 | |
| | | 10004746 | | | U-2242 | |
| | | 10010507 | | DP | | |
| | | 10016435 | | | U-1650 | |
| | | 10106548 | | DS DP | | |
| | | 10125140 | | DS DP | | |
| | | 7514444 | | DS DP | | |
| | | 8008309 | | DS DP | | |
| | | 8476284 | | | U-1456 | |
| | | 8476284 | | | U-1650 | |
| | | 8476284 | | | U-1946 | |
| | | 8476284 | | | U-1947 | |
| | | 8476284 | | | U-2241 | |
| | | 8497277 | | | U-1456 | |
| | | 8497277 | | | U-1491 | |
| | | 8497277 | | | U-1650 | |
| | | 8497277 | | | U-1946 | |
| | | 8497277 | | | U-1947 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>IBRUTINIB - IMBRUVICA</u> | | | | | | |
| N 210563 | 004 | 8497277 | Dec 28, 2026 | U-2241 | | |
| | | 8497277 | Dec 28, 2026 | U-2242 | | |
| | | 8563563 | Apr 26, 2027 | U-1491 | | |
| | | 8563563 | Apr 26, 2027 | U-1650 | | |
| | | 8563563 | Apr 26, 2027 | U-1946 | | |
| | | 8563563 | Apr 26, 2027 | U-2241 | | |
| | | 8563563 | Apr 26, 2027 | U-2242 | | |
| | | 8697711 | Dec 28, 2026 | DS DP | | |
| | | 8703780 | Dec 28, 2026 | U-1491 | | |
| | | 8703780 | Dec 28, 2026 | U-2242 | | |
| | | 8735403 | Dec 28, 2026 | DS DP | | |
| | | 8754090 | Jun 03, 2031 | U-1456 | | |
| | | 8754091 | Dec 28, 2026 | DP | | |
| | | 8952015 | Dec 28, 2026 | U-1456 | | |
| | | 8952015 | Dec 28, 2026 | U-1491 | | |
| | | 8952015 | Dec 28, 2026 | U-1650 | | |
| | | 8952015 | Dec 28, 2026 | U-1946 | | |
| | | 8952015 | Dec 28, 2026 | U-1947 | | |
| | | 8952015 | Dec 28, 2026 | U-2241 | | |
| | | 8952015 | Dec 28, 2026 | U-2242 | | |
| | | 8957079 | Dec 28, 2026 | DS DP | | |
| | | 8999999 | Jun 03, 2031 | U-1491 | | |
| | | 8999999 | Jun 03, 2031 | U-1946 | | |
| | | 8999999 | Jun 03, 2031 | U-2241 | | |
| | | 8999999 | Jun 03, 2031 | U-2242 | | |
| | | 9125889 | Jun 03, 2031 | U-1650 | | |
| | | 9181257 | Dec 28, 2026 | DS | | |
| | | 9296753 | Oct 30, 2033 | DS | | |
| | | 9655857 | Mar 03, 2036 | DP | | |
| | | 9725455 | Jun 03, 2033 | DS | | |
| | | 9795604 | Oct 24, 2034 | U-2150 | | |
| | | 9801881 | Jun 03, 2031 | U-1491 | | |
| | | 9801881 | Jun 03, 2031 | U-2242 | | |
| | | 9801883 | Jun 03, 2031 | U-2159 | | |
| | | 9801883 | Jun 03, 2031 | U-2243 | | |
| | | 9814721 | Jun 03, 2031 | U-1947 | | |
| <u>IBUPROFEN - CALDOLOR</u> | | | | | | |
| N 022348 | 001 | 6727286 | Nov 27, 2021 | DP U-981 | | |
| <u>IBUPROFEN - CALDOLOR</u> | | | | | | |
| N 022348 | 002 | 6727286 | Nov 27, 2021 | DP U-981 | | |
| | | 8735452 | Sep 30, 2029 | U-981 | | |
| | | 8871810 | Sep 30, 2029 | U-1599 | | |
| | | 9012508 | Sep 14, 2030 | U-981 | | |
| | | 9072661 | Mar 16, 2032 | U-2264 | | |
| | | 9072710 | Mar 16, 2032 | U-2266 | | |
| | | 9114068 | Sep 30, 2029 | U-1735 | | |
| | | 9138404 | Sep 30, 2029 | U-1756 | | |
| | | 9295639 | Sep 30, 2029 | U-1756 | | |
| | | 9649284 | Sep 30, 2029 | U-2018 | | |
| <u>IBUPROFEN LYSINE - NEOPROFEN</u> | | | | | | |
| N 021903 | 001 | 6342530 | Nov 14, 2020 | DP U-1127 | | |
| | | 6342530 | Nov 14, 2020 | DP U-794 | | |
| | | 6344479 | Mar 20, 2021 | DS DP U-794 | Y | |
| | | 8415337 | Mar 02, 2032 | DS DP | | |
| <u>IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE - CHILDREN'S MOTRIN COLD</u> | | | | | | |
| N 021128 | 001 | 6211246 | Jun 10, 2019 | | | |
| <u>ICATIBANT ACETATE - FIRAZYR</u> | | | | | | |
| N 022150 | 001 | 5648333 | Jul 15, 2019 | DS DP U-1187 | | |
| <u>ICOSAPENT ETHYL - VASCEPA</u> | | | | | | |
| N 202057 | 001 | 8188146 | Jan 27, 2020 | DS DP | | |
| | | 8293727 | Feb 09, 2030 | U-1287 | | |
| | | 8293728 | Feb 09, 2030 | U-1287 | | |
| | | 8298554 | Apr 29, 2030 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ICOSAPENT ETHYL - VASCEPA</u> | | | | | | |
| N 202057 001 | 8314086 | Feb 09, 2030 | U-1287 | | | |
| | 8318715 | Feb 09, 2030 | U-1287 | | | |
| | 8357677 | Feb 09, 2030 | U-1287 | | | |
| | 8367652 | Feb 09, 2030 | U-1287 | | | |
| | 8377920 | Feb 09, 2030 | U-1287 | | | |
| | 8399446 | Feb 09, 2030 | U-1287 | | | |
| | 8415335 | Feb 09, 2030 | U-1287 | | | |
| | 8426399 | Feb 09, 2030 | U-1287 | | | |
| | 8431560 | Feb 09, 2030 | U-1287 | | | |
| | 8440650 | Feb 09, 2030 | U-1287 | | | |
| | 8445003 | Apr 29, 2030 | U-1287 | | | |
| | 8445013 | Apr 29, 2030 | U-1287 | | | |
| | 8501225 | Apr 29, 2030 | U-1287 | | | |
| | 8518929 | Apr 29, 2030 | U-1287 | | | |
| | 8524698 | Apr 29, 2030 | U-1287 | | | |
| | 8546372 | Apr 29, 2030 | U-1287 | | | |
| | 8551521 | Apr 29, 2030 | U-1287 | | | |
| | 8563608 | Apr 29, 2030 | U-1287 | | | |
| | 8617593 | Apr 29, 2030 | U-1478 | | | |
| | 8617594 | Apr 29, 2030 | U-1287 | | | |
| | 8623406 | Apr 29, 2030 | U-1478 | | | |
| <u>ICOSAPENT ETHYL - VASCEPA</u> | | | | | | |
| N 202057 002 | 8188146 | Jan 27, 2020 | DS DP | | | |
| | 8293727 | Feb 09, 2030 | U-1287 | | | |
| | 8293728 | Feb 09, 2030 | U-1287 | | | |
| | 8298554 | Apr 29, 2030 | DP | | | |
| | 8314086 | Feb 09, 2030 | U-1287 | | | |
| | 8318715 | Feb 09, 2030 | U-1287 | | | |
| | 8357677 | Feb 09, 2030 | U-1287 | | | |
| | 8367652 | Feb 09, 2030 | U-1287 | | | |
| | 8377920 | Feb 09, 2030 | U-1287 | | | |
| | 8399446 | Feb 09, 2030 | U-1287 | | | |
| | 8415335 | Feb 09, 2030 | U-1287 | | | |
| | 8426399 | Feb 09, 2030 | U-1287 | | | |
| | 8440650 | Feb 09, 2030 | U-1287 | | | |
| | 8445003 | Apr 29, 2030 | U-1287 | | | |
| | 8445013 | Apr 29, 2030 | U-1287 | | | |
| | 8501225 | Apr 29, 2030 | U-1287 | | | |
| | 8518929 | Apr 29, 2030 | U-1287 | | | |
| | 8524698 | Apr 29, 2030 | U-1287 | | | |
| | 8546372 | Apr 29, 2030 | U-1287 | | | |
| | 8551521 | Apr 29, 2030 | U-1287 | | | |
| | 8563608 | Apr 29, 2030 | U-1287 | | | |
| | 8617593 | Apr 29, 2030 | U-1287 | | | |
| | 8617594 | Apr 29, 2030 | U-1287 | | | |
| | 8623406 | Apr 29, 2030 | U-1287 | | | |
| <u>IDELALISIB - ZYDELIG</u> | | | | | | |
| N 205858 001 | 6800620 | Apr 24, 2021 | DS U-1560 | | NCE | Jul 23, 2019 |
| | 6949535 | Apr 24, 2021 | DS U-1560 | | ODE-70 | Jul 23, 2021 |
| | 8138195 | Apr 24, 2021 | DS DP U-1549 | | ODE-71 | Jul 23, 2021 |
| | 8492389 | Apr 24, 2021 | DS DP | | | |
| | 8637533 | Apr 24, 2021 | DS DP | | | |
| | 8865730 | Mar 05, 2033 | DS DP U-1615 | | | |
| | 8980901 | May 12, 2025 | U-1678 | | | |
| | 9149477 | May 12, 2025 | U-1757 | | | |
| | 9469643 | Sep 02, 2033 | DS | | | |
| | 9492449 | Mar 11, 2030 | U-1914 | | | |
| | RE44599 | Jul 21, 2025 | U-1558 | | | |
| | RE44599 | Jul 21, 2025 | U-1615 | | | |
| | RE44638 | Aug 05, 2025 | DS DP | | | |
| <u>IDELALISIB - ZYDELIG</u> | | | | | | |
| N 205858 002 | 6800620 | Apr 24, 2021 | DS U-1560 | | NCE | Jul 23, 2019 |
| | 6949535 | Apr 24, 2021 | DS U-1560 | | ODE-70 | Jul 23, 2021 |
| | 8138195 | Apr 24, 2021 | DS DP U-1549 | | ODE-71 | Jul 23, 2021 |
| | 8492389 | Apr 24, 2021 | DS DP | | | |
| | 8637533 | Apr 24, 2021 | DS DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>IDELALISIB - ZYDELIG</u> | | | | | | |
| N 205858 | 002 | 8865730 | Mar 05, 2033 | DS DP U-1615 | | |
| | | 8980901 | May 12, 2025 | U-1678 | | |
| | | 9149477 | May 12, 2025 | U-1757 | | |
| | | 9469643 | Sep 02, 2033 | DS | | |
| | | 9492449 | Mar 11, 2030 | U-1914 | | |
| | | RE44599 | Jul 21, 2025 | U-1558 | | |
| | | RE44599 | Jul 21, 2025 | U-1615 | | |
| | | RE44638 | Aug 05, 2025 | DS DP | | |
| <u>ILOPERIDONE - FANAPT</u> | | | | | | |
| N 022192 | 001 | 8586610 | Nov 02, 2027 | U-1625 | M-180 | May 26, 2019 |
| | | 8652776 | Aug 31, 2030 | U-1685 | | |
| | | 8999638 | Oct 28, 2030 | U-1674 | | |
| | | 9072742 | Jan 16, 2031 | U-1674 | | |
| | | 9074254 | Dec 28, 2031 | U-1674 | | |
| | | 9074255 | Dec 17, 2030 | U-1674 | | |
| | | 9074256 | Feb 10, 2031 | U-1674 | | |
| | | 9138432 | Sep 30, 2025 | U-1737 | | |
| | | 9157121 | Apr 05, 2030 | U-1674 | | |
| <u>ILOPERIDONE - FANAPT</u> | | | | | | |
| N 022192 | 002 | 8586610 | Nov 02, 2027 | U-1625 | M-180 | May 26, 2019 |
| | | 8652776 | Aug 31, 2030 | U-1685 | | |
| | | 8999638 | Oct 28, 2030 | U-1674 | | |
| | | 9072742 | Jan 16, 2031 | U-1674 | | |
| | | 9074254 | Dec 28, 2031 | U-1674 | | |
| | | 9074255 | Dec 17, 2030 | U-1674 | | |
| | | 9074256 | Feb 10, 2031 | U-1674 | | |
| | | 9138432 | Sep 30, 2025 | U-1737 | | |
| | | 9157121 | Apr 05, 2030 | U-1674 | | |
| <u>ILOPERIDONE - FANAPT</u> | | | | | | |
| N 022192 | 003 | 8586610 | Nov 02, 2027 | U-1625 | M-180 | May 26, 2019 |
| | | 8652776 | Aug 31, 2030 | U-1685 | | |
| | | 8999638 | Oct 28, 2030 | U-1674 | | |
| | | 9072742 | Jan 16, 2031 | U-1674 | | |
| | | 9074254 | Dec 28, 2031 | U-1674 | | |
| | | 9074255 | Dec 17, 2030 | U-1674 | | |
| | | 9074256 | Feb 10, 2031 | U-1674 | | |
| | | 9138432 | Sep 30, 2025 | U-1737 | | |
| | | 9157121 | Apr 05, 2030 | U-1674 | | |
| <u>ILOPERIDONE - FANAPT</u> | | | | | | |
| N 022192 | 004 | 8586610 | Nov 02, 2027 | U-1625 | M-180 | May 26, 2019 |
| | | 8652776 | Aug 31, 2030 | U-1685 | | |
| | | 8999638 | Oct 28, 2030 | U-1674 | | |
| | | 9072742 | Jan 16, 2031 | U-1674 | | |
| | | 9074254 | Dec 28, 2031 | U-1674 | | |
| | | 9074255 | Dec 17, 2030 | U-1674 | | |
| | | 9074256 | Feb 10, 2031 | U-1674 | | |
| | | 9138432 | Sep 30, 2025 | U-1737 | | |
| | | 9157121 | Apr 05, 2030 | U-1674 | | |
| <u>ILOPERIDONE - FANAPT</u> | | | | | | |
| N 022192 | 005 | 8586610 | Nov 02, 2027 | U-1625 | M-180 | May 26, 2019 |
| | | 8652776 | Aug 31, 2030 | U-1685 | | |
| | | 8999638 | Oct 28, 2030 | U-1674 | | |
| | | 9072742 | Jan 16, 2031 | U-1674 | | |
| | | 9074254 | Dec 28, 2031 | U-1674 | | |
| | | 9074255 | Dec 17, 2030 | U-1674 | | |
| | | 9074256 | Feb 10, 2031 | U-1674 | | |
| | | 9138432 | Sep 30, 2025 | U-1737 | | |
| | | 9157121 | Apr 05, 2030 | U-1674 | | |
| <u>ILOPERIDONE - FANAPT</u> | | | | | | |
| N 022192 | 006 | 8586610 | Nov 02, 2027 | U-1625 | M-180 | May 26, 2019 |
| | | 8652776 | Aug 31, 2030 | U-1685 | | |
| | | 8999638 | Oct 28, 2030 | U-1674 | | |
| | | 9072742 | Jan 16, 2031 | U-1674 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ILOPERIDONE - FANAPT</u> | | | | | | |
| N 022192 | 006 | 9074254 | Dec 28, 2031 | U-1674 | | |
| | | 9074255 | Dec 17, 2030 | U-1674 | | |
| | | 9074256 | Feb 10, 2031 | U-1674 | | |
| | | 9138432 | Sep 30, 2025 | U-1737 | | |
| | | 9157121 | Apr 05, 2030 | U-1674 | | |
| <u>ILOPERIDONE - FANAPT</u> | | | | | | |
| N 022192 | 007 | 8586610 | Nov 02, 2027 | U-1625 | M-180 | May 26, 2019 |
| | | 8652776 | Aug 31, 2030 | U-1685 | | |
| | | 8999638 | Oct 28, 2030 | U-1674 | | |
| | | 9072742 | Jan 16, 2031 | U-1674 | | |
| | | 9074254 | Dec 28, 2031 | U-1674 | | |
| | | 9074255 | Dec 17, 2030 | U-1674 | | |
| | | 9074256 | Feb 10, 2031 | U-1674 | | |
| | | 9138432 | Sep 30, 2025 | U-1737 | | |
| | | 9157121 | Apr 05, 2030 | U-1674 | | |
| <u>IMATINIB MESYLATE - GLEEVEC</u> | | | | | | |
| N 021335 | 001 | 6894051 | May 23, 2019 | DS DP U-649 | | |
| | | 6958335 | Dec 19, 2021 | U-791 | | |
| | | RE43932*PED | Jan 16, 2019 | | | |
| <u>IMATINIB MESYLATE - GLEEVEC</u> | | | | | | |
| N 021335 | 002 | 6894051 | May 23, 2019 | DS DP U-649 | | |
| | | 6958335 | Dec 19, 2021 | U-791 | | |
| | | RE43932*PED | Jan 16, 2019 | | | |
| <u>IMATINIB MESYLATE - GLEEVEC</u> | | | | | | |
| N 021588 | 001 | 6894051 | May 23, 2019 | DS DP U-649 | ODE-40 | Jan 25, 2020 |
| | | 6958335 | Dec 19, 2021 | U-1883 | | |
| | | 6958335 | Dec 19, 2021 | U-791 | | |
| | | 6958335*PED | Jun 19, 2022 | | | |
| | | RE43932*PED | Jan 16, 2019 | | | |
| <u>IMATINIB MESYLATE - GLEEVEC</u> | | | | | | |
| N 021588 | 002 | 6894051 | May 23, 2019 | DS DP U-649 | ODE-40 | Jan 25, 2020 |
| | | 6958335 | Dec 19, 2021 | U-1883 | | |
| | | 6958335 | Dec 19, 2021 | U-791 | | |
| | | 6958335*PED | Jun 19, 2022 | | | |
| | | RE43932*PED | Jan 16, 2019 | | | |
| <u>IMIQUIMOD - ALDARA</u> | | | | | | |
| N 020723 | 001 | 7696159 | Apr 01, 2024 | DS U-1047 | | |
| | | 7696159 | Apr 01, 2024 | DS U-1048 | | |
| <u>IMIQUIMOD - ZYCLARA</u> | | | | | | |
| N 022483 | 001 | 8236816 | Dec 11, 2029 | U-68 | | |
| | | 8299109 | Dec 11, 2029 | U-68 | | |
| | | 8598196 | Aug 18, 2029 | U-1455 | | |
| | | 8598196 | Aug 18, 2029 | U-172 | | |
| <u>IMIQUIMOD - ZYCLARA</u> | | | | | | |
| N 022483 | 002 | 8222270 | Dec 11, 2029 | U-68 | | |
| <u>INDACATEROL MALEATE - ARCAPTA NEOHALER</u> | | | | | | |
| N 022383 | 001 | 6878721 | Feb 25, 2025 | DS DP U-1168 | | |
| | | 8067437 | Jun 02, 2020 | U-1168 | | |
| | | 8479730 | Oct 11, 2028 | DP | | |
| | | 8658673 | Jun 02, 2020 | DS DP U-1168 | | |
| | | 8796307 | Jun 02, 2020 | DS DP | | |
| <u>INDINAVIR SULFATE - CRIXIVAN</u> | | | | | | |
| N 020685 | 001 | 6689761 | Feb 10, 2021 | U-554 | | |
| <u>INDINAVIR SULFATE - CRIXIVAN</u> | | | | | | |
| N 020685 | 003 | 6689761 | Feb 10, 2021 | U-554 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>INDINAVIR SULFATE - CRIXIVAN</u> | | | | | | |
| N 020685 | 005 | 6689761 | Feb 10, 2021 | U-554 | | |
| <u>INDINAVIR SULFATE - CRIXIVAN</u> | | | | | | |
| N 020685 | 006 | 6689761 | Feb 10, 2021 | U-554 | | |
| <u>INDOCYANINE GREEN - SPY AGENT GREEN KIT</u> | | | | | | |
| N 211580 | 001 | 6915154 | Aug 01, 2021 | U-2460 | NP | Nov 21, 2021 |
| | | 7881777 | Sep 22, 2021 | U-2461 | | |
| | | 8185176 | Jun 04, 2028 | U-2462 | | |
| | | 8406860 | Apr 09, 2029 | U-2463 | | |
| | | 8647605 | Feb 11, 2029 | U-2464 | | |
| | | 8647605 | Feb 11, 2029 | U-2468 | | |
| | | 8892190 | Aug 11, 2020 | U-2465 | | |
| | | 9421280 | Nov 24, 2025 | U-2466 | | |
| | | 9421280 | Nov 24, 2025 | U-2467 | | |
| <u>INDOMETHACIN - TIVORBEX</u> | | | | | | |
| N 204768 | 001 | 8734847 | Apr 23, 2030 | DP | | |
| | | 8992982 | Apr 23, 2030 | DP | | |
| | | 9089471 | Apr 23, 2030 | U-55 | | |
| <u>INDOMETHACIN - TIVORBEX</u> | | | | | | |
| N 204768 | 002 | 8734847 | Apr 23, 2030 | DP | | |
| | | 8992982 | Apr 23, 2030 | DP | | |
| | | 9089471 | Apr 23, 2030 | U-55 | | |
| <u>INGENOL MEBUTATE - PICATO</u> | | | | | | |
| N 202833 | 001 | 7410656 | Oct 10, 2020 | U-1222 | | |
| | | 8278292 | Jul 06, 2027 | DP | | |
| | | 8372827 | Dec 18, 2026 | DP | | |
| | | 8372828 | Dec 18, 2026 | DP | | |
| | | 8377919 | Dec 18, 2026 | DP | | |
| | | 8536163 | Dec 18, 2026 | U-1440 | | |
| | | 8716271 | Dec 18, 2026 | U-1440 | | |
| | | 8735375 | Dec 18, 2026 | U-1440 | | |
| | | 9789078 | May 15, 2033 | U-2138 | | |
| | | 9820959 | Dec 18, 2026 | DP U-1440 | | |
| | | 9833428 | Dec 18, 2026 | DP | | |
| | | 9833429 | Dec 18, 2026 | DP | | |
| | | 9861603 | Dec 18, 2026 | U-1440 | | |
| <u>INGENOL MEBUTATE - PICATO</u> | | | | | | |
| N 202833 | 002 | 7410656 | Oct 10, 2020 | U-1222 | | |
| | | 8278292 | Jul 06, 2027 | DP | | |
| | | 8372827 | Dec 18, 2026 | DP | | |
| | | 8372828 | Dec 18, 2026 | DP | | |
| | | 8377919 | Dec 18, 2026 | DP | | |
| | | 8536163 | Dec 18, 2026 | U-1440 | | |
| | | 8716271 | Dec 18, 2026 | U-1440 | | |
| | | 8735375 | Dec 18, 2026 | U-1440 | | |
| | | 9820959 | Dec 18, 2026 | DP U-1440 | | |
| | | 9833428 | Dec 18, 2026 | DP | | |
| | | 9833429 | Dec 18, 2026 | DP | | |
| | | 9861603 | Dec 18, 2026 | U-1440 | | |
| <u>INOTERSEN SODIUM - TEGSEDI</u> | | | | | | |
| N 211172 | 001 | 7015315 | Mar 21, 2023 | DS | NCE | Oct 05, 2023 |
| | | 7101993 | Sep 05, 2023 | DS | ODE-212 | Oct 05, 2025 |
| | | 8101743 | Apr 01, 2025 | DS DP | | |
| | | 8697860 | Apr 29, 2031 | DP | | |
| | | 9061044 | Apr 29, 2031 | DS | | |
| | | 9399774 | Apr 29, 2031 | U-2430 | | |
| <u>INSULIN ASPART - FIASP</u> | | | | | | |
| N 208751 | 001 | 8324157 | Jun 25, 2030 | DP | NP | Sep 29, 2020 |
| <u>INSULIN ASPART - FIASP FLEXTOUCH</u> | | | | | | |
| N 208751 | 002 | 6899699 | Jan 02, 2022 | DP | NP | Sep 29, 2020 |
| | | 7686786 | Aug 03, 2026 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>INSULIN ASPART - FIASP FLEXTOUCH</u> | | | | | | |
| N 208751 002 | 8324157 | Jun 25, 2030 | DP | | | |
| | 8672898 | Jan 02, 2022 | DP | | | |
| | 8684969 | Oct 20, 2025 | DP | | | |
| | 8920383 | Jul 17, 2026 | DP | | | |
| | 9108002 | Jan 20, 2026 | DP | | | |
| | 9132239 | Feb 01, 2032 | DP | | | |
| | 9457154 | Jan 20, 2026 | DP | | | |
| | 9486588 | Jan 02, 2022 | DP | | | |
| | 9616180 | Jan 20, 2026 | DP | | | |
| | 9687611 | Feb 27, 2027 | DP | | | |
| | 9775953 | Jul 17, 2026 | DP | | | |
| | 9861757 | Jan 20, 2026 | DP | | | |
| <u>INSULIN ASPART PROTAMINE RECOMBINANT; INSULIN ASPART RECOMBINANT - NOVOLOG MIX 70/30 FLEXPEN</u> | | | | | | |
| N 021172 004 | 6004297 | Jan 28, 2019 | DP | | | |
| | 9265893 | Sep 23, 2032 | DP | | | |
| | RE41956 | Jan 21, 2021 | DP | | | |
| | RE43834 | Jan 28, 2019 | DP | | | |
| <u>INSULIN ASPART RECOMBINANT - NOVOLOG FLEXPEN</u> | | | | | | |
| N 020986 003 | 6004297 | Jan 28, 2019 | DP | | | |
| | 9265893 | Sep 23, 2032 | DP | | | |
| | RE41956 | Jan 21, 2021 | DP | | | |
| | RE43834 | Jan 28, 2019 | DP | | | |
| <u>INSULIN ASPART RECOMBINANT - NOVOLOG INNOLET</u> | | | | | | |
| N 020986 004 | RE41956 | Jan 21, 2021 | DP | | | |
| <u>INSULIN ASPART RECOMBINANT - NOVOLOG FLEXTOUCH</u> | | | | | | |
| N 020986 005 | 6899699 | Jan 02, 2022 | DP | | | |
| | 8672898 | Jan 02, 2022 | DP | | | |
| | 8684969 | Oct 20, 2025 | DP | | | |
| | 8920383 | Jul 17, 2026 | DP | | | |
| | 9108002 | Jan 20, 2026 | DP | | | |
| | 9132239 | Feb 01, 2032 | DP | | | |
| | 9457154 | Sep 27, 2027 | DP | | | |
| | 9486588 | Jan 02, 2022 | DP | | | |
| | 9616180 | Jan 20, 2026 | DP | | | |
| | 9687611 | Feb 27, 2027 | DP | | | |
| | 9775953 | Jul 17, 2026 | DP | | | |
| | 9861757 | Jan 20, 2026 | DP | | | |
| | RE46363 | Aug 03, 2026 | DP | | | |
| <u>INSULIN ASPART; INSULIN DEGLUDEC - RYZODEG 70/30</u> | | | | | | |
| N 203313 001 | 6899699 | Jan 02, 2022 | DP | | NCE | Sep 25, 2020 |
| | 7615532 | Jun 28, 2029 | DS DP | | NPP | Dec 16, 2019 |
| | 8672898 | Jan 02, 2022 | DP | | | |
| | 8684969 | Oct 20, 2025 | DP | | | |
| | 8920383 | Jul 17, 2026 | DP | | | |
| | 9108002 | Jan 20, 2026 | DP | | | |
| | 9132239 | Feb 01, 2032 | DP | | | |
| | 9457154 | Sep 27, 2027 | DP | | | |
| | 9486588 | Jan 02, 2022 | DP | | | |
| | 9616180 | Jan 20, 2026 | DP | | | |
| | 9687611 | Feb 27, 2027 | DP | | | |
| | 9775953 | Jul 17, 2026 | DP | | | |
| | 9861757 | Jan 20, 2026 | DP | | | |
| | 9884094 | May 01, 2033 | DP | U-2238 | | |
| | RE46363 | Aug 03, 2026 | DP | | | |
| <u>INSULIN DEGLUDEC - TRESIBA</u> | | | | | | |
| N 203314 001 | 6899699 | Jan 02, 2022 | DP | | NCE | Sep 25, 2020 |
| | 7615532 | Jun 28, 2029 | DS DP | | NPP | Dec 16, 2019 |
| | 8672898 | Jan 02, 2022 | DP | | | |
| | 8684969 | Oct 20, 2025 | DP | | | |
| | 8920383 | Jul 17, 2026 | DP | | | |
| | 9108002 | Jan 20, 2026 | DP | | | |
| | 9132239 | Feb 01, 2032 | DP | | | |
| | 9457154 | Sep 27, 2027 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>INSULIN DEGLUDEC - TRESIBA</u> | | | | | | |
| N 203314 001 | 9486588 | Jan 02, 2022 | DP | | | |
| | 9616180 | Jan 20, 2026 | DP | | | |
| | 9687611 | Feb 27, 2027 | DP | | | |
| | 9775953 | Jul 17, 2026 | DP | | | |
| | 9861757 | Jan 20, 2026 | DP | | | |
| | RE46363 | Aug 03, 2026 | DP | | | |
| <u>INSULIN DEGLUDEC - TRESIBA</u> | | | | | | |
| N 203314 002 | 6899699 | Jan 02, 2022 | DP | | NCE | Sep 25, 2020 |
| | 7615532 | Jun 28, 2029 | DS DP | | NFP | Dec 16, 2019 |
| | 8672898 | Jan 02, 2022 | DP | | | |
| | 8684969 | Oct 20, 2025 | DP | | | |
| | 8920383 | Jul 17, 2026 | DP | | | |
| | 9108002 | Jan 20, 2026 | DP | | | |
| | 9132239 | Feb 01, 2032 | DP | | | |
| | 9457154 | Sep 27, 2027 | DP | | | |
| | 9486588 | Jan 02, 2022 | DP | | | |
| | 9616180 | Jan 20, 2026 | DP | | | |
| | 9687611 | Feb 27, 2027 | DP | | | |
| | 9775953 | Jul 17, 2026 | DP | | | |
| | 9861757 | Jan 20, 2026 | DP | | | |
| | RE46363 | Aug 03, 2026 | DP | | | |
| <u>INSULIN DEGLUDEC; LIRAGLUTIDE - XULTOPHY 100/3.6</u> | | | | | | |
| N 208583 001 | 6268343 | Aug 22, 2022 | DS DP | | NC | Nov 21, 2019 |
| | 6899699 | Jan 02, 2022 | DP | | NCE | Sep 25, 2020 |
| | 7615532 | Jun 28, 2029 | DS DP | | | |
| | 8672898 | Jan 02, 2022 | DP | | | |
| | 8684969 | Oct 20, 2025 | DP | | | |
| | 8846618 | Jun 27, 2022 | DS DP | | | |
| | 8920383 | Jul 17, 2026 | DP | | | |
| | 8937042 | May 05, 2029 | DP | | | |
| | 9108002 | Jan 20, 2026 | DP | | | |
| | 9132239 | Feb 01, 2032 | DP | | | |
| | 9457154 | Sep 27, 2027 | DP | | | |
| | 9486588 | Jan 02, 2022 | DP | | | |
| | 9616180 | Jan 20, 2026 | DP | | | |
| | 9687611 | Feb 27, 2027 | DP | | | |
| | 9775953 | Jul 17, 2026 | DP | | | |
| | 9861757 | Jan 20, 2026 | DP | | | |
| | RE46363 | Aug 03, 2026 | DP | | | |
| <u>INSULIN DETEMIR RECOMBINANT - LEVEMIR</u> | | | | | | |
| N 021536 001 | 5750497 | Jun 16, 2019 | DS DP U-668 | | | |
| <u>INSULIN DETEMIR RECOMBINANT - LEVEMIR FLEXPEN</u> | | | | | | |
| N 021536 002 | 5750497 | Jun 16, 2019 | DS DP U-668 | | | |
| | 6004297 | Jan 28, 2019 | DP | | | |
| | 9265893 | Sep 23, 2032 | DP | | | |
| | RE41956 | Jan 21, 2021 | DP | | | |
| | RE43834 | Jan 28, 2019 | DP | | | |
| <u>INSULIN DETEMIR RECOMBINANT - LEVEMIR INNOLET</u> | | | | | | |
| N 021536 003 | 5750497 | Jun 16, 2019 | DS DP U-668 | | | |
| <u>INSULIN DETEMIR RECOMBINANT - LEVEMIR PENFILL</u> | | | | | | |
| N 021536 004 | 5750497 | Jun 16, 2019 | DS DP U-668 | | | |
| <u>INSULIN DETEMIR RECOMBINANT - LEVEMIR FLEXTOUCH</u> | | | | | | |
| N 021536 005 | 5750497 | Jun 16, 2019 | DS DP U-668 | | | |
| | 6899699 | Jan 02, 2022 | DP | | | |
| | 7686786 | Aug 03, 2026 | DP | | | |
| | 8672898 | Jan 02, 2022 | DP | | | |
| | 8684969 | Oct 20, 2025 | DP | | | |
| | 8920383 | Jul 17, 2026 | DP | | | |
| | 9108002 | Jan 20, 2026 | DP | | | |
| | 9132239 | Feb 01, 2032 | DP | | | |
| | 9457154 | Sep 27, 2027 | DP | | | |
| | 9486588 | Jan 02, 2022 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>INSULIN DETEMIR RECOMBINANT - LEVEMIR FLEXTOUCH</u> | | | | | | |
| N 021536 005 | 9616180 | Jan 20, 2026 | DP | | | |
| | 9687611 | Feb 27, 2027 | DP | | | |
| | 9775953 | Jul 17, 2026 | DP | | | |
| | 9861757 | Jan 20, 2026 | DP | | | |
| | RE46363 | Aug 03, 2026 | DP | | | |
| <u>INSULIN GLARGINE RECOMBINANT - LANTUS</u> | | | | | | |
| N 021081 001 | 7476652 | Jul 23, 2023 | DP | | | |
| | 7713930 | Jun 13, 2023 | DP | | | |
| | 7918833 | Sep 23, 2027 | DP | | | |
| <u>INSULIN GLARGINE RECOMBINANT - LANTUS SOLOSTAR</u> | | | | | | |
| N 021081 002 | 8512297 | Sep 15, 2024 | DP | | | |
| | 8556864 | Mar 03, 2024 | DP | | | |
| | 8603044 | Mar 02, 2024 | DP | | | |
| | 8679069 | Apr 12, 2025 | DP | | | |
| | 8992486 | Jun 05, 2024 | DP | | | |
| | 9011391 | Mar 26, 2024 | | U-1832 | | |
| | 9233211 | Mar 02, 2024 | DP | | | |
| | 9408979 | Mar 02, 2024 | DP | | | |
| | 9526844 | Mar 02, 2024 | DP | | | |
| | 9533105 | Aug 17, 2024 | DP | | | |
| | 9561331 | Aug 28, 2024 | DP | | | |
| | 9604008 | Mar 02, 2024 | DP | | | |
| | 9604009 | Aug 16, 2024 | DP | | | |
| | 9610409 | Mar 02, 2024 | DP | | | |
| | 9623189 | Aug 19, 2024 | DP | | | |
| | 9717852 | Apr 08, 2033 | DP | | | |
| | 9775954 | Mar 02, 2024 | DP | | | |
| | 9827379 | Mar 02, 2024 | DP | U-2146 | | |
| <u>INSULIN GLARGINE RECOMBINANT - TOUJEO SOLOSTAR</u> | | | | | | |
| N 206538 001 | 7918833 | Sep 23, 2027 | DP | | | |
| | 7918833*PED | Mar 23, 2028 | | | | |
| | 8512297 | Sep 15, 2024 | DP | | | |
| | 8556864 | Mar 03, 2024 | DP | | | |
| | 8603044 | Mar 02, 2024 | DP | | | |
| | 8679069 | Apr 12, 2025 | DP | | | |
| | 8992486 | Jun 05, 2024 | DP | | | |
| | 9011391 | Mar 26, 2024 | | U-1832 | | |
| | 9233211 | Mar 02, 2024 | DP | | | |
| | 9345750 | May 18, 2031 | DP | U-1855 | | |
| | 9408979 | Mar 02, 2024 | DP | | | |
| | 9526844 | Mar 02, 2024 | DP | | | |
| | 9533105 | Aug 17, 2024 | DP | | | |
| | 9561331 | Aug 28, 2024 | DP | | | |
| | 9604008 | Mar 02, 2024 | DP | | | |
| | 9604009 | Aug 16, 2024 | DP | | | |
| | 9610409 | Mar 02, 2024 | DP | | | |
| | 9623189 | Aug 19, 2024 | DP | | | |
| | 9775954 | Mar 02, 2024 | DP | | | |
| | 9827379 | Mar 02, 2024 | DP | U-2146 | | |
| <u>INSULIN GLARGINE RECOMBINANT - TOUJEO MAX SOLOSTAR</u> | | | | | | |
| N 206538 002 | 7918833 | Sep 23, 2027 | DP | | | |
| | 8512297 | Sep 15, 2024 | DP | | | |
| | 8556864 | Mar 03, 2024 | DP | | | |
| | 8603044 | Mar 02, 2024 | DP | | | |
| | 8679069 | Apr 12, 2025 | DP | | | |
| | 8992486 | Jun 05, 2024 | DP | | | |
| | 9011391 | Mar 26, 2024 | DP | U-1832 | | |
| | 9233211 | Mar 02, 2024 | DP | | | |
| | 9345750 | May 18, 2031 | DP | U-1855 | | |
| | 9408979 | Mar 02, 2024 | DP | | | |
| | 9526844 | Mar 02, 2024 | DP | | | |
| | 9533105 | Aug 17, 2024 | DP | | | |
| | 9561331 | Aug 28, 2024 | DP | | | |
| | 9604008 | Mar 02, 2024 | DP | | | |
| | 9604009 | Aug 16, 2024 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>INSULIN GLARGINE RECOMBINANT - TOUJEO MAX SOLOSTAR</u> | | | | | | |
| N 206538 | 002 | 9610409 | Mar 02, 2024 | DP | | |
| | | 9623189 | Aug 19, 2024 | DP | | |
| | | 9775954 | Mar 02, 2024 | DP | | |
| | | 9827379 | Mar 02, 2024 | DP | U-2146 | |
| <u>INSULIN GLARGINE; LIXISENATIDE - SOLIOUA 100/33</u> | | | | | | |
| N 208673 | 001 | 10029011 | Aug 02, 2032 | DP | NC | Nov 21, 2019 |
| | | 10117909 | Oct 09, 2029 | DP | NCE | Jul 27, 2021 |
| | | 7918833 | Sep 23, 2027 | DP | | |
| | | 8512297 | Sep 15, 2024 | DP | | |
| | | 8556864 | Mar 03, 2024 | DP | | |
| | | 8603044 | Mar 02, 2024 | DP | | |
| | | 8679069 | Apr 12, 2025 | DP | | |
| | | 8992486 | Jun 05, 2024 | DP | | |
| | | 9011391 | Mar 26, 2024 | | U-1923 | |
| | | 9233211 | Mar 02, 2024 | DP | | |
| | | 9408979 | Mar 02, 2024 | DP | | |
| | | 9526764 | Oct 09, 2029 | DP | | |
| | | 9526844 | Mar 02, 2024 | DP | | |
| | | 9533105 | Aug 17, 2024 | DP | | |
| | | 9561331 | Aug 28, 2024 | DP | | |
| | | 9604008 | Mar 02, 2024 | DP | | |
| | | 9604009 | Aug 16, 2024 | DP | | |
| | | 9610409 | Mar 02, 2024 | DP | | |
| | | 9623189 | Aug 19, 2024 | DP | | |
| | | 9707176 | Nov 11, 2030 | DP | | |
| | | 9717852 | Apr 08, 2033 | DP | | |
| | | 9775954 | Mar 02, 2024 | DP | | |
| | | 9821032 | May 09, 2032 | | U-2182 | |
| | | 9827379 | Mar 02, 2024 | DP | U-2146 | |
| | | 9950039 | Dec 10, 2035 | | U-2277 | |
| | | 9950039 | Dec 10, 2035 | | U-2278 | |
| | | 9950039 | Dec 10, 2035 | | U-2279 | |
| | | RE45313 | Jul 12, 2020 | DS DP | | |
| <u>INSULIN GLULISINE RECOMBINANT - APIDRA</u> | | | | | | |
| N 021629 | 001 | 6960561 | Jan 25, 2023 | DP | U-471 | |
| | | 7452860 | Mar 22, 2022 | DP | | |
| | | 7696162 | Mar 22, 2022 | DP | U-471 | |
| <u>INSULIN GLULISINE RECOMBINANT - APIDRA</u> | | | | | | |
| N 021629 | 002 | 6960561 | Jan 25, 2023 | DP | U-471 | |
| | | 7452860 | Mar 22, 2022 | DP | | |
| | | 7696162 | Mar 22, 2022 | DP | U-471 | |
| <u>INSULIN GLULISINE RECOMBINANT - APIDRA SOLOSTAR</u> | | | | | | |
| N 021629 | 003 | 6960561 | Jan 25, 2023 | DP | U-471 | |
| | | 7452860 | Mar 22, 2022 | DP | | |
| | | 7696162 | Mar 22, 2022 | DP | U-471 | |
| | | 7918833 | Sep 23, 2027 | DP | | |
| | | 8512297 | Sep 15, 2024 | DP | | |
| | | 8556864 | Mar 03, 2024 | DP | | |
| | | 8603044 | Mar 02, 2024 | DP | | |
| | | 8679069 | Apr 12, 2025 | DP | | |
| | | 8992486 | Jun 05, 2024 | DP | | |
| | | 9011391 | Mar 26, 2024 | | U-1832 | |
| | | 9233211 | Mar 02, 2024 | DP | | |
| | | 9408979 | Mar 02, 2024 | DP | | |
| | | 9526844 | Mar 02, 2024 | DP | | |
| | | 9533105 | Aug 17, 2024 | DP | | |
| | | 9561331 | Aug 28, 2024 | DP | | |
| | | 9604008 | Mar 02, 2024 | DP | | |
| | | 9604009 | Aug 16, 2024 | DP | | |
| | | 9610409 | Mar 02, 2024 | DP | | |
| | | 9623189 | Aug 19, 2024 | DP | | |
| | | 9717852 | Apr 08, 2033 | DP | | |
| | | 9775954 | Mar 02, 2024 | DP | | |
| | | 9827379 | Mar 02, 2024 | DP | U-2146 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>INSULIN HUMAN - HUMULIN R</u> | | | | | | |
| N 018780 | 004 | 7291132 | Aug 09, 2024 | DP | | |
| <u>INSULIN LISPRO - ADMELOG</u> | | | | | | |
| N 209196 | 001 | | | | NP | Dec 11, 2020 |
| <u>INSULIN LISPRO - ADMELOG SOLOSTAR</u> | | | | | | |
| N 209196 | 002 | 7918833 | Sep 23, 2027 | DP | NP | Dec 11, 2020 |
| | | 8512297 | Sep 15, 2024 | DP | | |
| | | 8556864 | Mar 03, 2024 | DP | | |
| | | 8603044 | Mar 02, 2024 | DP | | |
| | | 8679069 | Apr 12, 2025 | DP | | |
| | | 8992486 | Jun 05, 2024 | DP | | |
| | | 9011391 | Mar 26, 2024 | DP | U-1832 | |
| | | 9233211 | Mar 02, 2024 | DP | | |
| | | 9408979 | Mar 02, 2024 | DP | | |
| | | 9526844 | Mar 02, 2024 | DP | | |
| | | 9533105 | Aug 17, 2024 | DP | | |
| | | 9561331 | Aug 28, 2024 | DP | | |
| | | 9604008 | Mar 02, 2024 | DP | | |
| | | 9604009 | Aug 16, 2024 | DP | | |
| | | 9610409 | Mar 02, 2024 | DP | | |
| | | 9623189 | Aug 19, 2024 | DP | | |
| | | 9717852 | Apr 08, 2033 | DP | | |
| | | 9775954 | Mar 02, 2024 | DP | | |
| | | 9827379 | Mar 04, 2024 | DP | U-2146 | |
| <u>INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT - HUMALOG MIX 75/25 KWIKPEN</u> | | | | | | |
| N 021017 | 002 | 7291132 | Aug 09, 2024 | DP | | |
| <u>INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT - HUMALOG MIX 50/50 KWIKPEN</u> | | | | | | |
| N 021018 | 002 | 7291132 | Aug 09, 2024 | DP | | |
| <u>INSULIN LISPRO RECOMBINANT - HUMALOG KWIKPEN</u> | | | | | | |
| N 020563 | 003 | 7291132 | Aug 09, 2024 | DP | | |
| <u>INSULIN LISPRO RECOMBINANT - HUMALOG KWIKPEN</u> | | | | | | |
| N 205747 | 001 | 7291132 | Aug 09, 2024 | DP | | |
| <u>INSULIN RECOMBINANT HUMAN - EXUBERA</u> | | | | | | |
| N 021868 | 001 | 6257233 | May 14, 2019 | U-704 | | |
| | | 6546929 | May 14, 2019 | U-704 | | |
| | | 6582728 | Jun 24, 2020 | DP | | |
| <u>INSULIN RECOMBINANT HUMAN - EXUBERA</u> | | | | | | |
| N 021868 | 002 | 6257233 | May 14, 2019 | U-704 | | |
| | | 6546929 | May 14, 2019 | U-704 | | |
| | | 6582728 | Jun 24, 2020 | DP | | |
| <u>INSULIN RECOMBINANT HUMAN - AFREZZA</u> | | | | | | |
| N 022472 | 001 | 10046031 | Aug 11, 2029 | U-2383 | | |
| | | 6444226 | Jun 29, 2020 | DP | U-1534 | |
| | | 6652885 | Jun 29, 2020 | U-1535 | | |
| | | 7305986 | Jan 16, 2023 | DP | | |
| | | 7464706 | Mar 02, 2023 | DP | | |
| | | 7648960 | Jun 29, 2020 | U-1535 | | |
| | | 7943178 | Jun 29, 2020 | DP | U-1535 | |
| | | 7943572 | Aug 10, 2026 | U-1539 | | |
| | | 8119593 | Aug 11, 2029 | U-1537 | | |
| | | 8146588 | Apr 24, 2023 | DP | | |
| | | 8156936 | Jan 16, 2023 | DP | | |
| | | 8215300 | Nov 24, 2022 | DP | | |
| | | 8258095 | Aug 11, 2029 | U-1537 | | |
| | | 8389470 | Jun 29, 2020 | DP | U-1621 | |
| | | 8424518 | Oct 17, 2031 | DP | | |
| | | 8485180 | Mar 25, 2030 | DP | | |
| | | 8499757 | Feb 19, 2032 | DP | | |
| | | 8551528 | Jun 11, 2030 | DP | | |
| | | 8623817 | Sep 18, 2029 | U-1537 | | |
| | | 8636001 | Jul 12, 2032 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>INSULIN RECOMBINANT HUMAN - AFREZZA</u> | | | | | | |
| N 022472 001 | 8729019 | Dec 26, 2028 | DP | | | |
| | 8734845 | Jun 11, 2030 | DP | | | |
| | 8778403 | Jun 11, 2030 | DP | U-1538 | | |
| | 8889099 | Jun 29, 2020 | DP | U-1621 | | |
| | 8912193 | Jun 12, 2029 | DP | U-1538 | | |
| | 8950397 | Jul 20, 2021 | DP | | | |
| | 9192675 | Jun 12, 2029 | DP | U-1788 | | |
| | 9283193 | Sep 14, 2026 | DP | | | |
| | 9339615 | Oct 20, 2029 | DP | | | |
| | 9358352 | Feb 15, 2031 | DP | U-1861 | | |
| | 9393372 | Jul 04, 2029 | DP | | | |
| | 9446133 | Jun 12, 2029 | DP | U-1861 | | |
| | 9511198 | Feb 16, 2030 | | U-1929 | | |
| | 9511198 | Feb 16, 2030 | | U-1930 | | |
| | 9597374 | Oct 08, 2031 | | U-1987 | | |
| | 9662461 | Jun 12, 2029 | DP | U-2019 | | |
| | 9717689 | Sep 14, 2026 | DP | | | |
| | 9943571 | Aug 11, 2029 | | U-1537 | | |
| <u>INSULIN RECOMBINANT HUMAN - AFREZZA</u> | | | | | | |
| N 022472 002 | 10046031 | Aug 11, 2029 | | U-2383 | | |
| | 6444226 | Jun 29, 2020 | DP | U-1534 | | |
| | 6652885 | Jun 29, 2020 | | U-1535 | | |
| | 7305986 | Jan 16, 2023 | DP | | | |
| | 7464706 | Mar 02, 2023 | DP | | | |
| | 7648960 | Jun 29, 2020 | | U-1535 | | |
| | 7943178 | Jun 29, 2020 | DP | U-1535 | | |
| | 7943572 | Aug 10, 2026 | | U-1539 | | |
| | 8119593 | Aug 11, 2029 | | U-1537 | | |
| | 8146588 | Apr 24, 2023 | DP | | | |
| | 8156936 | Jan 16, 2023 | DP | | | |
| | 8215300 | Nov 24, 2022 | DP | | | |
| | 8258095 | Aug 11, 2029 | | U-1537 | | |
| | 8389470 | Jun 29, 2020 | DP | U-1621 | | |
| | 8424518 | Oct 17, 2031 | DP | | | |
| | 8485180 | Mar 25, 2030 | DP | | | |
| | 8499757 | Feb 19, 2032 | DP | | | |
| | 8551528 | Jun 11, 2030 | DP | | | |
| | 8623817 | Sep 18, 2029 | | U-1537 | | |
| | 8636001 | Jul 12, 2032 | DP | | | |
| | 8729019 | Dec 26, 2028 | DP | | | |
| | 8734845 | Jun 11, 2030 | DP | | | |
| | 8778403 | Jun 11, 2030 | DP | U-1538 | | |
| | 8889099 | Jun 29, 2020 | DP | U-1621 | | |
| | 8912193 | Jun 12, 2029 | DP | U-1538 | | |
| | 8950397 | Jul 20, 2021 | DP | | | |
| | 9192675 | Jun 12, 2029 | DP | U-1788 | | |
| | 9283193 | Sep 14, 2026 | DP | | | |
| | 9339615 | Oct 20, 2029 | DP | | | |
| | 9358352 | Feb 15, 2031 | DP | U-1861 | | |
| | 9393372 | Jul 04, 2029 | DP | | | |
| | 9446133 | Jun 12, 2029 | DP | U-1861 | | |
| | 9511198 | Feb 16, 2030 | | U-1929 | | |
| | 9511198 | Feb 16, 2030 | | U-1930 | | |
| | 9597374 | Oct 08, 2031 | | U-1987 | | |
| | 9662461 | Jun 12, 2029 | DP | U-2019 | | |
| | 9717689 | Sep 14, 2026 | DP | | | |
| | 9943571 | Aug 11, 2029 | | U-1537 | | |
| <u>INSULIN RECOMBINANT HUMAN - AFREZZA</u> | | | | | | |
| N 022472 003 | 10046031 | Aug 11, 2029 | | U-2383 | | |
| | 6444226 | Jun 29, 2020 | DP | U-1534 | | |
| | 6652885 | Jun 29, 2020 | | U-1535 | | |
| | 7305986 | Jan 16, 2023 | DP | | | |
| | 7464706 | Mar 02, 2023 | DP | | | |
| | 7648960 | Jun 29, 2020 | | U-1535 | | |
| | 7943178 | Jun 29, 2020 | DP | U-1535 | | |
| | 7943572 | Aug 10, 2026 | | U-1539 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>INSULIN RECOMBINANT HUMAN - AFREZZA</u> | | | | | | |
| N 022472 003 | 8119593 | Aug 11, 2029 | | U-1537 | | |
| | 8146588 | Apr 24, 2023 | DP | | | |
| | 8156936 | Jan 16, 2023 | DP | | | |
| | 8215300 | Nov 24, 2022 | DP | | | |
| | 8258095 | Aug 11, 2029 | | U-1537 | | |
| | 8389470 | Jun 29, 2020 | DP | U-1621 | | |
| | 8424518 | Oct 17, 2031 | DP | | | |
| | 8485180 | Mar 25, 2030 | DP | | | |
| | 8499757 | Feb 19, 2032 | DP | | | |
| | 8551528 | Jun 11, 2030 | DP | | | |
| | 8623817 | Sep 18, 2029 | | U-1537 | | |
| | 8636001 | Jul 12, 2032 | DP | | | |
| | 8729019 | Dec 26, 2028 | DP | | | |
| | 8734845 | Jun 11, 2030 | DP | | | |
| | 8778403 | Jun 11, 2030 | DP | U-1538 | | |
| | 8889099 | Jun 29, 2020 | DP | U-1621 | | |
| | 8912193 | Jun 12, 2029 | DP | U-1538 | | |
| | 8950397 | Jul 20, 2021 | DP | | | |
| | 9192675 | Jun 12, 2029 | DP | U-1788 | | |
| | 9283193 | Sep 14, 2026 | DP | | | |
| | 9339615 | Oct 20, 2029 | DP | | | |
| | 9358352 | Feb 15, 2031 | DP | U-1861 | | |
| | 9393372 | Jul 04, 2029 | DP | | | |
| | 9446133 | Jun 12, 2029 | DP | U-1861 | | |
| | 9511198 | Feb 16, 2030 | | U-1929 | | |
| | 9511198 | Feb 16, 2030 | | U-1930 | | |
| | 9597374 | Oct 08, 2031 | | U-1987 | | |
| | 9662461 | Jun 12, 2029 | DP | U-2019 | | |
| | 9717689 | Sep 14, 2026 | DP | | | |
| | 9943571 | Aug 11, 2029 | | U-1537 | | |
| <u>INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN - HUMULIN 70/30</u> | | | | | | |
| N 019717 001 | 7291132 | Aug 09, 2024 | DP | | | |
| <u>INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN - HUMULIN 70/30 PEN</u> | | | | | | |
| N 019717 002 | 7291132 | Aug 09, 2024 | DP | | | |
| <u>INSULIN SUSP ISOPHANE RECOMBINANT HUMAN - HUMULIN N</u> | | | | | | |
| N 018781 001 | 7291132 | Aug 09, 2024 | DP | | | |
| <u>ILOBENGUANE I-131 - AZEDRA</u> | | | | | | |
| N 209607 001 | | | | | NP ODE-204 | Jul 30, 2021 Jul 30, 2025 |
| <u>IODIXANOL - VISIPAQUE 320</u> | | | | | | |
| N 020351 002 | | | | | I-752 | Apr 05, 2020 |
| <u>IODIXANOL - VISIPAQUE 320</u> | | | | | | |
| N 020808 002 | | | | | I-752 | Apr 05, 2020 |
| <u>IPRATROPIUM BROMIDE - ATROVENT HFA</u> | | | | | | |
| N 021527 001 | 6739333 | May 26, 2020 | DP | | | |
| | 6983743 | May 26, 2020 | DP | | | |
| | 8474447 | Jan 17, 2030 | DP | | | |
| <u>IRINOTECAN HYDROCHLORIDE - CAMPTOSAR</u> | | | | | | |
| N 020571 001 | 6403569 | Apr 28, 2020 | | U-449 | | |
| | 6794370 | May 01, 2020 | | U-606 | | |
| <u>IRINOTECAN HYDROCHLORIDE - CAMPTOSAR</u> | | | | | | |
| N 020571 002 | 6403569 | Apr 28, 2020 | | U-449 | | |
| | 6794370 | May 01, 2020 | | U-606 | | |
| <u>IRINOTECAN HYDROCHLORIDE - ONIVYDE</u> | | | | | | |
| N 207793 001 | 8147867 | Aug 29, 2028 | DS DP | | ODE-99 | Oct 22, 2022 |
| | 8329213 | May 02, 2025 | DS DP | | | |
| | 8703181 | May 02, 2025 | | U-1434 | | |
| | 8992970 | May 02, 2025 | DS DP | | | |
| | 9339497 | Jun 12, 2033 | | U-1848 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>IRINOTECAN HYDROCHLORIDE - ONIVYDE</u> | | | | | | |
| N 207793 | 001 | 9364473 | Jun 12, 2033 | | U-1856 | |
| | | 9452162 | Jun 12, 2033 | | U-1899 | |
| | | 9492442 | Jun 12, 2033 | | U-1848 | |
| | | 9492442 | Jun 12, 2033 | | U-1899 | |
| | | 9492442 | Jun 12, 2033 | | U-1917 | |
| | | 9717724 | Jun 12, 2033 | | U-1848 | |
| | | 9717724 | Jun 12, 2033 | | U-2091 | |
| | | 9724303 | May 02, 2025 | DS DP | | |
| | | 9730891 | May 02, 2025 | | U-1848 | |
| | | 9782349 | May 02, 2025 | DS DP | | |
| <u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u> | | | | | | |
| N 207500 | 001 | 6812238 | Oct 31, 2020 | DS | NCE | Mar 06, 2020 |
| | | 7459561 | Oct 31, 2020 | DS | ODE-90 | Mar 06, 2022 |
| | | | | | GAIN | Mar 06, 2025 |
| | | | | | GAIN | Mar 06, 2027 |
| <u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u> | | | | | | |
| N 207501 | 001 | 6812238 | Oct 31, 2020 | DS | NCE | Mar 06, 2020 |
| | | 7459561 | Oct 31, 2020 | DS | ODE-90 | Mar 06, 2022 |
| | | | | | GAIN | Mar 06, 2025 |
| | | | | | GAIN | Mar 06, 2027 |
| <u>ISOTRETINOIN - ABSORICA</u> | | | | | | |
| N 021951 | 001 | 7435427 | Sep 21, 2021 | DP | | |
| | | 8367102 | Sep 21, 2021 | | U-1347 | |
| | | 8952064 | Sep 21, 2021 | DP | | |
| | | 9078925 | Sep 21, 2021 | DP | | |
| | | 9089534 | Sep 21, 2021 | DP | | |
| <u>ISOTRETINOIN - ABSORICA</u> | | | | | | |
| N 021951 | 002 | 7435427 | Sep 21, 2021 | DP | | |
| | | 8367102 | Sep 21, 2021 | | U-1347 | |
| | | 8952064 | Sep 21, 2021 | DP | | |
| | | 9078925 | Sep 21, 2021 | DP | | |
| | | 9089534 | Sep 21, 2021 | DP | | |
| <u>ISOTRETINOIN - ABSORICA</u> | | | | | | |
| N 021951 | 003 | 7435427 | Sep 21, 2021 | DP | | |
| | | 8367102 | Sep 21, 2021 | | U-1347 | |
| | | 8952064 | Sep 21, 2021 | DP | | |
| | | 9078925 | Sep 21, 2021 | DP | | |
| | | 9089534 | Sep 21, 2021 | DP | | |
| <u>ISOTRETINOIN - ABSORICA</u> | | | | | | |
| N 021951 | 004 | 7435427 | Sep 21, 2021 | DP | | |
| | | 8367102 | Sep 21, 2021 | | U-1347 | |
| | | 8952064 | Sep 21, 2021 | DP | | |
| | | 9078925 | Sep 21, 2021 | DP | | |
| | | 9089534 | Sep 21, 2021 | DP | | |
| <u>ISOTRETINOIN - ABSORICA</u> | | | | | | |
| N 021951 | 005 | 7435427 | Sep 21, 2021 | DP | | |
| | | 8367102 | Sep 21, 2021 | | U-1347 | |
| | | 8952064 | Sep 21, 2021 | DP | | |
| | | 9078925 | Sep 21, 2021 | DP | | |
| | | 9089534 | Sep 21, 2021 | DP | | |
| <u>ISOTRETINOIN - ABSORICA</u> | | | | | | |
| N 021951 | 006 | 7435427 | Sep 21, 2021 | DP | | |
| | | 8367102 | Sep 21, 2021 | | U-1347 | |
| | | 8952064 | Sep 21, 2021 | DP | | |
| | | 9078925 | Sep 21, 2021 | DP | | |
| | | 9089534 | Sep 21, 2021 | DP | | |
| <u>ITRACONAZOLE - ITRACONAZOLE</u> | | | | | | |
| A 205573 | 001 | | | | PC | Mar 17, 2019 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------------------|--------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ITRACONAZOLE - SPORANOX</u> | | | | | | |
| N 020657 | 001 6407079 | Jun 18, 2019 | | | | |
| <u>ITRACONAZOLE - SPORANOX</u> | | | | | | |
| N 020966 | 001 6407079 | Jun 18, 2019 | | | | |
| <u>ITRACONAZOLE - ONMEL</u> | | | | | | |
| N 022484 | 001 8486456 | Oct 03, 2028 | DP U-1054 | | | |
| <u>ITRACONAZOLE - TOLSURA</u> | | | | | | |
| N 208901 | 001 8771739 | Jul 25, 2023 | DP | | | |
| | 8921374 | Jun 21, 2033 | DP | | | |
| | 9272046 | Jun 21, 2033 | DP | | | |
| | 9713642 | Jun 21, 2033 | U-2453 | | | |
| <u>IVABRADINE HYDROCHLORIDE - CORLANOR</u> | | | | | | |
| N 206143 | 001 7361649 | Apr 17, 2026 | DS DP U-1694 | | NCE | Apr 15, 2020 |
| | 7361650 | Apr 14, 2026 | DS DP U-1694 | | | |
| | 7867996 | Feb 22, 2026 | DS DP U-1694 | | | |
| | 7879842 | Feb 22, 2026 | DS DP U-1694 | | | |
| <u>IVABRADINE HYDROCHLORIDE - CORLANOR</u> | | | | | | |
| N 206143 | 002 7361649 | Apr 17, 2026 | DS DP U-1694 | | NCE | Apr 15, 2020 |
| | 7361650 | Apr 14, 2026 | DS DP U-1694 | | | |
| | 7867996 | Feb 22, 2026 | DS DP U-1694 | | | |
| | 7879842 | Feb 22, 2026 | DS DP U-1694 | | | |
| <u>IVACAFTOR - KALYDECO</u> | | | | | | |
| N 203188 | 001 7495103 | May 20, 2027 | DS DP | | NPP | Jul 31, 2020 |
| | 8324242 | Aug 05, 2027 | U-1311 | | ODE-186 | Feb 21, 2021 |
| | 8324242 | Aug 05, 2027 | U-1906 | | ODE-187 | Dec 29, 2021 |
| | 8354427 | Jul 06, 2026 | U-1311 | | ODE-189 | Jul 31, 2024 |
| | 8354427 | Jul 06, 2026 | U-1905 | | ODE-190 | May 17, 2024 |
| | 8410274 | Dec 28, 2026 | DP | | ODE-199 | Aug 15, 2025 |
| | 8629162 | Jun 24, 2025 | U-2234 | | ODE-20 | Jan 31, 2019 |
| | 8754224 | Dec 28, 2026 | DS DP | | | |
| | 9670163 | Dec 28, 2026 | DP U-1311 | | | |
| <u>IVACAFTOR - KALYDECO</u> | | | | | | |
| N 207925 | 001 7495103 | May 20, 2027 | DS DP | | NPP | Jul 31, 2020 |
| | 8324242 | Aug 05, 2027 | U-1311 | | ODE-188 | Mar 17, 2022 |
| | 8324242 | Aug 05, 2027 | U-1906 | | ODE-189 | Jul 31, 2024 |
| | 8354427 | Jul 06, 2026 | U-1311 | | ODE-190 | May 17, 2024 |
| | 8354427 | Jul 06, 2026 | U-1905 | | ODE-199 | Aug 15, 2025 |
| | 8410274 | Dec 28, 2026 | DP | | ODE-20 | Jan 31, 2019 |
| | 8629162 | Jun 24, 2025 | U-2234 | | | |
| | 8754224 | Dec 28, 2026 | DS DP | | | |
| | 8883206 | Feb 27, 2033 | DP | | | |
| | 9670163 | Dec 28, 2026 | DP U-1311 | | | |
| <u>IVACAFTOR - KALYDECO</u> | | | | | | |
| N 207925 | 002 7495103 | May 20, 2027 | DS DP | | NPP | Jul 31, 2020 |
| | 8324242 | Aug 05, 2027 | U-1311 | | ODE-188 | Mar 17, 2022 |
| | 8324242 | Aug 05, 2027 | U-1906 | | ODE-189 | Jul 31, 2024 |
| | 8354427 | Jul 06, 2026 | U-1311 | | ODE-190 | May 17, 2024 |
| | 8354427 | Jul 06, 2026 | U-1905 | | ODE-199 | Aug 15, 2025 |
| | 8410274 | Dec 28, 2026 | DP | | ODE-20 | Jan 31, 2019 |
| | 8629162 | Jun 24, 2025 | U-2234 | | | |
| | 8754224 | Dec 28, 2026 | DS DP | | | |
| | 8883206 | Feb 27, 2033 | DP | | | |
| | 9670163 | Dec 28, 2026 | DP U-1311 | | | |
| <u>IVACAFTOR; IVACAFTOR, TEZACAFTOR - SYMDEKO (COPACKAGED)</u> | | | | | | |
| N 210491 | 001 10022352 | Apr 09, 2027 | DP U-2343 | | NCE | Feb 12, 2023 |
| | 10058546 | Jul 15, 2033 | U-2399 | | ODE-173 | Feb 12, 2025 |
| | 10081621 | Mar 25, 2031 | DP U-2420 | | | |
| | 7495103 | May 20, 2027 | DS DP | | | |
| | 7645789 | May 01, 2027 | DS DP | | | |
| | 7776905 | Jun 03, 2027 | DS DP | | | |
| | 8324242 | Aug 05, 2027 | U-2246 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------------------|-----------|------------------------|--------------|-------------------------|---------------------|-----------------------------|
| <u>IVACAFTOR; IVACAFTOR, TEZACAFTOR - SYMDEKO (COPACKAGED)</u> | | | | | | |
| N 210491 001 | 8410274 | Dec 28, 2026 | DP | | | |
| | 8415387 | Nov 12, 2027 | | U-2246 | | |
| | 8598181 | May 01, 2027 | | U-2246 | | |
| | 8623905 | May 01, 2027 | DS DP | | | |
| | 8629162 | Jun 24, 2025 | | U-2247 | | |
| | 8754224 | Dec 28, 2026 | DS DP | | | |
| | 9012496 | Jul 15, 2033 | | U-2248 | | |
| | 9670163 | Dec 28, 2026 | DP | U-2246 | | |
| | 9931334 | Dec 28, 2026 | DP | U-2275 | | |
| | 9974781 | Apr 09, 2027 | DP | U-2318 | | |
| <u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u> | | | | | | |
| N 206038 001 | 10076513 | Dec 04, 2028 | DP | U-2411 | M-218 | Jan 25, 2021 |
| | 7495103 | May 20, 2027 | DS DP | | NCE | Jul 02, 2020 |
| | 7973038 | Nov 08, 2026 | | U-1973 | ODE-123 | Sep 28, 2023 |
| | 8324242 | Aug 05, 2027 | | U-1311 | ODE-93 | Jul 02, 2022 |
| | 8324242 | Aug 05, 2027 | | U-1911 | | |
| | 8410274 | Dec 28, 2026 | DP | | | |
| | 8507534 | Sep 20, 2030 | DS DP | | | |
| | 8653103 | Dec 04, 2028 | DP | | | |
| | 8716338 | Sep 20, 2030 | DP | U-1718 | | |
| | 8716338 | Sep 20, 2030 | DP | U-1910 | | |
| | 8741933 | Nov 08, 2026 | | U-1717 | | |
| | 8741933 | Nov 08, 2026 | | U-1909 | | |
| | 8754224 | Dec 28, 2026 | DS DP | | | |
| | 8846718 | Dec 04, 2028 | | U-1717 | | |
| | 8846718 | Dec 04, 2028 | | U-1908 | | |
| | 8993600 | Dec 11, 2030 | DP | | | |
| | 9150552 | Dec 04, 2028 | | U-1908 | | |
| | 9192606 | Sep 29, 2029 | DP | U-1912 | | |
| | 9216969 | Nov 08, 2026 | DS DP | | | |
| | 9670163 | Dec 28, 2026 | DP | U-1911 | | |
| | 9931334 | Dec 28, 2026 | DP | U-2276 | | |
| <u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u> | | | | | | |
| N 206038 002 | 7495103 | May 20, 2027 | DS DP | | M-218 | Jan 25, 2021 |
| | 7973038 | Nov 08, 2026 | | U-1973 | NCE | Jul 02, 2020 |
| | 8324242 | Aug 05, 2027 | | U-1911 | NPP | Sep 28, 2019 |
| | 8410274 | Dec 28, 2026 | DP | | ODE-123 | Sep 28, 2023 |
| | 8507534 | Sep 20, 2030 | DS DP | | ODE-93 | Jul 02, 2022 |
| | 8653103 | Dec 04, 2028 | DP | | | |
| | 8716338 | Sep 20, 2030 | DP | U-1910 | | |
| | 8741933 | Nov 08, 2026 | | U-1909 | | |
| | 8754224 | Dec 28, 2026 | DS DP | | | |
| | 8846718 | Dec 04, 2028 | | U-1908 | | |
| | 8993600 | Dec 11, 2030 | DP | | | |
| | 9150552 | Dec 04, 2028 | | U-1908 | | |
| | 9192606 | Sep 29, 2029 | DP | U-1912 | | |
| | 9216969 | Nov 08, 2026 | DP | | | |
| | 9670163 | Dec 28, 2026 | DP | U-1911 | | |
| | 9931334 | Dec 28, 2026 | DP | U-2276 | | |
| <u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u> | | | | | | |
| N 211358 001 | 7495103 | May 20, 2027 | DS DP | | NCE | Jul 02, 2020 |
| | 7973038 | Nov 08, 2026 | | U-2374 | NP | Aug 07, 2021 |
| | 8324242 | Aug 05, 2027 | | U-2374 | ODE-195 | Aug 07, 2025 |
| | 8410274 | Dec 28, 2026 | DP | | | |
| | 8507534 | Sep 20, 2030 | DS DP | | | |
| | 8653103 | Dec 04, 2028 | DP | | | |
| | 8716338 | Sep 20, 2030 | DP | U-2396 | | |
| | 8741933 | Nov 08, 2026 | | U-2374 | | |
| | 8754224 | Dec 28, 2026 | DS DP | | | |
| | 8846718 | Dec 04, 2028 | | U-2375 | | |
| | 8993600 | Dec 11, 2030 | DP | | | |
| | 9150552 | Dec 04, 2028 | | U-2375 | | |
| | 9192606 | Sep 29, 2029 | DP | U-2397 | | |
| | 9216969 | Nov 08, 2026 | DP | | | |
| | 9670163 | Dec 28, 2026 | DP | U-2376 | | |
| | 9931334 | Dec 28, 2026 | DP | U-2376 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u> | | | | | | |
| N 211358 001 | 7495103 | May 20, 2027 | DS DP | | NCE | Jul 02, 2020 |
| | 7973038 | Nov 08, 2026 | | U-2374 | NP | Aug 07, 2021 |
| | 8324242 | Aug 05, 2027 | | U-2374 | ODE-195 | Aug 07, 2025 |
| | 8410274 | Dec 28, 2026 | | DP | | |
| | 8507534 | Sep 20, 2030 | DS DP | | | |
| | 8653103 | Dec 04, 2028 | | DP | | |
| | 8716338 | Sep 20, 2030 | | DP U-2396 | | |
| | 8741933 | Nov 08, 2026 | | U-2374 | | |
| | 8754224 | Dec 28, 2026 | DS DP | | | |
| | 8846718 | Dec 04, 2028 | | U-2375 | | |
| | 8993600 | Dec 11, 2030 | | DP | | |
| | 9150552 | Dec 04, 2028 | | U-2375 | | |
| | 9192606 | Sep 29, 2029 | | DP U-2397 | | |
| | 9216969 | Nov 08, 2026 | | DP | | |
| | 9670163 | Dec 28, 2026 | | DP U-2376 | | |
| | 9931334 | Dec 28, 2026 | | DP U-2376 | | |
| <u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u> | | | | | | |
| N 211358 002 | 7495103 | May 20, 2027 | DS DP | | NCE | Jul 02, 2020 |
| | 7973038 | Nov 08, 2026 | | U-2374 | NP | Aug 07, 2021 |
| | 8324242 | Aug 05, 2027 | | U-2374 | ODE-195 | Aug 07, 2025 |
| | 8410274 | Dec 28, 2026 | | DP | | |
| | 8507534 | Sep 20, 2030 | DS DP | | | |
| | 8653103 | Dec 04, 2028 | | DP | | |
| | 8716338 | Sep 20, 2030 | | DP U-2396 | | |
| | 8741933 | Nov 08, 2026 | | U-2374 | | |
| | 8754224 | Dec 28, 2026 | DS DP | | | |
| | 8846718 | Dec 04, 2028 | | U-2375 | | |
| | 8993600 | Dec 11, 2030 | | DP | | |
| | 9150552 | Dec 04, 2028 | | U-2375 | | |
| | 9192606 | Sep 29, 2029 | | DP U-2397 | | |
| | 9216969 | Nov 08, 2026 | | DP | | |
| | 9670163 | Dec 28, 2026 | | DP U-2376 | | |
| | 9931334 | Dec 28, 2026 | | DP U-2376 | | |
| <u>IVERMECTIN - SKLICE</u> | | | | | | |
| N 202736 001 | 8791153 | Oct 12, 2027 | | DP | | |
| | 8927595 | Oct 12, 2027 | | U-1782 | | |
| <u>IVERMECTIN - SOOLANTRA</u> | | | | | | |
| N 206255 001 | 6133310 | Apr 26, 2019 | | U-1631 | | |
| | 7550440 | Apr 22, 2024 | | DP U-1631 | | |
| | 8080530 | Apr 22, 2024 | | DP U-1631 | | |
| | 8093219 | Apr 22, 2024 | | DP U-1631 | | |
| | 8415311 | Apr 22, 2024 | | DP U-1631 | | |
| | 8470788 | Apr 22, 2024 | | DP U-1631 | | |
| | 8815816 | Apr 22, 2024 | | DP U-1631 | | |
| | 9089587 | Mar 13, 2034 | | U-1631 | | |
| | 9233117 | Mar 13, 2034 | | U-1631 | | |
| | 9233118 | Mar 13, 2034 | | U-1631 | | |
| | 9782425 | Mar 13, 2034 | | U-1631 | | |
| <u>IVOSIDENIB - TIBSOVO</u> | | | | | | |
| N 211192 001 | 9474779 | Aug 19, 2033 | DS DP | U-2350 | NCE | Jul 20, 2023 |
| | 9850277 | Jan 18, 2033 | DS DP | U-2350 | ODE-203 | Jul 20, 2025 |
| | 9968595 | Mar 13, 2035 | | DP U-2351 | | |
| <u>IXABEPILONE - IXEMPRA KIT</u> | | | | | | |
| N 022065 001 | 6670384 | Jan 23, 2022 | | DP U-959 | | |
| | 6670384 | Jan 23, 2022 | | DP U-960 | | |
| | 7022330 | Jan 23, 2022 | | DP U-958 | | |
| | 7312237 | Aug 21, 2024 | | U-965 | | |
| | RE41393 | Feb 08, 2022 | | U-961 | | |
| | RE41911 | Sep 28, 2020 | DS DP | U-961 | | |
| <u>IXABEPILONE - IXEMPRA KIT</u> | | | | | | |
| N 022065 002 | 6670384 | Jan 23, 2022 | | DP U-959 | | |
| | 6670384 | Jan 23, 2022 | | DP U-960 | | |
| | 7022330 | Jan 23, 2022 | | DP U-958 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>IXABEPILONE - IXEMPRA KIT</u> | | | | | | |
| N 022065 | 002 | 7312237 | Aug 21, 2024 | U-965 | | |
| | | RE41393 | Feb 08, 2022 | U-961 | | |
| | | RE41911 | Sep 28, 2020 | DS DP U-961 | | |
| <u>IXAZOMIB CITRATE - NINLARO</u> | | | | | | |
| N 208462 | 001 | 7442830 | Aug 06, 2027 | DS DP U-2434 | NCE | Nov 20, 2020 |
| | | 7687662 | Aug 06, 2027 | DS DP | ODE-103 | Nov 20, 2022 |
| | | 8003819 | Aug 06, 2027 | DS DP U-2434 | | |
| | | 8530694 | Aug 06, 2027 | DS DP U-2434 | | |
| | | 8546608 | Aug 12, 2024 | DS | | |
| | | 8859504 | Jun 16, 2029 | DS DP | | |
| | | 8871745 | Aug 06, 2027 | U-2434 | | |
| | | 9175017 | Jun 16, 2029 | U-2434 | | |
| | | 9233115 | Aug 12, 2024 | U-2434 | | |
| <u>IXAZOMIB CITRATE - NINLARO</u> | | | | | | |
| N 208462 | 002 | 7442830 | Aug 06, 2027 | DS DP U-2434 | NCE | Nov 20, 2020 |
| | | 7687662 | Aug 06, 2027 | DS DP | ODE-103 | Nov 20, 2022 |
| | | 8003819 | Aug 06, 2027 | DS DP U-2434 | | |
| | | 8530694 | Aug 06, 2027 | DS DP U-2434 | | |
| | | 8546608 | Aug 12, 2024 | DS | | |
| | | 8859504 | Jun 16, 2029 | DS DP | | |
| | | 8871745 | Aug 06, 2027 | U-2434 | | |
| | | 9175017 | Jun 16, 2029 | U-2434 | | |
| | | 9233115 | Aug 12, 2024 | U-2434 | | |
| <u>IXAZOMIB CITRATE - NINLARO</u> | | | | | | |
| N 208462 | 003 | 7442830 | Aug 06, 2027 | DS DP U-2434 | NCE | Nov 20, 2020 |
| | | 7687662 | Aug 06, 2027 | DS DP | ODE-103 | Nov 20, 2022 |
| | | 8003819 | Aug 06, 2027 | DS DP U-2434 | | |
| | | 8530694 | Aug 06, 2027 | DS DP U-2434 | | |
| | | 8546608 | Aug 12, 2024 | DS | | |
| | | 8859504 | Jun 16, 2029 | DS DP | | |
| | | 8871745 | Aug 06, 2027 | U-2434 | | |
| | | 9175017 | Jun 16, 2029 | U-2434 | | |
| | | 9233115 | Aug 12, 2024 | U-2434 | | |
| <u>KETOCONAZOLE - XOLEGEL</u> | | | | | | |
| N 021946 | 001 | 8232276 | Nov 24, 2020 | DP | | |
| <u>KETOROLAC TROMETHAMINE - ACULAR LS</u> | | | | | | |
| N 021528 | 001 | 8008338 | May 24, 2027 | DS DP U-1181 | | |
| | | 8207215 | May 28, 2024 | U-1251 | | |
| | | 8377982 | May 28, 2024 | U-1363 | | |
| | | 8377982*PED | Nov 28, 2024 | | | |
| | | 8541463 | May 28, 2024 | U-1441 | | |
| | | 8541463*PED | Nov 28, 2024 | | | |
| | | 8648107 | May 28, 2024 | DP | | |
| | | 8906950 | May 28, 2024 | U-1626 | | |
| | | 8946281 | May 28, 2024 | U-1662 | | |
| | | 9216167 | May 28, 2024 | U-1800 | | |
| <u>KETOROLAC TROMETHAMINE - ACUVAIL</u> | | | | | | |
| N 022427 | 001 | 7842714 | Aug 15, 2029 | DS DP | | |
| | | 8512717 | Mar 07, 2028 | DP | | |
| | | 8992952 | Aug 05, 2024 | DP | | |
| | | 9192571 | Mar 07, 2028 | DP | | |
| <u>KETOROLAC TROMETHAMINE; PHENYLEPHRINE HYDROCHLORIDE - OMIIDRIA</u> | | | | | | |
| N 205388 | 001 | 8173707 | Jul 30, 2023 | U-1518 | NPP | Dec 08, 2020 |
| | | 8173707*PED | Jan 30, 2024 | | PED | Jun 08, 2021 |
| | | 8586633 | Jul 30, 2023 | DP | | |
| | | 8586633*PED | Jan 30, 2024 | | | |
| | | 9066856 | Oct 23, 2033 | DP | | |
| | | 9066856*PED | Apr 23, 2034 | | | |
| | | 9278101 | Jul 30, 2023 | U-1518 | | |
| | | 9278101*PED | Jan 30, 2024 | | | |
| | | 9399040 | Jul 30, 2023 | DP | | |
| | | 9399040*PED | Jan 30, 2024 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>KETOROLAC TROMETHAMINE; PHENYLEPHRINE HYDROCHLORIDE - OMIIDRIA</u> | | | | | | |
| N 205388 | 001 | 9486406 | Oct 23, 2033 | DP | | |
| | | 9486406*PED | Apr 23, 2034 | | | |
| | | 9855246 | Oct 23, 2033 | DP | | |
| <u>L-GLUTAMINE - ENDARI</u> | | | | | | |
| N 208587 | 001 | | | | I-748 ODE-150 | Jul 07, 2020 Jul 07, 2024 |
| <u>LACOSAMIDE - VIMPAT</u> | | | | | | |
| N 022253 | 001 | RE38551 | Mar 17, 2022 | DS DP U-1567 | NPP | Nov 03, 2020 |
| | | RE38551 | Mar 17, 2022 | DS DP U-2140 | | |
| <u>LACOSAMIDE - VIMPAT</u> | | | | | | |
| N 022253 | 002 | RE38551 | Mar 17, 2022 | DS DP U-1567 | NPP | Nov 03, 2020 |
| | | RE38551 | Mar 17, 2022 | DS DP U-2140 | | |
| <u>LACOSAMIDE - VIMPAT</u> | | | | | | |
| N 022253 | 003 | RE38551 | Mar 17, 2022 | DS DP U-1567 | NPP | Nov 03, 2020 |
| | | RE38551 | Mar 17, 2022 | DS DP U-2140 | | |
| <u>LACOSAMIDE - VIMPAT</u> | | | | | | |
| N 022253 | 004 | RE38551 | Mar 17, 2022 | DS DP U-1567 | NPP | Nov 03, 2020 |
| | | RE38551 | Mar 17, 2022 | DS DP U-2140 | | |
| <u>LACOSAMIDE - VIMPAT</u> | | | | | | |
| N 022254 | 001 | RE38551 | Mar 17, 2022 | DS DP U-1565 | M-217 | Nov 03, 2020 |
| | | RE38551 | Mar 17, 2022 | DS DP U-1568 | | |
| <u>LACOSAMIDE - VIMPAT</u> | | | | | | |
| N 022255 | 001 | RE38551 | Mar 17, 2022 | DS DP U-1567 | NPP | Nov 03, 2020 |
| | | RE38551 | Mar 17, 2022 | DS DP U-2140 | | |
| <u>LAMIVUDINE; RALTEGRAVIR POTASSIUM - DUTREBIS</u> | | | | | | |
| N 206510 | 001 | 7169780 | Oct 03, 2023 | DS DP | | |
| | | 7169780*PED | Apr 03, 2024 | | | |
| | | 7217713 | Oct 21, 2022 | | U-1663 | |
| | | 7217713*PED | Apr 21, 2023 | | | |
| | | 7435734 | Oct 21, 2022 | | U-1663 | |
| | | 7435734*PED | Apr 21, 2023 | | | |
| | | 7754731 | Mar 11, 2029 | DS DP | U-1663 | |
| | | 7754731*PED | Sep 11, 2029 | | | |
| | | 7820660 | Apr 25, 2023 | DS | | |
| <u>LAMOTRIGINE - LAMICTAL XR</u> | | | | | | |
| N 022115 | 001 | 8637512 | Jun 14, 2028 | DP | | |
| | | 9144547 | Sep 22, 2023 | DP | | |
| <u>LAMOTRIGINE - LAMICTAL XR</u> | | | | | | |
| N 022115 | 002 | 8637512 | Jun 14, 2028 | DP | | |
| | | 9144547 | Sep 22, 2023 | DP | | |
| <u>LAMOTRIGINE - LAMICTAL XR</u> | | | | | | |
| N 022115 | 003 | 8637512 | Jun 14, 2028 | DP | | |
| | | 9144547 | Sep 22, 2023 | DP | | |
| <u>LAMOTRIGINE - LAMICTAL XR</u> | | | | | | |
| N 022115 | 004 | 8637512 | Jun 14, 2028 | DP | | |
| | | 9144547 | Sep 22, 2023 | DP | | |
| <u>LAMOTRIGINE - LAMICTAL XR</u> | | | | | | |
| N 022115 | 005 | 8637512 | Jun 14, 2028 | DP | | |
| | | 9144547 | Sep 22, 2023 | DP | | |
| <u>LAMOTRIGINE - LAMICTAL XR</u> | | | | | | |
| N 022115 | 006 | 8637512 | Jun 14, 2028 | DP | | |
| | | 9144547 | Sep 22, 2023 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|----------------------------|----------------------------------------------|
| <u>LAMOTRIGINE - LAMICTAL ODT</u> | | | | | | |
| N 022251 | 001 | 7919115 | Jan 04, 2029 | DS DP | | |
| | | 8840925 | Jul 02, 2028 | DP U-1596 | | |
| | | 9339504 | Jul 02, 2028 | DP U-1596 | | |
| <u>LAMOTRIGINE - LAMICTAL ODT</u> | | | | | | |
| N 022251 | 004 | 7919115 | Jan 04, 2029 | DS DP | | |
| | | 8840925 | Jul 02, 2028 | DP U-1596 | | |
| | | 9339504 | Jul 02, 2028 | DP U-1596 | | |
| <u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u> | | | | | | |
| N 022074 | 001 | 5595760 | Mar 08, 2020 | DP U-831 | I-754 ODE-156 ODE-82 | Sep 15, 2020 Sep 15, 2024 Dec 16, 2021 |
| <u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u> | | | | | | |
| N 022074 | 002 | 5595760 | Mar 08, 2020 | DP U-831 | I-754 ODE-156 ODE-82 | Sep 15, 2020 Sep 15, 2024 Dec 16, 2021 |
| <u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u> | | | | | | |
| N 022074 | 003 | 5595760 | Mar 08, 2020 | DP U-831 | I-754 ODE-156 ODE-82 | Sep 15, 2020 Sep 15, 2024 Dec 16, 2021 |
| <u>LANSOPRAZOLE - PREVACID</u> | | | | | | |
| N 021428 | 001 | 6328994 | May 17, 2019 | | | |
| | | 7431942 | May 17, 2019 | DP | | |
| | | 7875292 | May 17, 2019 | DP | | |
| | | 9901546 | May 17, 2019 | DP | | |
| | | 9901546*PED | Nov 17, 2019 | | | |
| <u>LANSOPRAZOLE - PREVACID</u> | | | | | | |
| N 021428 | 002 | 6328994 | May 17, 2019 | | | |
| | | 7431942 | May 17, 2019 | DP | | |
| | | 7875292 | May 17, 2019 | DP | | |
| | | 9901546 | May 17, 2019 | DP | | |
| | | 9901546*PED | Nov 17, 2019 | | | |
| <u>LANSOPRAZOLE - PREVACID IV</u> | | | | | | |
| N 021566 | 001 | 7396841 | Aug 17, 2021 | DP U-947 | | |
| <u>LANTHANUM CARBONATE - FOSRENOL</u> | | | | | | |
| N 021468 | 001 | 7381428 | Aug 26, 2024 | U-890 | | |
| | | 7465465 | Aug 26, 2024 | DP | | |
| <u>LANTHANUM CARBONATE - FOSRENOL</u> | | | | | | |
| N 021468 | 002 | 7381428 | Aug 26, 2024 | U-890 | | |
| | | 7465465 | Aug 26, 2024 | DP | | |
| <u>LANTHANUM CARBONATE - FOSRENOL</u> | | | | | | |
| N 021468 | 003 | 7381428 | Aug 26, 2024 | U-890 | | |
| | | 7465465 | Aug 26, 2024 | DP | | |
| <u>LANTHANUM CARBONATE - FOSRENOL</u> | | | | | | |
| N 021468 | 004 | 7381428 | Aug 26, 2024 | U-890 | | |
| | | 7465465 | Aug 26, 2024 | DP | | |
| <u>LANTHANUM CARBONATE - FOSRENOL</u> | | | | | | |
| N 204734 | 001 | 7465465 | Aug 26, 2024 | DP | | |
| | | 8980327 | Dec 01, 2030 | DP | | |
| | | 9023397 | Dec 01, 2030 | DP | | |
| <u>LANTHANUM CARBONATE - FOSRENOL</u> | | | | | | |
| N 204734 | 002 | 7465465 | Aug 26, 2024 | DP | | |
| | | 8980327 | Dec 01, 2030 | DP | | |
| | | 9023397 | Dec 01, 2030 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LAPATINIB DITOSYLATE - TYKERB</u> | | | | | | |
| N 022059 | 001 | 6713485 | Sep 29, 2020 | DS DP U-1429 | M-235 | Dec 06, 2021 |
| | | 6713485 | Sep 29, 2020 | DS DP U-800 | | |
| | | 6727256 | Jan 08, 2019 | DS DP U-1429 | | |
| | | 6727256 | Jan 08, 2019 | DS DP U-800 | | |
| | | 7157466 | Nov 19, 2021 | DS DP | | |
| | | 8513262 | Jan 08, 2019 | DS DP | | |
| | | 8821927 | Sep 18, 2029 | DS DP | | |
| <u>LAROTRECTINIB - VITRAKVI</u> | | | | | | |
| N 210861 | 001 | 10005783 | Oct 21, 2029 | U-2472 | ODE-215 | Nov 26, 2025 |
| | | 10047097 | Oct 21, 2029 | U-2474 | ODE-220 | Nov 26, 2025 |
| | | 8513263 | Dec 23, 2029 | DS DP | ODE-221 | Nov 26, 2025 |
| | | 8865698 | Oct 21, 2029 | U-2469 | | |
| | | 9127013 | Oct 21, 2029 | DS DP | | |
| | | 9447104 | Oct 21, 2029 | U-2470 | | |
| | | 9676783 | Oct 21, 2029 | U-2469 | | |
| | | 9782414 | Nov 16, 2035 | U-2475 | | |
| <u>LAROTRECTINIB - VITRAKVI</u> | | | | | | |
| N 210861 | 002 | 10005783 | Oct 21, 2029 | U-2472 | ODE-215 | Nov 26, 2025 |
| | | 10047097 | Oct 21, 2029 | U-2474 | ODE-220 | Nov 26, 2025 |
| | | 8513263 | Dec 23, 2029 | DS DP | ODE-221 | Nov 26, 2025 |
| | | 8865698 | Oct 21, 2029 | U-2469 | | |
| | | 9127013 | Oct 21, 2029 | DS DP | | |
| | | 9447104 | Oct 21, 2029 | U-2470 | | |
| | | 9676783 | Oct 21, 2029 | U-2469 | | |
| | | 9782414 | Nov 16, 2035 | U-2475 | | |
| <u>LAROTRECTINIB - VITRAKVI</u> | | | | | | |
| N 211710 | 001 | 10005783 | Oct 21, 2029 | U-2472 | ODE-215 | Nov 26, 2025 |
| | | 10045991 | Apr 04, 2037 | U-2473 | ODE-220 | Nov 26, 2025 |
| | | 10047097 | Oct 21, 2029 | U-2474 | ODE-221 | Nov 26, 2025 |
| | | 10137127 | Apr 04, 2037 | DP | | |
| | | 8513263 | Dec 23, 2029 | DS DP | | |
| | | 8865698 | Oct 21, 2029 | U-2469 | | |
| | | 9127013 | Oct 21, 2029 | DS DP | | |
| | | 9447104 | Oct 21, 2029 | U-2470 | | |
| | | 9676783 | Oct 21, 2029 | U-2469 | | |
| | | 9782414 | Nov 16, 2035 | U-2471 | | |
| <u>LATANOPROST - XELPROS</u> | | | | | | |
| N 206185 | 001 | 9539262 | Oct 15, 2028 | DP U-2400 | | |
| | | 9629852 | Sep 12, 2029 | DP | | |
| <u>LATANOPROSTENE BUNOD - VYZULTA</u> | | | | | | |
| N 207795 | 001 | 7273946 | Oct 03, 2025 | DS DP U-2144 | | |
| | | 7629345 | Jan 05, 2025 | DP U-2144 | | |
| | | 7910767 | Jan 05, 2025 | DS DP U-2144 | | |
| | | 8058467 | Jan 05, 2025 | DS U-2144 | | |
| <u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u> | | | | | | |
| N 205834 | 001 | 10039779 | Jan 30, 2034 | DS DP U-2369 | D-158 | Feb 12, 2019 |
| | | 10039779 | Jan 30, 2034 | DS DP U-2370 | D-159 | Feb 12, 2019 |
| | | 7964580 | Mar 26, 2029 | DS DP U-1470 | D-160 | Feb 12, 2019 |
| | | 8088368 | May 12, 2030 | DS DP | NCE | Oct 10, 2019 |
| | | 8273341 | May 12, 2030 | U-1470 | NPP | Apr 07, 2020 |
| | | 8334270 | Mar 21, 2028 | DS DP U-1470 | ODE-136 | Apr 07, 2024 |
| | | 8580765 | Mar 21, 2028 | DS DP U-1470 | | |
| | | 8618076 | Dec 11, 2030 | DS DP U-1470 | | |
| | | 8633309 | Mar 26, 2029 | DS DP U-1470 | | |
| | | 8735372 | Mar 21, 2028 | U-1470 | | |
| | | 8822430 | May 12, 2030 | DS DP U-1470 | | |
| | | 8841278 | May 12, 2030 | DP U-1470 | | |
| | | 8889159 | Mar 26, 2029 | DP U-1470 | | |
| | | 9085573 | Mar 21, 2028 | DS DP U-1470 | | |
| | | 9284342 | Sep 13, 2030 | DS DP U-1470 | | |
| | | 9393256 | Sep 14, 2032 | U-1470 | | |
| | | 9511056 | May 12, 2030 | DP U-1470 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LENALIDOMIDE - REVLIMID</u> | | | | | | |
| N 021880 001 | 5635517 | Oct 04, 2019 | DS | | ODE-131 | Feb 22, 2024 |
| | 6315720 | Oct 23, 2020 | U-1210 | | ODE-49 | Jun 05, 2020 |
| | 6561977 | Oct 23, 2020 | U-1210 | | ODE-88 | Feb 17, 2022 |
| | 6755784 | Oct 23, 2020 | U-1210 | | | |
| | 7189740 | Apr 11, 2023 | U-1215 | | | |
| | 7465800 | Apr 27, 2027 | DS DP | | | |
| | 7468363 | Oct 07, 2023 | U-1414 | | | |
| | 7855217 | Nov 24, 2024 | DS DP | | | |
| | 7968569 | Oct 07, 2023 | U-1216 | | | |
| | 8315886 | Oct 23, 2020 | U-1249 | | | |
| | 8404717 | Apr 11, 2023 | U-1215 | | | |
| | 8530498 | May 15, 2023 | U-1216 | | | |
| | 8626531 | Oct 23, 2020 | U-1210 | | | |
| | 8648095 | May 15, 2023 | U-1216 | | | |
| | 8741929 | Mar 08, 2028 | U-1414 | | | |
| | 9056120 | Apr 11, 2023 | U-1215 | | | |
| | 9101621 | May 15, 2023 | U-1216 | | | |
| | 9101622 | May 15, 2023 | U-1216 | | | |
| <u>LENALIDOMIDE - REVLIMID</u> | | | | | | |
| N 021880 002 | 5635517 | Oct 04, 2019 | DS | | ODE-131 | Feb 22, 2024 |
| | 6315720 | Oct 23, 2020 | U-1210 | | ODE-49 | Jun 05, 2020 |
| | 6561977 | Oct 23, 2020 | U-1210 | | ODE-88 | Feb 17, 2022 |
| | 6755784 | Oct 23, 2020 | U-1210 | | | |
| | 7189740 | Apr 11, 2023 | U-1215 | | | |
| | 7465800 | Apr 27, 2027 | DS DP | | | |
| | 7468363 | Oct 07, 2023 | U-1414 | | | |
| | 7855217 | Nov 24, 2024 | DS DP | | | |
| | 7968569 | Oct 07, 2023 | U-1216 | | | |
| | 8315886 | Oct 23, 2020 | U-1249 | | | |
| | 8404717 | Apr 11, 2023 | U-1215 | | | |
| | 8530498 | May 15, 2023 | U-1216 | | | |
| | 8626531 | Oct 23, 2020 | U-1210 | | | |
| | 8648095 | May 15, 2023 | U-1216 | | | |
| | 8741929 | Mar 08, 2028 | U-1414 | | | |
| | 9056120 | Apr 11, 2023 | U-1215 | | | |
| | 9101621 | May 15, 2023 | U-1216 | | | |
| | 9101622 | May 15, 2023 | U-1216 | | | |
| <u>LENALIDOMIDE - REVLIMID</u> | | | | | | |
| N 021880 003 | 5635517 | Oct 04, 2019 | DS | | ODE-131 | Feb 22, 2024 |
| | 6315720 | Oct 23, 2020 | U-1210 | | ODE-49 | Jun 05, 2020 |
| | 6561977 | Oct 23, 2020 | U-1210 | | ODE-88 | Feb 17, 2022 |
| | 6755784 | Oct 23, 2020 | U-1210 | | | |
| | 7189740 | Apr 11, 2023 | U-1215 | | | |
| | 7465800 | Apr 27, 2027 | DS DP | | | |
| | 7468363 | Oct 07, 2023 | U-1414 | | | |
| | 7855217 | Nov 24, 2024 | DS DP | | | |
| | 7968569 | Oct 07, 2023 | U-1216 | | | |
| | 8315886 | Oct 23, 2020 | U-1249 | | | |
| | 8404717 | Apr 11, 2023 | U-1215 | | | |
| | 8530498 | May 15, 2023 | U-1216 | | | |
| | 8626531 | Oct 23, 2020 | U-1210 | | | |
| | 8648095 | May 15, 2023 | U-1216 | | | |
| | 8741929 | Mar 08, 2028 | U-1414 | | | |
| | 9056120 | Apr 11, 2023 | U-1215 | | | |
| | 9101621 | May 15, 2023 | U-1216 | | | |
| | 9101622 | May 15, 2023 | U-1216 | | | |
| <u>LENALIDOMIDE - REVLIMID</u> | | | | | | |
| N 021880 004 | 5635517 | Oct 04, 2019 | DS | | ODE-131 | Feb 22, 2024 |
| | 6315720 | Oct 23, 2020 | U-1210 | | ODE-49 | Jun 05, 2020 |
| | 6561977 | Oct 23, 2020 | U-1210 | | ODE-88 | Feb 17, 2022 |
| | 6755784 | Oct 23, 2020 | U-1210 | | | |
| | 7189740 | Apr 11, 2023 | U-1215 | | | |
| | 7465800 | Apr 27, 2027 | DS DP | | | |
| | 7468363 | Oct 07, 2023 | U-1414 | | | |
| | 7855217 | Nov 24, 2024 | DS DP | | | |
| | 7968569 | Oct 07, 2023 | U-1216 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LENALIDOMIDE - REVLIMID</u> | | | | | | |
| N 021880 004 | 8315886 | Oct 23, 2020 | U-1249 | | | |
| | 8404717 | Apr 11, 2023 | U-1215 | | | |
| | 8530498 | May 15, 2023 | U-1216 | | | |
| | 8626531 | Oct 23, 2020 | U-1210 | | | |
| | 8648095 | May 15, 2023 | U-1216 | | | |
| | 8741929 | Mar 08, 2028 | U-1414 | | | |
| | 9056120 | Apr 11, 2023 | U-1215 | | | |
| | 9101621 | May 15, 2023 | U-1216 | | | |
| | 9101622 | May 15, 2023 | U-1216 | | | |
| <u>LENALIDOMIDE - REVLIMID</u> | | | | | | |
| N 021880 005 | 5635517 | Oct 04, 2019 | DS | | ODE-131 | Feb 22, 2024 |
| | 6315720 | Oct 23, 2020 | U-1210 | | ODE-49 | Jun 05, 2020 |
| | 6561977 | Oct 23, 2020 | U-1210 | | ODE-88 | Feb 17, 2022 |
| | 6755784 | Oct 23, 2020 | U-1210 | | | |
| | 7189740 | Apr 11, 2023 | U-1215 | | | |
| | 7465800 | Apr 27, 2027 | DS DP | | | |
| | 7468363 | Oct 07, 2023 | U-1414 | | | |
| | 7855217 | Nov 24, 2024 | DS DP | | | |
| | 7968569 | Oct 07, 2023 | U-1216 | | | |
| | 8315886 | Oct 23, 2020 | U-1249 | | | |
| | 8404717 | Apr 11, 2023 | U-1215 | | | |
| | 8530498 | May 15, 2023 | U-1216 | | | |
| | 8626531 | Oct 23, 2020 | U-1210 | | | |
| | 8648095 | May 15, 2023 | U-1216 | | | |
| | 8741929 | Mar 08, 2028 | U-1414 | | | |
| | 9056120 | Apr 11, 2023 | U-1215 | | | |
| | 9101621 | May 15, 2023 | U-1216 | | | |
| | 9101622 | May 15, 2023 | U-1216 | | | |
| <u>LENALIDOMIDE - REVLIMID</u> | | | | | | |
| N 021880 006 | 5635517 | Oct 04, 2019 | DS | | ODE-131 | Feb 22, 2024 |
| | 6315720 | Oct 23, 2020 | U-1210 | | ODE-49 | Jun 05, 2020 |
| | 6561977 | Oct 23, 2020 | U-1210 | | ODE-88 | Feb 17, 2022 |
| | 6755784 | Oct 23, 2020 | U-1210 | | | |
| | 7189740 | Apr 11, 2023 | U-1215 | | | |
| | 7465800 | Apr 27, 2027 | DS DP | | | |
| | 7468363 | Oct 07, 2023 | U-1414 | | | |
| | 7855217 | Nov 24, 2024 | DS DP | | | |
| | 7968569 | Oct 07, 2023 | U-1216 | | | |
| | 8315886 | Oct 23, 2020 | U-1249 | | | |
| | 8404717 | Apr 11, 2023 | U-1215 | | | |
| | 8530498 | May 15, 2023 | U-1216 | | | |
| | 8626531 | Oct 23, 2020 | U-1210 | | | |
| | 8648095 | May 15, 2023 | U-1216 | | | |
| | 8741929 | Mar 08, 2028 | U-1414 | | | |
| | 9056120 | Apr 11, 2023 | U-1215 | | | |
| | 9101621 | May 15, 2023 | U-1216 | | | |
| | 9101622 | May 15, 2023 | U-1216 | | | |
| <u>LENAVATINIB MESYLATE - LENVIMA</u> | | | | | | |
| N 206947 001 | 7253286 | Oct 19, 2021 | DS DP | | I-734 | May 13, 2019 |
| | 7612208 | Sep 19, 2026 | DS DP | | I-787 | Aug 15, 2021 |
| | 9006256 | Jul 27, 2027 | U-1695 | | NCE | Feb 13, 2020 |
| | | | | | ODE-196 | Aug 15, 2025 |
| | | | | | ODE-87 | Feb 13, 2022 |
| <u>LENAVATINIB MESYLATE - LENVIMA</u> | | | | | | |
| N 206947 002 | 7253286 | Oct 19, 2021 | DS DP | | I-734 | May 13, 2019 |
| | 7612208 | Sep 19, 2026 | DS DP | | I-787 | Aug 15, 2021 |
| | 9006256 | Jul 27, 2027 | U-1695 | | NCE | Feb 13, 2020 |
| | | | | | ODE-196 | Aug 15, 2025 |
| | | | | | ODE-87 | Feb 13, 2022 |
| <u>LESINURAD - ZURAMPIC</u> | | | | | | |
| N 207988 001 | 8003681 | Aug 25, 2025 | DS | | NCE | Dec 22, 2020 |
| | 8084483 | Aug 17, 2029 | U-1801 | | | |
| | 8283369 | Nov 26, 2028 | U-1802 | | | |
| | 8283369 | Nov 26, 2028 | U-1804 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LESINURAD - ZURAMPIC</u> | | | | | | |
| N 207988 | 001 | 8357713 | Nov 26, 2028 | DP U-1801 | | |
| | | 8357713 | Nov 26, 2028 | DP U-1802 | | |
| | | 8357713 | Nov 26, 2028 | DP U-1803 | | |
| | | 8546436 | Feb 29, 2032 | DS DP | | |
| | | 8546437 | Apr 29, 2029 | U-1803 | | |
| | | 9216179 | Aug 01, 2031 | U-1806 | | |
| | | 9956205 | Dec 28, 2031 | U-2311 | | |
| <u>LETERMOVIR - PREVYMIS</u> | | | | | | |
| N 209939 | 001 | 7196086 | May 22, 2024 | DS DP | NCE | Nov 08, 2022 |
| | | 8513255 | May 22, 2024 | DS DP | ODE-165 | Nov 08, 2024 |
| <u>LETERMOVIR - PREVYMIS</u> | | | | | | |
| N 209939 | 002 | 7196086 | May 22, 2024 | DS DP | NCE | Nov 08, 2022 |
| | | 8513255 | May 22, 2024 | DS DP | ODE-165 | Nov 08, 2024 |
| <u>LETERMOVIR - PREVYMIS</u> | | | | | | |
| N 209940 | 001 | 7196086 | May 22, 2024 | DS DP | NCE | Nov 08, 2022 |
| | | 8513255 | May 22, 2024 | DS DP | ODE-165 | Nov 08, 2024 |
| <u>LETERMOVIR - PREVYMIS</u> | | | | | | |
| N 209940 | 002 | 7196086 | May 22, 2024 | DS DP | NCE | Nov 08, 2022 |
| | | 8513255 | May 22, 2024 | DS DP | ODE-165 | Nov 08, 2024 |
| <u>LETROZOLE; RIBOCICLIB SUCCINATE - KISOALI FEMARA CO-PACK (COPACKAGED)</u> | | | | | | |
| N 209935 | 001 | 8324225 | Jun 17, 2028 | DS DP | NCE | Mar 13, 2022 |
| | | 8415355 | Feb 19, 2031 | DS DP | | |
| | | 8685980 | May 25, 2030 | DS DP | | |
| | | 8962630 | Dec 09, 2029 | U-1981 | | |
| | | 9193732 | Nov 09, 2031 | DS DP | | |
| | | 9416136 | Aug 20, 2029 | U-1981 | | |
| | | 9868739 | Nov 09, 2031 | U-1981 | | |
| <u>LEUPROLIDE ACETATE - LUPRON DEPOT</u> | | | | | | |
| N 020517 | 003 | 7429559 | Jan 13, 2019 | DP | | |
| | | 8815801 | Jun 28, 2022 | DP | | |
| | | 8921326 | Feb 05, 2031 | DP U-1666 | | |
| <u>LEUPROLIDE ACETATE - ELIGARD</u> | | | | | | |
| N 021343 | 001 | 6626870 | Mar 27, 2020 | DP | | |
| <u>LEUPROLIDE ACETATE - ELIGARD</u> | | | | | | |
| N 021379 | 001 | 6626870 | Mar 27, 2020 | DP | | |
| | | 8470359 | Oct 15, 2023 | DS DP U-621 | | |
| | | 8840916 | Nov 13, 2020 | DP | | |
| | | 9539333 | Nov 13, 2020 | DS DP U-621 | | |
| <u>LEUPROLIDE ACETATE - ELIGARD</u> | | | | | | |
| N 021488 | 001 | 6626870 | Mar 27, 2020 | DP | | |
| | | 8470359 | Oct 15, 2023 | DS DP U-621 | | |
| | | 8840916 | Nov 13, 2020 | DP | | |
| | | 9539333 | Nov 13, 2020 | DS DP U-621 | | |
| <u>LEUPROLIDE ACETATE - ELIGARD</u> | | | | | | |
| N 021731 | 001 | 6626870 | Mar 27, 2020 | DP | | |
| | | 8470359 | Oct 15, 2023 | DS DP U-621 | | |
| | | 8840916 | Nov 13, 2020 | DP | | |
| | | 9539333 | Nov 13, 2020 | DS DP U-621 | | |
| | | 9914802 | Nov 13, 2020 | DS DP U-1666 | | |
| <u>LEUPROLIDE ACETATE - LUTRATE DEPOT KIT</u> | | | | | | |
| N 205054 | 001 | 9789064 | Dec 15, 2020 | DP | | |
| <u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u> | | | | | | |
| N 020837 | 001 | 6451289 | Mar 21, 2021 | | | |
| <u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u> | | | | | | |
| N 020837 | 002 | 6451289 | Mar 21, 2021 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u> | | | | | | |
| N 020837 | 003 | 6451289 | Mar 21, 2021 | | | |
| <u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u> | | | | | | |
| N 020837 | 004 | 6451289 | Mar 21, 2021 | DP | | |
| <u>LEVALBUTEROL TARTRATE - XOPENEX HFA</u> | | | | | | |
| N 021730 | 001 | 7256310 | Oct 08, 2024 | DS DP | U-636 | |
| | | 8765153 | Dec 08, 2023 | DP | | |
| <u>LEVETIRACETAM - KEPPRA</u> | | | | | | |
| N 021035 | 001 | 8802142 | Jun 07, 2031 | DP | | |
| | | 8802142*PED | Dec 07, 2031 | | | |
| <u>LEVETIRACETAM - KEPPRA</u> | | | | | | |
| N 021035 | 002 | 8802142 | Jun 07, 2031 | DP | | |
| | | 8802142*PED | Dec 07, 2031 | | | |
| <u>LEVETIRACETAM - KEPPRA</u> | | | | | | |
| N 021035 | 003 | 8802142 | Jun 07, 2031 | DP | | |
| | | 8802142*PED | Dec 07, 2031 | | | |
| <u>LEVETIRACETAM - KEPPRA</u> | | | | | | |
| N 021035 | 004 | 8802142 | Jun 07, 2031 | DP | | |
| | | 8802142*PED | Dec 07, 2031 | | | |
| <u>LEVETIRACETAM - KEPPRA XR</u> | | | | | | |
| N 022285 | 001 | 7858122 | Sep 17, 2028 | DP | | |
| <u>LEVETIRACETAM - KEPPRA XR</u> | | | | | | |
| N 022285 | 002 | 7858122 | Sep 17, 2028 | DP | | |
| <u>LEVETIRACETAM - ELEPSIA XR</u> | | | | | | |
| N 204417 | 001 | 8163306 | Sep 03, 2027 | DP | | |
| | | 8425938 | Feb 22, 2026 | DP | | |
| | | 8431156 | Oct 31, 2027 | DP | | |
| | | 8470367 | Feb 22, 2026 | DP | | |
| | | 8535717 | Feb 22, 2026 | DP | | |
| <u>LEVETIRACETAM - ELEPSIA XR</u> | | | | | | |
| N 204417 | 002 | 8163306 | Sep 03, 2027 | DP | | |
| | | 8425938 | Feb 22, 2026 | DP | | |
| | | 8431156 | Oct 31, 2027 | DP | | |
| | | 8470367 | Feb 22, 2026 | DP | | |
| | | 8535717 | Feb 22, 2026 | DP | | |
| <u>LEVETIRACETAM - SPRITAM</u> | | | | | | |
| N 207958 | 001 | 9339489 | Mar 14, 2034 | DP | U-1850 | |
| | | 9669009 | Mar 14, 2034 | | U-1850 | |
| | | 9669009 | Mar 14, 2034 | | U-2021 | |
| | | 9669009 | Mar 14, 2034 | | U-2022 | |
| <u>LEVETIRACETAM - SPRITAM</u> | | | | | | |
| N 207958 | 002 | 9339489 | Mar 14, 2034 | DP | U-1850 | |
| | | 9669009 | Mar 14, 2034 | | U-1850 | |
| | | 9669009 | Mar 14, 2034 | | U-2021 | |
| | | 9669009 | Mar 14, 2034 | | U-2022 | |
| <u>LEVETIRACETAM - SPRITAM</u> | | | | | | |
| N 207958 | 003 | 9339489 | Mar 14, 2034 | DP | U-1850 | |
| | | 9669009 | Mar 14, 2034 | | U-1850 | |
| | | 9669009 | Mar 14, 2034 | | U-2021 | |
| | | 9669009 | Mar 14, 2034 | | U-2022 | |
| <u>LEVETIRACETAM - SPRITAM</u> | | | | | | |
| N 207958 | 004 | 9339489 | Mar 14, 2034 | DP | U-1850 | |
| | | 9669009 | Mar 14, 2034 | | U-1850 | |
| | | 9669009 | Mar 14, 2034 | | U-2021 | |
| | | 9669009 | Mar 14, 2034 | | U-2022 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LEVOCARNITINE - CARNITOR</u> | | | | | | |
| N 020182 | 001 | 6335369 | Jan 18, 2021 | U-433 | | |
| | | 6429230 | Jan 18, 2021 | U-433 | | |
| | | 6696493 | Jan 18, 2021 | U-433 | | |
| <u>LEVOCETIRIZINE DIHYDROCHLORIDE - XYZAL ALLERGY 24HR</u> | | | | | | |
| N 209090 | 001 | 8633194 | Oct 16, 2027 | DP | | |
| <u>LEVOFLOXACIN - LEVAQUIN</u> | | | | | | |
| N 021721 | 001 | 6806256 | Feb 26, 2022 | DP | | |
| <u>LEVOLEUCOVORIN CALCIUM - FUSILEV</u> | | | | | | |
| N 020140 | 001 | 6500829 | Mar 07, 2022 | DS DP | | |
| <u>LEVOLEUCOVORIN CALCIUM - FUSILEV</u> | | | | | | |
| N 020140 | 002 | 6500829 | Mar 07, 2022 | DS DP | | |
| <u>LEVOLEUCOVORIN CALCIUM - FUSILEV</u> | | | | | | |
| N 020140 | 003 | 6500829 | Mar 07, 2022 | DS DP | | |
| <u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u> | | | | | | |
| N 204168 | 001 | 8481598 | Mar 02, 2031 | U-839 | | |
| | | 8865937 | May 23, 2032 | DS DP | | |
| | | RE43879 | Jun 03, 2023 | U-839 | | |
| <u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u> | | | | | | |
| N 204168 | 002 | 8481598 | Mar 02, 2031 | U-839 | | |
| | | 8865937 | May 23, 2032 | DS DP | | |
| | | RE43879 | Jun 03, 2023 | U-839 | | |
| <u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u> | | | | | | |
| N 204168 | 003 | 8481598 | Mar 02, 2031 | U-839 | | |
| | | 8865937 | May 23, 2032 | DS DP | | |
| | | RE43879 | Jun 03, 2023 | U-839 | | |
| <u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u> | | | | | | |
| N 204168 | 004 | 8481598 | Mar 02, 2031 | U-839 | | |
| | | 8865937 | May 23, 2032 | DS DP | | |
| | | RE43879 | Jun 03, 2023 | U-839 | | |
| <u>LEVONORGESTREL - MIRENA</u> | | | | | | |
| N 021225 | 001 | 9615965 | Sep 16, 2029 | DP U-2003 | | |
| | | 9668912 | Apr 01, 2031 | DP | | |
| <u>LEVONORGESTREL - SKYLA</u> | | | | | | |
| N 203159 | 001 | 7252839 | Nov 13, 2023 | DP | | |
| | | 9615965 | Sep 16, 2029 | DP U-2003 | | |
| | | 9668912 | Apr 01, 2031 | DP | | |
| <u>LEVONORGESTREL - LILETTA</u> | | | | | | |
| N 206229 | 001 | 10028858 | Mar 22, 2034 | DP U-2348 | | |
| <u>LEVONORGESTREL - KYLEENA</u> | | | | | | |
| N 208224 | 001 | 7252839 | Nov 13, 2023 | DP | NP | Sep 16, 2019 |
| | | 9615965 | Sep 16, 2029 | DP U-2003 | | |
| | | 9668912 | Apr 01, 2031 | DP | | |
| <u>LEVORPHANOL TARTRATE - LEVORPHANOL TARTRATE</u> | | | | | | |
| A 211484 | 001 | | | | CGT | Jun 11, 2019 |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | | |
| N 021301 | 001 | 6555581 | Feb 15, 2022 | | | |
| | | 7067148 | Feb 15, 2022 | DP | | |
| | | 7101569 | Oct 02, 2023 | U-759 | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | | |
| N 021301 | 002 | 6555581 | Feb 15, 2022 | | | |
| | | 7067148 | Feb 15, 2022 | DP | | |
| | | 7101569 | Oct 02, 2023 | U-759 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | | |
| N 021301 003 | 6555581 | Feb 15, 2022 | | | | |
| | 7067148 | Feb 15, 2022 | DP | | | |
| | 7101569 | Oct 02, 2023 | | U-759 | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | | |
| N 021301 004 | 6555581 | Feb 15, 2022 | | | | |
| | 7067148 | Feb 15, 2022 | DP | | | |
| | 7101569 | Oct 02, 2023 | | U-759 | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | | |
| N 021301 005 | 6555581 | Feb 15, 2022 | | | | |
| | 7067148 | Feb 15, 2022 | DP | | | |
| | 7101569 | Oct 02, 2023 | | U-759 | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | | |
| N 021301 006 | 6555581 | Feb 15, 2022 | | | | |
| | 7067148 | Feb 15, 2022 | DP | | | |
| | 7101569 | Oct 02, 2023 | | U-759 | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | | |
| N 021301 007 | 6555581 | Feb 15, 2022 | | | | |
| | 7067148 | Feb 15, 2022 | DP | | | |
| | 7101569 | Oct 02, 2023 | | U-759 | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | | |
| N 021301 008 | 6555581 | Feb 15, 2022 | | | | |
| | 7067148 | Feb 15, 2022 | DP | | | |
| | 7101569 | Oct 02, 2023 | | U-759 | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | | |
| N 021301 009 | 6555581 | Feb 15, 2022 | | | | |
| | 7067148 | Feb 15, 2022 | DP | | | |
| | 7101569 | Oct 02, 2023 | | U-759 | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | | |
| N 021301 010 | 6555581 | Feb 15, 2022 | | | | |
| | 7067148 | Feb 15, 2022 | DP | | | |
| | 7101569 | Oct 02, 2023 | | U-759 | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | | |
| N 021301 011 | 6555581 | Feb 15, 2022 | | | | |
| | 7067148 | Feb 15, 2022 | DP | | | |
| | 7101569 | Oct 02, 2023 | | U-759 | | |
| <u>LEVOTHYROXINE SODIUM - LEVOXYL</u> | | | | | | |
| N 021301 012 | 6555581 | Feb 15, 2022 | | | | |
| | 7067148 | Feb 15, 2022 | DP | | | |
| | 7101569 | Oct 02, 2023 | | U-759 | | |
| <u>LEVOTHYROXINE SODIUM - LEVO-T</u> | | | | | | |
| N 021342 001 | 6399101 | Mar 30, 2020 | | | | |
| <u>LEVOTHYROXINE SODIUM - LEVO-T</u> | | | | | | |
| N 021342 002 | 6399101 | Mar 30, 2020 | | | | |
| <u>LEVOTHYROXINE SODIUM - LEVO-T</u> | | | | | | |
| N 021342 003 | 6399101 | Mar 30, 2020 | | | | |
| <u>LEVOTHYROXINE SODIUM - LEVO-T</u> | | | | | | |
| N 021342 004 | 6399101 | Mar 30, 2020 | | | | |
| <u>LEVOTHYROXINE SODIUM - LEVO-T</u> | | | | | | |
| N 021342 005 | 6399101 | Mar 30, 2020 | | | | |
| <u>LEVOTHYROXINE SODIUM - LEVO-T</u> | | | | | | |
| N 021342 006 | 6399101 | Mar 30, 2020 | | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LEVOTHYROXINE SODIUM - LEVO-T</u> | | | | | | |
| N 021342 007 | 6399101 | Mar 30, 2020 | | | | |
| <u>LEVOTHYROXINE SODIUM - LEVO-T</u> | | | | | | |
| N 021342 008 | 6399101 | Mar 30, 2020 | | | | |
| <u>LEVOTHYROXINE SODIUM - LEVO-T</u> | | | | | | |
| N 021342 009 | 6399101 | Mar 30, 2020 | | | | |
| <u>LEVOTHYROXINE SODIUM - LEVO-T</u> | | | | | | |
| N 021342 010 | 6399101 | Mar 30, 2020 | | | | |
| <u>LEVOTHYROXINE SODIUM - LEVO-T</u> | | | | | | |
| N 021342 011 | 6399101 | Mar 30, 2020 | | | | |
| <u>LEVOTHYROXINE SODIUM - TIROSINT</u> | | | | | | |
| N 021924 002 | 7691411 | Mar 14, 2024 | | DP | | |
| | 7723390 | Mar 14, 2024 | | DP | | |
| <u>LEVOTHYROXINE SODIUM - TIROSINT</u> | | | | | | |
| N 021924 003 | 7691411 | Mar 14, 2024 | | DP | | |
| | 7723390 | Mar 14, 2024 | | DP | | |
| <u>LEVOTHYROXINE SODIUM - TIROSINT</u> | | | | | | |
| N 021924 004 | 7691411 | Mar 14, 2024 | | DP | | |
| | 7723390 | Mar 14, 2024 | | DP | | |
| <u>LEVOTHYROXINE SODIUM - TIROSINT</u> | | | | | | |
| N 021924 005 | 7691411 | Mar 14, 2024 | | DP | | |
| | 7723390 | Mar 14, 2024 | | DP | | |
| <u>LEVOTHYROXINE SODIUM - TIROSINT</u> | | | | | | |
| N 021924 006 | 7691411 | Mar 14, 2024 | | DP | | |
| | 7723390 | Mar 14, 2024 | | DP | | |
| <u>LEVOTHYROXINE SODIUM - TIROSINT</u> | | | | | | |
| N 021924 007 | 7691411 | Mar 14, 2024 | | DP | | |
| | 7723390 | Mar 14, 2024 | | DP | | |
| <u>LEVOTHYROXINE SODIUM - TIROSINT</u> | | | | | | |
| N 021924 008 | 7691411 | Mar 14, 2024 | | DP | | |
| | 7723390 | Mar 14, 2024 | | DP | | |
| <u>LEVOTHYROXINE SODIUM - TIROSINT</u> | | | | | | |
| N 021924 009 | 7691411 | Mar 14, 2024 | | DP | | |
| | 7723390 | Mar 14, 2024 | | DP | | |
| <u>LEVOTHYROXINE SODIUM - TIROSINT</u> | | | | | | |
| N 021924 010 | 7691411 | Mar 14, 2024 | | DP | | |
| | 7723390 | Mar 14, 2024 | | DP | | |
| <u>LEVOTHYROXINE SODIUM - TIROSINT</u> | | | | | | |
| N 021924 013 | 7691411 | Mar 14, 2024 | | DP | | |
| | 7723390 | Mar 14, 2024 | | DP | | |
| <u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u> | | | | | | |
| N 202231 001 | 9006289 | Oct 03, 2032 | | DP | | |
| | 9168238 | Aug 29, 2032 | | DP | | |
| | 9168239 | Aug 29, 2032 | | DP | | |
| <u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u> | | | | | | |
| N 202231 002 | 9006289 | Oct 03, 2032 | | DP | | |
| | 9168238 | Aug 29, 2032 | | DP | | |
| | 9168239 | Aug 29, 2032 | | DP | | |
| <u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u> | | | | | | |
| N 202231 003 | 9006289 | Oct 03, 2032 | | DP | | |
| | 9168238 | Aug 29, 2032 | | DP | | |
| | 9168239 | Aug 29, 2032 | | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LIDOCAINE - ZTLIDO</u> | | | | | | |
| N 207962 | 001 | 9283174 | May 10, 2031 | DP | NP | Feb 28, 2021 |
| | | 9925264 | May 10, 2031 | DP U-2267 | | |
| | | 9931403 | May 10, 2031 | DP | | |
| <u>LIDOCAINE HYDROCHLORIDE - ZINGO</u> | | | | | | |
| N 022114 | 001 | 8540665 | Oct 22, 2029 | U-1438 | | |
| | | 9358338 | Apr 27, 2035 | U-1870 | | |
| | | 9370622 | Sep 28, 2035 | U-1870 | | |
| <u>LIDOCAINE HYDROCHLORIDE - AKTEN</u> | | | | | | |
| N 022221 | 001 | 8759401 | Jul 24, 2026 | DP U-1523 | | |
| <u>LIDOCAINE; TETRACAINE - SYNERA</u> | | | | | | |
| N 021623 | 001 | 6465709 | Jul 07, 2020 | DP | | |
| <u>LIDOCAINE; TETRACAINE - PLIAGLIS</u> | | | | | | |
| N 021717 | 001 | 6528086 | Sep 28, 2019 | DP | | |
| <u>LIFITEGRAST - XIIDRA</u> | | | | | | |
| N 208073 | 001 | 10124000 | Nov 05, 2024 | U-1900 | NCE | Jul 11, 2021 |
| | | 7314938 | Mar 10, 2025 | DS DP | | |
| | | 7745460 | Nov 05, 2024 | DS DP U-1880 | | |
| | | 7790743 | Nov 05, 2024 | U-1880 | | |
| | | 7928122 | Nov 05, 2024 | DS DP | | |
| | | 8084047 | May 17, 2026 | DS DP | | |
| | | 8168655 | May 09, 2029 | U-1880 | | |
| | | 8367701 | Apr 15, 2029 | DP U-1880 | | |
| | | 8592450 | May 17, 2026 | U-1880 | | |
| | | 8927574 | Nov 12, 2030 | DP | | |
| | | 9085553 | Jul 25, 2033 | DP | | |
| | | 9216174 | Nov 05, 2024 | DP | | |
| | | 9353088 | Oct 21, 2030 | DP | | |
| | | 9447077 | Apr 15, 2029 | U-1900 | | |
| | | 9890141 | Oct 21, 2030 | DS | | |
| <u>LINACLOTIDE - LINZESS</u> | | | | | | |
| N 202811 | 001 | 7304036 | Aug 30, 2026 | DS DP U-1278 | | |
| | | 7304036 | Aug 30, 2026 | DS DP U-1516 | | |
| | | 7371727 | Jan 28, 2024 | DS | | |
| | | 7704947 | Jan 28, 2024 | DS DP | | |
| | | 7745409 | Jan 28, 2024 | DS DP | | |
| | | 8080526 | Jan 28, 2024 | DS DP | | |
| | | 8110553 | Jan 28, 2024 | U-1278 | | |
| | | 8748573 | Oct 30, 2031 | U-1515 | | |
| | | 8748573 | Oct 30, 2031 | U-1516 | | |
| | | 8802628 | Nov 17, 2031 | DP | | |
| | | 8933030 | Feb 17, 2031 | DP | | |
| | | 9708371 | Aug 16, 2033 | DP U-1515 | | |
| | | 9708371 | Aug 16, 2033 | DP U-1516 | | |
| <u>LINACLOTIDE - LINZESS</u> | | | | | | |
| N 202811 | 002 | 7304036 | Aug 30, 2026 | DS DP U-1278 | | |
| | | 7304036 | Aug 30, 2026 | DS DP U-1516 | | |
| | | 7371727 | Jan 28, 2024 | DS | | |
| | | 7704947 | Jan 28, 2024 | DS DP | | |
| | | 7745409 | Jan 28, 2024 | DS DP | | |
| | | 8080526 | Jan 28, 2024 | DS DP | | |
| | | 8110553 | Jan 28, 2024 | U-1278 | | |
| | | 8748573 | Oct 30, 2031 | U-1515 | | |
| | | 8748573 | Oct 30, 2031 | U-1516 | | |
| | | 8802628 | Nov 17, 2031 | DP | | |
| | | 8933030 | Feb 17, 2031 | DP | | |
| | | 9708371 | Aug 16, 2033 | DP U-1515 | | |
| <u>LINACLOTIDE - LINZESS</u> | | | | | | |
| N 202811 | 003 | 7304036 | Aug 30, 2026 | DS DP U-1516 | NS | Jan 25, 2020 |
| | | 7371727 | Jan 28, 2024 | DS | | |
| | | 7704947 | Jan 28, 2024 | DS DP | | |
| | | 7745409 | Jan 28, 2024 | DS DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LINACLOTIDE - LINZESS</u> | | | | | | |
| N 202811 003 | 8080526 | Jan 28, 2024 | DS DP | | | |
| | 8110553 | Jan 28, 2024 | U-1516 | | | |
| | 8933030 | Feb 17, 2031 | DP U-1516 | | | |
| | 9708371 | Aug 16, 2033 | DP U-1516 | | | |
| <u>LINAGLIPTIN - TRADJENTA</u> | | | | | | |
| N 201280 001 | 10034877 | Aug 05, 2029 | U-2347 | | | |
| | 6890898 | Feb 02, 2019 | U-1270 | | | |
| | 6890898 | Feb 02, 2019 | U-493 | | | |
| | 7078381 | Feb 02, 2019 | U-1270 | | | |
| | 7078381 | Feb 02, 2019 | U-493 | | | |
| | 7407955 | May 02, 2025 | DS DP | | | |
| | 7459428 | Feb 02, 2019 | U-1270 | | | |
| | 7459428 | Feb 02, 2019 | U-493 | | | |
| | 8119648 | Aug 12, 2023 | U-1270 | | | |
| | 8119648 | Aug 12, 2023 | U-774 | | | |
| | 8178541 | Aug 12, 2023 | U-1244 | | | |
| | 8178541 | Aug 12, 2023 | U-1245 | | | |
| | 8178541 | Aug 12, 2023 | U-1270 | | | |
| | 8178541 | Aug 12, 2023 | U-775 | | | |
| | 8673927 | May 04, 2027 | U-1503 | | | |
| | 8846695 | Jun 04, 2030 | U-1503 | Y | | |
| | 8853156 | Mar 05, 2031 | U-1642 | | | |
| | 8883805 | Nov 26, 2025 | DP | | | |
| | 9173859 | May 04, 2027 | DP U-1503 | | | |
| | 9173859 | May 04, 2027 | DP U-1768 | | | |
| | 9486526 | Aug 05, 2029 | U-1915 | | | |
| <u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u> | | | | | | |
| N 201281 001 | 10022379 | Apr 02, 2029 | U-2339 | | | |
| | 6890898 | Feb 02, 2019 | U-1039 | | | |
| | 7078381 | Feb 02, 2019 | U-1039 | | | |
| | 7407955 | May 02, 2025 | DS DP | | | |
| | 7459428 | Feb 02, 2019 | U-1039 | | | |
| | 8119648 | Aug 12, 2023 | U-802 | | | |
| | 8178541 | Aug 12, 2023 | DP U-775 | | | |
| | 8673927 | May 04, 2027 | U-1503 | | | |
| | 8846695 | Jun 04, 2030 | U-1503 | | | |
| | 8883805 | Nov 26, 2025 | DP | | | |
| | 9155705 | May 21, 2030 | DP | | | |
| | 9173859 | May 04, 2027 | DP U-1503 | | | |
| | 9415016 | Apr 02, 2029 | DP | | | |
| <u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u> | | | | | | |
| N 201281 002 | 10022379 | Apr 02, 2029 | U-2339 | | | |
| | 6890898 | Feb 02, 2019 | U-1039 | | | |
| | 7078381 | Feb 02, 2019 | U-1039 | | | |
| | 7407955 | May 02, 2025 | DS DP | | | |
| | 7459428 | Feb 02, 2019 | U-1039 | | | |
| | 8119648 | Aug 12, 2023 | U-802 | | | |
| | 8178541 | Aug 12, 2023 | DP U-775 | | | |
| | 8673927 | May 04, 2027 | U-1503 | | | |
| | 8846695 | Jun 04, 2030 | U-1503 | | | |
| | 8883805 | Nov 26, 2025 | DP | | | |
| | 9155705 | May 21, 2030 | DP | | | |
| | 9173859 | May 04, 2027 | DP U-1503 | | | |
| | 9415016 | Apr 02, 2029 | DP | | | |
| <u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u> | | | | | | |
| N 201281 003 | 10022379 | Apr 02, 2029 | U-2339 | | | |
| | 6890898 | Feb 02, 2019 | U-1039 | | | |
| | 7078381 | Feb 02, 2019 | U-1039 | | | |
| | 7407955 | May 02, 2025 | DS DP | | | |
| | 7459428 | Feb 02, 2019 | U-1039 | | | |
| | 8119648 | Aug 12, 2023 | U-802 | | | |
| | 8178541 | Aug 12, 2023 | DP U-775 | | | |
| | 8673927 | May 04, 2027 | U-1503 | | | |
| | 8846695 | Jun 04, 2030 | U-1503 | | | |
| | 8883805 | Nov 26, 2025 | DP | | | |
| | 9155705 | May 21, 2030 | DP | | | |
| | 9173859 | May 04, 2027 | DP U-1503 | | | |
| | 9415016 | Apr 02, 2029 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO</u> | | | | | | |
| N 201281 003 | 9155705 | May 21, 2030 | DP | | | |
| | 9173859 | May 04, 2027 | DP | U-1503 | | |
| | 9415016 | Apr 02, 2029 | DP | | | |
| <u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</u> | | | | | | |
| N 208026 001 | 10022379 | Apr 02, 2029 | | U-2339 | | |
| | 6488962 | Jun 20, 2020 | DP | | | |
| | 6890898 | Feb 02, 2019 | | U-803 | | |
| | 7078381 | Feb 02, 2019 | | U-803 | | |
| | 7407955 | May 02, 2025 | DS DP | | | |
| | 7459428 | Feb 02, 2019 | | U-803 | | |
| | 8119648 | Aug 12, 2023 | | U-802 | | |
| | 8178541 | Aug 12, 2023 | DP | U-1853 | | |
| | 8673927 | May 04, 2027 | | U-1503 | | |
| | 8883805 | Nov 26, 2025 | DP | | | |
| | 9155705 | May 21, 2030 | DP | | | |
| | 9173859 | May 04, 2027 | DP | U-1503 | | |
| | 9415016 | Apr 02, 2029 | DP | | | |
| | 9555001 | Mar 06, 2033 | DP | U-1967 | | |
| | 9555001 | Mar 06, 2033 | DP | U-1968 | | |
| <u>LINAGLIPTIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</u> | | | | | | |
| N 208026 002 | 10022379 | Apr 02, 2029 | | U-2339 | | |
| | 6488962 | Jun 20, 2020 | DP | | | |
| | 6890898 | Feb 02, 2019 | | U-803 | | |
| | 7078381 | Feb 02, 2019 | | U-803 | | |
| | 7407955 | May 02, 2025 | DS DP | | | |
| | 7459428 | Feb 02, 2019 | | U-803 | | |
| | 8119648 | Aug 12, 2023 | | U-802 | | |
| | 8178541 | Aug 12, 2023 | DP | U-1853 | | |
| | 8673927 | May 04, 2027 | | U-1503 | | |
| | 8883805 | Nov 26, 2025 | DP | | | |
| | 9155705 | May 21, 2030 | DP | | | |
| | 9173859 | May 04, 2027 | DP | U-1503 | | |
| | 9415016 | Apr 02, 2029 | DP | | | |
| | 9555001 | Mar 06, 2033 | DP | U-1967 | | |
| | 9555001 | Mar 06, 2033 | DP | U-1968 | | |
| <u>LINEZOLID - ZYVOX</u> | | | | | | |
| N 021130 001 | 6514529 | Mar 15, 2021 | DP | | | |
| | 6559305 | Jan 29, 2021 | DS | | | |
| <u>LINEZOLID - ZYVOX</u> | | | | | | |
| N 021130 002 | 6514529 | Mar 15, 2021 | DP | | | |
| | 6559305 | Jan 29, 2021 | DS | | | |
| <u>LINEZOLID - ZYVOX</u> | | | | | | |
| N 021131 001 | 6559305 | Jan 29, 2021 | DS | | | |
| <u>LINEZOLID - ZYVOX</u> | | | | | | |
| N 021131 002 | 6559305 | Jan 29, 2021 | DS | | | |
| | 6559305*PED | Jul 29, 2021 | | | | |
| <u>LINEZOLID - ZYVOX</u> | | | | | | |
| N 021131 003 | 6559305 | Jan 29, 2021 | DS | | | |
| | 6559305*PED | Jul 29, 2021 | | | | |
| <u>LINEZOLID - ZYVOX</u> | | | | | | |
| N 021132 001 | 6559305 | Jan 29, 2021 | DS | | | |
| <u>LIRAGLUTIDE RECOMBINANT - VICTOZA</u> | | | | | | |
| N 022341 001 | 6004297 | Jan 28, 2019 | DP | | I-750 | Aug 25, 2020 |
| | 6268343 | Aug 22, 2022 | DS DP | U-968 | M-176 | Apr 22, 2019 |
| | 8114833 | Aug 13, 2025 | DP | | | |
| | 8846618 | Jun 27, 2022 | DP | | | |
| | 9265893 | Sep 23, 2032 | DP | | | |
| | 9968659 | Jan 09, 2037 | | U-2313 | | |
| | RE41956 | Jan 21, 2021 | DP | | | |
| | RE43834 | Jan 28, 2019 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LIRAGLUTIDE RECOMBINANT - VICTOZA</u> | | | | | | |
| N 022341 001 | 6004297 | Jan 28, 2019 | DP | | I-750 | Aug 25, 2020 |
| | 6268343 | Aug 22, 2022 | DS DP U-968 | | M-176 | Apr 22, 2019 |
| | 8114833 | Aug 13, 2025 | DP | | | |
| | 8846618 | Jun 27, 2022 | DP | | | |
| | 9265893 | Sep 23, 2032 | DP | | | |
| | 9968659 | Jan 09, 2037 | | U-2313 | | |
| | RE41956 | Jan 21, 2021 | DP | | | |
| | RE43834 | Jan 28, 2019 | DP | | | |
| <u>LIRAGLUTIDE RECOMBINANT - SAXENDA</u> | | | | | | |
| N 206321 001 | 6268343 | Aug 22, 2022 | DS DP U-1255 | | | |
| | 6899699 | Jan 01, 2022 | DP | | | |
| | 8114833 | Aug 13, 2025 | DP | | | |
| | 8672898 | Jan 02, 2022 | DP | | | |
| | 8684969 | Oct 20, 2025 | DP | | | |
| | 8846618 | Jun 27, 2022 | DP | | | |
| | 8920383 | Jul 17, 2026 | DP | | | |
| | 9108002 | Jan 26, 2026 | DP | | | |
| | 9132239 | Feb 01, 2032 | DP | | | |
| | 9457154 | Sep 27, 2027 | DP | | | |
| | 9486588 | Jan 02, 2022 | DP | | | |
| | 9616180 | Jan 20, 2026 | DP | | | |
| | 9687611 | Feb 27, 2027 | DP | | | |
| | 9775953 | Jul 17, 2026 | DP | | | |
| | 9861757 | Jan 20, 2026 | DP | | | |
| | 9968659 | Jan 09, 2037 | | U-2438 | | |
| | RE46363 | Aug 03, 2026 | DP | | | |
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 021977 001 | 7105486 | Feb 24, 2023 | | U-727 | M-188 | Oct 14, 2019 |
| | 7223735 | Feb 24, 2023 | DP | | | |
| | 7655630 | Feb 24, 2023 | DS | | | |
| | 7659253 | Feb 24, 2023 | DS DP U-727 | | | |
| | 7659254 | Feb 24, 2023 | | U-1034 | | |
| | 7662787 | Feb 24, 2023 | DS | | | |
| | 7662788 | Feb 24, 2023 | | U-727 | | |
| | 7671030 | Feb 24, 2023 | DP U-727 | | | |
| | 7671031 | Feb 24, 2023 | | U-727 | | |
| | 7674774 | Feb 24, 2023 | DP U-842 | | | |
| | 7678770 | Feb 24, 2023 | | U-842 | | |
| | 7678771 | Feb 24, 2023 | DP U-842 | | | |
| | 7687466 | Feb 24, 2023 | DP | | | |
| | 7687467 | Feb 24, 2023 | DP U-842 | | | |
| | 7700561 | Feb 24, 2023 | DP | | | |
| | 7713936 | Feb 24, 2023 | | U-727 | | |
| | 7718619 | Feb 24, 2023 | DP U-842 | | | |
| | 7723305 | Feb 24, 2023 | DP U-842 | | | |
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 021977 002 | 7105486 | Feb 24, 2023 | | U-727 | M-188 | Oct 14, 2019 |
| | 7223735 | Feb 24, 2023 | DP | | | |
| | 7655630 | Feb 24, 2023 | DS | | | |
| | 7659253 | Feb 24, 2023 | DS DP U-727 | | | |
| | 7659254 | Feb 24, 2023 | | U-1034 | | |
| | 7662787 | Feb 24, 2023 | DS | | | |
| | 7662788 | Feb 24, 2023 | | U-727 | | |
| | 7671030 | Feb 24, 2023 | DP U-727 | | | |
| | 7671031 | Feb 24, 2023 | | U-727 | | |
| | 7674774 | Feb 24, 2023 | DP U-842 | | | |
| | 7678770 | Feb 24, 2023 | | U-842 | | |
| | 7678771 | Feb 24, 2023 | DP U-842 | | | |
| | 7687466 | Feb 24, 2023 | DP | | | |
| | 7687467 | Feb 24, 2023 | DP U-842 | | | |
| | 7700561 | Feb 24, 2023 | DP | | | |
| | 7713936 | Feb 24, 2023 | | U-727 | | |
| | 7718619 | Feb 24, 2023 | DP U-842 | | | |
| | 7723305 | Feb 24, 2023 | DP U-842 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 021977 003 | 7105486 | Feb 24, 2023 | U-727 | | M-188 | Oct 14, 2019 |
| | 7223735 | Feb 24, 2023 | DP | | | |
| | 7655630 | Feb 24, 2023 | DS | | | |
| | 7659253 | Feb 24, 2023 | DS DP U-727 | | | |
| | 7659254 | Feb 24, 2023 | U-1034 | | | |
| | 7662787 | Feb 24, 2023 | DS | | | |
| | 7662788 | Feb 24, 2023 | U-727 | | | |
| | 7671030 | Feb 24, 2023 | DP U-727 | | | |
| | 7671031 | Feb 24, 2023 | U-727 | | | |
| | 7674774 | Feb 24, 2023 | DP U-842 | | | |
| | 7678770 | Feb 24, 2023 | U-842 | | | |
| | 7678771 | Feb 24, 2023 | DP U-842 | | | |
| | 7687466 | Feb 24, 2023 | DP | | | |
| | 7687467 | Feb 24, 2023 | DP U-842 | | | |
| | 7700561 | Feb 24, 2023 | DP | | | |
| | 7713936 | Feb 24, 2023 | U-727 | | | |
| | 7718619 | Feb 24, 2023 | DP U-842 | | | |
| | 7723305 | Feb 24, 2023 | DP U-842 | | | |
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 021977 004 | 7105486 | Feb 24, 2023 | U-727 | | M-188 | Oct 14, 2019 |
| | 7105486 | Feb 24, 2023 | U-842 | | | |
| | 7223735 | Feb 24, 2023 | DP | | | |
| | 7655630 | Feb 24, 2023 | DS | | | |
| | 7659253 | Feb 24, 2023 | DS DP U-727 | | | |
| | 7659254 | Feb 24, 2023 | U-1034 | | | |
| | 7662787 | Feb 24, 2023 | DS | | | |
| | 7662788 | Feb 24, 2023 | U-727 | | | |
| | 7671030 | Feb 24, 2023 | DP U-727 | | | |
| | 7671031 | Feb 24, 2023 | U-727 | | | |
| | 7674774 | Feb 24, 2023 | DP U-842 | | | |
| | 7678770 | Feb 24, 2023 | U-842 | | | |
| | 7678771 | Feb 24, 2023 | DP U-842 | | | |
| | 7687466 | Feb 24, 2023 | DP | | | |
| | 7687467 | Feb 24, 2023 | DP U-842 | | | |
| | 7700561 | Feb 24, 2023 | DP | | | |
| | 7713936 | Feb 24, 2023 | U-727 | | | |
| | 7718619 | Feb 24, 2023 | DP U-842 | | | |
| | 7723305 | Feb 24, 2023 | DP U-842 | | | |
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 021977 005 | 7105486 | Feb 24, 2023 | U-842 | | M-188 | Oct 14, 2019 |
| | 7223735 | Feb 24, 2023 | DP | | | |
| | 7655630 | Feb 24, 2023 | DS | | | |
| | 7659253 | Feb 24, 2023 | DS DP U-727 | | | |
| | 7659254 | Feb 24, 2023 | U-1034 | | | |
| | 7662787 | Feb 24, 2023 | DS | | | |
| | 7662788 | Feb 24, 2023 | U-727 | | | |
| | 7671030 | Feb 24, 2023 | DP U-727 | | | |
| | 7671031 | Feb 24, 2023 | U-727 | | | |
| | 7674774 | Feb 24, 2023 | DP U-842 | | | |
| | 7678770 | Feb 24, 2023 | U-842 | | | |
| | 7678771 | Feb 24, 2023 | DP U-842 | | | |
| | 7687466 | Feb 24, 2023 | DP | | | |
| | 7687467 | Feb 24, 2023 | DP U-842 | | | |
| | 7700561 | Feb 24, 2023 | DP | | | |
| | 7713936 | Feb 24, 2023 | U-727 | | | |
| | 7718619 | Feb 24, 2023 | DP U-842 | | | |
| | 7723305 | Feb 24, 2023 | DP U-842 | | | |
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 021977 006 | 7105486 | Feb 24, 2023 | U-727 | | M-188 | Oct 14, 2019 |
| | 7105486 | Feb 24, 2023 | U-842 | | | |
| | 7223735 | Feb 24, 2023 | DP | | | |
| | 7655630 | Feb 24, 2023 | DS | | | |
| | 7659253 | Feb 24, 2023 | DS DP U-727 | | | |
| | 7659254 | Feb 24, 2023 | U-1034 | | | |
| | 7662787 | Feb 24, 2023 | DS | | | |
| | 7662788 | Feb 24, 2023 | U-727 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 021977 006 | 7671030 | Feb 24, 2023 | DP U-727 | | | |
| | 7671031 | Feb 24, 2023 | U-727 | | | |
| | 7674774 | Feb 24, 2023 | DP U-842 | | | |
| | 7678770 | Feb 24, 2023 | U-842 | | | |
| | 7678771 | Feb 24, 2023 | DP U-842 | | | |
| | 7687466 | Feb 24, 2023 | DP | | | |
| | 7687467 | Feb 24, 2023 | DP U-842 | | | |
| | 7700561 | Feb 24, 2023 | DP | | | |
| | 7713936 | Feb 24, 2023 | U-727 | | | |
| | 7718619 | Feb 24, 2023 | DP U-842 | | | |
| | 7723305 | Feb 24, 2023 | DP U-842 | | | |
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 021977 007 | 7223735 | Feb 24, 2023 | DP | | M-188 | Oct 14, 2019 |
| | 7655630 | Feb 24, 2023 | DS | | | |
| | 7659253 | Feb 24, 2023 | DS DP U-727 | | | |
| | 7659254 | Feb 24, 2023 | U-1034 | | | |
| | 7662787 | Feb 24, 2023 | DS | | | |
| | 7662788 | Feb 24, 2023 | U-727 | | | |
| | 7671030 | Feb 24, 2023 | DP U-727 | | | |
| | 7671031 | Feb 24, 2023 | U-727 | | | |
| | 7674774 | Feb 24, 2023 | DP U-842 | | | |
| | 7678770 | Feb 24, 2023 | U-842 | | | |
| | 7678771 | Feb 24, 2023 | DP | | | |
| | 7687466 | Feb 24, 2023 | DP | | | |
| | 7687467 | Feb 24, 2023 | DP U-842 | | | |
| | 7700561 | Feb 24, 2023 | DP | | | |
| | 7713936 | Feb 24, 2023 | U-727 | | | |
| | 7718619 | Feb 24, 2023 | DP U-842 | | | |
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 208510 001 | 7105486 | Feb 24, 2023 | U-727 | | M-188 | Oct 14, 2019 |
| | 7223735 | Feb 24, 2023 | DP | | | |
| | 7655630 | Feb 24, 2023 | DS DP | | | |
| | 7659253 | Feb 24, 2023 | DS DP U-727 | | | |
| | 7659254 | Feb 24, 2023 | U-727 | | | |
| | 7662787 | Feb 24, 2023 | DS | | | |
| | 7662788 | Feb 24, 2023 | U-727 | | | |
| | 7671030 | Feb 24, 2023 | DP U-727 | | | |
| | 7671031 | Feb 24, 2023 | U-727 | | | |
| | 7674774 | Feb 24, 2023 | DP U-727 | | | |
| | 7678770 | Feb 24, 2023 | U-727 | | | |
| | 7678771 | Feb 24, 2023 | DP U-727 | | | |
| | 7687466 | Feb 24, 2023 | DP | | | |
| | 7687467 | Feb 24, 2023 | DP U-727 | | | |
| | 7713936 | Feb 24, 2023 | U-727 | | | |
| | 7718619 | Feb 24, 2023 | DP U-727 | | | |
| | 7723305 | Feb 24, 2023 | DP U-727 | | | |
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 208510 002 | 7105486 | Feb 24, 2023 | U-727 | | M-188 | Oct 14, 2019 |
| | 7223735 | Feb 24, 2023 | DP | | | |
| | 7655630 | Feb 24, 2023 | DS DP | | | |
| | 7659253 | Feb 24, 2023 | DS DP U-727 | | | |
| | 7659254 | Feb 24, 2023 | U-727 | | | |
| | 7662787 | Feb 24, 2023 | DS | | | |
| | 7662788 | Feb 24, 2023 | U-727 | | | |
| | 7671030 | Feb 24, 2023 | DP U-727 | | | |
| | 7671031 | Feb 24, 2023 | U-727 | | | |
| | 7674774 | Feb 24, 2023 | DP U-727 | | | |
| | 7678770 | Feb 24, 2023 | U-727 | | | |
| | 7678771 | Feb 24, 2023 | DP U-727 | | | |
| | 7687466 | Feb 24, 2023 | DP | | | |
| | 7687467 | Feb 24, 2023 | DP U-727 | | | |
| | 7713936 | Feb 24, 2023 | U-727 | | | |
| | 7718619 | Feb 24, 2023 | DP U-727 | | | |
| | 7723305 | Feb 24, 2023 | DP U-727 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 208510 003 | 7105486 | Feb 24, 2023 | U-727 | | M-188 | Oct 14, 2019 |
| | 7223735 | Feb 24, 2023 | DP | | | |
| | 7655630 | Feb 24, 2023 | DS DP | | | |
| | 7659253 | Feb 24, 2023 | DS DP U-727 | | | |
| | 7659254 | Feb 24, 2023 | U-727 | | | |
| | 7662787 | Feb 24, 2023 | DS | | | |
| | 7662788 | Feb 24, 2023 | U-727 | | | |
| | 7671030 | Feb 24, 2023 | DP U-727 | | | |
| | 7671031 | Feb 24, 2023 | U-727 | | | |
| | 7674774 | Feb 24, 2023 | DP U-727 | | | |
| | 7678770 | Feb 24, 2023 | U-727 | | | |
| | 7678771 | Feb 24, 2023 | DP U-727 | | | |
| | 7687466 | Feb 24, 2023 | DP | | | |
| | 7687467 | Feb 24, 2023 | DP U-727 | | | |
| | 7713936 | Feb 24, 2023 | U-727 | | | |
| | 7718619 | Feb 24, 2023 | DP U-727 | | | |
| | 7723305 | Feb 24, 2023 | DP U-727 | | | |
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 208510 004 | 7105486 | Feb 24, 2023 | U-727 | | M-188 | Oct 14, 2019 |
| | 7223735 | Feb 24, 2023 | DP | | | |
| | 7655630 | Feb 24, 2023 | DS DP | | | |
| | 7659253 | Feb 24, 2023 | DS DP U-727 | | | |
| | 7659254 | Feb 24, 2023 | U-727 | | | |
| | 7662787 | Feb 24, 2023 | DS | | | |
| | 7662788 | Feb 24, 2023 | U-727 | | | |
| | 7671030 | Feb 24, 2023 | DP U-727 | | | |
| | 7671031 | Feb 24, 2023 | U-727 | | | |
| | 7674774 | Feb 24, 2023 | DP U-727 | | | |
| | 7678770 | Feb 24, 2023 | U-727 | | | |
| | 7678771 | Feb 24, 2023 | DP U-727 | | | |
| | 7687466 | Feb 24, 2023 | DP | | | |
| | 7687467 | Feb 24, 2023 | DP U-727 | | | |
| | 7713936 | Feb 24, 2023 | U-727 | | | |
| | 7718619 | Feb 24, 2023 | DP U-727 | | | |
| | 7723305 | Feb 24, 2023 | DP U-727 | | | |
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 208510 005 | 7105486 | Feb 24, 2023 | U-727 | | M-188 | Oct 14, 2019 |
| | 7223735 | Feb 24, 2023 | DP | | | |
| | 7655630 | Feb 24, 2023 | DS DP | | | |
| | 7659253 | Feb 24, 2023 | DS DP U-727 | | | |
| | 7659254 | Feb 24, 2023 | U-727 | | | |
| | 7662787 | Feb 24, 2023 | DS | | | |
| | 7662788 | Feb 24, 2023 | U-727 | | | |
| | 7671030 | Feb 24, 2023 | DP U-727 | | | |
| | 7671031 | Feb 24, 2023 | U-727 | | | |
| | 7674774 | Feb 24, 2023 | DP U-727 | | | |
| | 7678770 | Feb 24, 2023 | U-727 | | | |
| | 7678771 | Feb 24, 2023 | DP U-727 | | | |
| | 7687466 | Feb 24, 2023 | DP | | | |
| | 7687467 | Feb 24, 2023 | DP U-727 | | | |
| | 7713936 | Feb 24, 2023 | U-727 | | | |
| | 7718619 | Feb 24, 2023 | DP U-727 | | | |
| | 7723305 | Feb 24, 2023 | DP U-727 | | | |
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 208510 006 | 7105486 | Feb 24, 2023 | U-727 | | M-188 | Oct 14, 2019 |
| | 7223735 | Feb 24, 2023 | DP | | | |
| | 7655630 | Feb 24, 2023 | DS DP | | | |
| | 7659253 | Feb 24, 2023 | DS DP U-727 | | | |
| | 7659254 | Feb 24, 2023 | U-727 | | | |
| | 7662787 | Feb 24, 2023 | DS | | | |
| | 7662788 | Feb 24, 2023 | U-727 | | | |
| | 7671030 | Feb 24, 2023 | DP U-727 | | | |
| | 7671031 | Feb 24, 2023 | U-727 | | | |
| | 7674774 | Feb 24, 2023 | DP U-727 | | | |
| | 7678770 | Feb 24, 2023 | U-727 | | | |
| | 7678771 | Feb 24, 2023 | DP U-727 | | | |
| | 7687466 | Feb 24, 2023 | DP | | | |
| | 7687467 | Feb 24, 2023 | DP U-727 | | | |
| | 7713936 | Feb 24, 2023 | U-727 | | | |
| | 7718619 | Feb 24, 2023 | DP U-727 | | | |
| | 7678771 | Feb 24, 2023 | DP U-727 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u> | | | | | | |
| N 208510 006 | 7687466 | Feb 24, 2023 | DP | | | |
| | 7687467 | Feb 24, 2023 | DP U-727 | | | |
| | 7713936 | Feb 24, 2023 | U-727 | | | |
| | 7718619 | Feb 24, 2023 | DP U-727 | | | |
| | 7723305 | Feb 24, 2023 | DP U-727 | | | |
| <u>LISINAPRIL - OBRELIS</u> | | | | | | |
| N 208401 001 | 10039800 | Nov 06, 2035 | U-1723 | | | |
| | 10039800 | Nov 06, 2035 | U-185 | | | |
| | 10039800 | Nov 06, 2035 | U-1864 | | | |
| | 10039800 | Nov 06, 2035 | U-1991 | | | |
| | 10039800 | Nov 06, 2035 | U-3 | | | |
| | 10039800 | Nov 06, 2035 | U-71 | | | |
| | 10039800 | Nov 06, 2035 | U-8 | | | |
| | 9463183 | Nov 06, 2035 | DP | | | |
| | 9616096 | Nov 06, 2035 | U-1723 | | | |
| | 9616096 | Nov 06, 2035 | U-185 | | | |
| | 9616096 | Nov 06, 2035 | U-1864 | | | |
| | 9616096 | Nov 06, 2035 | U-1991 | | | |
| | 9616096 | Nov 06, 2035 | U-3 | | | |
| | 9616096 | Nov 06, 2035 | U-71 | | | |
| | 9616096 | Nov 06, 2035 | U-8 | | | |
| | 9814751 | Nov 06, 2035 | DP | | | |
| <u>LIXISENATIDE - ADLYXIN</u> | | | | | | |
| N 208471 001 | 10028910 | Nov 11, 2030 | DP | | NCE | Jul 27, 2021 |
| | 8475414 | Dec 28, 2030 | DP U-1881 | | | |
| | 8882721 | Jun 28, 2031 | DP | | | |
| | 8915888 | Jun 08, 2030 | DP U-1881 | | | |
| | 9072836 | Mar 15, 2032 | DP | | | |
| | 9084853 | Oct 05, 2031 | DP | | | |
| | 9308329 | Dec 28, 2030 | DP U-1881 | | | |
| | 9408893 | Aug 27, 2032 | U-1894 | | | |
| | 9440029 | Jan 30, 2032 | DP | | | |
| | 9511193 | Jan 19, 2032 | DP | | | |
| | 9707176 | Nov 11, 2030 | DP | | | |
| | 9821032 | May 09, 2032 | U-2200 | | | |
| | 9855388 | Apr 24, 2029 | DP U-1881 | | | |
| | 9981013 | Aug 30, 2030 | U-2297 | | | |
| | RE45313 | Jul 12, 2020 | DS DP | | | |
| <u>LIXISENATIDE - ADLYXIN</u> | | | | | | |
| N 208471 002 | 10028910 | Nov 11, 2030 | DP | | NCE | Jul 27, 2021 |
| | 8475414 | Dec 28, 2030 | DP U-1881 | | | |
| | 8882721 | Jun 28, 2031 | DP | | | |
| | 8915888 | Jun 08, 2030 | DP U-1881 | | | |
| | 9072836 | Mar 15, 2032 | DP | | | |
| | 9084853 | Oct 05, 2031 | DP | | | |
| | 9308329 | Dec 28, 2030 | DP U-1881 | | | |
| | 9408893 | Aug 27, 2032 | U-1894 | | | |
| | 9440029 | Jan 30, 2032 | DP | | | |
| | 9511193 | Jan 19, 2032 | DP | | | |
| | 9707176 | Nov 11, 2030 | DP | | | |
| | 9821032 | May 09, 2032 | U-2200 | | | |
| | 9855388 | Apr 24, 2029 | DP U-1881 | | | |
| | 9981013 | Aug 30, 2030 | U-2297 | | | |
| | RE45313 | Jul 12, 2020 | DS DP | | | |
| <u>LOFEXIDINE HYDROCHLORIDE - LUCEMYRA</u> | | | | | | |
| N 209229 001 | | | | | NCE | May 16, 2023 |
| <u>LOMITAPIDE MESYLATE - JUXTAPID</u> | | | | | | |
| N 203858 001 | 10016404 | Mar 07, 2025 | U-1316 | | ODE-36 | Dec 21, 2019 |
| | 5712279 | Feb 21, 2020 | DS DP U-1317 | | | |
| | 6492365 | Dec 10, 2019 | U-1318 | | | |
| | 7932268 | Aug 19, 2027 | U-1316 | | | |
| | 8618135 | Mar 07, 2025 | U-1316 | | | |
| | 9265758 | Mar 07, 2025 | U-1316 | | | |
| | 9364470 | Mar 07, 2025 | U-1851 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LOMITAPIDE MESYLATE - JUXTAPID</u> | | | | | | |
| N 203858 001 | 9433617 | Mar 07, 2025 | U-1316 | | | |
| | 9861622 | Mar 07, 2025 | U-1316 | | | |
| <u>LOMITAPIDE MESYLATE - JUXTAPID</u> | | | | | | |
| N 203858 002 | 10016404 | Mar 07, 2025 | U-1316 | | ODE-36 | Dec 21, 2019 |
| | 5712279 | Feb 21, 2020 | DS DP U-1317 | | | |
| | 6492365 | Dec 10, 2019 | U-1318 | | | |
| | 7932268 | Aug 19, 2027 | U-1316 | | | |
| | 8618135 | Mar 07, 2025 | U-1316 | | | |
| | 9265758 | Mar 07, 2025 | U-1316 | | | |
| | 9364470 | Mar 07, 2025 | U-1851 | | | |
| | 9433617 | Mar 07, 2025 | U-1316 | | | |
| | 9861622 | Mar 07, 2025 | U-1316 | | | |
| <u>LOMITAPIDE MESYLATE - JUXTAPID</u> | | | | | | |
| N 203858 003 | 10016404 | Mar 07, 2025 | U-1316 | | ODE-36 | Dec 21, 2019 |
| | 5712279 | Feb 21, 2020 | DS DP U-1317 | | | |
| | 6492365 | Dec 10, 2019 | U-1318 | | | |
| | 7932268 | Aug 19, 2027 | U-1316 | | | |
| | 8618135 | Mar 07, 2025 | U-1316 | | | |
| | 9265758 | Mar 07, 2025 | U-1316 | | | |
| | 9364470 | Mar 07, 2025 | U-1851 | | | |
| | 9433617 | Mar 07, 2025 | U-1316 | | | |
| | 9861622 | Mar 07, 2025 | U-1316 | | | |
| <u>LOMITAPIDE MESYLATE - JUXTAPID</u> | | | | | | |
| N 203858 004 | 10016404 | Mar 07, 2025 | U-1316 | | ODE-36 | Dec 21, 2019 |
| | 5712279 | Feb 21, 2020 | DS DP U-1317 | | | |
| | 6492365 | Dec 10, 2019 | U-1318 | | | |
| | 7932268 | Aug 19, 2027 | U-1316 | | | |
| | 8618135 | Mar 07, 2025 | U-1316 | | | |
| | 9265758 | Mar 07, 2025 | U-1316 | | | |
| | 9364470 | Mar 07, 2025 | U-1851 | | | |
| | 9433617 | Mar 07, 2025 | U-1316 | | | |
| | 9861622 | Mar 07, 2025 | U-1316 | | | |
| <u>LOMITAPIDE MESYLATE - JUXTAPID</u> | | | | | | |
| N 203858 005 | 10016404 | Mar 07, 2025 | U-1316 | | ODE-36 | Dec 21, 2019 |
| | 5712279 | Feb 21, 2020 | DS DP U-1317 | | | |
| | 6492365 | Dec 10, 2019 | U-1318 | | | |
| | 7932268 | Aug 19, 2027 | U-1316 | | | |
| | 8618135 | Mar 07, 2025 | U-1316 | | | |
| | 9265758 | Mar 07, 2025 | U-1316 | | | |
| | 9364470 | Mar 07, 2025 | U-1851 | | | |
| | 9433617 | Mar 07, 2025 | U-1316 | | | |
| | 9861622 | Mar 07, 2025 | U-1316 | | | |
| <u>LOMITAPIDE MESYLATE - JUXTAPID</u> | | | | | | |
| N 203858 006 | 10016404 | Mar 07, 2025 | U-1316 | | ODE-36 | Dec 21, 2019 |
| | 5712279 | Feb 21, 2020 | DS DP U-1317 | | | |
| | 6492365 | Dec 10, 2019 | U-1318 | | | |
| | 7932268 | Aug 19, 2027 | U-1316 | | | |
| | 8618135 | Mar 07, 2025 | U-1316 | | | |
| | 9265758 | Mar 07, 2025 | U-1316 | | | |
| | 9364470 | Mar 07, 2025 | U-1851 | | | |
| | 9433617 | Mar 07, 2025 | U-1316 | | | |
| | 9861622 | Mar 07, 2025 | U-1316 | | | |
| <u>LOPERAMIDE HYDROCHLORIDE - IMODIUM A-D EZ CHEWS</u> | | | | | | |
| N 020448 001 | 6814978 | Aug 26, 2021 | DP | | | |
| <u>LOPINAVIR; RITONAVIR - KALETRA</u> | | | | | | |
| N 021226 001 | 7141593 | May 22, 2020 | DP | | | |
| | 7432294 | May 22, 2020 | DP | | | |
| <u>LOPINAVIR; RITONAVIR - KALETRA</u> | | | | | | |
| N 021251 001 | 6911214 | Nov 28, 2021 | DP U-895 | | | |
| | 8501219 | Nov 28, 2021 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------|------------------------------------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LOPINAVIR; RITONAVIR - KALETRA</u> | | | | | | |
| N 021906 001 | 7148359 | Jul 19, 2019 | DP | | | |
| | 7364752 | Nov 10, 2020 | DP U-688 | | | |
| | 8025899 | Dec 14, 2027 | DP | | | |
| | 8025899*PED | Jun 14, 2028 | | | | |
| | 8268349 | Aug 25, 2024 | DP | | | |
| | 8309613 | Dec 24, 2024 | U-688 | | | |
| | 8377952 | Oct 22, 2027 | U-1372 | | | |
| | 8377952*PED | Apr 22, 2028 | | | | |
| | 8399015 | Aug 25, 2024 | DP | | | |
| | 8399015*PED | Feb 25, 2025 | | | | |
| | 8470347 | Sep 17, 2026 | DP | | | |
| | 8470347*PED | Mar 17, 2027 | | | | |
| | 8691878 | Aug 25, 2024 | U-1513 | | | |
| | 8691878*PED | Feb 25, 2025 | | | | |
| | <u>LOPINAVIR; RITONAVIR - KALETRA</u> | | | | | |
| N 021906 002 | 7148359 | Jul 19, 2019 | DP | | | |
| | 7364752 | Nov 10, 2020 | DP U-688 | | | |
| | 8025899 | Dec 14, 2027 | DP | | | |
| | 8025899*PED | Jun 14, 2028 | | | | |
| | 8268349 | Aug 25, 2024 | DP | | | |
| | 8309613 | Dec 24, 2024 | U-688 | | | |
| | 8377952 | Oct 22, 2027 | U-1372 | | | |
| | 8377952*PED | Apr 22, 2028 | | | | |
| | 8399015 | Aug 25, 2024 | DP | | | |
| | 8399015*PED | Feb 25, 2025 | | | | |
| | 8470347 | Sep 17, 2026 | DP | | | |
| | 8470347*PED | Mar 17, 2027 | | | | |
| | 8691878 | Aug 25, 2024 | U-1513 | | | |
| | 8691878*PED | Feb 25, 2025 | | | | |
| | <u>LORCASERIN HYDROCHLORIDE - BELVIO</u> | | | | | |
| N 022529 001 | 6953787 | Apr 10, 2023 | DS DP U-1252 | | | |
| | 6953787 | Apr 10, 2023 | DS DP U-1253 | | | |
| | 6953787 | Apr 10, 2023 | DS DP U-1254 | | | |
| | 6953787 | Apr 10, 2023 | DS DP U-1255 | | | |
| | 7514422 | Apr 10, 2023 | U-1252 | | | |
| | 7514422 | Apr 10, 2023 | U-1253 | | | |
| | 7514422 | Apr 10, 2023 | U-1254 | | | |
| | 7514422 | Apr 10, 2023 | U-1255 | | | |
| | 7977329 | Apr 10, 2023 | DS DP U-1252 | | | |
| | 7977329 | Apr 10, 2023 | DS DP U-1253 | | | |
| | 7977329 | Apr 10, 2023 | DS DP U-1254 | | | |
| | 7977329 | Apr 10, 2023 | DS DP U-1255 | | | |
| | 8168624 | Apr 18, 2029 | DS DP | | | |
| | 8207158 | Apr 10, 2023 | U-1252 | | | |
| | 8207158 | Apr 10, 2023 | U-1253 | | | |
| | 8207158 | Apr 10, 2023 | U-1254 | | | |
| | 8207158 | Apr 10, 2023 | U-1255 | | | |
| | 8273734 | Apr 10, 2023 | U-1254 | | | |
| | 8273734 | Apr 10, 2023 | U-1255 | | | |
| | 8367657 | Apr 10, 2023 | DS DP U-1252 | | | |
| | 8367657 | Apr 10, 2023 | DS DP U-1253 | | | |
| | 8367657 | Apr 10, 2023 | DS DP U-1254 | | | |
| | 8367657 | Apr 10, 2023 | DS DP U-1255 | | | |
| | 8546379 | Apr 10, 2023 | DS DP U-1252 | | | |
| | 8546379 | Apr 10, 2023 | DS DP U-1253 | | | |
| | 8546379 | Apr 10, 2023 | DS DP U-1254 | | | |
| | 8546379 | Apr 10, 2023 | DS DP U-1255 | | | |
| | 8575149 | Apr 10, 2023 | U-1452 | | | |
| | 8697686 | Dec 20, 2025 | DS DP | | | |
| | 8946207 | Jun 16, 2024 | DP | | | |
| | 8980881 | Dec 20, 2025 | U-1252 | | | |
| | 8980881 | Dec 20, 2025 | U-1253 | | | |
| | 8980881 | Dec 20, 2025 | U-1254 | | | |
| | 8980881 | Dec 20, 2025 | U-1255 | | | |
| 8999970 | Feb 07, 2033 | U-1688 | | | | |
| 8999970 | Feb 07, 2033 | U-1689 | | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LORCASERIN HYDROCHLORIDE - BELVIO</u> | | | | | | |
| N 022529 001 | 8999970 | Feb 07, 2033 | U-1692 | | | |
| | 9169213 | Dec 06, 2032 | U-1762 | | | |
| | 9169213 | Dec 06, 2032 | U-1763 | | | |
| | 9169213 | Dec 06, 2032 | U-1764 | | | |
| | 9169213 | Dec 06, 2032 | U-1765 | | | |
| | 9770455 | Aug 31, 2031 | U-2110 | | | |
| <u>LORCASERIN HYDROCHLORIDE - BELVIO XR</u> | | | | | | |
| N 208524 001 | 6953787 | Apr 10, 2023 | DS DP U-1252 | | | |
| | 6953787 | Apr 10, 2023 | DS DP U-1253 | | | |
| | 6953787 | Apr 10, 2023 | DS DP U-1254 | | | |
| | 6953787 | Apr 10, 2023 | DS DP U-1255 | | | |
| | 7514422 | Apr 10, 2023 | U-1252 | | | |
| | 7514422 | Apr 10, 2023 | U-1253 | | | |
| | 7514422 | Apr 10, 2023 | U-1254 | | | |
| | 7514422 | Apr 10, 2023 | U-1255 | | | |
| | 7977329 | Apr 10, 2023 | DS DP U-1252 | | | |
| | 7977329 | Apr 10, 2023 | DS DP U-1253 | | | |
| | 7977329 | Apr 10, 2023 | DS DP U-1254 | | | |
| | 7977329 | Apr 10, 2023 | DS DP U-1255 | | | |
| | 8168624 | Apr 18, 2029 | DS DP | | | |
| | 8207158 | Apr 10, 2023 | U-1252 | | | |
| | 8207158 | Apr 10, 2023 | U-1253 | | | |
| | 8207158 | Apr 10, 2023 | U-1254 | | | |
| | 8207158 | Apr 10, 2023 | U-1255 | | | |
| | 8273734 | Apr 10, 2023 | U-1254 | | | |
| | 8273734 | Apr 10, 2023 | U-1255 | | | |
| | 8367657 | Apr 10, 2023 | DS DP U-1252 | | | |
| | 8367657 | Apr 10, 2023 | DS DP U-1253 | | | |
| | 8367657 | Apr 10, 2023 | DS DP U-1254 | | | |
| | 8367657 | Apr 10, 2023 | DS DP U-1255 | | | |
| | 8546379 | Apr 10, 2023 | DS DP U-1252 | | | |
| | 8546379 | Apr 10, 2023 | DS DP U-1253 | | | |
| | 8546379 | Apr 10, 2023 | DS DP U-1254 | | | |
| | 8546379 | Apr 10, 2023 | DS DP U-1255 | | | |
| | 8575149 | Apr 10, 2023 | U-1452 | | | |
| | 8697686 | Dec 20, 2025 | DS DP | | | |
| | 8946207 | Jun 16, 2024 | DP | | | |
| | 8980881 | Dec 20, 2025 | U-1252 | | | |
| | 8980881 | Dec 20, 2025 | U-1253 | | | |
| | 8980881 | Dec 20, 2025 | U-1254 | | | |
| | 8980881 | Dec 20, 2025 | U-1255 | | | |
| | 8999970 | Feb 07, 2033 | U-1688 | | | |
| | 8999970 | Feb 07, 2033 | U-1689 | | | |
| | 8999970 | Feb 07, 2033 | U-1692 | | | |
| | 9169213 | Dec 06, 2032 | U-1884 | | | |
| | 9169213 | Dec 06, 2032 | U-1885 | | | |
| | 9169213 | Dec 06, 2032 | U-1886 | | | |
| | 9169213 | Dec 06, 2032 | U-1887 | | | |
| | 9770455 | Aug 31, 2031 | U-2110 | | | |
| <u>LORLATINIB - LORBRENA</u> | | | | | | |
| N 210868 001 | 8680111 | Mar 05, 2033 | DS DP | | NCE | Nov 02, 2023 |
| | | | | | ODE-217 | Nov 02, 2025 |
| | | | | | ODE-218 | Nov 02, 2025 |
| | | | | | ODE-219 | Nov 02, 2025 |
| <u>LORLATINIB - LORBRENA</u> | | | | | | |
| N 210868 002 | 8680111 | Mar 05, 2033 | DS DP | | NCE | Nov 02, 2023 |
| | | | | | ODE-217 | Nov 02, 2025 |
| | | | | | ODE-218 | Nov 02, 2025 |
| | | | | | ODE-219 | Nov 02, 2025 |
| <u>LOTEPREDNOL ETABONATE - LOTEMAX</u> | | | | | | |
| N 202872 001 | | | | | M-229 | Jul 20, 2021 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LOTEPREDNOL ETABONATE - INVELTYS</u> | | | | | | |
| N 210565 | 001 | | | | NP | Aug 22, 2021 |
| <u>LOXAPINE - ADASUVE</u> | | | | | | |
| N 022549 | 001 | 6716416 | May 20, 2022 | DP | | |
| | | 7052679 | Oct 26, 2021 | DP | | |
| | | 7078020 | Oct 26, 2021 | DP | U-1375 | |
| | | 7090830 | Oct 26, 2021 | DP | | |
| | | 7458374 | Aug 18, 2024 | DP | | |
| | | 7537009 | Oct 28, 2024 | DP | | |
| | | 7585493 | Oct 26, 2021 | DP | | |
| | | 7601337 | Oct 26, 2021 | DP | | |
| | | 8074644 | Jul 25, 2022 | DP | | |
| | | 8173107 | Oct 26, 2021 | DP | | |
| | | 8235037 | Oct 26, 2021 | DP | | |
| | | 8387612 | Oct 23, 2026 | DP | | |
| | | 8955512 | Oct 26, 2021 | DP | | |
| | | 8991387 | May 21, 2024 | DP | | |
| | | 9370629 | May 20, 2024 | DP | | |
| | | 9439907 | Oct 26, 2021 | DP | | |
| | | 9440034 | Oct 26, 2021 | DP | | |
| | | 9687487 | Oct 26, 2021 | DS DP | | |
| <u>LUBIPROSTONE - AMITIZA</u> | | | | | | |
| N 021908 | 001 | 6414016 | Sep 05, 2020 | U-1392 | M-225 | Apr 26, 2021 |
| | | 6414016 | Sep 05, 2020 | U-717 | | |
| | | 6583174 | Oct 16, 2020 | DP | | |
| | | 6982283 | Dec 04, 2022 | U-1391 | | |
| | | 7064148 | Aug 30, 2022 | U-1404 | | |
| | | 7064148 | Aug 30, 2022 | U-739 | | |
| | | 7417067 | Oct 16, 2020 | DP | | |
| | | 8026393 | Oct 25, 2027 | DP | | |
| | | 8071613 | Sep 05, 2020 | U-1203 | | |
| | | 8071613 | Sep 05, 2020 | U-1393 | | |
| | | 8088934 | May 18, 2021 | DS | | |
| | | 8097649 | Oct 16, 2020 | DP | | |
| | | 8097653 | Nov 14, 2022 | U-1214 | | |
| | | 8097653 | Nov 14, 2022 | U-1394 | | |
| | | 8114890 | Sep 05, 2020 | DP | | |
| | | 8338639 | Jan 23, 2027 | DP | | |
| | | 8389542 | Nov 14, 2022 | DP | U-1345 | |
| | | 8389542 | Nov 14, 2022 | DP | U-1395 | |
| | | 8748481 | Sep 01, 2025 | U-1520 | | |
| | | 8779187 | Jul 23, 2027 | DP | | |
| <u>LUBIPROSTONE - AMITIZA</u> | | | | | | |
| N 021908 | 002 | 6414016 | Sep 05, 2020 | U-874 | M-225 | Apr 26, 2021 |
| | | 6583174 | Oct 16, 2020 | DP | | |
| | | 7064148 | Aug 30, 2022 | U-739 | | |
| | | 7064148 | Aug 30, 2022 | U-873 | | |
| | | 7417067 | Oct 16, 2020 | DP | | |
| | | 7795312 | Sep 17, 2024 | U-1085 | | |
| | | 8026393 | Oct 25, 2027 | DP | | |
| | | 8071613 | Sep 05, 2020 | U-1202 | | |
| | | 8088934 | May 18, 2021 | DS | | |
| | | 8097649 | Oct 16, 2020 | DP | | |
| | | 8114890 | Sep 05, 2020 | DP | | |
| | | 8338639 | Jan 23, 2027 | DP | | |
| | | 8748481 | Sep 01, 2025 | U-1519 | | |
| | | 8779187 | Jan 23, 2027 | DP | | |
| <u>LULICONAZOLE - LUZU</u> | | | | | | |
| N 204153 | 001 | 5900488 | Jan 18, 2020 | DS DP | NPP | Feb 20, 2021 |
| | | 8980931 | Apr 28, 2034 | DP | | |
| | | 9012484 | Sep 06, 2033 | DS DP | U-540 | |
| | | 9199977 | Sep 06, 2033 | DS DP | | |
| | | 9453006 | Sep 06, 2033 | DS | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LURASIDONE HYDROCHLORIDE - LATUDA</u> | | | | | | |
| N 200603 001 | 5532372*PED | Jan 02, 2019 | | | M-195 | Jan 27, 2020 |
| | 8729085 | May 26, 2026 | DP | | NPP | Jan 27, 2020 |
| | 8729085*PED | Nov 26, 2026 | | | NPP | Mar 05, 2021 |
| | 8883794 | May 26, 2026 | DP | | PED | Jul 27, 2020 |
| | 8883794*PED | Nov 26, 2026 | | | PED | Jul 27, 2020 |
| | 9174975 | Feb 20, 2024 | | U-1770 | | |
| | 9174975*PED | Aug 20, 2024 | | | | |
| | 9259423 | May 23, 2031 | | U-1822 | | |
| | 9259423*PED | Nov 23, 2031 | | | | |
| | 9555027 | May 26, 2026 | DP | U-543 | | |
| | 9815827 | Feb 20, 2024 | | U-2166 | | |
| | 9815827 | Feb 20, 2024 | | U-543 | | |
| | 9827242 | May 23, 2031 | | U-2199 | | |
| | 9827242 | May 23, 2031 | | U-2201 | | |
| | 9907794 | May 26, 2026 | | DP | | |
| | RE45573 | Jun 23, 2025 | DS | | | |
| | RE45573*PED | Dec 23, 2025 | | | | |
| <u>LURASIDONE HYDROCHLORIDE - LATUDA</u> | | | | | | |
| N 200603 002 | 5532372*PED | Jan 02, 2019 | | | NPP | Jan 27, 2020 |
| | 8729085 | May 26, 2026 | DP | | NPP | Mar 05, 2021 |
| | 8729085*PED | Nov 26, 2026 | | | PED | Jul 27, 2020 |
| | 8883794 | May 26, 2026 | DP | | | |
| | 8883794*PED | Nov 26, 2026 | | | | |
| | 9174975 | Feb 20, 2024 | | U-1770 | | |
| | 9174975*PED | Aug 20, 2024 | | | | |
| | 9259423 | May 23, 2031 | | U-1822 | | |
| | 9259423*PED | Nov 23, 2031 | | | | |
| | 9555027 | May 26, 2026 | DP | U-543 | | |
| | 9815827 | Feb 20, 2024 | | U-2166 | | |
| | 9815827 | Feb 20, 2024 | | U-543 | | |
| | 9827242 | May 23, 2031 | | U-2199 | | |
| | 9827242 | May 23, 2031 | | U-2201 | | |
| | 9907794 | May 26, 2026 | | DP | | |
| | RE45573 | Jun 23, 2025 | DS | | | |
| | RE45573*PED | Dec 23, 2025 | | | | |
| <u>LURASIDONE HYDROCHLORIDE - LATUDA</u> | | | | | | |
| N 200603 003 | 5532372*PED | Jan 02, 2019 | | | M-195 | Jan 27, 2020 |
| | 8729085 | May 26, 2026 | DP | | NPP | Jan 27, 2020 |
| | 8729085*PED | Nov 26, 2026 | | | NPP | Mar 05, 2021 |
| | 8883794 | May 26, 2026 | DP | | PED | Jul 27, 2020 |
| | 8883794*PED | Nov 26, 2026 | | | PED | Jul 27, 2020 |
| | 9174975 | Feb 20, 2024 | | U-1770 | | |
| | 9174975*PED | Aug 20, 2024 | | | | |
| | 9259423 | May 23, 2031 | | U-1822 | | |
| | 9259423*PED | Nov 23, 2031 | | | | |
| | 9555027 | May 26, 2026 | DP | U-543 | | |
| | 9815827 | Feb 20, 2024 | | U-2166 | | |
| | 9815827 | Feb 20, 2024 | | U-543 | | |
| | 9827242 | May 23, 2031 | | U-2199 | | |
| | 9827242 | May 23, 2031 | | U-2201 | | |
| | 9907794 | May 26, 2026 | | DP | | |
| | RE45573 | Jun 23, 2025 | DS | | | |
| | RE45573*PED | Dec 23, 2025 | | | | |
| <u>LURASIDONE HYDROCHLORIDE - LATUDA</u> | | | | | | |
| N 200603 004 | 5532372*PED | Jan 02, 2019 | | | | |
| | 8729085 | May 26, 2026 | DP | | | |
| | 8729085*PED | Nov 26, 2026 | | | | |
| | 8883794 | May 26, 2026 | DP | | | |
| | 8883794*PED | Nov 26, 2026 | | | | |
| | 9174975 | Feb 20, 2024 | | U-1770 | | |
| | 9174975*PED | Aug 20, 2024 | | | | |
| | 9259423 | May 23, 2031 | | U-1822 | | |
| | 9259423*PED | Nov 23, 2031 | | | | |
| | 9555027 | May 26, 2026 | DP | U-543 | | |
| | 9815827 | Feb 20, 2024 | | U-2166 | | |
| | 9815827 | Feb 20, 2024 | | U-543 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>LURASIDONE HYDROCHLORIDE - LATUDA</u> | | | | | | |
| N 200603 004 | 9827242 | May 23, 2031 | | U-2199 | | |
| | 9827242 | May 23, 2031 | | U-2201 | | |
| | 9907794 | May 26, 2026 | DP | | | |
| | RE45573 | Jun 23, 2025 | DS | | | |
| | RE45573*PED | Dec 23, 2025 | | | | |
| <u>LURASIDONE HYDROCHLORIDE - LATUDA</u> | | | | | | |
| N 200603 005 | 5532372*PED | Jan 02, 2019 | | | M-195 | Jan 27, 2020 |
| | 8729085 | May 26, 2026 | DP | | NPP | Jan 27, 2020 |
| | 8729085*PED | Nov 26, 2026 | | | NPP | Mar 05, 2021 |
| | 8883794 | May 26, 2026 | DP | | PED | Jul 27, 2020 |
| | 8883794*PED | Nov 26, 2026 | | | PED | Jul 27, 2020 |
| | 9174975 | Feb 20, 2024 | | U-1770 | | |
| | 9174975*PED | Aug 20, 2024 | | | | |
| | 9259423 | May 23, 2031 | | U-1822 | | |
| | 9259423*PED | Nov 23, 2031 | | | | |
| | 9555027 | May 26, 2026 | DP | U-543 | | |
| | 9815827 | Feb 20, 2024 | | U-2166 | | |
| | 9815827 | Feb 20, 2024 | | U-543 | | |
| | 9827242 | May 23, 2031 | | U-2199 | | |
| | 9827242 | May 23, 2031 | | U-2201 | | |
| | 9907794 | May 26, 2026 | DP | | | |
| | RE45573 | Jun 23, 2025 | DS | | | |
| | RE45573*PED | Dec 23, 2025 | | | | |
| <u>LUSUTROMBOPAG - MULPLETA</u> | | | | | | |
| N 210923 001 | 7601746 | Sep 05, 2024 | DS DP | U-2344 | NCE | Jul 31, 2023 |
| | 8530668 | Jan 21, 2030 | DS DP | | | |
| | 8889722 | Jul 29, 2028 | DS DP | | | |
| | 9427402 | Sep 29, 2031 | DP | | | |
| <u>LUTETIUM DOTATATE LU-177 - LUTATHERA</u> | | | | | | |
| N 208700 001 | | | | | NCE | Jan 26, 2023 |
| | | | | | ODE-166 | Jan 26, 2025 |
| <u>MACIMORELIN ACETATE - MACRILEN</u> | | | | | | |
| N 205598 001 | 6861409 | Aug 01, 2022 | DS DP | U-2220 | NCE | Dec 20, 2022 |
| | 8192719 | Oct 12, 2027 | | U-2220 | ODE-170 | Dec 20, 2024 |
| <u>MACITENTAN - OPSUMIT</u> | | | | | | |
| N 204410 001 | 7094781 | Dec 05, 2025 | DS DP | | ODE-54 | Oct 18, 2020 |
| | 8268847 | Apr 18, 2029 | | U-1446 | | |
| | 8367685 | Oct 04, 2028 | DP | U-1445 | | |
| | 9265762 | May 29, 2027 | DP | U-1820 | | |
| <u>MAGNESIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE - NORMOCARB HF 25</u> | | | | | | |
| N 021910 001 | 7300674 | Mar 04, 2023 | DP | U-785 | | |
| <u>MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE - SUPREP BOWEL PREP KIT</u> | | | | | | |
| N 022372 001 | 6946149 | Mar 07, 2023 | DP | U-837 | | |
| <u>MALATHION - OVIDE</u> | | | | | | |
| N 018613 001 | 7560445 | Feb 01, 2027 | DS DP | U-986 | | |
| | 7977324 | Aug 14, 2026 | DP | | | |
| <u>MARAVIROC - SELZENTRY</u> | | | | | | |
| N 022128 001 | 6586430 | Dec 01, 2019 | DS DP | U-824 | NPP | Nov 04, 2019 |
| | 6667314 | Aug 06, 2021 | DS DP | U-824 | NS | Nov 04, 2019 |
| | 7368460 | Nov 25, 2022 | | U-824 | | |
| | 7576097 | May 25, 2021 | DS | | | |
| <u>MARAVIROC - SELZENTRY</u> | | | | | | |
| N 022128 002 | 6586430 | Dec 01, 2019 | DS DP | U-824 | NPP | Nov 04, 2019 |
| | 6667314 | Aug 06, 2021 | DS DP | U-824 | NS | Nov 04, 2019 |
| | 7368460 | Nov 25, 2022 | | U-824 | | |
| | 7576097 | May 25, 2021 | DS | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>MARAVIROC - SELZENTRY</u> | | | | | | |
| N 022128 | 003 | 6586430 | Dec 01, 2019 | DS DP U-824 | NPP | Nov 04, 2019 |
| | | 6667314 | Aug 06, 2021 | DS DP U-824 | NS | Nov 04, 2019 |
| | | 7368460 | Nov 25, 2022 | U-824 | | |
| | | 7576097 | May 25, 2021 | DS | | |
| <u>MARAVIROC - SELZENTRY</u> | | | | | | |
| N 022128 | 004 | 6586430 | Dec 01, 2019 | DS DP U-824 | NPP | Nov 04, 2019 |
| | | 6667314 | Aug 06, 2021 | DS DP U-824 | NS | Nov 04, 2019 |
| | | 7368460 | Nov 25, 2022 | U-824 | | |
| | | 7576097 | May 25, 2021 | DS | | |
| <u>MARAVIROC - SELZENTRY</u> | | | | | | |
| N 208984 | 001 | 6586430 | Dec 01, 2019 | DS DP U-824 | NP | Nov 04, 2019 |
| | | 6667314 | Aug 06, 2021 | DS DP U-824 | | |
| | | 7368460 | Nov 25, 2022 | U-824 | | |
| | | 7576097 | May 25, 2021 | DS | | |
| <u>MEBENDAZOLE - VERMOX</u> | | | | | | |
| N 208398 | 001 | | | | NS | Oct 19, 2019 |
| <u>MECHLORETHAMINE HYDROCHLORIDE - VALCHLOR</u> | | | | | | |
| N 202317 | 001 | 7838564 | Mar 07, 2026 | DP | ODE-51 | Aug 23, 2020 |
| | | 7872050 | Jul 08, 2029 | U-1427 | | |
| | | 8450375 | Mar 07, 2026 | DP | | |
| | | 8501818 | Mar 07, 2026 | DP | | |
| | | 8501819 | Mar 07, 2026 | U-1427 | | |
| | | 9382191 | Mar 07, 2026 | DP | | |
| <u>MEDROXYPROGESTERONE ACETATE - DEPO-SUBQ PROVERA 104</u> | | | | | | |
| N 021583 | 001 | 6495534 | May 15, 2020 | DP | | |
| <u>MEGESTROL ACETATE - MEGACE ES</u> | | | | | | |
| N 021778 | 001 | 6592903 | Sep 21, 2020 | DP | | |
| | | 7101576 | Apr 22, 2024 | U-755 | | |
| | | 9040088 | Apr 22, 2024 | U-755 | | |
| | | 9101540 | Apr 22, 2024 | DP U-755 | | |
| | | 9101549 | Apr 22, 2024 | U-755 | | |
| | | 9107827 | Apr 22, 2024 | U-755 | | |
| <u>MELOXICAM - MOBIC</u> | | | | | | |
| N 021530 | 001 | 6184220 | Mar 25, 2019 | DP | | |
| <u>MELOXICAM - VIVLODEX</u> | | | | | | |
| N 207233 | 001 | 9526734 | Mar 31, 2033 | DP | | |
| | | 9649318 | Mar 31, 2035 | DP | | |
| | | 9808468 | Mar 31, 2035 | U-2160 | | |
| | | 9808468 | Mar 31, 2035 | U-2165 | | |
| <u>MELOXICAM - VIVLODEX</u> | | | | | | |
| N 207233 | 002 | 9526734 | Mar 31, 2033 | DP | | |
| | | 9649318 | Mar 31, 2035 | DP | | |
| | | 9808468 | Mar 31, 2035 | U-2160 | | |
| | | 9808468 | Mar 31, 2035 | U-2165 | | |
| <u>MELOXICAM - OMIIZ ODT</u> | | | | | | |
| N 211210 | 001 | 8545879 | Aug 31, 2030 | DP | | |
| <u>MELOXICAM - OMIIZ ODT</u> | | | | | | |
| N 211210 | 002 | 8545879 | Aug 31, 2030 | DP | | |
| <u>MELPHALAN HYDROCHLORIDE - EVOMELA</u> | | | | | | |
| N 207155 | 001 | 10040872 | Jan 30, 2034 | DP | ODE-110 | Mar 10, 2023 |
| | | 8410077 | Mar 13, 2029 | DP | | |
| | | 9200088 | Mar 13, 2029 | DP | | |
| | | 9493582 | Feb 27, 2033 | DP | | |
| <u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u> | | | | | | |
| N 022525 | 001 | 8039009 | Mar 24, 2029 | U-539 | | |
| | | 8039009*PED | Sep 24, 2029 | | | |
| | | 8168209 | Nov 22, 2025 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u> | | | | | | |
| N 022525 001 | 8168209*PED | May 22, 2026 | | | | |
| | 8173708 | Nov 22, 2025 | U-539 | | | |
| | 8173708*PED | May 22, 2026 | | | | |
| | 8283379 | Nov 22, 2025 | U-539 | | | |
| | 8283379*PED | May 22, 2026 | | | | |
| | 8329752 | Nov 22, 2025 | DP | | | |
| | 8329752*PED | May 22, 2026 | | | | |
| | 8362085 | Nov 22, 2025 | U-539 | | | |
| | 8362085*PED | May 22, 2026 | | | | |
| <u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u> | | | | | | |
| N 022525 002 | 8039009 | Mar 24, 2029 | U-539 | | | |
| | 8039009*PED | Sep 24, 2029 | | | | |
| | 8168209 | Nov 22, 2025 | DP | | | |
| | 8168209*PED | May 22, 2026 | | | | |
| | 8173708 | Nov 22, 2025 | U-539 | | | |
| | 8173708*PED | May 22, 2026 | | | | |
| | 8283379 | Nov 22, 2025 | U-539 | | | |
| | 8283379*PED | May 22, 2026 | | | | |
| | 8329752 | Nov 22, 2025 | DP | | | |
| | 8329752*PED | May 22, 2026 | | | | |
| | 8362085 | Nov 22, 2025 | U-539 | | | |
| | 8362085*PED | May 22, 2026 | | | | |
| <u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u> | | | | | | |
| N 022525 003 | 8039009 | Mar 24, 2029 | U-539 | | | |
| | 8039009*PED | Sep 24, 2029 | | | | |
| | 8168209 | Nov 22, 2025 | DP | | | |
| | 8168209*PED | May 22, 2026 | | | | |
| | 8173708 | Nov 22, 2025 | U-539 | | | |
| | 8173708*PED | May 22, 2026 | | | | |
| | 8283379 | Nov 22, 2025 | U-539 | | | |
| | 8283379*PED | May 22, 2026 | | | | |
| | 8329752 | Nov 22, 2025 | DP | | | |
| | 8329752*PED | May 22, 2026 | | | | |
| | 8362085 | Nov 22, 2025 | U-539 | | | |
| | 8362085*PED | May 22, 2026 | | | | |
| <u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u> | | | | | | |
| N 022525 004 | 8039009 | Mar 24, 2029 | U-539 | | | |
| | 8039009*PED | Sep 24, 2029 | | | | |
| | 8168209 | Nov 22, 2025 | DP | | | |
| | 8168209*PED | May 22, 2026 | | | | |
| | 8173708 | Nov 22, 2025 | U-539 | | | |
| | 8173708*PED | May 22, 2026 | | | | |
| | 8283379 | Nov 22, 2025 | U-539 | | | |
| | 8283379*PED | May 22, 2026 | | | | |
| | 8329752 | Nov 22, 2025 | DP | | | |
| | 8329752*PED | May 22, 2026 | | | | |
| | 8362085 | Nov 22, 2025 | U-539 | | | |
| | 8362085*PED | May 22, 2026 | | | | |
| | 8598233 | Nov 22, 2025 | DP | | | |
| | 8598233*PED | May 22, 2026 | | | | |
| <u>MENTHOL; METHYL SALICYLATE - SALONPAS</u> | | | | | | |
| N 022029 001 | 8809615 | Jan 03, 2030 | DP | | | |
| | 9233184 | Aug 01, 2027 | DP | | | |
| <u>MENTHOL; METHYL SALICYLATE - SALONPAS</u> | | | | | | |
| N 022029 002 | 8809615 | Jan 03, 2030 | DP | | | |
| | 9233184 | Aug 01, 2027 | DP | | | |
| <u>MEQUINOL; TRETINOLIN - SOLAGE</u> | | | | | | |
| N 020922 001 | 6353029 | Aug 24, 2020 | | | | |
| <u>MERCAPTOPYRINE - PURIXAN</u> | | | | | | |
| N 205919 001 | | | | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>MEROPENEM; VABORBACTAM - VABOMERE</u> | | | | | | |
| N 209776 | 001 8680136 | Aug 17, 2031 | DS DP | | | |
| | 9694025 | Aug 08, 2031 | | U-2120 | | |
| <u>MESALAMINE - SFROWASA</u> | | | | | | |
| N 019618 | 002 7645801 | Jul 24, 2027 | DS DP | | | |
| <u>MESALAMINE - CANASA</u> | | | | | | |
| N 021252 | 001 | | | | M-187 | Sep 02, 2019 |
| <u>MESALAMINE - CANASA</u> | | | | | | |
| N 021252 | 002 8217083 | Jun 06, 2028 | DP | | | |
| | 8436051 | Jun 06, 2028 | DP | | | |
| <u>MESALAMINE - ASACOL HD</u> | | | | | | |
| N 021830 | 001 6893662 | Nov 15, 2021 | DP | U-141 | | |
| | 8580302 | Nov 15, 2021 | DP | | | |
| | 9089492 | Nov 15, 2021 | DP | | | |
| <u>MESALAMINE - LIALDA</u> | | | | | | |
| N 022000 | 001 6773720 | Jun 08, 2020 | DP | | | |
| <u>MESALAMINE - APRISO</u> | | | | | | |
| N 022301 | 001 8865688 | May 01, 2030 | | U-1310 | | |
| <u>MESALAMINE - DELZICOL</u> | | | | | | |
| N 204412 | 001 6649180 | Apr 13, 2020 | DP | | | |
| <u>METAXALONE - SKELAXIN</u> | | | | | | |
| N 013217 | 003 7122566 | Feb 06, 2026 | | U-915 | | |
| | 7714006 | Dec 03, 2021 | | U-1050 | | |
| <u>METFORMIN HYDROCHLORIDE - FORTAMET</u> | | | | | | |
| N 021574 | 001 6790459 | Mar 17, 2021 | | U-604 | | |
| | 6866866 | Mar 17, 2021 | DP | | | |
| <u>METFORMIN HYDROCHLORIDE - FORTAMET</u> | | | | | | |
| N 021574 | 002 6790459 | Mar 17, 2021 | | U-604 | | |
| | 6866866 | Mar 17, 2021 | DP | | | |
| <u>METFORMIN HYDROCHLORIDE - RIOMET</u> | | | | | | |
| N 021591 | 001 6890957 | Sep 14, 2023 | DP | | | |
| <u>METFORMIN HYDROCHLORIDE - GLUMETZA</u> | | | | | | |
| N 021748 | 001 6488962 | Jun 20, 2020 | DS DP | | | |
| | 6723340 | Oct 25, 2021 | DS DP | | | |
| <u>METFORMIN HYDROCHLORIDE - GLUMETZA</u> | | | | | | |
| N 021748 | 002 7780987 | Mar 23, 2025 | DS DP | | | |
| | 8323692 | Mar 30, 2023 | DP | | | |
| <u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET</u> | | | | | | |
| N 021842 | 001 9101660 | Jan 22, 2027 | DP | | | |
| | 9320714 | Feb 03, 2029 | DP | | | |
| <u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET</u> | | | | | | |
| N 021842 | 002 9101660 | Jan 22, 2027 | DP | | | |
| | 9320714 | Feb 03, 2029 | DP | | | |
| <u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</u> | | | | | | |
| N 022024 | 001 6790459 | Mar 17, 2021 | | U-974 | | |
| | 6866866 | Mar 17, 2021 | DP | | | |
| | 7785627 | Jul 31, 2026 | DP | | | |
| | 7959946 | Jul 31, 2026 | DP | | | |
| | 8470368 | Sep 19, 2023 | DP | | | |
| | 8668931 | Sep 19, 2023 | DP | | | |
| | 9060941 | Sep 19, 2023 | DP | | | |
| <u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</u> | | | | | | |
| N 022024 | 002 6790459 | Mar 17, 2021 | | U-974 | | |
| | 6866866 | Mar 17, 2021 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</u> | | | | | | |
| N 022024 | 002 | 7785627 | Jul 31, 2026 | DP | | |
| | | 7959946 | Jul 31, 2026 | DP | | |
| | | 8470368 | Sep 19, 2023 | DP | | |
| | | 8668931 | Sep 19, 2023 | DP | | |
| | | 9060941 | Sep 19, 2023 | DP | | |
| <u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u> | | | | | | |
| N 021410 | 001 | 8236345 | Oct 07, 2022 | DP | | |
| <u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u> | | | | | | |
| N 021410 | 002 | 8236345 | Oct 07, 2022 | DP | | |
| <u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u> | | | | | | |
| N 021410 | 003 | 8236345 | Oct 07, 2022 | DP | | |
| <u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u> | | | | | | |
| N 021410 | 004 | 8236345 | Oct 07, 2022 | DP | | |
| <u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u> | | | | | | |
| N 021410 | 005 | 8236345 | Oct 07, 2022 | DP | | |
| <u>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</u> | | | | | | |
| N 200678 | 001 | 8628799 | Jul 13, 2025 | DP | M-175 | Apr 05, 2019 |
| | | 9339472 | Jul 13, 2025 | DP | M-198 | Feb 27, 2020 |
| | | RE44186 | Jul 31, 2023 | DS DP U-1097 | | |
| | | RE44186 | Jul 31, 2023 | DS DP U-1838 | | |
| <u>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</u> | | | | | | |
| N 200678 | 002 | 9339472 | Jul 13, 2025 | DP | M-175 | Apr 05, 2019 |
| | | RE44186 | Jul 31, 2023 | DS DP U-1097 | M-198 | Feb 27, 2020 |
| | | RE44186 | Jul 31, 2023 | DS DP U-1838 | | |
| <u>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</u> | | | | | | |
| N 200678 | 003 | 9339472 | Jul 13, 2025 | DP | M-175 | Apr 05, 2019 |
| | | RE44186 | Jul 31, 2023 | DS DP U-1097 | M-198 | Feb 27, 2020 |
| | | RE44186 | Jul 31, 2023 | DS DP U-1838 | | |
| <u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</u> | | | | | | |
| N 022044 | 001 | 6699871 | Jul 26, 2022 | DS DP U-802 | | |
| | | 6890898 | Feb 02, 2019 | U-1996 | | |
| | | 7078381 | Feb 02, 2019 | U-1996 | | |
| | | 7125873 | Jul 26, 2022 | DP U-1036 | | |
| | | 7125873 | Jul 26, 2022 | DP U-1038 | | |
| | | 7125873 | Jul 26, 2022 | DP U-803 | | |
| | | 7326708 | Nov 24, 2026 | DS DP U-802 | | |
| | | 7459428 | Feb 02, 2019 | U-1996 | | |
| | | 8414921 | Jul 21, 2028 | DP U-1036 | | |
| <u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</u> | | | | | | |
| N 022044 | 002 | 6699871 | Jul 26, 2022 | DS DP U-802 | | |
| | | 6890898 | Feb 02, 2019 | U-1996 | | |
| | | 7078381 | Feb 02, 2019 | U-1996 | | |
| | | 7125873 | Jul 26, 2022 | DP U-1036 | | |
| | | 7125873 | Jul 26, 2022 | DP U-1038 | | |
| | | 7125873 | Jul 26, 2022 | DP U-803 | | |
| | | 7326708 | Nov 24, 2026 | DS DP U-802 | | |
| | | 7459428 | Feb 02, 2019 | U-1996 | | |
| | | 8414921 | Jul 21, 2028 | DP U-1036 | | |
| <u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u> | | | | | | |
| N 202270 | 001 | 6699871 | Jul 26, 2022 | DS DP U-1227 | | |
| | | 6890898 | Feb 02, 2019 | U-1996 | | |
| | | 7078381 | Feb 02, 2019 | U-1996 | | |
| | | 7125873 | Jul 26, 2022 | DP U-1227 | | |
| | | 7326708 | Nov 24, 2026 | DS DP U-1227 | | |
| | | 7459428 | Feb 02, 2019 | U-1996 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u> | | | | | | |
| N 202270 002 | 6699871 | Jul 26, 2022 | DS DP U-1227 | | | |
| | 6890898 | Feb 02, 2019 | U-1996 | | | |
| | 7078381 | Feb 02, 2019 | U-1996 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1227 | | | |
| | 7326708 | Nov 24, 2026 | DS DP U-1227 | | | |
| | 7459428 | Feb 02, 2019 | U-1996 | | | |
| <u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u> | | | | | | |
| N 202270 003 | 6699871 | Jul 26, 2022 | DS DP U-1227 | | | |
| | 6890898 | Feb 02, 2019 | U-1996 | | | |
| | 7078381 | Feb 02, 2019 | U-1996 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1227 | | | |
| | 7326708 | Nov 24, 2026 | DS DP U-1227 | | | |
| | 7459428 | Feb 02, 2019 | U-1996 | | | |
| <u>METHOTREXATE - OTREXUP</u> | | | | | | |
| N 204824 001 | 6746429 | Apr 12, 2020 | DP | | | |
| | 7744582 | Aug 10, 2019 | DP U-1442 | | | |
| | 7776015 | Aug 10, 2019 | DP | | | |
| | 8021335 | Oct 04, 2026 | DP | | | |
| | 8480631 | Mar 19, 2030 | DP U-1442 | | | |
| | 8562564 | Jan 24, 2026 | DP | | | |
| | 8579865 | Mar 19, 2030 | DP U-1442 | | | |
| | 8945063 | Mar 19, 2030 | DP U-1442 | | | |
| | 9421333 | Mar 19, 2030 | DP U-1442 | | | |
| | 9533102 | Jan 24, 2026 | DP | | | |
| | 9629959 | Jan 24, 2026 | DP | | | |
| | RE44846 | Aug 10, 2019 | DP | | | |
| | RE44847 | Aug 10, 2019 | DP U-1442 | | | |
| <u>METHOTREXATE - OTREXUP</u> | | | | | | |
| N 204824 002 | 6746429 | Apr 12, 2020 | DP | | | |
| | 7744582 | Aug 10, 2019 | DP U-1442 | | | |
| | 7776015 | Aug 10, 2019 | DP | | | |
| | 8021335 | Oct 04, 2026 | DP | | | |
| | 8480631 | Mar 19, 2030 | DP U-1442 | | | |
| | 8562564 | Jan 24, 2026 | DP | | | |
| | 8579865 | Mar 19, 2030 | DP U-1442 | | | |
| | 8945063 | Mar 19, 2030 | DP U-1442 | | | |
| | 9421333 | Mar 19, 2030 | DP U-1442 | | | |
| | 9533102 | Jan 24, 2026 | DP | | | |
| | 9629959 | Jan 24, 2026 | DP | | | |
| | RE44846 | Aug 10, 2019 | DP | | | |
| | RE44847 | Aug 10, 2019 | DP U-1442 | | | |
| <u>METHOTREXATE - OTREXUP</u> | | | | | | |
| N 204824 003 | 6746429 | Apr 12, 2020 | DP | | | |
| | 7744582 | Aug 10, 2019 | DP U-1442 | | | |
| | 7776015 | Aug 10, 2019 | DP | | | |
| | 8021335 | Oct 04, 2026 | DP | | | |
| | 8480631 | Mar 19, 2030 | DP U-1442 | | | |
| | 8562564 | Jan 24, 2026 | DP | | | |
| | 8579865 | Mar 19, 2030 | DP U-1442 | | | |
| | 8945063 | Mar 19, 2030 | DP U-1442 | | | |
| | 9421333 | Mar 19, 2030 | DP U-1442 | | | |
| | 9533102 | Jan 24, 2026 | DP | | | |
| | 9629959 | Jan 24, 2026 | DP | | | |
| | RE44846 | Aug 10, 2019 | DP | | | |
| | RE44847 | Aug 10, 2019 | DP U-1442 | | | |
| <u>METHOTREXATE - OTREXUP</u> | | | | | | |
| N 204824 004 | 6746429 | Apr 12, 2020 | DP | | | |
| | 7744582 | Aug 10, 2019 | DP U-1442 | | | |
| | 7776015 | Aug 10, 2019 | DP | | | |
| | 8021335 | Oct 04, 2026 | DP | | | |
| | 8480631 | Mar 19, 2030 | DP U-1442 | | | |
| | 8562564 | Jan 24, 2026 | DP | | | |
| | 8579865 | Mar 19, 2030 | DP U-1442 | | | |
| | 8945063 | Mar 19, 2030 | DP U-1442 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>METHOTREXATE - OTREXUP</u> | | | | | | |
| N 204824 004 | 9421333 | Mar 19, 2030 | DP U-1442 | | | |
| | 9533102 | Jan 24, 2026 | DP | | | |
| | 9629959 | Jan 24, 2026 | DP | | | |
| | RE44846 | Aug 10, 2019 | DP | | | |
| | RE44847 | Aug 10, 2019 | DP U-1442 | | | |
| <u>METHOTREXATE - OTREXUP</u> | | | | | | |
| N 204824 005 | 6746429 | Apr 12, 2020 | DP | | | |
| | 8021335 | Oct 04, 2026 | DP | | | |
| | 8480631 | Mar 19, 2030 | DP U-1442 | | | |
| | 8562564 | Jan 24, 2026 | DP | | | |
| | 8579865 | Mar 19, 2030 | DP U-1442 | | | |
| | 8945063 | Mar 19, 2030 | DP U-1442 | | | |
| | 9421333 | Mar 19, 2030 | DP U-1442 | | | |
| | 9533102 | Jan 24, 2026 | DP | | | |
| | 9629959 | Jan 24, 2026 | DP | | | |
| | RE44846 | Aug 10, 2019 | DP | | | |
| | RE44847 | Aug 10, 2019 | DP U-1442 | | | |
| <u>METHOTREXATE - OTREXUP</u> | | | | | | |
| N 204824 006 | 6746429 | Apr 12, 2020 | DP | | | |
| | 8021335 | Oct 04, 2026 | DP | | | |
| | 8480631 | Mar 19, 2030 | DP U-1442 | | | |
| | 8562564 | Jan 24, 2026 | DP | | | |
| | 8579865 | Mar 19, 2030 | DP U-1442 | | | |
| | 8945063 | Mar 19, 2030 | DP U-1442 | | | |
| | 9421333 | Mar 19, 2030 | DP U-1442 | | | |
| | 9533102 | Jan 24, 2026 | DP | | | |
| | 9629959 | Jan 24, 2026 | DP | | | |
| | RE44846 | Aug 10, 2019 | DP | | | |
| | RE44847 | Aug 10, 2019 | DP U-1442 | | | |
| <u>METHOTREXATE - OTREXUP</u> | | | | | | |
| N 204824 007 | 6746429 | Apr 12, 2020 | DP | | | |
| | 8021335 | Oct 04, 2026 | DP | | | |
| | 8480631 | Mar 19, 2030 | DP U-1442 | | | |
| | 8562564 | Jan 24, 2026 | DP | | | |
| | 8579865 | Mar 19, 2030 | DP U-1442 | | | |
| | 8945063 | Mar 19, 2030 | DP U-1442 | | | |
| | 9421333 | Mar 19, 2030 | DP U-1442 | | | |
| | 9533102 | Jan 24, 2026 | DP | | | |
| | 9629959 | Jan 24, 2026 | DP | | | |
| | RE44846 | Aug 10, 2019 | DP | | | |
| | RE44847 | Aug 10, 2019 | DP U-1442 | | | |
| <u>METHOTREXATE - OTREXUP</u> | | | | | | |
| N 204824 008 | 6746429 | Apr 12, 2020 | DP | | | |
| | 8021335 | Oct 04, 2026 | DP | | | |
| | 8480631 | Mar 19, 2030 | DP U-1442 | | | |
| | 8562564 | Jan 24, 2026 | DP | | | |
| | 8579865 | Mar 19, 2030 | DP U-1442 | | | |
| | 8945063 | Mar 19, 2030 | DP U-1442 | | | |
| | 9421333 | Mar 19, 2030 | DP U-1442 | | | |
| | 9533102 | Jan 24, 2026 | DP | | | |
| | 9629959 | Jan 24, 2026 | DP | | | |
| | RE44846 | Aug 10, 2019 | DP | | | |
| | RE44847 | Aug 10, 2019 | DP U-1442 | | | |
| <u>METHOTREXATE - RASUVQ</u> | | | | | | |
| N 205776 001 | 8664231 | Jun 01, 2029 | U-1442 | | | |
| <u>METHOTREXATE - RASUVQ</u> | | | | | | |
| N 205776 002 | 8664231 | Jun 01, 2029 | U-1442 | | | |
| <u>METHOTREXATE - RASUVQ</u> | | | | | | |
| N 205776 003 | 8664231 | Jun 01, 2029 | U-1442 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>METHOTREXATE - RASUVO</u> | | | | | | |
| N 205776 | 004 8664231 | Jun 01, 2029 | U-1442 | | | |
| <u>METHOTREXATE - RASUVO</u> | | | | | | |
| N 205776 | 005 8664231 | Jun 01, 2029 | U-1442 | | | |
| <u>METHOTREXATE - RASUVO</u> | | | | | | |
| N 205776 | 006 8664231 | Jun 01, 2029 | U-1442 | | | |
| <u>METHOTREXATE - RASUVO</u> | | | | | | |
| N 205776 | 007 8664231 | Jun 01, 2029 | U-1442 | | | |
| <u>METHOTREXATE - RASUVO</u> | | | | | | |
| N 205776 | 008 8664231 | Jun 01, 2029 | U-1442 | | | |
| <u>METHOTREXATE - RASUVO</u> | | | | | | |
| N 205776 | 009 8664231 | Jun 01, 2029 | U-1442 | | | |
| <u>METHOTREXATE - RASUVO</u> | | | | | | |
| N 205776 | 010 8664231 | Jun 01, 2029 | U-1442 | | | |
| <u>METHOTREXATE SODIUM - XATMEP</u> | | | | | | |
| N 208400 | 001 9259427 | Jan 02, 2033 | DP | | ODE-137 | Apr 25, 2024 |
| | 9855215 | Jan 02, 2033 | DP | | ODE-138 | Apr 25, 2024 |
| <u>METHYLENE BLUE - PROVAYBLUE</u> | | | | | | |
| N 204630 | 001 | | | | ODE-113 | Apr 08, 2023 |
| <u>METHYLERGONOVINE MALEATE - METHYLERGONOVINE MALEATE</u> | | | | | | |
| A 211483 | 001 | | | | CGT | Mar 20, 2019 |
| <u>METHYLNALTREXONE BROMIDE - RELISTOR</u> | | | | | | |
| N 021964 | 001 8247425 | Dec 31, 2030 | U-1185 | | | |
| | 8420663 | Sep 30, 2029 | U-1185 | | | |
| | 8552025 | Apr 08, 2024 | DP | | | |
| | 8822490 | Sep 30, 2029 | DP U-1185 | | | |
| | 9180125 | Sep 30, 2029 | DP U-1185 | | | |
| | 9492445 | Sep 30, 2029 | DP U-1185 | | | |
| | 9669096 | Apr 08, 2024 | DP | | | |
| <u>METHYLNALTREXONE BROMIDE - RELISTOR</u> | | | | | | |
| N 021964 | 002 8247425 | Dec 31, 2030 | U-1185 | | | |
| | 8420663 | Sep 30, 2029 | U-1185 | | | |
| | 8552025 | Apr 08, 2024 | DP | | | |
| | 8822490 | Sep 30, 2029 | DP U-1185 | | | |
| | 9180125 | Sep 30, 2029 | DP U-1185 | | | |
| | 9492445 | Sep 30, 2029 | DP U-1185 | | | |
| | 9669096 | Apr 08, 2024 | DP | | | |
| <u>METHYLNALTREXONE BROMIDE - RELISTOR</u> | | | | | | |
| N 021964 | 003 8247425 | Dec 31, 2030 | U-1185 | | | |
| | 8420663 | Sep 30, 2029 | U-1185 | | | |
| | 8552025 | Apr 08, 2024 | DP | | | |
| | 8822490 | Sep 30, 2029 | DP U-1185 | | | |
| | 9180125 | Sep 30, 2029 | DP U-1185 | | | |
| | 9492445 | Sep 30, 2029 | DP U-1185 | | | |
| | 9669096 | Apr 08, 2024 | DP | | | |
| <u>METHYLNALTREXONE BROMIDE - RELISTOR</u> | | | | | | |
| N 208271 | 001 8420663 | Sep 30, 2029 | U-1185 | | NP | Jul 19, 2019 |
| | 8524276 | Mar 10, 2031 | DP | | | |
| | 8956651 | Mar 10, 2031 | DP | | | |
| | 9180125 | Sep 30, 2029 | DP U-1185 | | | |
| | 9314461 | Mar 10, 2031 | DP | | | |
| | 9492445 | Sep 30, 2029 | DP U-1185 | | | |
| | 9724343 | Sep 30, 2029 | DP U-1185 | | | |
| <u>METHYLPHENIDATE - DAYTRANA</u> | | | | | | |
| N 021514 | 001 8632802 | Oct 07, 2025 | DP | | | |
| | 9034370 | Oct 07, 2025 | DP | | | |
| | 9668981 | Oct 07, 2025 | U-2024 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>METHYLPHENIDATE - DAYTRANA</u> | | | | | | |
| N 021514 001 | 8632802 | Oct 07, 2025 | DP | | | |
| | 9034370 | Oct 07, 2025 | DP | | | |
| | 9668981 | Oct 07, 2025 | | | U-2024 | |
| <u>METHYLPHENIDATE - DAYTRANA</u> | | | | | | |
| N 021514 002 | 8632802 | Oct 07, 2025 | DP | | | |
| | 9034370 | Oct 07, 2025 | DP | | | |
| | 9668981 | Oct 07, 2025 | | | U-2024 | |
| <u>METHYLPHENIDATE - DAYTRANA</u> | | | | | | |
| N 021514 003 | 8632802 | Oct 07, 2025 | DP | | | |
| | 9034370 | Oct 07, 2025 | DP | | | |
| | 9668981 | Oct 07, 2025 | | | U-2024 | |
| <u>METHYLPHENIDATE - DAYTRANA</u> | | | | | | |
| N 021514 004 | 8632802 | Oct 07, 2025 | DP | | | |
| | 9034370 | Oct 07, 2025 | DP | | | |
| | 9668981 | Oct 07, 2025 | | | U-2024 | |
| <u>METHYLPHENIDATE - COTEMPLA XR-ODT</u> | | | | | | |
| N 205489 001 | 8840924 | Jun 05, 2026 | DP | | NP | Jun 19, 2020 |
| | 9072680 | Jun 28, 2032 | DP | | | |
| | 9089496 | Jun 28, 2032 | DP | | | |
| <u>METHYLPHENIDATE - COTEMPLA XR-ODT</u> | | | | | | |
| N 205489 002 | 8840924 | Jun 05, 2026 | DP | | NP | Jun 19, 2020 |
| | 9072680 | Jun 28, 2032 | DP | | | |
| | 9089496 | Jun 28, 2032 | DP | | | |
| <u>METHYLPHENIDATE - COTEMPLA XR-ODT</u> | | | | | | |
| N 205489 003 | 8840924 | Jun 05, 2026 | DP | | NP | Jun 19, 2020 |
| | 9072680 | Jun 28, 2032 | DP | | | |
| | 9089496 | Jun 28, 2032 | DP | | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u> | | | | | | |
| N 021259 001 | 6344215 | Oct 27, 2020 | DP | | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u> | | | | | | |
| N 021259 002 | 6344215 | Oct 27, 2020 | DP | | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u> | | | | | | |
| N 021259 003 | 6344215 | Oct 27, 2020 | DP | | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u> | | | | | | |
| N 021259 004 | 6344215 | Oct 27, 2020 | DP | | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - RITALIN LA</u> | | | | | | |
| N 021284 001 | 6228398 | Nov 01, 2019 | DP | | U-472 | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - RITALIN LA</u> | | | | | | |
| N 021284 002 | 6228398 | Nov 01, 2019 | DP | | U-472 | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - RITALIN LA</u> | | | | | | |
| N 021284 003 | 6228398 | Nov 01, 2019 | DP | | U-472 | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - RITALIN LA</u> | | | | | | |
| N 021284 004 | 6228398 | Nov 01, 2019 | DP | | U-472 | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - METHYLIN</u> | | | | | | |
| N 021419 001 | 7691880 | Oct 07, 2024 | DP | | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - METHYLIN</u> | | | | | | |
| N 021419 002 | 7691880 | Oct 07, 2024 | DP | | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - QUILLIVANT XR</u> | | | | | | |
| N 202100 001 | 8062667 | Mar 29, 2029 | DP | | | |
| | 8287903 | Feb 15, 2031 | DP | | | |
| | 8465765 | Feb 15, 2031 | DP | | U-1415 | |
| | 8563033 | Feb 15, 2031 | DP | | U-1415 | |
| | 8778390 | Feb 15, 2031 | DP | | U-1543 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>METHYLPHENIDATE HYDROCHLORIDE - OUIILLIVANT XR</u> | | | | | | |
| N 202100 | 001 | 8956649 | Feb 15, 2031 | DP U-1665 | | |
| | | 9040083 | Feb 15, 2031 | DP | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u> | | | | | | |
| N 205831 | 001 | 10039719 | Dec 16, 2019 | U-2357 | | |
| | | 6419960 | Dec 16, 2019 | DP | | |
| | | 7083808 | Dec 16, 2019 | DP | | |
| | | 7247318 | Dec 16, 2019 | DP | | |
| | | 7438930 | Dec 16, 2019 | DP | Y | |
| | | 8580310 | Dec 16, 2019 | DP | | |
| | | 9066869 | Dec 16, 2019 | DP | | |
| | | 9801823 | Dec 16, 2019 | DP | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u> | | | | | | |
| N 205831 | 002 | 10039719 | Dec 16, 2019 | U-2357 | | |
| | | 6419960 | Dec 16, 2019 | DP | | |
| | | 7083808 | Dec 16, 2019 | DP | | |
| | | 7247318 | Dec 16, 2019 | DP | | |
| | | 7438930 | Dec 16, 2019 | DP | Y | |
| | | 8580310 | Dec 16, 2019 | DP | | |
| | | 9066869 | Dec 16, 2019 | DP | | |
| | | 9801823 | Dec 16, 2019 | DP | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u> | | | | | | |
| N 205831 | 003 | 10039719 | Dec 16, 2019 | U-2357 | | |
| | | 6419960 | Dec 16, 2019 | DP | | |
| | | 7083808 | Dec 16, 2019 | DP | | |
| | | 7247318 | Dec 16, 2019 | DP | | |
| | | 7438930 | Dec 16, 2019 | DP | Y | |
| | | 8580310 | Dec 16, 2019 | DP | | |
| | | 9066869 | Dec 16, 2019 | DP | | |
| | | 9801823 | Dec 16, 2019 | DP | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u> | | | | | | |
| N 205831 | 004 | 10039719 | Dec 16, 2019 | U-2357 | | |
| | | 6419960 | Dec 16, 2019 | DP | | |
| | | 7083808 | Dec 16, 2019 | DP | | |
| | | 7247318 | Dec 16, 2019 | DP | | |
| | | 7438930 | Dec 16, 2019 | DP | Y | |
| | | 8580310 | Dec 16, 2019 | DP | | |
| | | 9066869 | Dec 16, 2019 | DP | | |
| | | 9801823 | Dec 16, 2019 | DP | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u> | | | | | | |
| N 205831 | 005 | 10039719 | Dec 16, 2019 | U-2357 | | |
| | | 6419960 | Dec 16, 2019 | DP | | |
| | | 7083808 | Dec 16, 2019 | DP | | |
| | | 7247318 | Dec 16, 2019 | DP | | |
| | | 7438930 | Dec 16, 2019 | DP | Y | |
| | | 8580310 | Dec 16, 2019 | DP | | |
| | | 9066869 | Dec 16, 2019 | DP | | |
| | | 9801823 | Dec 16, 2019 | DP | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u> | | | | | | |
| N 205831 | 006 | 10039719 | Dec 16, 2019 | U-2357 | | |
| | | 6419960 | Dec 16, 2019 | DP | | |
| | | 7083808 | Dec 16, 2019 | DP | | |
| | | 7247318 | Dec 16, 2019 | DP | | |
| | | 7438930 | Dec 16, 2019 | DP | Y | |
| | | 8580310 | Dec 16, 2019 | DP | | |
| | | 9066869 | Dec 16, 2019 | DP | | |
| | | 9801823 | Dec 16, 2019 | DP | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u> | | | | | | |
| N 205831 | 007 | 10039719 | Dec 16, 2019 | U-2357 | | |
| | | 6419960 | Dec 16, 2019 | DP | | |
| | | 7083808 | Dec 16, 2019 | DP | | |
| | | 7247318 | Dec 16, 2019 | DP | | |
| | | 7438930 | Dec 16, 2019 | DP | Y | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u> | | | | | | |
| N 205831 | 007 | 8580310 | Dec 16, 2019 | DP | | |
| | | 9066869 | Dec 16, 2019 | DP | | |
| | | 9801823 | Dec 16, 2019 | DP | | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - QUILLICHEW ER</u> | | | | | | |
| N 207960 | 001 | 8202537 | Mar 15, 2027 | DP | | |
| | | 8287903 | Feb 15, 2031 | DP | | |
| | | 8999386 | Aug 14, 2033 | DP | | |
| | | 9295642 | Aug 14, 2033 | DP | U-1827 | |
| | | 9545399 | Aug 14, 2033 | DP | U-1827 | |
| | | 9844544 | Aug 14, 2033 | DP | U-2203 | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - QUILLICHEW ER</u> | | | | | | |
| N 207960 | 002 | 8202537 | Mar 15, 2027 | DP | | |
| | | 8287903 | Feb 15, 2031 | DP | | |
| | | 8999386 | Aug 14, 2033 | DP | | |
| | | 9295642 | Aug 14, 2033 | DP | U-1827 | |
| | | 9545399 | Aug 14, 2033 | DP | U-1827 | |
| | | 9844544 | Aug 14, 2033 | DP | U-2203 | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - QUILLICHEW ER</u> | | | | | | |
| N 207960 | 003 | 8202537 | Mar 15, 2027 | DP | | |
| | | 8287903 | Feb 15, 2031 | DP | | |
| | | 8999386 | Aug 14, 2033 | DP | | |
| | | 9295642 | Aug 14, 2033 | DP | U-1827 | |
| | | 9545399 | Aug 14, 2033 | DP | U-1827 | |
| | | 9844544 | Aug 14, 2033 | DP | U-2203 | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u> | | | | | | |
| N 209311 | 001 | 8916588 | Mar 23, 2032 | | U-2357 | NP Aug 08, 2021 |
| | | 8927010 | Mar 23, 2032 | DP | | |
| | | 9023389 | Mar 23, 2032 | DP | | |
| | | 9028868 | Mar 23, 2032 | | U-2357 | |
| | | 9034902 | Mar 23, 2032 | | U-2357 | |
| | | 9283214 | Mar 23, 2032 | DP | | |
| | | 9498447 | Mar 23, 2032 | DP | | |
| | | 9603809 | Mar 23, 2032 | | U-2357 | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u> | | | | | | |
| N 209311 | 002 | 8916588 | Mar 23, 2032 | | U-2357 | NP Aug 08, 2021 |
| | | 8927010 | Mar 23, 2032 | DP | | |
| | | 9023389 | Mar 23, 2032 | DP | | |
| | | 9028868 | Mar 23, 2032 | | U-2357 | |
| | | 9034902 | Mar 23, 2032 | | U-2357 | |
| | | 9283214 | Mar 23, 2032 | DP | | |
| | | 9498447 | Mar 23, 2032 | DP | | |
| | | 9603809 | Mar 23, 2032 | | U-2357 | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u> | | | | | | |
| N 209311 | 003 | 8916588 | Mar 23, 2032 | | U-2357 | NP Aug 08, 2021 |
| | | 8927010 | Mar 23, 2032 | DP | | |
| | | 9023389 | Mar 23, 2032 | DP | | |
| | | 9028868 | Mar 23, 2032 | | U-2357 | |
| | | 9034902 | Mar 23, 2032 | | U-2357 | |
| | | 9283214 | Mar 23, 2032 | DP | | |
| | | 9498447 | Mar 23, 2032 | DP | | |
| | | 9603809 | Mar 23, 2032 | | U-2357 | |
| <u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u> | | | | | | |
| N 209311 | 004 | 8916588 | Mar 23, 2032 | | U-2357 | NP Aug 08, 2021 |
| | | 8927010 | Mar 23, 2032 | DP | | |
| | | 9023389 | Mar 23, 2032 | DP | | |
| | | 9028868 | Mar 23, 2032 | | U-2357 | |
| | | 9034902 | Mar 23, 2032 | | U-2357 | |
| | | 9283214 | Mar 23, 2032 | DP | | |
| | | 9498447 | Mar 23, 2032 | DP | | |
| | | 9603809 | Mar 23, 2032 | | U-2357 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>METHYLPHENIDATE HYDROCHLORIDE - JORNAY PM</u> | | | | | | |
| N 209311 | 005 | 8916588 | Mar 23, 2032 | U-2357 | NP | Aug 08, 2021 |
| | | 8927010 | Mar 23, 2032 | DP | | |
| | | 9023389 | Mar 23, 2032 | DP | | |
| | | 9028868 | Mar 23, 2032 | U-2357 | | |
| | | 9034902 | Mar 23, 2032 | U-2357 | | |
| | | 9283214 | Mar 23, 2032 | DP | | |
| | | 9498447 | Mar 23, 2032 | DP | | |
| | | 9603809 | Mar 23, 2032 | U-2357 | | |
| <u>METOPROLOL SUCCINATE - KAPSPARGO SPRINKLE</u> | | | | | | |
| N 210428 | 001 | 9504655 | Jul 09, 2035 | DP | | |
| | | 9700530 | Jul 09, 2035 | DP | | |
| <u>METOPROLOL SUCCINATE - KAPSPARGO SPRINKLE</u> | | | | | | |
| N 210428 | 002 | 9504655 | Jul 09, 2035 | DP | | |
| | | 9700530 | Jul 09, 2035 | DP | | |
| <u>METOPROLOL SUCCINATE - KAPSPARGO SPRINKLE</u> | | | | | | |
| N 210428 | 003 | 9504655 | Jul 09, 2035 | DP | | |
| | | 9700530 | Jul 09, 2035 | DP | | |
| <u>METOPROLOL SUCCINATE - KAPSPARGO SPRINKLE</u> | | | | | | |
| N 210428 | 004 | 9504655 | Jul 09, 2035 | DP | | |
| | | 9700530 | Jul 09, 2035 | DP | | |
| <u>METRONIDAZOLE - METROGEL</u> | | | | | | |
| N 021789 | 001 | 6881726 | Feb 21, 2022 | DP U-743 | | |
| | | 7348317 | Feb 21, 2022 | DP U-743 | | |
| <u>METRONIDAZOLE - VANDAZOLE</u> | | | | | | |
| N 021806 | 001 | 7456207 | Sep 22, 2024 | DP | | |
| <u>METRONIDAZOLE - NUVESSA</u> | | | | | | |
| N 205223 | 001 | 7893097 | Feb 19, 2028 | DP | | |
| | | 8658678 | Jun 27, 2028 | U-1682 | | |
| | | 8877792 | Feb 02, 2028 | DP | | |
| | | 8946276 | Jun 28, 2032 | U-1664 | | |
| | | 9198858 | Jun 28, 2032 | U-1664 | | |
| <u>MICAFUNGIN SODIUM - MYCAMINE</u> | | | | | | |
| N 021506 | 002 | 6107458 | Mar 16, 2019 | DS DP U-650 | | |
| | | 6107458 | Mar 16, 2019 | DS DP U-845 | | |
| | | 6774104 | Jan 08, 2021 | DP U-650 | | |
| | | 6774104 | Jan 08, 2021 | DP U-845 | | |
| <u>MICAFUNGIN SODIUM - MYCAMINE</u> | | | | | | |
| N 021506 | 003 | 6107458 | Mar 16, 2019 | DS DP U-650 | | |
| | | 6107458 | Mar 16, 2019 | DS DP U-845 | | |
| | | 6774104 | Jan 08, 2021 | DP U-650 | | |
| | | 6774104 | Jan 08, 2021 | DP U-845 | | |
| <u>MICONAZOLE - ORAVIG</u> | | | | | | |
| N 022404 | 001 | 6916485 | Sep 11, 2022 | DP U-1051 | | |
| | | 7651698 | Sep 11, 2022 | U-1051 | | |
| | | 8518442 | Sep 11, 2022 | DP | | |
| <u>MICONAZOLE NITRATE; MICONAZOLE NITRATE - MONISTAT 1 COMBINATION PACK</u> | | | | | | |
| N 021308 | 001 | 6153635 | Nov 28, 2020 | | Y | |
| <u>MICONAZOLE NITRATE; PETROLATUM, WHITE; ZINC OXIDE - VUSION</u> | | | | | | |
| N 021026 | 001 | 8147852 | Mar 30, 2028 | U-1426 | | |
| <u>MIDAZOLAM HYDROCHLORIDE - SEIZALAM</u> | | | | | | |
| N 209566 | 001 | | | | ODE-207 | Sep 14, 2025 |
| <u>MIDOSTAURIN - RYDAPT</u> | | | | | | |
| N 207997 | 001 | 7973031 | Oct 17, 2024 | U-2007 | NCE | Apr 28, 2022 |
| | | 8222244 | Oct 29, 2022 | U-2007 | ODE-140 | Apr 28, 2024 |
| | | 8575146 | Dec 02, 2030 | U-2008 | ODE-141 | Apr 28, 2024 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>MIFEPRISTONE - KORLYM</u> | | | | | | |
| N 202107 | 001 | 10006924 | Aug 12, 2036 | U-1643 | ODE-22 | Feb 17, 2019 |
| | | 10151763 | Jan 18, 2037 | U-1643 | | |
| | | 10166242 | Apr 20, 2036 | U-1643 | | |
| | | 10166243 | Apr 20, 2036 | U-1643 | | |
| | | 8921348 | Aug 27, 2028 | U-1643 | | |
| | | 9829495 | Aug 15, 2036 | U-1643 | | |
| | | 9943526 | Apr 20, 2036 | U-1643 | | |
| <u>MIGALASTAT HYDROCHLORIDE - GALAFOLD</u> | | | | | | |
| N 208623 | 001 | 10076514 | Mar 15, 2037 | U-2371 | NCE | Aug 10, 2023 |
| | | 8592362 | Feb 12, 2029 | U-2371 | ODE-205 | Aug 10, 2025 |
| | | 9000011 | May 16, 2027 | U-2371 | | |
| | | 9095584 | Feb 12, 2029 | U-2371 | | |
| | | 9480682 | May 16, 2027 | U-2371 | | |
| | | 9987263 | May 16, 2027 | U-2371 | | |
| | | 9999618 | Apr 28, 2028 | U-2372 | | |
| | | 9999618 | Apr 28, 2028 | U-2373 | | |
| <u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u> | | | | | | |
| N 022256 | 001 | 6602911 | Jan 14, 2023 | U-882 | | |
| | | 6992110 | Nov 05, 2021 | U-882 | | |
| | | 7888342 | Nov 05, 2021 | U-882 | | |
| | | 7994220 | Sep 19, 2029 | U-819 | | |
| <u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u> | | | | | | |
| N 022256 | 002 | 6602911 | Jan 14, 2023 | U-882 | | |
| | | 6992110 | Nov 05, 2021 | U-882 | | |
| | | 7888342 | Nov 05, 2021 | U-882 | | |
| | | 7994220 | Sep 19, 2029 | U-819 | | |
| <u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u> | | | | | | |
| N 022256 | 003 | 6602911 | Jan 14, 2023 | U-882 | | |
| | | 6992110 | Nov 05, 2021 | U-882 | | |
| | | 7888342 | Nov 05, 2021 | U-882 | | |
| | | 7994220 | Sep 19, 2029 | U-819 | | |
| <u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u> | | | | | | |
| N 022256 | 004 | 6602911 | Jan 14, 2023 | U-882 | | |
| | | 6992110 | Nov 05, 2021 | U-882 | | |
| | | 7888342 | Nov 05, 2021 | U-882 | | |
| | | 7994220 | Sep 19, 2029 | U-819 | | |
| <u>MILTEFOSINE - IMPAVIDO</u> | | | | | | |
| N 204684 | 001 | | | | NCE | Mar 19, 2019 |
| | | | | | ODE-63 | Mar 19, 2021 |
| <u>MINOCYCLINE HYDROCHLORIDE - MINOCIN</u> | | | | | | |
| N 050444 | 001 | 9084802 | May 12, 2031 | U-282 | | |
| | | 9278105 | May 12, 2031 | U-282 | | |
| <u>MINOCYCLINE HYDROCHLORIDE - ARESTIN</u> | | | | | | |
| N 050781 | 001 | 6682348 | Mar 29, 2022 | DP | | |
| | | 7699609 | Mar 29, 2022 | DP | | |
| <u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u> | | | | | | |
| N 050808 | 001 | 7790705 | Jun 24, 2025 | U-1078 | | |
| | | 7919483 | Mar 07, 2027 | U-1078 | | |
| | | 8252776 | Jun 24, 2025 | U-124 | | |
| | | 8268804 | Jun 24, 2025 | U-1078 | | |
| <u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u> | | | | | | |
| N 050808 | 002 | 7541347 | Apr 02, 2027 | U-917 | | |
| | | 7544373 | Apr 02, 2027 | DP | | |
| | | 7790705 | Jun 24, 2025 | U-1078 | | |
| | | 7919483 | Mar 07, 2027 | U-1078 | | |
| | | 8252776 | Jun 24, 2025 | U-124 | | |
| | | 8268804 | Jun 24, 2025 | U-1078 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u> | | | | | | |
| N 050808 003 | 7790705 | Jun 24, 2025 | U-1078 | | | |
| | 7919483 | Mar 07, 2027 | U-1078 | | | |
| | 8252776 | Jun 24, 2025 | U-124 | | | |
| | 8268804 | Jun 24, 2025 | U-1078 | | | |
| <u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u> | | | | | | |
| N 050808 004 | 7790705 | Jun 24, 2025 | U-1078 | | | |
| | 7919483 | Mar 07, 2027 | U-1078 | | | |
| | 8252776 | Jun 24, 2025 | U-124 | | | |
| | 8268804 | Jun 24, 2025 | U-1078 | | | |
| | 9192615 | Nov 17, 2031 | DP | | | |
| <u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u> | | | | | | |
| N 050808 005 | 7790705 | Jun 24, 2025 | U-1078 | | | |
| | 7919483 | Mar 07, 2027 | U-1078 | | | |
| | 8252776 | Jun 24, 2025 | U-124 | | | |
| | 8268804 | Jun 24, 2025 | U-1078 | | | |
| | 9192615 | Nov 17, 2031 | DP | | | |
| <u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u> | | | | | | |
| N 050808 006 | 7790705 | Jun 24, 2025 | U-1078 | | | |
| | 7919483 | Mar 07, 2027 | U-1078 | | | |
| | 8252776 | Jun 24, 2025 | U-124 | | | |
| | 8268804 | Jun 24, 2025 | U-1078 | | | |
| | 8722650 | Jun 24, 2025 | U-1078 | | | |
| <u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u> | | | | | | |
| N 050808 007 | 7790705 | Jun 24, 2025 | U-1078 | | | |
| | 7919483 | Mar 07, 2027 | U-1078 | | | |
| | 8252776 | Jun 24, 2025 | U-124 | | | |
| | 8268804 | Jun 24, 2025 | U-1078 | | | |
| | 8722650 | Jun 24, 2025 | U-1078 | | | |
| <u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u> | | | | | | |
| N 050808 008 | 7790705 | Jun 24, 2025 | U-1078 | | | |
| | 7919483 | Mar 07, 2027 | U-1078 | | | |
| | 8252776 | Jun 24, 2025 | U-124 | | | |
| | 8268804 | Jun 24, 2025 | U-1078 | | | |
| | 8722650 | Jun 24, 2025 | U-1078 | | | |
| <u>MINOCYCLINE HYDROCHLORIDE - XIMINO</u> | | | | | | |
| N 201922 001 | 7541347 | Apr 02, 2027 | U-917 | | | |
| | 7544373 | Apr 02, 2027 | DP | | | |
| | 7790705 | Jun 24, 2025 | U-124 | | | |
| | 7919483 | Mar 07, 2027 | U-124 | | | |
| | 8252776 | Jun 24, 2025 | U-124 | | | |
| | 8268804 | Jun 24, 2025 | U-124 | | | |
| <u>MINOCYCLINE HYDROCHLORIDE - XIMINO</u> | | | | | | |
| N 201922 003 | 7541347 | Apr 02, 2027 | U-917 | | | |
| | 7544373 | Apr 02, 2027 | DP | | | |
| | 7790705 | Jun 24, 2025 | U-124 | | | |
| | 7919483 | Mar 07, 2027 | U-124 | | | |
| | 8252776 | Jun 24, 2025 | U-124 | | | |
| | 8268804 | Jun 24, 2025 | U-124 | | | |
| <u>MINOCYCLINE HYDROCHLORIDE - XIMINO</u> | | | | | | |
| N 201922 005 | 7541347 | Apr 02, 2027 | U-917 | | | |
| | 7544373 | Apr 02, 2027 | DP | | | |
| | 7790705 | Jun 24, 2025 | U-124 | | | |
| | 7919483 | Mar 07, 2027 | U-124 | | | |
| | 8252776 | Jun 24, 2025 | U-124 | | | |
| | 8268804 | Jun 24, 2025 | U-124 | | | |
| <u>MINOXIDIL - MEN'S ROGAINE</u> | | | | | | |
| N 021812 001 | 6946120 | Apr 20, 2019 | DP U-702 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>MINOXIDIL - WOMEN'S ROGAINE</u> | | | | | | |
| N 021812 | 002 6946120 | Apr 20, 2019 | DP | U-702 | | |
| <u>MIPOMERSEN SODIUM - KYNAMRO</u> | | | | | | |
| N 203568 | 001 7015315 | Mar 21, 2023 | DS | | ODE-41 | Jan 29, 2020 |
| | 7101993 | Sep 05, 2023 | DS | | | |
| | 7407943 | Aug 01, 2021 | | U-1353 | | |
| | 7511131 | Jan 29, 2027 | DS | | | |
| <u>MIRABEGRON - MYRBETRIO</u> | | | | | | |
| N 202611 | 001 6346532 | Mar 27, 2022 | DS DP | | I-777 | Apr 27, 2021 |
| | 6562375 | Aug 01, 2020 | DP | | | |
| | 7342117 | Nov 04, 2023 | DS | | | |
| | 7982049 | Nov 04, 2023 | DP | | | |
| | 8772315 | Oct 30, 2028 | | U-2300 | | |
| | 8835474 | Nov 04, 2023 | | U-1527 | | |
| | RE44872 | Nov 04, 2023 | | U-1527 | | |
| <u>MIRABEGRON - MYRBETRIO</u> | | | | | | |
| N 202611 | 002 6346532 | Mar 27, 2022 | DS DP | | I-777 | Apr 27, 2021 |
| | 6562375 | Aug 01, 2020 | DP | | | |
| | 7342117 | Nov 04, 2023 | DS | | | |
| | 7982049 | Nov 04, 2023 | DP | | | |
| | 8772315 | Oct 30, 2028 | | U-2300 | | |
| | 8835474 | Nov 04, 2023 | | U-1527 | | |
| | RE44872 | Nov 04, 2023 | | U-1527 | | |
| <u>MITOMYCIN - MITOSOL</u> | | | | | | |
| N 022572 | 001 7806265 | Feb 01, 2029 | DP | | ODE-21 | Feb 07, 2019 |
| | 8186511 | Jul 19, 2026 | DP | | | |
| | 9205075 | Jul 19, 2026 | DP | | | |
| | 9539241 | Jan 02, 2028 | DS DP | U-2095 | | |
| | 9649428 | May 21, 2029 | | U-2095 | | |
| <u>MODAFINIL - PROVIGIL</u> | | | | | | |
| N 020717 | 001 7297346 | Nov 29, 2023 | DP | | | |
| <u>MODAFINIL - PROVIGIL</u> | | | | | | |
| N 020717 | 002 7297346 | Nov 29, 2023 | DP | | | |
| <u>MOMETASONE FUROATE - SINUVA</u> | | | | | | |
| N 209310 | 001 7544192 | Nov 29, 2026 | | U-2272 | NP | Dec 08, 2020 |
| | 7662141 | Mar 12, 2024 | | U-2272 | | |
| | 7713255 | Mar 12, 2024 | | U-2272 | | |
| | 7951130 | Mar 12, 2024 | | U-2272 | | |
| | 7951131 | Mar 12, 2024 | | U-2272 | | |
| | 7951133 | Mar 12, 2024 | | U-2272 | | |
| | 8025635 | Jun 12, 2027 | DP | U-2272 | | |
| | 8109918 | Mar 12, 2024 | | U-2272 | | |
| | 8763222 | Feb 08, 2032 | DP | | | |
| | 9585681 | Apr 04, 2026 | | U-2272 | | |
| <u>MONTELUKAST SODIUM - SINGULAIR</u> | | | | | | |
| N 021409 | 001 8007830 | Oct 24, 2022 | DP | | | |
| <u>MORPHINE SULFATE - MORPHINE SULFATE</u> | | | | | | |
| N 204223 | 001 9072781 | Mar 12, 2034 | DP | | | |
| | 9192608 | Mar 12, 2034 | | U-43 | | |
| | 9192608 | Mar 12, 2034 | | U-55 | | |
| | 9248229 | Mar 12, 2034 | DP | | | |
| <u>MORPHINE SULFATE - MORPHINE SULFATE</u> | | | | | | |
| N 204223 | 002 9072781 | Mar 12, 2034 | DP | | | |
| | 9192608 | Mar 12, 2034 | | U-43 | | |
| | 9192608 | Mar 12, 2034 | | U-55 | | |
| | 9248229 | Mar 12, 2034 | DP | | | |
| <u>MORPHINE SULFATE - MORPHINE SULFATE</u> | | | | | | |
| N 204223 | 003 9072781 | Mar 12, 2034 | DP | | | |
| | 9192608 | Mar 12, 2034 | | U-43 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>MORPHINE SULFATE - MORPHINE SULFATE</u> | | | | | | |
| N 204223 | 003 | 9192608 | Mar 12, 2034 | U-55 | | |
| | | 9248229 | Mar 12, 2034 | DP | | |
| <u>MORPHINE SULFATE - MORPHINE SULFATE</u> | | | | | | |
| N 204223 | 004 | 9072781 | Mar 12, 2034 | DP | | |
| | | 9192608 | Mar 12, 2034 | U-43 | | |
| | | 9192608 | Mar 12, 2034 | U-55 | | |
| | | 9248229 | Mar 12, 2034 | DP | | |
| <u>MORPHINE SULFATE - MORPHINE SULFATE</u> | | | | | | |
| N 204223 | 005 | 9072781 | Mar 12, 2034 | DP | | |
| | | 9192608 | Mar 12, 2034 | U-43 | | |
| | | 9192608 | Mar 12, 2034 | U-55 | | |
| | | 9248229 | Mar 12, 2034 | DP | | |
| <u>MORPHINE SULFATE - MORPHABOND ER</u> | | | | | | |
| N 206544 | 001 | 7955619 | Aug 12, 2028 | DP | | |
| <u>MORPHINE SULFATE - MORPHABOND ER</u> | | | | | | |
| N 206544 | 002 | 7955619 | Aug 12, 2028 | DP | | |
| <u>MORPHINE SULFATE - MORPHABOND ER</u> | | | | | | |
| N 206544 | 003 | 7955619 | Aug 12, 2028 | DP | | |
| <u>MORPHINE SULFATE - MORPHABOND ER</u> | | | | | | |
| N 206544 | 004 | 7955619 | Aug 12, 2028 | DP | | |
| <u>MORPHINE SULFATE - ARYMO ER</u> | | | | | | |
| N 208603 | 001 | 9044402 | Jul 01, 2033 | DP U-1556 | | |
| | | 9549899 | Jul 01, 2033 | DP U-1556 | | |
| <u>MORPHINE SULFATE - ARYMO ER</u> | | | | | | |
| N 208603 | 002 | 9044402 | Jul 01, 2033 | DP U-1556 | | |
| | | 9549899 | Jul 01, 2033 | DP U-1556 | | |
| <u>MORPHINE SULFATE - ARYMO ER</u> | | | | | | |
| N 208603 | 003 | 9044402 | Jul 01, 2033 | DP U-1556 | | |
| | | 9549899 | Jul 01, 2033 | DP U-1556 | | |
| <u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u> | | | | | | |
| N 022321 | 001 | 7682633 | Jun 19, 2027 | U-1510 | | |
| | | 7682634 | Jun 19, 2027 | DP | | |
| | | 7815934 | Dec 12, 2027 | DP | | |
| | | 8158156 | Jun 19, 2027 | U-1510 | | |
| | | 8623418 | Nov 07, 2029 | U-1640 | | |
| | | 8685443 | Jul 03, 2025 | U-1508 | | |
| | | 8685444 | Jul 03, 2025 | DP | | |
| | | 8846104 | Jun 19, 2027 | DP | | |
| | | 8877247 | Jun 19, 2027 | DP | | |
| <u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u> | | | | | | |
| N 022321 | 002 | 7682633 | Jun 19, 2027 | U-1510 | | |
| | | 7682634 | Jun 19, 2027 | DP | | |
| | | 7815934 | Dec 12, 2027 | DP | | |
| | | 8158156 | Jun 19, 2027 | U-1510 | | |
| | | 8623418 | Nov 07, 2029 | U-1640 | | |
| | | 8685443 | Jul 03, 2025 | U-1508 | | |
| | | 8685444 | Jul 03, 2025 | DP | | |
| | | 8846104 | Jun 19, 2027 | DP | | |
| | | 8877247 | Jun 19, 2027 | DP | | |
| <u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u> | | | | | | |
| N 022321 | 003 | 7682633 | Jun 19, 2027 | U-1510 | | |
| | | 7682634 | Jun 19, 2027 | DP | | |
| | | 7815934 | Dec 12, 2027 | DP | | |
| | | 8158156 | Jun 19, 2027 | U-1510 | | |
| | | 8623418 | Nov 07, 2029 | U-1640 | | |
| | | 8685443 | Jul 03, 2025 | U-1508 | | |
| | | 8685444 | Jul 03, 2025 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u> | | | | | | |
| N 022321 003 | 8846104 | Jun 19, 2027 | DP | | | |
| | 8877247 | Jun 19, 2027 | DP | | | |
| <u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u> | | | | | | |
| N 022321 004 | 7682633 | Jun 19, 2027 | | U-1510 | | |
| | 7682634 | Jun 19, 2027 | DP | | | |
| | 7815934 | Dec 12, 2027 | DP | | | |
| | 8158156 | Jun 19, 2027 | | U-1510 | | |
| | 8623418 | Nov 07, 2029 | | U-1640 | | |
| | 8685443 | Jul 03, 2025 | | U-1508 | | |
| | 8685444 | Jul 03, 2025 | DP | | | |
| | 8846104 | Jun 19, 2027 | DP | | | |
| | 8877247 | Jun 19, 2027 | DP | | | |
| <u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u> | | | | | | |
| N 022321 005 | 7682633 | Jun 19, 2027 | | U-1510 | | |
| | 7682634 | Jun 19, 2027 | DP | | | |
| | 7815934 | Dec 12, 2027 | DP | | | |
| | 8158156 | Jun 19, 2027 | | U-1510 | | |
| | 8623418 | Nov 07, 2029 | | U-1640 | | |
| | 8685443 | Jul 03, 2025 | | U-1508 | | |
| | 8685444 | Jul 03, 2025 | DP | | | |
| | 8846104 | Jun 19, 2027 | DP | | | |
| | 8877247 | Jun 19, 2027 | DP | | | |
| <u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u> | | | | | | |
| N 022321 006 | 7682633 | Jun 19, 2027 | | U-1510 | | |
| | 7682634 | Jun 19, 2027 | DP | | | |
| | 7815934 | Dec 12, 2027 | DP | | | |
| | 8158156 | Jun 19, 2027 | | U-1510 | | |
| | 8623418 | Nov 07, 2029 | | U-1640 | | |
| | 8685443 | Jul 03, 2025 | | U-1508 | | |
| | 8685444 | Jul 03, 2025 | DP | | | |
| | 8846104 | Jun 19, 2027 | DP | | | |
| | 8877247 | Jun 19, 2027 | DP | | | |
| <u>MOXIDECTIN - MOXIDECTIN</u> | | | | | | |
| N 210867 001 | | | | | NCE ODE-193 | Jun 13, 2023 Jun 13, 2025 |
| <u>MOXIFLOXACIN HYDROCHLORIDE - AVELOX</u> | | | | | | |
| N 021085 001 | 6610327 | Oct 29, 2019 | DP U-298 | | M-185 | Sep 27, 2019 |
| <u>MOXIFLOXACIN HYDROCHLORIDE - AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER</u> | | | | | | |
| N 021277 001 | 6548079 | Jul 25, 2020 | DP U-298 | | M-185 | Sep 27, 2019 |
| <u>MOXIFLOXACIN HYDROCHLORIDE - VIGAMOX</u> | | | | | | |
| N 021598 001 | 6716830 | Sep 29, 2019 | DP | | | |
| | 7671070 | Sep 29, 2019 | | U-709 | | |
| <u>MOXIFLOXACIN HYDROCHLORIDE - MOXEZA</u> | | | | | | |
| N 022428 001 | 6716830 | Sep 29, 2019 | DP | | | |
| | 7671070 | Sep 29, 2019 | DP U-709 | | | |
| | 8450311 | May 29, 2029 | DP | | | |
| | 9114168 | May 29, 2029 | DP | | | |
| <u>NAFTIFINE HYDROCHLORIDE - NAFTIN</u> | | | | | | |
| N 019599 002 | | | | | M-191 | Nov 10, 2019 |
| <u>NAFTIFINE HYDROCHLORIDE - NAFTIN</u> | | | | | | |
| N 204286 001 | 8778365 | Jan 31, 2033 | DP | | | |
| | 9161914 | Jan 31, 2033 | | U-540 | | |
| <u>NALDEMEDINE TOSYLATE - SYMPROIC</u> | | | | | | |
| N 208854 001 | 9108975 | Nov 11, 2031 | DS DP | | NCE | Mar 23, 2022 |
| | RE46365 | Jan 11, 2028 | DS DP | | | |
| | RE46375 | Oct 05, 2026 | DS DP U-1185 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>NALOXEGOL OXALATE - MOVANTIK</u> | | | | | | |
| N 204760 001 | 7056500 | Jun 29, 2024 | DP | U-1185 | NCE | Sep 16, 2019 |
| | 7662365 | Oct 18, 2022 | DS DP | | | |
| | 7786133 | Dec 19, 2027 | DS DP | | | |
| | 8067431 | Dec 16, 2024 | | U-1185 | | |
| | 8617530 | Oct 18, 2022 | | U-1185 | | |
| | 9012469 | Apr 02, 2032 | DS DP | | | |
| <u>NALOXEGOL OXALATE - MOVANTIK</u> | | | | | | |
| N 204760 002 | 7056500 | Jun 29, 2024 | DP | U-1185 | NCE | Sep 16, 2019 |
| | 7662365 | Oct 18, 2022 | DS DP | | | |
| | 7786133 | Dec 19, 2027 | DS DP | | | |
| | 8067431 | Dec 16, 2024 | | U-1185 | | |
| | 8617530 | Oct 18, 2022 | | U-1185 | | |
| | 9012469 | Apr 02, 2032 | DS DP | | | |
| <u>NALOXONE HYDROCHLORIDE - EVZIO</u> | | | | | | |
| N 205787 001 | 10143972 | May 24, 2031 | | U-2476 | | |
| | 7731686 | Jun 10, 2026 | DP | | | |
| | 7731690 | Jan 15, 2025 | DP | | | |
| | 7749194 | Oct 30, 2028 | DP | | | |
| | 7918823 | Nov 23, 2024 | DP | | | |
| | 7947017 | Mar 12, 2028 | DP | | | |
| | 8016788 | Mar 21, 2025 | DP | | | |
| | 8021344 | Nov 02, 2029 | DP | | | |
| | 8206360 | Feb 27, 2027 | DP | | | |
| | 8226610 | Apr 10, 2029 | DP | | | |
| | 8231573 | Nov 25, 2028 | DP | | | |
| | 8313466 | Nov 23, 2024 | DP | | | |
| | 8361029 | Nov 23, 2024 | DP | | | |
| | 8425462 | Nov 23, 2024 | DP | | | |
| | 8608698 | Nov 23, 2024 | DP | | | |
| | 8627816 | Feb 04, 2032 | DP | | | |
| | 8926594 | Mar 31, 2026 | DP | | | |
| | 8939943 | Feb 28, 2031 | DP | | | |
| | 9022022 | Feb 28, 2031 | DP | | | |
| | 9056170 | Nov 23, 2024 | DP | | | |
| | 9238108 | Feb 20, 2027 | DP | | | |
| | 9278182 | Feb 01, 2026 | DP | | | |
| | 9474869 | Feb 28, 2031 | DP | U-1907 | | |
| | 9517307 | Jul 18, 2034 | DP | U-1925 | | |
| | 9724471 | May 23, 2027 | DP | U-2092 | | |
| | 9737669 | Nov 23, 2024 | DP | | | |
| <u>NALOXONE HYDROCHLORIDE - NARCAN</u> | | | | | | |
| N 208411 001 | 10085937 | Mar 16, 2035 | | U-1903 | | |
| | 9211253 | Mar 16, 2035 | DP | | | |
| | 9468747 | Mar 16, 2035 | DP | U-1903 | | |
| | 9561177 | Mar 16, 2035 | DP | U-1903 | | |
| | 9629965 | Mar 16, 2035 | DP | U-1903 | | |
| | 9775838 | Mar 16, 2035 | | U-1903 | | |
| <u>NALOXONE HYDROCHLORIDE - NARCAN</u> | | | | | | |
| N 208411 002 | 9480644 | Mar 16, 2035 | DP | U-1903 | | |
| | 9707226 | Mar 16, 2035 | DP | U-1903 | | |
| <u>NALOXONE HYDROCHLORIDE - EVZIO</u> | | | | | | |
| N 209862 001 | 10143972 | May 24, 2031 | | U-2476 | | |
| | 7731686 | Jun 01, 2026 | DP | | | |
| | 7731690 | Jan 15, 2025 | DP | | | |
| | 7749194 | Oct 30, 2028 | DP | | | |
| | 7918823 | Nov 23, 2024 | DP | | | |
| | 7947017 | Mar 12, 2028 | DP | | | |
| | 8016788 | Mar 21, 2025 | DP | | | |
| | 8021344 | Nov 02, 2029 | DP | | | |
| | 8206360 | Feb 27, 2027 | DP | | | |
| | 8226610 | Apr 10, 2029 | DP | | | |
| | 8231573 | Nov 25, 2028 | DP | | | |
| | 8313466 | Nov 23, 2024 | DP | | | |
| | 8361029 | Nov 23, 2024 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>NALOXONE HYDROCHLORIDE - EVZIO</u> | | | | | | |
| N 209862 001 | 8425462 | Nov 23, 2024 | DP | | | |
| | 8608698 | Nov 23, 2024 | DP | | | |
| | 8627816 | Feb 04, 2032 | DP | | | |
| | 8926594 | Mar 31, 2026 | DP | | | |
| | 8939943 | Feb 28, 2031 | DP | | | |
| | 9022022 | Feb 28, 2031 | DP | | | |
| | 9056170 | Nov 23, 2024 | DP | | | |
| | 9238108 | Feb 20, 2027 | DP | | | |
| | 9278182 | Feb 01, 2026 | DP | | | |
| | 9474869 | Feb 28, 2031 | DP U-1907 | | | |
| | 9517307 | Jul 18, 2034 | DP U-1925 | | | |
| | 9724471 | May 23, 2027 | DP U-2092 | | | |
| | 9737669 | Nov 23, 2024 | DP | | | |
| <u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u> | | | | | | |
| N 205777 001 | 8846090 | Apr 04, 2023 | DP | | | |
| | 8846091 | Apr 04, 2023 | DP | | | |
| | 8969369 | May 10, 2022 | DP U-1556 | | | |
| | 9056051 | May 10, 2022 | DP U-1556 | | | |
| | 9073933 | Mar 30, 2025 | DS | | | |
| | 9084729 | May 10, 2022 | DP U-1556 | | | |
| | 9161937 | May 10, 2022 | DP U-1556 | | | |
| | 9168252 | May 10, 2022 | DP U-1556 | | | |
| | 9283216 | May 10, 2022 | DP U-1819 | | | |
| | 9283221 | May 10, 2022 | DP U-1819 | | | |
| | 9345701 | May 10, 2022 | DP U-1819 | | | |
| | 9511066 | May 10, 2022 | U-1921 | | | |
| | 9522919 | Mar 30, 2025 | DS DP | | | |
| | 9555000 | Apr 04, 2023 | DP U-1556 | | | |
| | 9907793 | Apr 04, 2023 | DP U-1556 | | | |
| <u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u> | | | | | | |
| N 205777 002 | 8846090 | Apr 04, 2023 | DP | | | |
| | 8846091 | Apr 04, 2023 | DP | | | |
| | 8969369 | May 10, 2022 | DP U-1556 | | | |
| | 9056051 | May 10, 2022 | DP U-1556 | | | |
| | 9073933 | Mar 30, 2025 | DS | | | |
| | 9084729 | May 10, 2022 | DP U-1556 | | | |
| | 9161937 | May 10, 2022 | DP U-1556 | | | |
| | 9168252 | May 10, 2022 | DP U-1556 | | | |
| | 9283216 | May 10, 2022 | DP U-1819 | | | |
| | 9283221 | May 10, 2022 | DP U-1819 | | | |
| | 9345701 | May 10, 2022 | DP U-1819 | | | |
| | 9511066 | May 10, 2022 | U-1921 | | | |
| | 9522919 | Mar 30, 2025 | DS DP | | | |
| | 9555000 | Apr 04, 2023 | DP U-1556 | | | |
| | 9907793 | Apr 04, 2023 | DP U-1556 | | | |
| <u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u> | | | | | | |
| N 205777 003 | 8846090 | Apr 04, 2023 | DP | | | |
| | 8846091 | Apr 04, 2023 | DP | | | |
| | 8969369 | May 10, 2022 | DP U-1556 | | | |
| | 9056051 | May 10, 2022 | DP U-1556 | | | |
| | 9073933 | Mar 30, 2025 | DS | | | |
| | 9084729 | May 10, 2022 | DP U-1556 | | | |
| | 9161937 | May 10, 2022 | DP U-1556 | | | |
| | 9168252 | May 10, 2022 | DP U-1556 | | | |
| | 9283216 | May 10, 2022 | DP U-1819 | | | |
| | 9283221 | May 10, 2022 | DP U-1819 | | | |
| | 9345701 | May 10, 2022 | DP U-1819 | | | |
| | 9511066 | May 10, 2022 | U-1921 | | | |
| | 9522919 | Mar 30, 2025 | DS DP | | | |
| <u>NALTREXONE - VIVITROL</u> | | | | | | |
| N 021897 001 | 6264987 | May 19, 2020 | DP | | | |
| | 6331317 | Nov 12, 2019 | DP | | | |
| | 6379704 | May 19, 2020 | DP | | | |
| | 6395304 | Nov 12, 2019 | DP | | | |
| | 6495164 | May 25, 2020 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>NALTREXONE - VIVITROL</u> | | | | | | |
| N 021897 | 001 | 6495166 | Nov 12, 2019 | DP | | |
| | | 6534092 | May 19, 2020 | DP | | |
| | | 6537586 | Nov 12, 2019 | DP | | |
| | | 6667061 | May 25, 2020 | DP | | |
| | | 6713090 | Nov 12, 2019 | DP | | |
| | | 6939033 | Nov 12, 2019 | DP | | |
| | | 7799345 | May 25, 2020 | DP | | |
| | | 7919499 | Oct 15, 2029 | U-1123 | | |
| | | 7919499 | Oct 15, 2029 | U-1124 | | |
| <u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u> | | | | | | |
| N 207621 | 001 | 7815934 | Dec 12, 2027 | DP | NC | Aug 19, 2019 |
| | | 8685443 | Jul 03, 2025 | U-1508 | | |
| <u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u> | | | | | | |
| N 207621 | 002 | 7815934 | Dec 12, 2027 | DP | NC | Aug 19, 2019 |
| | | 8685443 | Jul 03, 2025 | U-1508 | | |
| <u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u> | | | | | | |
| N 207621 | 003 | 7815934 | Dec 12, 2027 | DP | NC | Aug 19, 2019 |
| | | 8685443 | Jul 03, 2025 | U-1508 | | |
| <u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u> | | | | | | |
| N 207621 | 004 | 7815934 | Dec 12, 2027 | DP | NC | Aug 19, 2019 |
| | | 8685443 | Jul 03, 2025 | U-1508 | | |
| <u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u> | | | | | | |
| N 207621 | 005 | 7815934 | Dec 12, 2027 | DP | NC | Aug 19, 2019 |
| | | 8685443 | Jul 03, 2025 | U-1508 | | |
| <u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u> | | | | | | |
| N 207621 | 006 | 7815934 | Dec 12, 2027 | DP | NC | Aug 19, 2019 |
| | | 8685443 | Jul 03, 2025 | U-1508 | | |
| <u>NAPROXEN SODIUM - NAPROXEN SODIUM</u> | | | | | | |
| N 021920 | 001 | 10022344 | Mar 03, 2026 | DP U-1731 | | |
| | | 10022344 | Mar 03, 2026 | DP U-1732 | | |
| | | 10028925 | Mar 03, 2026 | DP U-1731 | | |
| | | 10028925 | Mar 03, 2026 | DP U-1732 | | |
| | | 9693978 | Mar 03, 2026 | DP | | |
| | | 9693979 | Mar 03, 2026 | DP | | |
| <u>NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE - TREXIMET</u> | | | | | | |
| N 021926 | 001 | 7332183 | Oct 02, 2025 | DP U-867 | | |
| | | 7332183*PED | Apr 02, 2026 | | | |
| <u>NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE - TREXIMET</u> | | | | | | |
| N 021926 | 002 | 7332183 | Oct 02, 2025 | DP U-1719 | | |
| | | 7332183*PED | Apr 02, 2026 | | | |
| <u>NATEGLINIDE - STARLIX</u> | | | | | | |
| N 021204 | 001 | 6559188 | Sep 15, 2020 | DP U-827 | | |
| | | 6878749 | Sep 15, 2020 | DP | | |
| <u>NATEGLINIDE - STARLIX</u> | | | | | | |
| N 021204 | 002 | 6559188 | Sep 15, 2020 | DP U-827 | | |
| | | 6878749 | Sep 15, 2020 | DP | | |
| <u>NEBIVOLOL HYDROCHLORIDE - BYSTOLIC</u> | | | | | | |
| N 021742 | 002 | 6545040 | Dec 17, 2021 | DP U-3 | | |
| <u>NEBIVOLOL HYDROCHLORIDE - BYSTOLIC</u> | | | | | | |
| N 021742 | 003 | 6545040 | Dec 17, 2021 | DP U-3 | | |
| <u>NEBIVOLOL HYDROCHLORIDE - BYSTOLIC</u> | | | | | | |
| N 021742 | 004 | 6545040 | Dec 17, 2021 | DP U-3 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------------------------------------|--------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>NEBIVOLOL HYDROCHLORIDE - BYSTOLIC</u> | | | | | | |
| N 021742 | 005 6545040 | Dec 17, 2021 | DP U-3 | | | |
| <u>NEBIVOLOL HYDROCHLORIDE; VALSARTAN - BYVALSON</u> | | | | | | |
| N 206302 | 001 7803838 | Aug 29, 2026 | DP | | NC | Jun 03, 2019 |
| | 7838552 | Oct 04, 2027 | U-185 | | | |
| <u>NEPAFENAC - NEVANAC</u> | | | | | | |
| N 021862 | 001 7834059 | Jan 31, 2027 | U-1095 | | | |
| | 8071648 | Dec 02, 2025 | DP | | | |
| | 8324281 | Dec 02, 2025 | DP | | | |
| <u>NEPAFENAC - ILEVRO</u> | | | | | | |
| N 203491 | 001 7947295 | Jun 08, 2024 | DP | | | |
| | 8921337 | Mar 31, 2032 | DP | | | |
| | 9662398 | Dec 01, 2030 | DP | | | |
| <u>NERATINIB MALEATE - NERLYNX</u> | | | | | | |
| N 208051 | 001 10035788 | Oct 15, 2028 | U-2043 | | NCE | Jul 17, 2022 |
| | 6288082 | Sep 24, 2019 | DS DP U-2043 | | | |
| | 7399865 | Dec 29, 2025 | DS DP | | | |
| | 7982043 | Oct 08, 2025 | U-2043 | | | |
| | 8518446 | Nov 20, 2030 | DP U-2043 | | | |
| | 8790708 | Nov 05, 2030 | DP U-2043 | | | |
| | 9139558 | Oct 15, 2028 | U-2043 | | | |
| | 9211291 | Mar 24, 2030 | U-2043 | | | |
| | 9630946 | Oct 15, 2028 | U-2043 | | | |
| <u>NETARSUDIL DIMESYLATE - RHOPRESSA</u> | | | | | | |
| N 208254 | 001 8394826 | Nov 10, 2030 | DS DP U-1524 | | NCE | Dec 18, 2022 |
| | 8450344 | Jul 11, 2026 | DS DP U-1524 | | | |
| | 9096569 | Jul 11, 2026 | DS DP U-1524 | | | |
| | 9415043 | Mar 14, 2034 | DS | | | |
| | 9931336 | Mar 14, 2034 | DS DP U-1524 | | | |
| <u>NETUPITANT; PALONOSETRON HYDROCHLORIDE - AKYNZEO</u> | | | | | | |
| N 205718 | 001 6297375 | Feb 22, 2020 | DS | | NCE | Oct 10, 2019 |
| | 8623826 | Nov 18, 2030 | U-2293 | | | |
| | 8951969 | Nov 18, 2030 | DP | | | |
| | 9186357 | Nov 18, 2030 | U-2293 | | | |
| | 9271975 | Sep 09, 2031 | U-2293 | | | |
| | 9943515 | Nov 18, 2030 | U-2293 | | | |
| | 9951016 | Sep 25, 2035 | DS DP | | | |
| <u>NEVIRAPINE - VIRAMUNE XR</u> | | | | | | |
| N 201152 | 001 8460704 | Mar 12, 2029 | U-1409 | | | |
| <u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER</u> | | | | | | |
| N 019734 | 002 7612102 | Dec 26, 2027 | DP | | | |
| | 7659291 | Apr 18, 2027 | U-1029 | | | |
| | 8455524 | Apr 18, 2027 | U-1029 | | | |
| | 9364564 | Dec 26, 2027 | DP | | | |
| <u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER</u> | | | | | | |
| N 019734 | 003 7612102 | Dec 26, 2027 | DP | | | |
| | 7659291 | Apr 18, 2027 | U-1029 | | | |
| | 8455524 | Apr 18, 2027 | U-1029 | | | |
| | 9364564 | Dec 26, 2027 | DP | | | |
| <u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER</u> | | | | | | |
| N 019734 | 004 7612102 | Dec 26, 2027 | DP | | | |
| | 7659291 | Apr 18, 2027 | U-1029 | | | |
| | 8455524 | Apr 18, 2027 | U-1029 | | | |
| | 9364564 | Dec 26, 2027 | DP | | | |
| <u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 5.0% DEXTROSE IN PLASTIC CONTAINER</u> | | | | | | |
| N 019734 | 005 7612102 | Dec 26, 2027 | DP | | | |
| | 7659291 | Apr 18, 2027 | U-1029 | | | |
| | 8455524 | Apr 18, 2027 | U-1029 | | | |
| | 9364564 | Dec 26, 2027 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>NICOTINE - NICODERM CO</u> | | | | | | |
| N 020165 004 | 8075911 | May 22, 2021 | DP | | | |
| | 8663680 | Feb 13, 2020 | DP | | | |
| | 8999379 | Feb 13, 2020 | | U-1686 | | |
| | 9205059 | Dec 15, 2019 | DP | | | |
| <u>NICOTINE - NICODERM CO</u> | | | | | | |
| N 020165 005 | 8075911 | May 22, 2021 | DP | | | |
| | 8663680 | Feb 13, 2020 | DP | | | |
| | 8999379 | Feb 13, 2020 | | U-1686 | | |
| | 9205059 | Dec 15, 2019 | DP | | | |
| <u>NICOTINE - NICODERM CO</u> | | | | | | |
| N 020165 006 | 8075911 | May 22, 2021 | DP | | | |
| | 8663680 | Feb 13, 2020 | DP | | | |
| | 8999379 | Feb 13, 2020 | | U-1686 | | |
| | 9205059 | Dec 15, 2019 | DP | | | |
| <u>NICOTINE POLACRILEX - NICORETTE</u> | | | | | | |
| N 018612 002 | 8323683 | Apr 30, 2028 | | | | |
| <u>NICOTINE POLACRILEX - NICORETTE</u> | | | | | | |
| N 020066 002 | 8323683 | Apr 30, 2028 | DP | | | |
| <u>NICOTINE POLACRILEX - NICORETTE</u> | | | | | | |
| N 022360 001 | 8501164 | Jun 14, 2029 | DP | | | |
| | 8940772 | Apr 30, 2029 | DP | | | |
| <u>NICOTINE POLACRILEX - NICORETTE</u> | | | | | | |
| N 022360 002 | 8501164 | Jun 14, 2029 | DP | | | |
| | 8940772 | Apr 30, 2029 | DP | | | |
| <u>NILOTINIB HYDROCHLORIDE - TASIGNA</u> | | | | | | |
| N 022068 001 | 7169791 | Jul 04, 2023 | DS DP U-836 | | D-170 | Dec 22, 2020 |
| | 7169791*PED | Jan 04, 2024 | | | NPP | Mar 22, 2021 |
| | 8163904 | Aug 23, 2028 | DS DP | | ODE-171 | Mar 22, 2025 |
| | 8163904*PED | Feb 23, 2029 | | | ODE-172 | Mar 22, 2025 |
| | 8293756 | Sep 25, 2027 | | DP | PED | Jun 22, 2021 |
| | 8293756*PED | Mar 25, 2028 | | | PED | Sep 22, 2021 |
| | 8389537 | Jul 18, 2026 | DS DP U-1374 | | PED | Sep 22, 2025 |
| | 8389537*PED | Jan 18, 2027 | | | PED | Sep 22, 2025 |
| | 8415363 | Jul 18, 2026 | DS DP U-1407 | | | |
| | 8415363*PED | Jan 18, 2027 | | | | |
| | 8501760 | Jul 18, 2026 | DS DP | | | |
| | 8501760*PED | Jan 18, 2027 | | | | |
| | 9061029 | Apr 07, 2032 | | DP U-1374 | | |
| | 9061029*PED | Oct 07, 2032 | | | | |
| <u>NILOTINIB HYDROCHLORIDE - TASIGNA</u> | | | | | | |
| N 022068 002 | 7169791 | Jul 04, 2023 | DS DP U-836 | | D-170 | Dec 22, 2020 |
| | 7169791*PED | Jan 04, 2024 | | | NPP | Mar 22, 2021 |
| | 8163904 | Aug 23, 2028 | DS DP | | ODE-171 | Mar 22, 2025 |
| | 8163904*PED | Feb 23, 2029 | | | ODE-172 | Mar 22, 2025 |
| | 8293756 | Sep 25, 2027 | | DP | PED | Jun 22, 2021 |
| | 8293756*PED | Mar 25, 2028 | | | PED | Sep 22, 2021 |
| | 8389537 | Jul 18, 2026 | DS DP U-1374 | | PED | Sep 22, 2025 |
| | 8389537*PED | Jan 18, 2027 | | | PED | Sep 22, 2025 |
| | 8415363 | Jul 18, 2026 | DS DP U-1407 | | | |
| | 8415363*PED | Jan 18, 2027 | | | | |
| | 8501760 | Jul 18, 2026 | DS DP | | | |
| | 8501760*PED | Jan 18, 2027 | | | | |
| | 9061029 | Apr 07, 2032 | | DP U-1374 | | |
| | 9061029*PED | Oct 07, 2032 | | | | |
| <u>NILOTINIB HYDROCHLORIDE - TASIGNA</u> | | | | | | |
| N 022068 003 | 7169791 | Jul 04, 2023 | DS DP U-836 | | NPP | Mar 22, 2021 |
| | 7169791*PED | Jan 04, 2024 | | | ODE-171 | Mar 22, 2025 |
| | 8163904 | Aug 23, 2028 | DS DP | | ODE-172 | Mar 22, 2025 |
| | 8163904*PED | Feb 23, 2029 | | | PED | Sep 22, 2021 |
| | 8293756 | Sep 25, 2027 | | DP | PED | Sep 22, 2025 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>NILOTINIB HYDROCHLORIDE - TASIGNA</u> | | | | | | |
| N 022068 | 003 | 8293756*PED | Mar 25, 2028 | | PED | Sep 22, 2025 |
| | | 8389537 | Jul 18, 2026 | DS DP U-1374 | | |
| | | 8389537*PED | Jan 18, 2027 | | | |
| | | 8415363 | Jul 18, 2026 | DS DP U-1407 | | |
| | | 8415363*PED | Jan 18, 2027 | | | |
| | | 8501760 | Jul 18, 2026 | DS DP | | |
| | | 8501760*PED | Jan 18, 2027 | | | |
| | | 9061029 | Apr 07, 2032 | DS DP U-1374 | | |
| | | 9061029*PED | Oct 07, 2032 | | | |
| <u>NIMODIPINE - NYMALIZE</u> | | | | | | |
| N 203340 | 001 | | | | ODE-46 | May 10, 2020 |
| <u>NINTEDANIB ESYLATE - OFEV</u> | | | | | | |
| N 205832 | 001 | 10105323 | Jun 04, 2029 | DP | NCE | Oct 15, 2019 |
| | | 6762180 | Oct 03, 2020 | DS DP | ODE-77 | Oct 15, 2021 |
| | | 7119093 | Feb 21, 2024 | DS DP | | |
| | | 7989474 | Apr 06, 2024 | U-1677 | | |
| | | 9907756 | Jun 07, 2029 | DP | | |
| <u>NINTEDANIB ESYLATE - OFEV</u> | | | | | | |
| N 205832 | 002 | 10105323 | Jun 04, 2029 | DP | NCE | Oct 15, 2019 |
| | | 6762180 | Oct 03, 2020 | DS DP | ODE-77 | Oct 15, 2021 |
| | | 7119093 | Feb 21, 2024 | DS DP | | |
| | | 7989474 | Apr 06, 2024 | U-1677 | | |
| | | 9907756 | Jun 07, 2029 | DP | | |
| <u>NIRAPARIB TOSYLATE - ZEJULA</u> | | | | | | |
| N 208447 | 001 | 8071623 | Mar 22, 2030 | DS DP | NCE | Mar 27, 2022 |
| | | 8436185 | Apr 24, 2029 | DS | ODE-133 | Mar 27, 2024 |
| <u>NITISINONE - ORFADIN</u> | | | | | | |
| N 021232 | 001 | | | | D-169 | Sep 01, 2020 |
| <u>NITISINONE - ORFADIN</u> | | | | | | |
| N 021232 | 002 | | | | D-169 | Sep 01, 2020 |
| <u>NITISINONE - ORFADIN</u> | | | | | | |
| N 021232 | 003 | | | | D-169 | Sep 01, 2020 |
| <u>NITISINONE - ORFADIN</u> | | | | | | |
| N 021232 | 004 | | | | D-169 | Sep 01, 2020 |
| <u>NITISINONE - ORFADIN</u> | | | | | | |
| N 206356 | 001 | 9301932 | Feb 28, 2033 | DP U-1836 | D-169 | Sep 01, 2020 |
| <u>NITRIC OXIDE - INOMAX</u> | | | | | | |
| N 020845 | 002 | 8282966 | Jun 30, 2029 | U-1286 | | |
| | | 8291904 | Jan 06, 2031 | DP U-1226 | | |
| | | 8293284 | Jun 30, 2029 | U-1286 | | |
| | | 8431163 | Jun 30, 2029 | U-1286 | | |
| | | 8431163*PED | Dec 30, 2029 | | | |
| | | 8573209 | Jan 06, 2031 | DP | | |
| | | 8573209*PED | Jul 06, 2031 | | | |
| | | 8573210 | Jan 06, 2031 | DP U-1453 | | |
| | | 8573210*PED | Jul 06, 2031 | | | |
| | | 8776794 | Jan 06, 2031 | DP U-1226 | | |
| | | 8776794*PED | Jul 06, 2031 | | | |
| | | 8776795 | Jan 06, 2031 | DP U-1226 | | |
| | | 8776795*PED | Jul 06, 2031 | | | |
| | | 8795741 | Jun 30, 2029 | U-1286 | | |
| | | 8795741*PED | Dec 30, 2029 | | | |
| | | 8846112 | Jun 30, 2029 | U-1286 | | |
| | | 8846112*PED | Dec 30, 2029 | | | |
| <u>NITRIC OXIDE - INOMAX</u> | | | | | | |
| N 020845 | 003 | 8282966 | Jun 30, 2029 | U-1286 | | |
| | | 8291904 | Jan 06, 2031 | DP U-1226 | | |
| | | 8293284 | Jun 30, 2029 | U-1286 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>NITRIC OXIDE - INOMAX</u> | | | | | | |
| N 020845 003 | 8431163 | Jun 30, 2029 | | U-1286 | | |
| | 8431163*PED | Dec 30, 2029 | | | | |
| | 8573209 | Jan 06, 2031 | DP | | | |
| | 8573209*PED | Jul 06, 2031 | | | | |
| | 8573210 | Jan 06, 2031 | DP | U-1453 | | |
| | 8573210*PED | Jul 06, 2031 | | | | |
| | 8776794 | Jan 06, 2031 | DP | U-1226 | | |
| | 8776794*PED | Jul 06, 2031 | | | | |
| | 8776795 | Jan 06, 2031 | DP | U-1226 | | |
| | 8776795*PED | Jul 06, 2031 | | | | |
| | 8795741 | Jun 30, 2029 | | U-1286 | | |
| | 8795741*PED | Dec 30, 2029 | | | | |
| | 8846112 | Jun 30, 2029 | | U-1286 | | |
| | 8846112*PED | Dec 30, 2029 | | | | |
| | 9265911 | Jan 06, 2031 | DP | U-1824 | | |
| | 9265911*PED | Jul 06, 2031 | | | | |
| | 9279794 | Feb 19, 2034 | DP | U-1823 | | |
| | 9279794*PED | Aug 19, 2034 | | | | |
| | 9295802 | Jan 06, 2031 | DP | U-1226 | | |
| | 9295802*PED | Jul 06, 2031 | | | | |
| | 9408993 | Jan 06, 2031 | DP | U-1824 | | |
| | 9408993*PED | Jul 06, 2031 | | | | |
| | 9770570 | May 03, 2036 | | U-2148 | | |
| | 9770570*PED | Nov 03, 2036 | | | | |
| <u>NITROGLYCERIN - NITROLINGUAL PUMPSPRAY</u> | | | | | | |
| N 018705 002 | 7872049 | Mar 12, 2029 | DP | U-2223 | | |
| <u>NITROGLYCERIN - GONITRO</u> | | | | | | |
| N 208424 001 | 9101592 | Mar 11, 2032 | DP | | | |
| <u>NIZATIDINE - AXID</u> | | | | | | |
| N 021494 001 | 6930119 | Jul 17, 2022 | DP | | | |
| <u>NUSINERSEN SODIUM - SPINRAZA</u> | | | | | | |
| N 209531 001 | 7101993 | Sep 05, 2023 | DS | | M-226 | May 14, 2021 |
| | 7838657 | Jul 11, 2027 | DS | | NCE | Dec 23, 2021 |
| | 8110560 | Dec 05, 2025 | | U-1942 | ODE-127 | Dec 23, 2023 |
| | 8110560 | Dec 05, 2025 | | U-1943 | | |
| | 8110560 | Dec 05, 2025 | | U-1944 | | |
| | 8361977 | May 27, 2030 | DS DP | | | |
| | 8980853 | Nov 24, 2030 | | U-1941 | | |
| | 9717750 | Jun 17, 2030 | | U-1942 | | |
| | 9717750 | Jun 17, 2030 | | U-1943 | | |
| | 9717750 | Jun 17, 2030 | | U-2093 | | |
| | 9717750 | Jun 17, 2030 | | U-2094 | | |
| | 9926559 | Jan 09, 2034 | | U-1943 | | |
| <u>OBETICHOLIC ACID - OCALIVA</u> | | | | | | |
| N 207999 001 | 10047117 | Sep 06, 2033 | | U-1854 | NCE | May 27, 2021 |
| | 10052337 | Apr 26, 2036 | DP | | ODE-119 | May 27, 2023 |
| | 10174073 | Jun 17, 2033 | DS | | | |
| | 7138390 | Nov 16, 2022 | DS DP | | | |
| | 8058267 | Feb 21, 2022 | | U-1854 | | |
| | 8377916 | Feb 21, 2022 | | U-1854 | | |
| | 9238673 | Jun 17, 2033 | DP | | | |
| <u>OBETICHOLIC ACID - OCALIVA</u> | | | | | | |
| N 207999 002 | 10047117 | Sep 06, 2033 | | U-1854 | NCE | May 27, 2021 |
| | 10052337 | Apr 26, 2036 | DP | | ODE-119 | May 27, 2023 |
| | 10174073 | Jun 17, 2033 | DS | | | |
| | 7138390 | Nov 16, 2022 | DS DP | | | |
| | 8058267 | Feb 21, 2022 | | U-1854 | | |
| | 8377916 | Feb 21, 2022 | | U-1854 | | |
| | 9238673 | Jun 17, 2033 | DP | | | |
| <u>OLAPARIB - LYNPARZA</u> | | | | | | |
| N 206162 001 | 7151102 | Apr 29, 2022 | DS DP | | NCE | Dec 19, 2019 |
| | 7449464 | Oct 11, 2024 | DS DP | | ODE-83 | Dec 19, 2021 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------------------|--------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>OLAPARIB - LYNPARZA</u> | | | | | | |
| N 206162 001 | 7981889 | Oct 11, 2024 | DS DP | | | |
| | 8143241 | Aug 12, 2027 | | U-1634 | | |
| | 8247416 | Sep 24, 2028 | DS | | | |
| | 8859562 | Aug 04, 2031 | | U-1634 | | |
| | 8912187 | Mar 12, 2024 | | U-1634 | | |
| <u>OLAPARIB - LYNPARZA</u> | | | | | | |
| N 208558 001 | 7151102 | Apr 29, 2022 | DS DP | | I-762 | Jan 12, 2021 |
| | 7449464 | Oct 11, 2024 | DS DP | | I-776 | Dec 19, 2021 |
| | 7981889 | Oct 11, 2024 | DS DP | | NCE | Dec 19, 2019 |
| | 8143241 | Aug 12, 2027 | | U-2101 | NP | Aug 17, 2020 |
| | 8143241 | Aug 12, 2027 | | U-2102 | ODE-180 | Aug 17, 2024 |
| | 8143241 | Aug 12, 2027 | | U-2103 | ODE-181 | Aug 17, 2024 |
| | 8143241 | Aug 12, 2027 | | U-2358 | ODE-83 | Dec 19, 2021 |
| | 8143241 | Aug 12, 2027 | | U-2359 | | |
| | 8475842 | Dec 31, 2029 | DP | | | |
| | 8859562 | Aug 04, 2031 | | U-2101 | | |
| | 8859562 | Aug 04, 2031 | | U-2102 | | |
| | 8859562 | Aug 04, 2031 | | U-2358 | | |
| | 8859562 | Aug 04, 2031 | | U-2359 | | |
| | 8912187 | Mar 12, 2024 | | U-2101 | | |
| | 8912187 | Mar 12, 2024 | | U-2102 | | |
| | 8912187 | Mar 12, 2024 | | U-2358 | | |
| | 8912187 | Mar 12, 2024 | | U-2359 | | |
| <u>OLAPARIB - LYNPARZA</u> | | | | | | |
| N 208558 002 | 7151102 | Apr 29, 2022 | DS DP | | I-762 | Jan 12, 2021 |
| | 7449464 | Oct 11, 2024 | DS DP | | I-776 | Dec 19, 2021 |
| | 7981889 | Oct 11, 2024 | DS DP | | NCE | Dec 19, 2019 |
| | 8143241 | Aug 12, 2027 | | U-2101 | NP | Aug 17, 2020 |
| | 8143241 | Aug 12, 2027 | | U-2102 | ODE-180 | Aug 17, 2024 |
| | 8143241 | Aug 12, 2027 | | U-2103 | ODE-181 | Aug 17, 2024 |
| | 8143241 | Aug 12, 2027 | | U-2358 | ODE-83 | Dec 19, 2021 |
| | 8143241 | Aug 12, 2027 | | U-2359 | | |
| | 8475842 | Dec 31, 2029 | DP | | | |
| | 8859562 | Aug 04, 2031 | | U-2101 | | |
| | 8859562 | Aug 04, 2031 | | U-2102 | | |
| | 8859562 | Aug 04, 2031 | | U-2358 | | |
| | 8859562 | Aug 04, 2031 | | U-2359 | | |
| | 8912187 | Mar 12, 2024 | | U-2101 | | |
| | 8912187 | Mar 12, 2024 | | U-2102 | | |
| | 8912187 | Mar 12, 2024 | | U-2358 | | |
| | 8912187 | Mar 12, 2024 | | U-2359 | | |
| <u>OLODATEROL HYDROCHLORIDE - STRIVERDI RESPIMAT</u> | | | | | | |
| N 203108 001 | 6988496 | Feb 23, 2020 | | DP U-1547 | NCE | Jul 31, 2019 |
| | 7056916 | Dec 07, 2023 | DS DP | | | |
| | 7220742 | May 12, 2025 | DS DP | U-1547 | | |
| | 7284474 | Aug 26, 2024 | | DP | | |
| | 7396341 | Oct 10, 2026 | | DP U-1547 | | |
| | 7491719 | Nov 10, 2023 | DS DP | | | |
| | 7727984 | Nov 10, 2023 | DS | | | |
| | 7786111 | Nov 10, 2023 | | DP | | |
| | 7802568 | Feb 26, 2019 | | DP | | |
| | 7837235 | Mar 13, 2028 | | DP | | |
| | 7896264 | May 26, 2025 | | DP | | |
| | 7988001 | Aug 04, 2021 | | DP | | |
| | 8034809 | May 12, 2025 | | U-1547 | | |
| | 8044046 | Nov 10, 2023 | | U-1547 | | |
| | 8733341 | Oct 16, 2030 | | DP | | |
| 9027967 | Mar 31, 2027 | | DP | | | |
| <u>OLODATEROL HYDROCHLORIDE; TIOTROPIUM BROMIDE - STIOLTO RESPIMAT</u> | | | | | | |
| N 206756 001 | 6846413*PED | Feb 28, 2019 | | | M-173 | Mar 18, 2019 |
| | 6977042*PED | Feb 28, 2019 | | | M-233 | Oct 05, 2021 |
| | 6988496 | Feb 23, 2020 | DP | | NCE | Jul 31, 2019 |
| | 6988496*PED | Aug 23, 2020 | | | | |
| | 7056916 | Dec 07, 2023 | DS DP | | | |
| | 7220742 | May 12, 2025 | DS DP | U-1703 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------------|--------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>OLOPATADINE HYDROCHLORIDE; TIOTROPIUM BROMIDE - STIOLTO RESPIMAT</u> | | | | | | |
| N 206756 | 001 7284474 | Aug 26, 2024 | DP | | | |
| | 7284474*PED | Feb 26, 2025 | | | | |
| | 7396341 | Oct 10, 2026 | DP | | | |
| | 7396341*PED | Apr 10, 2027 | | | | |
| | 7491719 | Nov 10, 2023 | DS DP | | | |
| | 7727984 | Nov 10, 2023 | DS | | | |
| | 7786111 | Nov 10, 2023 | DP | | | |
| | 7802568 | Feb 26, 2019 | DP | | | |
| | 7802568*PED | Aug 26, 2019 | | | | |
| | 7837235 | Mar 13, 2028 | DP | | | |
| | 7837235*PED | Sep 13, 2028 | | | | |
| | 7896264 | May 26, 2025 | DP | | | |
| | 7988001 | Aug 04, 2021 | DP | | | |
| | 8034809 | May 12, 2025 | | U-1702 | | |
| | 8044046 | Nov 10, 2023 | | U-1702 | | |
| | 8733341 | Oct 16, 2030 | DP | | | |
| | 9027967 | Mar 31, 2027 | DP | | | |
| <u>OLOPATADINE HYDROCHLORIDE - PATADAY</u> | | | | | | |
| N 021545 | 001 6995186 | Nov 12, 2023 | DP U-765 | | | |
| | 7402609 | Jun 19, 2022 | DP | | | |
| <u>OLOPATADINE HYDROCHLORIDE - PATANASE</u> | | | | | | |
| N 021861 | 001 7977376 | Feb 02, 2023 | DP | | | |
| | 8399508 | Sep 17, 2022 | | U-726 | | |
| | 8399508*PED | Mar 17, 2023 | | | | |
| <u>OLOPATADINE HYDROCHLORIDE - PAZEO</u> | | | | | | |
| N 206276 | 001 8791154 | May 19, 2032 | DP U-1680 | | | |
| | 9533053 | May 19, 2032 | DP | | | |
| <u>OMACETAXINE MEPESUCCINATE - SYNRIBO</u> | | | | | | |
| N 203585 | 001 6987103 | Oct 26, 2026 | | U-1300 | ODE-32 | Oct 26, 2019 |
| | RE45128 | Mar 16, 2019 | DS DP | U-1576 | | |
| <u>OMADACYCLINE TOSYLATE - NUZYRA</u> | | | | | | |
| N 209816 | 001 10111890 | Aug 03, 2037 | | U-2444 | NCE | Oct 02, 2023 |
| | 10124014 | Mar 05, 2029 | | U-2449 | GAIN | Oct 02, 2028 |
| <u>OMADACYCLINE TOSYLATE - NUZYRA</u> | | | | | | |
| N 209817 | 001 10124014 | Mar 05, 2029 | | U-2449 | NCE | Oct 02, 2023 |
| | | | | | GAIN | Oct 02, 2028 |
| <u>OMBITASVIR; PARITAPREVIR; RITONAVIR - TECHNIVIE</u> | | | | | | |
| N 207931 | 001 7148359 | Jul 19, 2019 | DP | | NCE | Dec 19, 2019 |
| | 7148359*PED | Jan 19, 2020 | | | | |
| | 7364752 | Nov 10, 2020 | DP | | | |
| | 7364752*PED | May 10, 2021 | | | | |
| | 8268349 | Aug 25, 2024 | DP | | | |
| | 8268349*PED | Feb 25, 2025 | | | | |
| | 8399015 | Aug 25, 2024 | DP | | | |
| | 8399015*PED | Feb 25, 2025 | | | | |
| | 8420596 | Apr 10, 2031 | DS DP | | | |
| | 8420596*PED | Oct 10, 2031 | | | | |
| | 8642538 | Sep 10, 2029 | DS DP | U-1638 | | |
| | 8686026 | Jun 09, 2031 | DP | | | |
| | 8691938 | Apr 13, 2032 | DS DP | | | |
| | 9006387 | Jun 10, 2030 | | U-1687 | | |
| | 9044480 | Apr 10, 2031 | | U-1638 | | |
| <u>OMEGA-3-ACID ETHYL ESTERS TYPE A - OMTRYG</u> | | | | | | |
| N 204977 | 001 | | | | NCE | Apr 23, 2019 |
| <u>OMEGA-3-CARBOXYLIC ACIDS - EPANOVA</u> | | | | | | |
| N 205060 | 001 10117844 | Jan 04, 2033 | | U-2447 | NCE | May 05, 2019 |
| | 5792795 | May 13, 2019 | DP | | | |
| | 5948818 | May 13, 2019 | DP | | | |
| | 7960370 | Feb 07, 2025 | DP | | | |
| | 8383678 | Feb 07, 2025 | DP | U-1511 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>OMEGA-3-CARBOXYLIC ACIDS - EPANOVA</u> | | | | | | |
| N 205060 001 | 9012501 | Feb 07, 2025 | DP U-1511 | | | |
| | 9050308 | Jan 04, 2033 | U-1511 | | | |
| | 9050309 | Jan 04, 2033 | DS | | | |
| | 9132112 | Feb 07, 2025 | DP U-1511 | | | |
| <u>OMEPRAZOLE - OMEPRAZOLE</u> | | | | | | |
| N 022032 001 | 9023391 | Aug 16, 2025 | DP | | | |
| <u>OMEPRAZOLE - OMEPRAZOLE</u> | | | | | | |
| N 209400 001 | 10076494 | Dec 08, 2036 | DP | | | |
| <u>OMEPRAZOLE MAGNESIUM - PRILOSEC OTC</u> | | | | | | |
| N 021229 001 | 6403616 | Nov 15, 2019 | | | | |
| | 6428810 | Nov 03, 2019 | | | | |
| <u>OMEPRAZOLE MAGNESIUM - PRILOSEC</u> | | | | | | |
| N 022056 001 | 6428810 | Nov 03, 2019 | DP U-1817 | | | |
| | 6428810 | Nov 03, 2019 | DP U-864 | | | |
| <u>OMEPRAZOLE MAGNESIUM - PRILOSEC</u> | | | | | | |
| N 022056 002 | 6428810 | Nov 03, 2019 | DP U-864 | | | |
| <u>ONDANSETRON - ZUPLENZ</u> | | | | | | |
| N 022524 001 | 8580830 | Nov 23, 2029 | DP | | | |
| | 9095577 | Jul 13, 2030 | DP | | | |
| <u>ONDANSETRON - ZUPLENZ</u> | | | | | | |
| N 022524 002 | 8580830 | Nov 23, 2029 | DP | | | |
| | 9095577 | Jul 13, 2030 | DP | | | |
| <u>ORITAVANCIN DIPHOSPHATE - ORBACTIV</u> | | | | | | |
| N 206334 001 | 5840684 | Nov 24, 2020 | DS DP U-1569 | | NCE | Aug 06, 2019 |
| | 8420592 | Aug 29, 2029 | U-1570 | | GAIN | Aug 06, 2024 |
| | 9649352 | Jul 16, 2035 | DP | | | |
| | 9682061 | Apr 26, 2030 | U-1569 | | | |
| <u>OSIMERTINIB MESYLATE - TAGRISSO</u> | | | | | | |
| N 208065 001 | 8946235 | Aug 08, 2032 | DS DP U-1777 | | I-774 | Apr 18, 2021 |
| | 8946235 | Aug 08, 2032 | DS DP U-2289 | | NCE | Nov 13, 2020 |
| | 9732058 | Jul 25, 2032 | DS DP U-1777 | | ODE-102 | Nov 13, 2022 |
| | 9732058 | Jul 25, 2032 | DS DP U-2289 | | ODE-176 | Apr 18, 2025 |
| <u>OSIMERTINIB MESYLATE - TAGRISSO</u> | | | | | | |
| N 208065 002 | 8946235 | Aug 08, 2032 | DS DP U-1777 | | I-774 | Apr 18, 2021 |
| | 8946235 | Aug 08, 2032 | DS DP U-2289 | | NCE | Nov 13, 2020 |
| | 9732058 | Jul 25, 2032 | DS DP U-1777 | | ODE-102 | Nov 13, 2022 |
| | 9732058 | Jul 25, 2032 | DS DP U-2289 | | ODE-176 | Apr 18, 2025 |
| <u>OSPEMIFENE - OSPHENA</u> | | | | | | |
| N 203505 001 | 6245819 | Jul 21, 2025 | U-1370 | | | |
| | 8236861 | Aug 11, 2026 | U-1369 | | | |
| | 8236861 | Aug 11, 2026 | U-1370 | | | |
| | 8470890 | Feb 13, 2024 | U-1369 | | | |
| | 8470890 | Feb 13, 2024 | U-1370 | | | |
| | 8642079 | Jul 09, 2028 | DP | | | |
| | 8772353 | Feb 13, 2024 | U-1369 | | | |
| | 8772353 | Feb 13, 2024 | U-1370 | | | |
| | 9241915 | Feb 13, 2024 | U-1369 | | | |
| | 9241915 | Feb 13, 2024 | U-1370 | | | |
| | 9566252 | Jul 21, 2020 | U-1370 | | | |
| | 9855224 | Feb 13, 2024 | U-1369 | | | |
| | 9855224 | Feb 13, 2024 | U-1370 | | | |
| <u>OXCARBAZEPINE - TRILEPTAL</u> | | | | | | |
| N 021285 001 | 8119148 | Dec 19, 2020 | DP U-724 | | | |
| <u>OXCARBAZEPINE - OXTELLAR XR</u> | | | | | | |
| N 202810 001 | 7722898 | Apr 13, 2027 | DP | | | |
| | 7910131 | Apr 13, 2027 | U-1298 | | | |
| | 8617600 | Apr 13, 2027 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>OXCARBAZEPINE - OXTELLAR XR</u> | | | | | | |
| N 202810 001 | 8821930 | Apr 13, 2027 | DP | | | |
| | 9119791 | Apr 13, 2027 | U-1298 | | | |
| | 9351975 | Apr 13, 2027 | DP | | | |
| | 9370525 | Apr 13, 2027 | DP | | | |
| | 9855278 | Apr 13, 2027 | DP | | | |
| <u>OXCARBAZEPINE - OXTELLAR XR</u> | | | | | | |
| N 202810 002 | 7722898 | Apr 13, 2027 | DP | | | |
| | 7910131 | Apr 13, 2027 | U-1298 | | | |
| | 8617600 | Apr 13, 2027 | DP | | | |
| | 8821930 | Apr 13, 2027 | DP | | | |
| | 9119791 | Apr 13, 2027 | U-1298 | | | |
| | 9351975 | Apr 13, 2027 | DP | | | |
| | 9370525 | Apr 13, 2027 | DP | | | |
| | 9855278 | Apr 13, 2027 | DP | | | |
| <u>OXCARBAZEPINE - OXTELLAR XR</u> | | | | | | |
| N 202810 003 | 7722898 | Apr 13, 2027 | DP | | | |
| | 7910131 | Apr 13, 2027 | U-1298 | | | |
| | 8617600 | Apr 13, 2027 | DP | | | |
| | 8821930 | Apr 13, 2027 | DP | | | |
| | 9119791 | Apr 13, 2027 | U-1298 | | | |
| | 9351975 | Apr 13, 2027 | DP | | | |
| | 9370525 | Apr 13, 2027 | DP | | | |
| | 9855278 | Apr 13, 2027 | DP | | | |
| <u>OXYBUTYNYNIN - OXYTROL</u> | | | | | | |
| N 021351 002 | 6743441 | Apr 26, 2020 | DP U-318 | | | |
| | 7081249 | Apr 26, 2020 | DP U-318 | | | |
| | 7081250 | Apr 26, 2020 | DP U-318 | | | |
| | 7081251 | Apr 26, 2020 | DP U-318 | | | |
| | 7081252 | Apr 26, 2020 | DP U-318 | | | |
| | 7179483 | Apr 26, 2020 | DS DP U-318 | | | |
| <u>OXYBUTYNYNIN - OXYTROL FOR WOMEN</u> | | | | | | |
| N 202211 001 | 6743441 | Apr 26, 2020 | DP U-1329 | | | |
| | 7081249 | Apr 26, 2020 | DP U-1329 | | | |
| | 7081250 | Apr 26, 2020 | DP U-1329 | | | |
| | 7081251 | Apr 26, 2020 | DP U-1329 | | | |
| | 7081252 | Apr 26, 2020 | DP U-1329 | | | |
| | 7179483 | Apr 26, 2020 | U-1329 | | | |
| <u>OXYBUTYNYNIN - GELNIQUE 3%</u> | | | | | | |
| N 202513 001 | 7029694 | Apr 26, 2020 | DP U-318 | | | |
| | 7179483 | Apr 26, 2020 | U-318 | | | |
| | 7198801 | Jun 25, 2022 | DP | | | |
| | 8241662 | Apr 26, 2020 | U-318 | | | |
| <u>OXYBUTYNYNIN CHLORIDE - GELNIQUE</u> | | | | | | |
| N 022204 001 | 7029694 | Apr 26, 2020 | DP U-318 | | | |
| | 7179483 | Apr 26, 2020 | U-318 | | | |
| | 8241662 | Apr 26, 2020 | U-318 | | | |
| | 8920392 | Mar 26, 2031 | U-1644 | | | |
| | 9259388 | Nov 06, 2029 | U-1644 | | | |
| <u>OXYCODONE - XTAMPZA ER</u> | | | | | | |
| N 208090 001 | 10004729 | Dec 10, 2030 | DP U-1556 | | NP | Apr 26, 2019 |
| | 7399488 | Mar 24, 2025 | DP | | | |
| | 7771707 | Mar 24, 2025 | DP | | | |
| | 8449909 | Mar 24, 2025 | DP | | | |
| | 8557291 | Mar 21, 2025 | DP | | | |
| | 8758813 | Jun 10, 2025 | U-1556 | | | |
| | 8840928 | Jul 07, 2023 | DP U-1556 | | | |
| | 9044398 | Jul 07, 2023 | DP | | | |
| | 9248195 | Jul 07, 2023 | U-1556 | | | |
| | 9592200 | Jul 07, 2023 | DP | | | |
| | 9682075 | Dec 10, 2030 | DP U-1556 | | | |
| | 9737530 | Sep 02, 2036 | DP U-1556 | | | |
| | 9763883 | Jul 07, 2023 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>OXYCODONE - XTAMPZA ER</u> | | | | | | |
| N 208090 | 001 | 9968598 | Sep 02, 2036 | DP U-1556 | | |
| <u>OXYCODONE - XTAMPZA ER</u> | | | | | | |
| N 208090 | 002 | 10004729 | Dec 10, 2030 | DP U-1556 | NP | Apr 26, 2019 |
| | | 7399488 | Mar 24, 2025 | DP | | |
| | | 7771707 | Mar 24, 2025 | DP | | |
| | | 8449909 | Mar 24, 2025 | DP | | |
| | | 8557291 | Mar 21, 2025 | DP | | |
| | | 8758813 | Jun 10, 2025 | U-1556 | | |
| | | 8840928 | Jul 07, 2023 | DP U-1556 | | |
| | | 9044398 | Jul 07, 2023 | DP | | |
| | | 9248195 | Jul 07, 2023 | U-1556 | | |
| | | 9592200 | Jul 07, 2023 | DP | | |
| | | 9682075 | Dec 10, 2030 | DP U-1556 | | |
| | | 9737530 | Sep 02, 2036 | DP U-1556 | | |
| | | 9763883 | Jul 07, 2023 | DP | | |
| | | 9968598 | Sep 02, 2036 | DP U-1556 | | |
| <u>OXYCODONE - XTAMPZA ER</u> | | | | | | |
| N 208090 | 003 | 10004729 | Dec 10, 2030 | DP U-1556 | NP | Apr 26, 2019 |
| | | 7399488 | Mar 24, 2025 | DP | | |
| | | 7771707 | Mar 24, 2025 | DP | | |
| | | 8449909 | Mar 24, 2025 | DP | | |
| | | 8557291 | Mar 21, 2025 | DP | | |
| | | 8758813 | Jun 10, 2025 | U-1556 | | |
| | | 8840928 | Jul 07, 2023 | DP U-1556 | | |
| | | 9044398 | Jul 07, 2023 | DP | | |
| | | 9248195 | Jul 07, 2023 | U-1556 | | |
| | | 9592200 | Jul 07, 2023 | DP | | |
| | | 9682075 | Dec 10, 2030 | DP U-1556 | | |
| | | 9737530 | Sep 02, 2036 | DP U-1556 | | |
| | | 9763883 | Jul 07, 2023 | DP | | |
| | | 9968598 | Sep 02, 2036 | DP U-1556 | | |
| <u>OXYCODONE - XTAMPZA ER</u> | | | | | | |
| N 208090 | 004 | 10004729 | Dec 10, 2030 | DP U-1556 | NP | Apr 26, 2019 |
| | | 7399488 | Mar 24, 2025 | DP | | |
| | | 7771707 | Mar 24, 2025 | DP | | |
| | | 8449909 | Mar 24, 2025 | DP | | |
| | | 8557291 | Mar 21, 2025 | DP | | |
| | | 8758813 | Jun 10, 2025 | U-1556 | | |
| | | 8840928 | Jul 07, 2023 | DP U-1556 | | |
| | | 9044398 | Jul 07, 2023 | DP | | |
| | | 9248195 | Jul 07, 2023 | U-1556 | | |
| | | 9592200 | Jul 07, 2023 | DP | | |
| | | 9682075 | Dec 10, 2030 | DP U-1556 | | |
| | | 9737530 | Sep 02, 2036 | DP U-1556 | | |
| | | 9763883 | Jul 07, 2023 | DP | | |
| | | 9968598 | Sep 02, 2036 | DP U-1556 | | |
| <u>OXYCODONE - XTAMPZA ER</u> | | | | | | |
| N 208090 | 005 | 10004729 | Dec 10, 2030 | DP U-1556 | NP | Apr 26, 2019 |
| | | 7399488 | Mar 24, 2025 | DP | | |
| | | 7771707 | Mar 24, 2025 | DP | | |
| | | 8449909 | Mar 24, 2025 | DP | | |
| | | 8557291 | Mar 21, 2025 | DP | | |
| | | 8758813 | Jun 10, 2025 | U-1556 | | |
| | | 8840928 | Jul 07, 2023 | DP U-1556 | | |
| | | 9044398 | Jul 07, 2023 | DP | | |
| | | 9248195 | Jul 07, 2023 | U-1556 | | |
| | | 9592200 | Jul 07, 2023 | DP | | |
| | | 9682075 | Dec 10, 2030 | DP U-1556 | | |
| | | 9737530 | Sep 02, 2036 | DP U-1556 | | |
| | | 9763883 | Jul 07, 2023 | DP | | |
| | | 9968598 | Sep 02, 2036 | DP U-1556 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u> | | | | | | |
| N 022272 001 | 10130591 | Nov 20, 2023 | DP U-1819 | | | |
| | 7674799 | Mar 30, 2025 | DP | Y | | |
| | 7674800 | Mar 30, 2025 | DS | Y | | |
| | 7683072 | Mar 30, 2025 | DS | Y | | |
| | 7776314 | Apr 19, 2025 | DP | Y | | |
| | 8309060 | Nov 20, 2023 | DP U-1556 | | | |
| | 8808741 | Aug 24, 2027 | U-1556 | | | |
| | 8894987 | Mar 29, 2030 | DP | | | |
| | 8894988 | Aug 24, 2027 | DP | | | |
| | 9060976 | Aug 06, 2022 | DP | | | |
| | 9073933 | Mar 30, 2025 | DS | | | |
| | 9492389 | Aug 24, 2027 | DP | | | |
| | 9492391 | Aug 24, 2027 | U-1556 | | | |
| | 9492392 | Aug 24, 2027 | DP | | | |
| | 9492393 | Aug 24, 2027 | U-1556 | | | |
| | 9522919 | Mar 30, 2025 | DS DP | | | |
| | 9675610 | Jun 16, 2023 | DP | | | |
| | 9763933 | Aug 24, 2027 | DP | | | |
| | 9770416 | Aug 24, 2027 | DP | | | |
| | 9775808 | Aug 24, 2027 | DP | | | |
| <u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u> | | | | | | |
| N 022272 002 | 10130591 | Nov 20, 2023 | DP U-1819 | | | |
| | 7674799 | Mar 30, 2025 | DP | Y | | |
| | 7674800 | Mar 30, 2025 | DS | Y | | |
| | 7683072 | Mar 30, 2025 | DS | Y | | |
| | 7776314 | Apr 19, 2025 | DP | Y | | |
| | 8309060 | Nov 20, 2023 | DP U-1556 | | | |
| | 8808741 | Aug 24, 2027 | U-1556 | | | |
| | 8894987 | Mar 29, 2030 | DP | | | |
| | 8894988 | Aug 24, 2027 | DP | | | |
| | 9060976 | Aug 06, 2022 | DP | | | |
| | 9073933 | Mar 30, 2025 | DS | | | |
| | 9492389 | Aug 24, 2027 | DP | | | |
| | 9492391 | Aug 24, 2027 | U-1556 | | | |
| | 9492392 | Aug 24, 2027 | DP | | | |
| | 9492393 | Aug 24, 2027 | U-1556 | | | |
| | 9522919 | Mar 30, 2025 | DS DP | | | |
| | 9675610 | Jun 16, 2023 | DP | | | |
| | 9763933 | Aug 24, 2027 | DP | | | |
| | 9770416 | Aug 24, 2027 | DP | | | |
| | 9775808 | Aug 24, 2027 | DP | | | |
| <u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u> | | | | | | |
| N 022272 003 | 10130591 | Nov 20, 2023 | DP U-1819 | | | |
| | 7674799 | Mar 30, 2025 | DP | Y | | |
| | 7674800 | Mar 30, 2025 | DS | Y | | |
| | 7683072 | Mar 30, 2025 | DS | Y | | |
| | 7776314 | Apr 19, 2025 | DP | Y | | |
| | 8309060 | Nov 20, 2023 | DP U-1556 | | | |
| | 8808741 | Aug 24, 2027 | U-1556 | | | |
| | 8894987 | Mar 29, 2030 | DP | | | |
| | 8894988 | Aug 24, 2027 | DP | | | |
| | 9060976 | Aug 06, 2022 | DP | | | |
| | 9073933 | Mar 30, 2025 | DS | | | |
| | 9492389 | Aug 24, 2027 | DP | | | |
| | 9492391 | Aug 24, 2027 | U-1556 | | | |
| | 9492392 | Aug 24, 2027 | DP | | | |
| | 9492393 | Aug 24, 2027 | U-1556 | | | |
| | 9522919 | Mar 30, 2025 | DS DP | | | |
| | 9675610 | Jun 16, 2023 | DP | | | |
| | 9763933 | Aug 24, 2027 | DP | | | |
| | 9770416 | Aug 24, 2027 | DP | | | |
| | 9775808 | Aug 24, 2027 | DP | | | |
| <u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u> | | | | | | |
| N 022272 004 | 10130591 | Nov 20, 2023 | DP U-1819 | | | |
| | 8309060 | Nov 20, 2023 | DP U-1556 | | | |
| | 8808741 | Aug 24, 2027 | U-1556 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-----------|------------------------|--------------|-------------------------|---------------------|-----------------------------|
| <u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u> | | | | | | |
| N 022272 004 | 8894987 | Mar 29, 2030 | DP | | | |
| | 8894988 | Aug 24, 2027 | DP | | | |
| | 9060976 | Aug 06, 2022 | DP | | | |
| | 9073933 | Mar 30, 2025 | DS | | | |
| | 9492389 | Aug 24, 2027 | DP | | | |
| | 9492391 | Aug 24, 2027 | | U-1556 | | |
| | 9492392 | Aug 24, 2027 | DP | | | |
| | 9492393 | Aug 24, 2027 | | U-1556 | | |
| | 9522919 | Mar 30, 2025 | DS DP | | | |
| | 9675610 | Jun 16, 2023 | DP | | | |
| | 9763933 | Aug 24, 2027 | DP | | | |
| | 9770416 | Aug 24, 2027 | DP | | | |
| | 9775808 | Aug 24, 2027 | DP | | | |
| <u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u> | | | | | | |
| N 022272 005 | 10130591 | Nov 20, 2023 | DP | U-1819 | | |
| | 8309060 | Nov 20, 2023 | DP | U-1556 | | |
| | 8808741 | Aug 24, 2027 | | U-1556 | | |
| | 8894988 | Aug 24, 2027 | DP | | | |
| | 9060976 | Aug 06, 2022 | DP | | | |
| | 9073933 | Mar 30, 2025 | DS | | | |
| | 9492389 | Aug 24, 2027 | DP | | | |
| | 9492391 | Aug 24, 2027 | | U-1556 | | |
| | 9492392 | Aug 24, 2027 | DP | | | |
| | 9492393 | Aug 24, 2027 | | U-1556 | | |
| | 9522919 | Mar 30, 2025 | DS DP | | | |
| | 9675610 | Jun 16, 2023 | DP | | | |
| | 9763933 | Aug 24, 2027 | DP | | | |
| | 9770416 | Aug 24, 2027 | DP | | | |
| | 9775808 | Aug 24, 2027 | DP | | | |
| <u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u> | | | | | | |
| N 022272 006 | 10130591 | Nov 20, 2023 | DP | U-1819 | | |
| | 8309060 | Nov 20, 2023 | DP | U-1556 | | |
| | 8808741 | Aug 24, 2027 | | U-1556 | | |
| | 8894988 | Aug 24, 2027 | DP | | | |
| | 9060976 | Aug 06, 2022 | DP | | | |
| | 9073933 | Mar 30, 2025 | DS | | | |
| | 9492389 | Aug 24, 2027 | DP | | | |
| | 9492391 | Aug 24, 2027 | | U-1556 | | |
| | 9492392 | Aug 24, 2027 | DP | | | |
| | 9492393 | Aug 24, 2027 | | U-1556 | | |
| | 9522919 | Mar 30, 2025 | DS DP | | | |
| | 9675610 | Jun 16, 2023 | DP | | | |
| | 9763933 | Aug 24, 2027 | DP | | | |
| | 9770416 | Aug 24, 2027 | DP | | | |
| | 9775808 | Aug 24, 2027 | DP | | | |
| <u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u> | | | | | | |
| N 022272 007 | 10130591 | Nov 20, 2023 | DP | U-1819 | | |
| | 8309060 | Nov 20, 2023 | DP | U-1556 | | |
| | 8808741 | Aug 24, 2027 | | U-1556 | | |
| | 8894988 | Aug 24, 2027 | DP | | | |
| | 9060976 | Aug 06, 2022 | DP | | | |
| | 9073933 | Mar 30, 2025 | DS | | | |
| | 9492389 | Aug 24, 2027 | DP | | | |
| | 9492391 | Aug 24, 2027 | | U-1556 | | |
| | 9492392 | Aug 24, 2027 | DP | | | |
| | 9492393 | Aug 24, 2027 | | U-1556 | | |
| | 9522919 | Mar 30, 2025 | DS DP | | | |
| | 9675610 | Jun 16, 2023 | DP | | | |
| | 9763933 | Aug 24, 2027 | DP | | | |
| | 9770416 | Aug 24, 2027 | DP | | | |
| | 9775808 | Aug 24, 2027 | DP | | | |
| <u>OXYCODONE HYDROCHLORIDE - OXAYDO</u> | | | | | | |
| N 202080 001 | 7201920 | Mar 16, 2025 | DP | | | |
| | 7510726 | Nov 26, 2023 | DP | | | |
| | 7981439 | Nov 26, 2023 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>OXYCODONE HYDROCHLORIDE - OXAYDO</u> | | | | | | |
| N 202080 001 | 8409616 | Nov 26, 2023 | DP | | | |
| | 8637540 | Nov 26, 2023 | DP | | | |
| | 9492443 | May 26, 2024 | DP | | | |
| <u>OXYCODONE HYDROCHLORIDE - OXAYDO</u> | | | | | | |
| N 202080 002 | 7201920 | Mar 16, 2025 | DP | | | |
| | 7510726 | Nov 26, 2023 | DP | | | |
| | 7981439 | Nov 26, 2023 | DP | | | |
| | 8409616 | Nov 26, 2023 | DP | | | |
| | 8637540 | Nov 26, 2023 | DP | | | |
| | 9492443 | May 26, 2024 | DP | | | |
| <u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u> | | | | | | |
| N 209777 001 | 7955619 | Aug 12, 2028 | DP | | NP | Apr 20, 2020 |
| <u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u> | | | | | | |
| N 209777 002 | 7955619 | Aug 12, 2028 | DP | | NP | Apr 20, 2020 |
| <u>OXYCODONE HYDROCHLORIDE - ROXYBOND</u> | | | | | | |
| N 209777 003 | 7955619 | Aug 12, 2028 | DP | | NP | Apr 20, 2020 |
| <u>OXYMETAZOLINE HYDROCHLORIDE - RHOFADÉ</u> | | | | | | |
| N 208552 001 | 7812049 | May 02, 2028 | | U-1959 | NP | Jan 18, 2020 |
| | 8420688 | Aug 02, 2024 | | U-1959 | | |
| | 8815929 | Jan 22, 2024 | | U-1959 | | |
| | 8883838 | Dec 01, 2031 | DP | | | |
| | 9974773 | Jun 11, 2035 | | U-2306 | | |
| <u>OXYMETAZOLINE HYDROCHLORIDE; TETRACAINE HYDROCHLORIDE - KOVANAZE</u> | | | | | | |
| N 208032 001 | 6413499 | Mar 20, 2020 | | U-1876 | NC | Jun 29, 2019 |
| | 8580282 | Apr 02, 2030 | DP | U-1876 | | |
| | 9308191 | Apr 02, 2030 | DP | U-1876 | | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 021610 001 | 7276250 | Feb 04, 2023 | DP | U-826 | | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 021610 002 | 7276250 | Feb 04, 2023 | DP | U-826 | | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 021610 003 | 7276250 | Feb 04, 2023 | DP | U-826 | | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 021610 004 | 7276250 | Feb 04, 2023 | DP | U-826 | | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 021610 005 | 7276250 | Feb 04, 2023 | DP | U-826 | | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 021610 006 | 7276250 | Feb 04, 2023 | DP | U-826 | | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 021610 007 | 7276250 | Feb 04, 2023 | DP | U-826 | | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 201655 001 | 7851482 | Jul 10, 2029 | DS | | | |
| | 8075872 | Nov 20, 2023 | DP | | | |
| | 8114383 | Aug 08, 2024 | DP | | | |
| | 8192722 | Sep 15, 2025 | DP | | | |
| | 8309060 | Nov 20, 2023 | DP | | | |
| | 8309122 | Feb 04, 2023 | DP | | | |
| | 8329216 | Feb 04, 2023 | DP | | | |
| | 8808737 | Jun 21, 2027 | | U-1598 | | |
| | 8871779 | Nov 22, 2029 | DS | | | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 201655 002 | 7851482 | Jul 10, 2029 | DS | | | |
| | 8075872 | Nov 20, 2023 | DP | | | |
| | 8114383 | Aug 08, 2024 | DP | | | |
| | 8192722 | Sep 15, 2025 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 201655 002 | 8309060 | Nov 20, 2023 | DP | | | |
| | 8309122 | Feb 04, 2023 | DP | | | |
| | 8329216 | Feb 04, 2023 | DP | | | |
| | 8808737 | Jun 21, 2027 | | U-1598 | | |
| | 8871779 | Nov 22, 2029 | DS | | | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 201655 003 | 7851482 | Jul 10, 2029 | DS | | | |
| | 8075872 | Nov 20, 2023 | DP | | | |
| | 8114383 | Aug 08, 2024 | DP | | | |
| | 8192722 | Sep 15, 2025 | DP | | | |
| | 8309060 | Nov 20, 2023 | DP | | | |
| | 8309122 | Feb 04, 2023 | DP | | | |
| | 8329216 | Feb 04, 2023 | DP | | | |
| | 8808737 | Jun 21, 2027 | | U-1598 | | |
| | 8871779 | Nov 22, 2029 | DS | | | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 201655 004 | 7851482 | Jul 10, 2029 | DS | | | |
| | 8075872 | Nov 20, 2023 | DP | | | |
| | 8114383 | Aug 08, 2024 | DP | | | |
| | 8192722 | Sep 15, 2025 | DP | | | |
| | 8309060 | Nov 20, 2023 | DP | | | |
| | 8309122 | Feb 04, 2023 | DP | | | |
| | 8329216 | Feb 04, 2023 | DP | | | |
| | 8808737 | Jun 21, 2027 | | U-1598 | | |
| | 8871779 | Nov 22, 2029 | DS | | | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 201655 005 | 7851482 | Jul 10, 2029 | DS | | | |
| | 8075872 | Nov 20, 2023 | DP | | | |
| | 8114383 | Aug 08, 2024 | DP | | | |
| | 8192722 | Sep 15, 2025 | DP | | | |
| | 8309060 | Nov 20, 2023 | DP | | | |
| | 8309122 | Feb 04, 2023 | DP | | | |
| | 8329216 | Feb 04, 2023 | DP | | | |
| | 8808737 | Jun 21, 2027 | | U-1598 | | |
| | 8871779 | Nov 22, 2029 | DS | | | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 201655 006 | 7851482 | Jul 10, 2029 | DS | | | |
| | 8075872 | Nov 20, 2023 | DP | | | |
| | 8114383 | Aug 08, 2024 | DP | | | |
| | 8192722 | Sep 15, 2025 | DP | | | |
| | 8309060 | Nov 20, 2023 | DP | | | |
| | 8309122 | Feb 04, 2023 | DP | | | |
| | 8329216 | Feb 04, 2023 | DP | | | |
| | 8808737 | Jun 21, 2027 | | U-1598 | | |
| | 8871779 | Nov 22, 2029 | DS | | | |
| <u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u> | | | | | | |
| N 201655 007 | 7851482 | Jul 10, 2029 | DS | | | |
| | 8075872 | Nov 20, 2023 | DP | | | |
| | 8114383 | Aug 08, 2024 | DP | | | |
| | 8192722 | Sep 15, 2025 | DP | | | |
| | 8309060 | Nov 20, 2023 | DP | | | |
| | 8309122 | Feb 04, 2023 | DP | | | |
| | 8329216 | Feb 04, 2023 | DP | | | |
| | 8808737 | Jun 21, 2027 | | U-1598 | | |
| | 8871779 | Nov 22, 2029 | DS | | | |
| <u>OZENOXACIN - XEPI</u> | | | | | | |
| N 208945 001 | 6335447 | Apr 06, 2019 | DS | | NCE | Dec 11, 2022 |
| | 9180200 | Jan 29, 2032 | DP | U-805 | | |
| | 9399014 | Dec 15, 2029 | | U-805 | | |
| <u>PACLITAXEL - ABRAXANE</u> | | | | | | |
| N 021660 001 | 7758891 | Feb 21, 2026 | | U-1434 | ODE-52 | Sep 06, 2020 |
| | 7820788 | Oct 27, 2024 | DP | U-1092 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PACLITAXEL - ABRAXANE</u> | | | | | | |
| N 021660 001 | 7820788 | Oct 27, 2024 | DP U-1290 | | | |
| | 7820788 | Oct 27, 2024 | DP U-1434 | | | |
| | 7923536 | Dec 09, 2023 | U-1117 | | | |
| | 7923536 | Dec 09, 2023 | U-1290 | | | |
| | 7923536 | Dec 09, 2023 | U-1434 | | | |
| | 8034375 | Aug 13, 2026 | U-1290 | | | |
| | 8138229 | Dec 09, 2023 | DP U-1092 | | | |
| | 8138229 | Dec 09, 2023 | DP U-1290 | | | |
| | 8138229 | Dec 09, 2023 | DP U-1434 | | | |
| | 8268348 | Feb 21, 2026 | U-1290 | | | |
| | 8314156 | Dec 09, 2023 | U-1290 | | | |
| | 8314156 | Dec 09, 2023 | U-1434 | | | |
| | 8853260 | Oct 10, 2020 | DP U-1092 | | | |
| | 8853260 | Oct 10, 2020 | DP U-1290 | | | |
| | 8853260 | Oct 10, 2020 | DP U-1434 | | | |
| | 9101543 | Feb 21, 2026 | U-1434 | | | |
| | 9393318 | Mar 04, 2032 | U-1290 | | | |
| | 9511046 | Jan 12, 2034 | U-1434 | | | |
| | 9597409 | Mar 04, 2032 | U-1290 | | | |
| <u>PALBOCICLIB - IBRANCE</u> | | | | | | |
| N 207103 001 | 6936612 | Jan 22, 2023 | DS DP | | I-725 | Feb 19, 2019 |
| | 7208489 | Jan 16, 2023 | DS DP | | NCE | Feb 03, 2020 |
| | 7456168 | Jan 16, 2023 | U-1998 | | | |
| <u>PALBOCICLIB - IBRANCE</u> | | | | | | |
| N 207103 002 | 6936612 | Jan 22, 2023 | DS DP | | I-725 | Feb 19, 2019 |
| | 7208489 | Jan 16, 2023 | DS DP | | NCE | Feb 03, 2020 |
| | 7456168 | Jan 16, 2023 | U-1998 | | | |
| <u>PALBOCICLIB - IBRANCE</u> | | | | | | |
| N 207103 003 | 6936612 | Jan 22, 2023 | DS DP | | I-725 | Feb 19, 2019 |
| | 7208489 | Jan 16, 2023 | DS DP | | NCE | Feb 03, 2020 |
| | 7456168 | Jan 16, 2023 | U-1998 | | | |
| <u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u> | | | | | | |
| N 022264 001 | 9439906 | Jan 26, 2031 | U-1901 | | M-215 | Dec 20, 2020 |
| | 9439906 | Jan 26, 2031 | U-543 | | | |
| <u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u> | | | | | | |
| N 022264 002 | 9439906 | Jan 26, 2031 | U-1901 | | M-215 | Dec 20, 2020 |
| | 9439906 | Jan 26, 2031 | U-543 | | | |
| <u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u> | | | | | | |
| N 022264 003 | 9439906 | Jan 26, 2031 | U-1901 | | M-215 | Dec 20, 2020 |
| | 9439906 | Jan 26, 2031 | U-543 | | | |
| <u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u> | | | | | | |
| N 022264 004 | 9439906 | Jan 26, 2031 | U-1901 | | M-215 | Dec 20, 2020 |
| | 9439906 | Jan 26, 2031 | U-543 | | | |
| <u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u> | | | | | | |
| N 022264 005 | 9439906 | Jan 26, 2031 | U-1901 | | M-215 | Dec 20, 2020 |
| | 9439906 | Jan 26, 2031 | U-543 | | | |
| <u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u> | | | | | | |
| N 207946 001 | 10143693 | Apr 05, 2036 | U-2457 | | | |
| | 10143693 | Apr 05, 2036 | U-2458 | | | |
| <u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u> | | | | | | |
| N 207946 002 | 10143693 | Apr 05, 2036 | U-2457 | | | |
| | 10143693 | Apr 05, 2036 | U-2458 | | | |
| <u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u> | | | | | | |
| N 207946 003 | 10143693 | Apr 05, 2036 | U-2457 | | | |
| | 10143693 | Apr 05, 2036 | U-2458 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u> | | | | | | |
| N 207946 004 | 10143693 | Apr 05, 2036 | | U-2457 | | |
| | 10143693 | Apr 05, 2036 | | U-2458 | | |
| <u>PALONOSETRON HYDROCHLORIDE - ALOXI</u> | | | | | | |
| N 021372 001 | 7947724 | Jan 30, 2024 | | DP | | |
| | 7947724*PED | Jul 30, 2024 | | | | |
| | 7947725 | Jan 30, 2024 | | DP | | |
| | 7947725*PED | Jul 30, 2024 | | | | |
| | 7960424 | Jan 30, 2024 | | DP | | |
| | 7960424*PED | Jul 30, 2024 | | | | |
| | 8518981 | Jan 30, 2024 | | DP | | |
| | 8518981*PED | Jul 30, 2024 | | | | |
| | 8598218 | Jan 30, 2024 | | DP | | |
| | 8598218*PED | Jul 30, 2024 | | | | |
| | 8598219 | Jan 30, 2024 | | DP | | |
| | 8598219*PED | Jul 30, 2024 | | | | |
| | 8729094 | Jan 30, 2024 | | DP U-528 | | |
| | 8729094*PED | Jul 30, 2024 | | | | |
| | 9066980 | Jan 30, 2024 | | DP U-528 | | |
| | 9066980*PED | Jul 30, 2024 | | | | |
| | 9125905 | Jan 30, 2024 | | DP | | |
| | 9125905*PED | Jul 30, 2024 | | | | |
| | 9173942 | Jan 30, 2024 | | DP | | |
| | 9173942*PED | Jul 30, 2024 | | | | |
| | 9439854 | Jan 30, 2024 | | DP | | |
| | 9439854*PED | Jul 30, 2024 | | | | |
| | 9457020 | Jan 30, 2024 | | DP | | |
| | 9457020*PED | Jul 30, 2024 | | | | |
| | 9457021 | Jan 30, 2024 | | DP | | |
| | 9457021*PED | Jul 30, 2024 | | | | |
| <u>PALONOSETRON HYDROCHLORIDE - ALOXI</u> | | | | | | |
| N 021372 002 | 7947724 | Jan 30, 2024 | | DP | | |
| | 7947724*PED | Jul 30, 2024 | | | | |
| | 7947725 | Jan 30, 2024 | | DP | | |
| | 7947725*PED | Jul 30, 2024 | | | | |
| | 7960424 | Jan 30, 2024 | | DP | | |
| | 7960424*PED | Jul 30, 2024 | | | | |
| | 8518981 | Jan 30, 2024 | | DP | | |
| | 8518981*PED | Jul 30, 2024 | | | | |
| | 8598218 | Jan 30, 2024 | | DP | | |
| | 8598218*PED | Jul 30, 2024 | | | | |
| | 9173942 | Jan 30, 2024 | | DP | | |
| | 9173942*PED | Jul 30, 2024 | | | | |
| | 9439854 | Jan 30, 2024 | | DP | | |
| | 9439854*PED | Jul 30, 2024 | | | | |
| | 9457020 | Jan 30, 2024 | | DP | | |
| | 9457020*PED | Jul 30, 2024 | | | | |
| <u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u> | | | | | | |
| N 022210 001 | 7658918 | Feb 20, 2028 | | DP | | |
| | 8221747 | Feb 20, 2028 | | DP | | |
| | 8246950 | Feb 20, 2028 | | U-1274 | | |
| | 8562978 | Feb 20, 2028 | | DP | | |
| | 8562979 | Feb 20, 2028 | | DP U-1274 | | |
| | 8562980 | Feb 20, 2028 | | DP U-1274 | | |
| | 8562981 | Feb 20, 2028 | | U-1274 | | |
| <u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u> | | | | | | |
| N 022210 002 | 7658918 | Feb 20, 2028 | | DP | | |
| | 8221747 | Feb 20, 2028 | | DP | | |
| | 8246950 | Feb 20, 2028 | | U-1274 | | |
| | 8562978 | Feb 20, 2028 | | DP | | |
| | 8562979 | Feb 20, 2028 | | DP U-1274 | | |
| | 8562980 | Feb 20, 2028 | | DP U-1274 | | |
| | 8562981 | Feb 20, 2028 | | U-1274 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------------------------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u> | | | | | | |
| N 022210 003 | 7658918 | Feb 20, 2028 | DP | | | |
| | 8221747 | Feb 20, 2028 | DP | | | |
| | 8246950 | Feb 20, 2028 | | U-1274 | | |
| | 8562978 | Feb 20, 2028 | DP | | | |
| | 8562979 | Feb 20, 2028 | DP | U-1274 | | |
| | 8562980 | Feb 20, 2028 | DP | U-1274 | | |
| | 8562981 | Feb 20, 2028 | | | | |
| <u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u> | | | | | | |
| N 022210 004 | 7658918 | Feb 20, 2028 | DP | | | |
| | 8221747 | Feb 20, 2028 | DP | | | |
| | 8246950 | Feb 20, 2028 | | U-1274 | | |
| | 8562978 | Feb 20, 2028 | DP | | | |
| | 8562979 | Feb 20, 2028 | DP | U-1274 | | |
| | 8562980 | Feb 20, 2028 | DP | U-1274 | | |
| | 8562981 | Feb 20, 2028 | | U-1274 | | |
| <u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u> | | | | | | |
| N 022210 005 | 8221747 | Feb 20, 2028 | DP | | | |
| | 8562978 | Feb 20, 2028 | DP | | | |
| | 8562979 | Feb 20, 2028 | DP | U-1274 | | |
| | 8562980 | Feb 20, 2028 | DP | U-1274 | | |
| | 8562981 | Feb 20, 2028 | | U-1274 | | |
| <u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u> | | | | | | |
| N 022210 006 | 8221747 | Feb 20, 2028 | DP | | | |
| | 8562978 | Feb 20, 2028 | DP | | | |
| | 8562979 | Feb 20, 2028 | DP | U-1274 | | |
| | 8562980 | Feb 20, 2028 | DP | U-1274 | | |
| | 8562981 | Feb 20, 2028 | | U-1274 | | |
| <u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u> | | | | | | |
| N 022210 007 | 8221747 | Feb 20, 2028 | DP | | | |
| | 8562978 | Feb 20, 2028 | DP | | | |
| | 8562979 | Feb 20, 2028 | DP | U-1274 | | |
| | 8562980 | Feb 20, 2028 | DP | U-1274 | | |
| | 8562981 | Feb 20, 2028 | | U-1274 | | |
| <u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - PANCREAZE</u> | | | | | | |
| N 022523 005 | 8221747 | Feb 20, 2028 | DP | | | |
| | 8562978 | Feb 20, 2028 | DP | | | |
| | 8562979 | Feb 20, 2028 | DP | U-1274 | | |
| | 8562980 | Feb 20, 2028 | DP | U-1274 | | |
| | 8562981 | Feb 20, 2028 | DP | U-1274 | | |
| <u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u> | | | | | | |
| N 020725 001 | 9198871 | Feb 07, 2030 | DP | U-1787 | | |
| <u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u> | | | | | | |
| N 020725 002 | 9198871 | Feb 07, 2030 | DP | U-1787 | | |
| <u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u> | | | | | | |
| N 020725 003 | 9198871 | Feb 07, 2030 | DP | U-1787 | | |
| <u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u> | | | | | | |
| N 020725 004 | 9198871 | Feb 07, 2030 | DP | U-1787 | M-93 | Jul 29, 2019 |
| <u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u> | | | | | | |
| N 020725 005 | 9198871 | Feb 07, 2030 | DP | U-1787 | M-93 | Jul 29, 2019 |
| <u>PANOBINOSTAT LACTATE - FARYDAK</u> | | | | | | |
| N 205353 001 | 6552065 | Aug 31, 2021 | DS DP | | NCE | Feb 23, 2020 |
| | 6833384 | Sep 30, 2021 | DS DP | U-1669 | ODE-89 | Feb 23, 2022 |
| | 7067551 | Aug 31, 2021 | | U-1669 | | |
| | 7989494 | Jan 17, 2028 | DS DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PANOBINOSTAT LACTATE - FARYDAK</u> | | | | | | |
| N 205353 | 001 8883842 | Jun 13, 2028 | U-1669 | | | |
| <u>PANOBINOSTAT LACTATE - FARYDAK</u> | | | | | | |
| N 205353 | 002 6552065 | Aug 31, 2021 | DS DP | | NCE | Feb 23, 2020 |
| | 6833384 | Sep 30, 2021 | DS DP | U-1669 | ODE-89 | Feb 23, 2022 |
| | 7067551 | Aug 31, 2021 | | U-1669 | | |
| | 7989494 | Jan 17, 2028 | DS DP | | | |
| | 8883842 | Jun 13, 2028 | U-1669 | | | |
| <u>PANOBINOSTAT LACTATE - FARYDAK</u> | | | | | | |
| N 205353 | 003 6552065 | Aug 31, 2021 | DS DP | | NCE | Feb 23, 2020 |
| | 6833384 | Sep 30, 2021 | DS DP | U-1669 | ODE-89 | Feb 23, 2022 |
| | 7067551 | Aug 31, 2021 | | U-1669 | | |
| | 7989494 | Jan 17, 2028 | DS DP | | | |
| | 8883842 | Jun 13, 2028 | U-1669 | | | |
| <u>PANTOPRAZOLE SODIUM - PROTONIX IV</u> | | | | | | |
| N 020988 | 001 6780881 | Nov 17, 2021 | DP | | | |
| | 7351723 | Nov 17, 2021 | DP | | | |
| | 8754108 | Nov 17, 2021 | DP | | | |
| | 8754108*PED | May 17, 2022 | | | | |
| <u>PANTOPRAZOLE SODIUM - PROTONIX</u> | | | | | | |
| N 022020 | 001 7544370 | Jun 07, 2026 | DP | | | |
| | 7550153 | Sep 30, 2024 | | U-859 | | |
| | 7553498 | Sep 30, 2024 | | U-859 | | |
| | 7838027 | Sep 30, 2024 | DP | U-859 | | |
| <u>PARICALCITOL - ZEMPLAR</u> | | | | | | |
| N 021606 | 001 | | | | NPP | Oct 18, 2019 |
| | | | | | NPP | Oct 18, 2019 |
| | | | | | ODE-125 | Oct 18, 2023 |
| <u>PARICALCITOL - ZEMPLAR</u> | | | | | | |
| N 021606 | 002 | | | | NPP | Oct 18, 2019 |
| | | | | | NPP | Oct 18, 2019 |
| | | | | | ODE-125 | Oct 18, 2023 |
| <u>PARICALCITOL - ZEMPLAR</u> | | | | | | |
| N 021606 | 003 | | | | NPP | Oct 18, 2019 |
| | | | | | NPP | Oct 18, 2019 |
| | | | | | ODE-125 | Oct 18, 2023 |
| <u>PAROXETINE HYDROCHLORIDE - PAXIL</u> | | | | | | |
| N 020885 | 001 6063927 | Apr 23, 2019 | | | | |
| <u>PAROXETINE HYDROCHLORIDE - PAXIL</u> | | | | | | |
| N 020885 | 002 6063927 | Apr 23, 2019 | | | | |
| <u>PAROXETINE HYDROCHLORIDE - PAXIL</u> | | | | | | |
| N 020885 | 003 6063927 | Apr 23, 2019 | | | | |
| <u>PAROXETINE HYDROCHLORIDE - PAXIL</u> | | | | | | |
| N 020885 | 004 6063927 | Apr 23, 2019 | | | | |
| <u>PAROXETINE MESYLATE - PEXEVA</u> | | | | | | |
| N 021299 | 001 7598271 | May 04, 2025 | DS | | | |
| <u>PAROXETINE MESYLATE - PEXEVA</u> | | | | | | |
| N 021299 | 002 7598271 | May 04, 2025 | DS | | | |
| <u>PAROXETINE MESYLATE - PEXEVA</u> | | | | | | |
| N 021299 | 003 7598271 | May 04, 2025 | DS | | | |
| <u>PAROXETINE MESYLATE - PEXEVA</u> | | | | | | |
| N 021299 | 004 7598271 | May 04, 2025 | DS | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PAROXETINE MESYLATE - BRISDELLE</u> | | | | | | |
| N 204516 001 | 7598271 | May 04, 2025 | DS | | | |
| | 8658663 | Apr 06, 2029 | DS DP | U-904 | | |
| | 8946251 | Aug 04, 2026 | DS DP | U-904 | | |
| | 9393237 | Aug 04, 2026 | | U-904 | | |
| <u>PASIREOTIDE DIASPARTATE - SIGNIFOR</u> | | | | | | |
| N 200677 001 | 7473761 | Dec 14, 2026 | DS DP | | ODE-34 | Dec 14, 2019 |
| | 8299209 | Dec 27, 2025 | DS DP | | | |
| <u>PASIREOTIDE DIASPARTATE - SIGNIFOR</u> | | | | | | |
| N 200677 002 | 7473761 | Dec 14, 2026 | DS DP | | ODE-34 | Dec 14, 2019 |
| | 8299209 | Dec 27, 2025 | DS DP | | | |
| <u>PASIREOTIDE DIASPARTATE - SIGNIFOR</u> | | | | | | |
| N 200677 003 | 7473761 | Dec 14, 2026 | DS DP | | ODE-34 | Dec 14, 2019 |
| | 8299209 | Dec 27, 2025 | DS DP | | | |
| <u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u> | | | | | | |
| N 203255 001 | 7473761 | Dec 14, 2026 | DS DP | | I-785 | Jun 29, 2021 |
| | 7759308 | Oct 25, 2026 | DP | | ODE-81 | Dec 15, 2021 |
| | 8822637 | Aug 06, 2023 | | U-1629 | | |
| | 9351923 | May 23, 2028 | DP | | | |
| <u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u> | | | | | | |
| N 203255 002 | 7473761 | Dec 14, 2026 | DS DP | | I-785 | Jun 29, 2021 |
| | 7759308 | Oct 25, 2026 | DP | | ODE-81 | Dec 15, 2021 |
| | 8822637 | Aug 06, 2023 | | U-1629 | | |
| | 9351923 | May 23, 2028 | DP | | | |
| <u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u> | | | | | | |
| N 203255 003 | 7473761 | Dec 14, 2026 | DS DP | | I-785 | Jun 29, 2021 |
| | 7759308 | Oct 25, 2026 | DP | | ODE-81 | Dec 15, 2021 |
| | 8822637 | Aug 06, 2023 | | U-1629 | | |
| | 9351923 | May 23, 2028 | DP | | | |
| <u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u> | | | | | | |
| N 203255 004 | | | | | I-785 | Jun 29, 2021 |
| <u>PASIREOTIDE PAMOATE - SIGNIFOR LAR KIT</u> | | | | | | |
| N 203255 005 | | | | | I-785 | Jun 29, 2021 |
| <u>PATIOROMER SORBITEX CALCIUM - VELTASSA</u> | | | | | | |
| N 205739 001 | 7556799 | Feb 27, 2025 | | U-1766 | NCE | Oct 21, 2020 |
| | 8147873 | Mar 11, 2026 | DP | | | |
| | 8216560 | Mar 14, 2027 | | U-1766 | | |
| | 8282913 | Mar 11, 2026 | DP | | | |
| | 8287847 | Mar 30, 2024 | | U-1766 | | |
| | 8337824 | May 29, 2030 | DS | U-1766 | | |
| | 8475780 | Mar 30, 2024 | | U-1766 | | |
| | 8778324 | Mar 30, 2024 | | U-1766 | | |
| | 8889115 | Mar 30, 2024 | | U-1766 | | |
| | 9492476 | Oct 08, 2033 | | U-1766 | | |
| | 9925212 | Oct 08, 2033 | | U-1766 | | |
| <u>PATIOROMER SORBITEX CALCIUM - VELTASSA</u> | | | | | | |
| N 205739 002 | 7556799 | Feb 27, 2025 | | U-1766 | NCE | Oct 21, 2020 |
| | 8147873 | Mar 11, 2026 | DP | | | |
| | 8216560 | Mar 14, 2027 | | U-1766 | | |
| | 8282913 | Mar 11, 2026 | DP | | | |
| | 8287847 | Mar 30, 2024 | | U-1766 | | |
| | 8337824 | May 29, 2030 | DS | U-1766 | | |
| | 8475780 | Mar 30, 2024 | | U-1766 | | |
| | 8778324 | Mar 30, 2024 | | U-1766 | | |
| | 8889115 | Mar 30, 2024 | | U-1766 | | |
| | 9492476 | Oct 08, 2033 | | U-1766 | | |
| | 9925212 | Oct 08, 2033 | | U-1766 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PATIROMER SORBITE X CALCIUM - VELTASSA</u> | | | | | | |
| N 205739 | 003 | 7556799 | Feb 27, 2025 | U-1766 | NCE | Oct 21, 2020 |
| | | 8147873 | Mar 11, 2026 | DP | | |
| | | 8216560 | Mar 14, 2027 | U-1766 | | |
| | | 8282913 | Mar 11, 2026 | DP | | |
| | | 8287847 | Mar 30, 2024 | U-1766 | | |
| | | 8337824 | May 29, 2030 | DS U-1766 | | |
| | | 8475780 | Mar 30, 2024 | U-1766 | | |
| | | 8778324 | Mar 30, 2024 | U-1766 | | |
| | | 8889115 | Mar 30, 2024 | U-1766 | | |
| | | 9492476 | Oct 08, 2033 | U-1766 | | |
| | | 9925212 | Oct 08, 2033 | U-1766 | | |
| <u>PATISIRAN SODIUM - ONPATTRO</u> | | | | | | |
| N 210922 | 001 | 8058069 | Apr 15, 2029 | DP | NCE | Aug 10, 2023 |
| | | 8158601 | Nov 10, 2030 | DP U-2378 | ODE-197 | Aug 10, 2025 |
| | | 8168775 | Oct 20, 2029 | DS DP U-2378 | | |
| | | 8334373 | May 27, 2025 | DS DP | | |
| | | 8362231 | Mar 30, 2021 | DS DP | | |
| | | 8372968 | Mar 30, 2021 | DS DP | | |
| | | 8492359 | Apr 15, 2029 | DP | | |
| | | 8552171 | Mar 30, 2021 | DS DP | | |
| | | 8642076 | Oct 03, 2027 | DP | | |
| | | 8741866 | Oct 20, 2029 | U-2378 | | |
| | | 8778902 | Mar 30, 2021 | U-2378 | | |
| | | 8802644 | Oct 21, 2030 | DP U-2378 | | |
| | | 8822668 | Apr 15, 2029 | DP U-2378 | | |
| | | 8895718 | Mar 30, 2021 | DS DP | | |
| | | 8895721 | Mar 30, 2021 | DS DP | | |
| | | 9193753 | Mar 30, 2021 | U-2378 | | |
| | | 9234196 | Oct 20, 2029 | DP U-2378 | | |
| | | 9364435 | Apr 15, 2029 | DP U-2378 | | |
| | | 9567582 | Mar 30, 2021 | DS DP | | |
| | | 9943538 | Nov 04, 2023 | DP | | |
| | | 9943539 | Nov 04, 2023 | DP | | |
| <u>PAZOPANIB HYDROCHLORIDE - VOTRIENT</u> | | | | | | |
| N 022465 | 001 | 7105530 | Oct 19, 2023 | DS DP | ODE-23 | Apr 26, 2019 |
| | | 7262203 | Dec 19, 2021 | DS DP | | |
| | | 8114885 | Dec 19, 2021 | DS DP | | |
| <u>PAZOPANIB HYDROCHLORIDE - VOTRIENT</u> | | | | | | |
| N 022465 | 002 | 7105530 | Oct 19, 2023 | DS DP | ODE-23 | Apr 26, 2019 |
| | | 7262203 | Dec 19, 2021 | DS DP | | |
| | | 8114885 | Dec 19, 2021 | DS DP | | |
| <u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u> | | | | | | |
| N 202799 | 001 | 7084245 | May 12, 2024 | DS DP U-1238 | | |
| | | 7414105 | May 12, 2024 | DS DP U-1238 | | |
| | | 7528104 | May 12, 2024 | DS DP | | |
| | | 7550433 | Jun 02, 2026 | U-1238 | | |
| | | 7919118 | May 12, 2024 | DS DP | | |
| | | 7919461 | Jun 02, 2026 | U-1238 | | |
| <u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u> | | | | | | |
| N 202799 | 002 | 7084245 | May 12, 2024 | DS DP U-1238 | | |
| | | 7414105 | May 12, 2024 | DS DP U-1238 | | |
| | | 7528104 | May 12, 2024 | DS DP | | |
| | | 7550433 | Jun 02, 2026 | U-1238 | | |
| | | 7919118 | May 12, 2024 | DS DP | | |
| | | 7919461 | Jun 02, 2026 | U-1238 | | |
| <u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u> | | | | | | |
| N 202799 | 003 | 7084245 | May 12, 2024 | DS DP U-1238 | | |
| | | 7414105 | May 12, 2024 | DS DP U-1238 | | |
| | | 7528104 | May 12, 2024 | DS DP | | |
| | | 7550433 | Jun 02, 2026 | U-1238 | | |
| | | 7919118 | May 12, 2024 | DS DP | | |
| | | 7919461 | Jun 02, 2026 | U-1238 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u> | | | | | | |
| N 202799 004 | 7084245 | May 12, 2024 | DS DP U-1238 | | | |
| | 7414105 | May 12, 2024 | DS DP U-1238 | | | |
| | 7528104 | May 12, 2024 | DS DP | | | |
| | 7550433 | Jun 02, 2026 | | | U-1238 | |
| | 7919118 | May 12, 2024 | DS DP | | | |
| | 7919461 | Jun 02, 2026 | | | U-1238 | |
| <u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u> | | | | | | |
| N 202799 005 | 7084245 | May 12, 2024 | DS DP U-1238 | | | |
| | 7414105 | May 12, 2024 | DS DP U-1238 | | | |
| | 7528104 | May 12, 2024 | DS DP | | | |
| | 7550433 | Jun 02, 2026 | | | U-1238 | |
| | 7919118 | May 12, 2024 | DS DP | | | |
| | 7919461 | Jun 02, 2026 | | | U-1238 | |
| <u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u> | | | | | | |
| N 202799 006 | 7084245 | May 12, 2024 | DS DP U-1238 | | | |
| | 7414105 | May 12, 2024 | DS DP U-1238 | | | |
| | 7528104 | May 12, 2024 | DS DP | | | |
| | 7550433 | Jun 02, 2026 | | | U-1238 | |
| | 7919118 | May 12, 2024 | DS DP | | | |
| | 7919461 | Jun 02, 2026 | | | U-1238 | |
| <u>PEGINESATIDE ACETATE - OMONTYS</u> | | | | | | |
| N 202799 007 | 7084245 | May 12, 2024 | DS DP U-1238 | | | |
| | 7414105 | May 12, 2024 | DS DP U-1238 | | | |
| | 7528104 | May 12, 2024 | DS DP | | | |
| | 7550433 | Jun 02, 2026 | | | U-1238 | |
| | 7919118 | May 12, 2024 | DS DP | | | |
| | 7919461 | Jun 02, 2026 | | | U-1238 | |
| <u>PEGINESATIDE ACETATE - OMONTYS</u> | | | | | | |
| N 202799 008 | 7084245 | May 12, 2024 | DS DP U-1238 | | | |
| | 7414105 | May 12, 2024 | DS DP U-1238 | | | |
| | 7528104 | May 12, 2024 | DS DP | | | |
| | 7550433 | Jun 02, 2026 | | | U-1238 | |
| | 7919118 | May 12, 2024 | DS DP | | | |
| | 7919461 | Jun 02, 2026 | | | U-1238 | |
| <u>PEMETREXED DISODIUM - ALIMTA</u> | | | | | | |
| N 021462 001 | 7772209 | Nov 24, 2021 | | | U-1296 | |
| <u>PEMETREXED DISODIUM - ALIMTA</u> | | | | | | |
| N 021462 002 | 7772209 | Nov 24, 2021 | | | U-1296 | |
| <u>PENCICLOVIR - DENAVIR</u> | | | | | | |
| N 020629 001 | 6469015 | Oct 22, 2019 | | | U-501 | |
| | 6579981 | Jun 17, 2020 | | | U-501 | |
| <u>PERAMIVIR - RAPIVAB</u> | | | | | | |
| N 206426 001 | 6503745 | Nov 05, 2019 | DS | | NCE | Dec 19, 2019 |
| | 6562861 | Dec 17, 2019 | DS | | NPP | Sep 20, 2020 |
| | 8778997 | May 07, 2027 | | | U-1627 | |
| <u>PERAMPANEL - FYCOMPA</u> | | | | | | |
| N 202834 001 | 6949571 | Jun 08, 2021 | DS DP U-106 | | | |
| | 6949571 | Jun 08, 2021 | DS DP U-2088 | | | |
| | 6949571 | Jun 08, 2021 | DS DP U-2089 | | | |
| | 6949571 | Jun 08, 2021 | DS DP U-2428 | | | |
| | 6949571 | Jun 08, 2021 | DS DP U-2429 | | | |
| | 8772497 | Jul 01, 2026 | DS | | | |
| <u>PERAMPANEL - FYCOMPA</u> | | | | | | |
| N 202834 002 | 6949571 | Jun 08, 2021 | DS DP U-106 | | | |
| | 6949571 | Jun 08, 2021 | DS DP U-2088 | | | |
| | 6949571 | Jun 08, 2021 | DS DP U-2089 | | | |
| | 6949571 | Jun 08, 2021 | DS DP U-2428 | | | |
| | 6949571 | Jun 08, 2021 | DS DP U-2429 | | | |
| | 8772497 | Jul 01, 2026 | DS | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PERAMPANEL - FYCOMPA</u> | | | | | | |
| N 202834 | 002 | 6949571 | Jun 08, 2021 | DS DP U-106 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2088 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2089 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2428 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2429 | | |
| | | 8772497 | Jul 01, 2026 | DS | | |
| <u>PERAMPANEL - FYCOMPA</u> | | | | | | |
| N 202834 | 003 | 6949571 | Jun 08, 2021 | DS DP U-106 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2088 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2089 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2428 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2429 | | |
| | | 8772497 | Jul 01, 2026 | DS | | |
| <u>PERAMPANEL - FYCOMPA</u> | | | | | | |
| N 202834 | 004 | 6949571 | Jun 08, 2021 | DS DP U-106 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2088 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2089 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2428 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2429 | | |
| | | 8772497 | Jul 01, 2026 | DS | | |
| <u>PERAMPANEL - FYCOMPA</u> | | | | | | |
| N 202834 | 005 | 6949571 | Jun 08, 2021 | DS DP U-106 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2088 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2089 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2428 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2429 | | |
| | | 8772497 | Jul 01, 2026 | DS | | |
| <u>PERAMPANEL - FYCOMPA</u> | | | | | | |
| N 202834 | 006 | 6949571 | Jun 08, 2021 | DS DP U-106 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2088 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2089 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2428 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2429 | | |
| | | 8772497 | Jul 01, 2026 | DS | | |
| <u>PERAMPANEL - FYCOMPA</u> | | | | | | |
| N 208277 | 001 | 6949571 | Jun 08, 2021 | DS DP U-106 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2088 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2089 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2428 | | |
| | | 6949571 | Jun 08, 2021 | DS DP U-2429 | | |
| | | 8772497 | Jul 01, 2026 | DS | | |
| <u>PERFLUTREN - DEFINITY</u> | | | | | | |
| N 021064 | 001 | 8658205 | Jun 18, 2019 | DP | | |
| | | 8685441 | Jan 13, 2019 | U-665 | | |
| | | 9545457 | Jan 13, 2019 | U-665 | | |
| | | 9789210 | Mar 16, 2037 | U-665 | | |
| <u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u> | | | | | | |
| N 202088 | 001 | 8440170 | Mar 14, 2029 | DP | | |
| <u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u> | | | | | | |
| N 202088 | 002 | 8440170 | Mar 14, 2029 | DP | | |
| <u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u> | | | | | | |
| N 202088 | 003 | 8440170 | Mar 14, 2029 | DP | | |
| <u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u> | | | | | | |
| N 022580 | 001 | 7056890 | Jun 14, 2020 | DP U-1262 | | |
| | | 7553818 | Jun 14, 2020 | U-1262 | | |
| | | 7659256 | Jun 14, 2020 | DP U-1262 | | |
| | | 7674776 | Jun 14, 2020 | DP U-1262 | | |
| | | 8580298 | May 15, 2029 | DP | | |
| | | 8580299 | Jun 14, 2029 | U-1262 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - OSYMLA</u> | | | | | | |
| N 022580 001 | 8895057 | Jun 09, 2028 | | | U-1262 | |
| | 8895058 | Jun 09, 2028 | DP | | | |
| | 9011905 | Jun 09, 2028 | DP | | | |
| | 9011906 | Jun 09, 2028 | | | U-1262 | |
| <u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - OSYMLA</u> | | | | | | |
| N 022580 002 | 7056890 | Jun 14, 2020 | DP | | U-1262 | |
| | 7553818 | Jun 14, 2020 | | | U-1262 | |
| | 7659256 | Jun 14, 2020 | DP | | U-1262 | |
| | 7674776 | Jun 14, 2020 | DP | | U-1262 | |
| | 8580298 | May 15, 2029 | DP | | | |
| | 8580299 | Jun 14, 2029 | | | U-1262 | |
| | 8895057 | Jun 09, 2028 | | | U-1262 | |
| | 8895058 | Jun 09, 2028 | DP | | | |
| | 9011905 | Jun 09, 2028 | DP | | | |
| | 9011906 | Jun 09, 2028 | | | U-1262 | |
| <u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - OSYMLA</u> | | | | | | |
| N 022580 003 | 7056890 | Jun 14, 2020 | DP | | U-1262 | |
| | 7553818 | Jun 14, 2020 | | | U-1262 | |
| | 7659256 | Jun 14, 2020 | DP | | U-1262 | |
| | 7674776 | Jun 14, 2020 | DP | | U-1262 | |
| | 8580298 | May 15, 2029 | DP | | | |
| | 8580299 | Jun 14, 2029 | | | U-1262 | |
| | 8895057 | Jun 09, 2028 | | | U-1262 | |
| | 8895058 | Jun 09, 2028 | DP | | | |
| | 9011905 | Jun 09, 2028 | DP | | | |
| | 9011906 | Jun 09, 2028 | | | U-1262 | |
| <u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - OSYMLA</u> | | | | | | |
| N 022580 004 | 7056890 | Jun 14, 2020 | DP | | U-1262 | |
| | 7553818 | Jun 14, 2020 | | | U-1262 | |
| | 7659256 | Jun 14, 2020 | DP | | U-1262 | |
| | 7674776 | Jun 14, 2020 | DP | | U-1262 | |
| | 8580298 | May 15, 2029 | DP | | | |
| | 8580299 | Jun 14, 2029 | | | U-1262 | |
| | 8895057 | Jun 09, 2028 | | | U-1262 | |
| | 8895058 | Jun 09, 2028 | DP | | | |
| | 9011905 | Jun 09, 2028 | DP | | | |
| | 9011906 | Jun 09, 2028 | | | U-1262 | |
| <u>PHENTOLAMINE MESYLATE - ORAVERSE</u> | | | | | | |
| N 022159 001 | 6764678 | May 11, 2021 | | | U-967 | |
| | 6872390 | May 11, 2021 | DP | | | |
| | 7229630 | Jun 20, 2023 | DP | | | |
| | 7569230 | Oct 17, 2023 | | | U-967 | |
| | 7575757 | Apr 21, 2025 | DP | | | |
| <u>PHENYLEPHRINE HYDROCHLORIDE - PHENYLEPHRINE HYDROCHLORIDE</u> | | | | | | |
| N 203510 001 | 8859623 | Nov 14, 2033 | | | U-1594 | |
| <u>PHENYLEPHRINE HYDROCHLORIDE - PHENYLEPHRINE HYDROCHLORIDE</u> | | | | | | |
| N 203510 002 | 8859623 | Nov 14, 2033 | | | U-1594 | |
| <u>PIMAVANSERIN TARTRATE - NUPLAZID</u> | | | | | | |
| N 207318 001 | 6756393 | Mar 06, 2021 | DS DP | | | |
| | 6815458 | Mar 06, 2021 | DS DP | | U-1843 | |
| | 7115634 | Oct 06, 2021 | DS DP | | | |
| | 7601740 | Jun 17, 2027 | DS DP | | | |
| | 7659285 | Aug 24, 2026 | | | U-1844 | |
| | 7732615 | Jun 03, 2028 | DS DP | | | |
| | 7858789 | Dec 13, 2020 | DS DP | | | |
| | 7923564 | Sep 26, 2025 | DS DP | | | |
| | 8110574 | Dec 13, 2020 | DS DP | | | |
| | 8618130 | Jan 15, 2024 | | | U-1845 | |
| | 8921393 | Jan 15, 2024 | | | U-1846 | |
| | 9296694 | Mar 06, 2021 | DS DP | | | |
| | 9566271 | Jan 15, 2024 | | | U-1974 | |
| | 9765053 | Jul 27, 2022 | | | U-1974 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PIMAVANSERIN TARTRATE - NUPLAZID</u> | | | | | | |
| N 207318 001 | 6756393 | Mar 06, 2021 | DS DP | | NCE | Apr 29, 2021 |
| | 6815458 | Mar 06, 2021 | DS DP | U-1843 | | |
| | 7115634 | Oct 06, 2021 | DS DP | | | |
| | 7601740 | Jun 17, 2027 | DS DP | | | |
| | 7659285 | Aug 24, 2026 | | U-1844 | | |
| | 7732615 | Jun 03, 2028 | DS DP | | | |
| | 7858789 | Dec 13, 2020 | DS DP | | | |
| | 7923564 | Sep 26, 2025 | DS DP | | | |
| | 8110574 | Dec 13, 2020 | DS DP | | | |
| | 8618130 | Jan 15, 2024 | | U-1845 | | |
| | 8921393 | Jan 15, 2024 | | U-1846 | | |
| | 9296694 | Mar 06, 2021 | DS DP | | | |
| | 9566271 | Jan 15, 2024 | | U-1974 | | |
| | 9765053 | Jul 27, 2022 | | U-1974 | | |
| <u>PIMAVANSERIN TARTRATE - NUPLAZID</u> | | | | | | |
| N 207318 002 | 10028944 | Jan 15, 2024 | | U-1974 | | |
| | 6756393 | Mar 06, 2021 | DS DP | | | |
| | 6815458 | Mar 06, 2021 | DS DP | U-1843 | | |
| | 7115634 | Oct 06, 2021 | DS DP | | | |
| | 7601740 | Jun 17, 2027 | DS DP | | | |
| | 7659285 | Aug 24, 2026 | | U-1844 | | |
| | 7732615 | Jun 03, 2028 | DS DP | | | |
| | 7858789 | Dec 13, 2020 | DS DP | | | |
| | 7923564 | Sep 26, 2025 | DS DP | | | |
| | 8110574 | Dec 13, 2020 | DS DP | | | |
| | 8618130 | Jan 15, 2024 | | U-1845 | | |
| | 8921393 | Jan 15, 2024 | | U-1846 | | |
| | 9296694 | Mar 06, 2021 | DS DP | | | |
| | 9566271 | Jan 15, 2024 | | U-1974 | | |
| | 9765053 | Jul 27, 2022 | | U-1974 | | |
| <u>PIMAVANSERIN TARTRATE - NUPLAZID</u> | | | | | | |
| N 210793 001 | 10028944 | Jan 15, 2024 | | | NCE | Apr 29, 2021 |
| | 6756393 | Mar 06, 2021 | DS DP | | | |
| | 6815458 | Mar 06, 2021 | DS DP | U-1843 | | |
| | 7115634 | Oct 06, 2021 | DS DP | | | |
| | 7601740 | Jun 17, 2027 | DS DP | | | |
| | 7659285 | Aug 24, 2026 | | U-1844 | | |
| | 7732615 | Jun 03, 2028 | DS DP | | | |
| | 7858789 | Dec 13, 2020 | DS DP | | | |
| | 7923564 | Sep 26, 2025 | DS DP | | | |
| | 8110574 | Dec 13, 2020 | DS DP | | | |
| | 8618130 | Jan 15, 2024 | | U-1845 | | |
| | 8921393 | Jan 15, 2024 | | U-1846 | | |
| | 9296694 | Mar 06, 2021 | DS DP | | | |
| | 9566271 | Jan 15, 2024 | | U-1974 | | |
| | 9765053 | Jul 27, 2022 | | U-1974 | | |
| <u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u> | | | | | | |
| N 050684 001 | 6900184 | Apr 14, 2023 | DP | U-282 | | |
| | 7915229 | Apr 14, 2023 | DP | | | |
| | 8133883 | Apr 14, 2023 | DP | U-282 | | |
| <u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u> | | | | | | |
| N 050684 002 | 6900184 | Apr 14, 2023 | DP | U-282 | | |
| | 7915229 | Apr 14, 2023 | DP | | | |
| | 8133883 | Apr 14, 2023 | DP | U-282 | | |
| <u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u> | | | | | | |
| N 050684 003 | 6900184 | Apr 14, 2023 | DP | U-282 | | |
| | 7915229 | Apr 14, 2023 | DP | | | |
| | 8133883 | Apr 14, 2023 | DP | U-282 | | |
| <u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u> | | | | | | |
| N 050684 004 | 6900184 | Apr 14, 2023 | DP | U-282 | | |
| | 7915229 | Apr 14, 2023 | DP | | | |
| | 8133883 | Apr 14, 2023 | DP | U-282 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u> | | | | | | |
| N 050750 001 | 6207661 | Feb 22, 2019 | DP | | | |
| | 6900184 | Apr 14, 2023 | DP U-282 | | | |
| | 7915229 | Apr 14, 2023 | DP | | | |
| | 8133883 | Apr 14, 2023 | DP U-282 | | | |
| <u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u> | | | | | | |
| N 050750 002 | 6207661 | Feb 22, 2019 | DP | | | |
| | 6900184 | Apr 14, 2023 | DP U-282 | | | |
| | 7915229 | Apr 14, 2023 | DP | | | |
| | 8133883 | Apr 14, 2023 | DP U-282 | | | |
| <u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u> | | | | | | |
| N 050750 003 | 6207661 | Feb 22, 2019 | DP | | | |
| | 6900184 | Apr 14, 2023 | DP U-282 | | | |
| | 7915229 | Apr 14, 2023 | DP | | | |
| | 8133883 | Apr 14, 2023 | DP U-282 | | | |
| <u>PIRFENIDONE - ESBRIET</u> | | | | | | |
| N 022535 001 | 7566729 | Apr 22, 2029 | U-1600 | | NCE | Oct 15, 2019 |
| | 7635707 | Apr 22, 2029 | U-1609 | | ODE-77 | Oct 15, 2021 |
| | 7696236 | Dec 18, 2027 | U-1601 | | | |
| | 7767225 | Sep 22, 2026 | DP U-1602 | | | |
| | 7767700 | Dec 18, 2027 | U-1601 | | | |
| | 7816383 | Jan 08, 2030 | U-1603 | | | |
| | 7910610 | Jan 08, 2030 | U-1604 | | | |
| | 7988994 | Sep 22, 2026 | DP U-1602 | | | |
| | 8013002 | Jan 08, 2030 | U-1603 | | | |
| | 8084475 | Jan 08, 2030 | U-1605 | | | |
| | 8318780 | Jan 08, 2030 | U-1606 | | | |
| | 8383150 | Sep 22, 2026 | DP U-1607 | | | |
| | 8383150 | Sep 22, 2026 | DP U-2361 | | | |
| | 8420674 | Dec 18, 2027 | DP U-1608 | | | |
| | 8592462 | Apr 22, 2029 | U-1609 | | | |
| | 8609701 | Apr 22, 2029 | U-1610 | | | |
| | 8648098 | Jan 08, 2030 | U-1611 | | | |
| | 8753679 | Sep 22, 2026 | DP U-1602 | | | |
| | 8754109 | Jan 08, 2030 | U-1612 | | | |
| | 8778947 | Aug 30, 2033 | U-1613 | | | |
| <u>PIRFENIDONE - ESBRIET</u> | | | | | | |
| N 208780 001 | 7566729 | Apr 22, 2029 | U-2077 | | NCE | Oct 15, 2019 |
| | 7566729 | Apr 22, 2029 | U-2078 | | ODE-77 | Oct 15, 2021 |
| | 7635707 | Apr 22, 2029 | U-2072 | | | |
| | 7635707 | Apr 22, 2029 | U-2073 | | | |
| | 7635707 | Apr 22, 2029 | U-2074 | | | |
| | 7635707 | Apr 22, 2029 | U-2075 | | | |
| | 7635707 | Apr 22, 2029 | U-2076 | | | |
| | 7635707 | Apr 22, 2029 | U-2083 | | | |
| | 7767700 | Dec 18, 2027 | U-2080 | | | |
| | 7816383 | Jan 08, 2030 | U-2042 | | | |
| | 7816383 | Jan 08, 2030 | U-2050 | | | |
| | 7910610 | Jan 08, 2030 | U-2048 | | | |
| | 7910610 | Jan 08, 2030 | U-2049 | | | |
| | 8013002 | Jan 08, 2030 | U-2047 | | | |
| | 8013002 | Jan 08, 2030 | U-2082 | | | |
| | 8084475 | Jan 08, 2030 | U-2052 | | | |
| | 8084475 | Jan 08, 2030 | U-2054 | | | |
| | 8318780 | Jan 08, 2030 | U-2046 | | | |
| | 8318780 | Jan 08, 2030 | U-2081 | | | |
| | 8383150 | Sep 22, 2026 | DP U-2361 | | | |
| | 8420674 | Dec 18, 2027 | U-2079 | | | |
| | 8592462 | Apr 22, 2029 | U-2055 | | | |
| | 8592462 | Apr 22, 2029 | U-2056 | | | |
| | 8592462 | Apr 22, 2029 | U-2057 | | | |
| | 8592462 | Apr 22, 2029 | U-2058 | | | |
| | 8592462 | Apr 22, 2029 | U-2059 | | | |
| | 8592462 | Apr 22, 2029 | U-2060 | | | |
| | 8592462 | Apr 22, 2029 | U-2061 | | | |
| | 8592462 | Apr 22, 2029 | U-2062 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PIRFENIDONE - ESBRIET</u> | | | | | | |
| N 208780 001 | 8592462 | Apr 22, 2029 | U-2063 | | | |
| | 8609701 | Apr 22, 2029 | U-2064 | | | |
| | 8609701 | Apr 22, 2029 | U-2065 | | | |
| | 8609701 | Apr 22, 2029 | U-2066 | | | |
| | 8609701 | Apr 22, 2029 | U-2067 | | | |
| | 8609701 | Apr 22, 2029 | U-2068 | | | |
| | 8609701 | Apr 22, 2029 | U-2069 | | | |
| | 8609701 | Apr 22, 2029 | U-2070 | | | |
| | 8648098 | Jan 08, 2030 | U-2051 | | | |
| | 8648098 | Jan 08, 2030 | U-2052 | | | |
| | 8754109 | Jan 08, 2030 | U-2053 | | | |
| | 8778947 | Aug 30, 2033 | U-2044 | | | |
| | 8778947 | Aug 30, 2033 | U-2045 | | | |
| | 9561217 | Jan 25, 2022 | | DP | | |
| <u>PIRFENIDONE - ESBRIET</u> | | | | | | |
| N 208780 002 | 7566729 | Apr 22, 2029 | U-2269 | | NCE | Oct 15, 2019 |
| | 7566729 | Apr 22, 2029 | U-2270 | | ODE-77 | Oct 15, 2021 |
| | 7635707 | Apr 22, 2029 | U-2072 | | | |
| | 7635707 | Apr 22, 2029 | U-2073 | | | |
| | 7635707 | Apr 22, 2029 | U-2074 | | | |
| | 7635707 | Apr 22, 2029 | U-2075 | | | |
| | 7635707 | Apr 22, 2029 | U-2076 | | | |
| | 7635707 | Apr 22, 2029 | U-2083 | | | |
| | 7767700 | Dec 18, 2027 | U-2080 | | | |
| | 7816383 | Jan 08, 2030 | U-2042 | | | |
| | 7816383 | Jan 08, 2030 | U-2050 | | | |
| | 7910610 | Jan 08, 2030 | U-2048 | | | |
| | 7910610 | Jan 08, 2030 | U-2049 | | | |
| | 8013002 | Jan 08, 2030 | U-2047 | | | |
| | 8013002 | Jan 08, 2030 | U-2082 | | | |
| | 8084475 | Jan 08, 2030 | U-2054 | | | |
| | 8084475 | Jan 08, 2030 | U-2268 | | | |
| | 8318780 | Jan 08, 2030 | U-2046 | | | |
| | 8318780 | Jan 08, 2030 | U-2081 | | | |
| | 8383150 | Sep 22, 2026 | | DP | | |
| | 8420674 | Dec 18, 2027 | U-2079 | | | |
| | 8592462 | Apr 22, 2029 | U-2055 | | | |
| | 8592462 | Apr 22, 2029 | U-2056 | | | |
| | 8592462 | Apr 22, 2029 | U-2057 | | | |
| | 8592462 | Apr 22, 2029 | U-2058 | | | |
| | 8592462 | Apr 22, 2029 | U-2059 | | | |
| | 8592462 | Apr 22, 2029 | U-2060 | | | |
| | 8592462 | Apr 22, 2029 | U-2061 | | | |
| | 8592462 | Apr 22, 2029 | U-2062 | | | |
| | 8592462 | Apr 22, 2029 | U-2063 | | | |
| | 8609701 | Apr 22, 2029 | U-2064 | | | |
| | 8609701 | Apr 22, 2029 | U-2065 | | | |
| | 8609701 | Apr 22, 2029 | U-2066 | | | |
| | 8609701 | Apr 22, 2029 | U-2067 | | | |
| | 8609701 | Apr 22, 2029 | U-2068 | | | |
| | 8609701 | Apr 22, 2029 | U-2069 | | | |
| | 8609701 | Apr 22, 2029 | U-2070 | | | |
| | 8648098 | Jan 08, 2030 | U-2051 | | | |
| | 8648098 | Jan 08, 2030 | U-2052 | | | |
| | 8754109 | Jan 08, 2030 | U-2053 | | | |
| | 8778947 | Aug 30, 2033 | U-2044 | | | |
| | 8778947 | Aug 30, 2033 | U-2045 | | | |
| | 9561217 | Jan 25, 2022 | | DP | | |
| <u>PIRFENIDONE - ESBRIET</u> | | | | | | |
| N 208780 003 | 7566729 | Apr 22, 2029 | U-2077 | | NCE | Oct 15, 2019 |
| | 7566729 | Apr 22, 2029 | U-2078 | | ODE-77 | Oct 15, 2021 |
| | 7635707 | Apr 22, 2029 | U-2072 | | | |
| | 7635707 | Apr 22, 2029 | U-2073 | | | |
| | 7635707 | Apr 22, 2029 | U-2074 | | | |
| | 7635707 | Apr 22, 2029 | U-2075 | | | |
| | 7635707 | Apr 22, 2029 | U-2076 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PIR Fenidone - ESBRIET</u> | | | | | | |
| N 208780 003 | 7635707 | Apr 22, 2029 | | | | |
| | 7767700 | Dec 18, 2027 | | | | |
| | 7816383 | Jan 08, 2030 | | | | |
| | 7816383 | Jan 08, 2030 | | | | |
| | 7910610 | Jan 08, 2030 | | | | |
| | 7910610 | Jan 08, 2030 | | | | |
| | 8013002 | Jan 08, 2030 | | | | |
| | 8013002 | Jan 08, 2030 | | | | |
| | 8084475 | Jan 08, 2030 | | | | |
| | 8084475 | Jan 08, 2030 | | | | |
| | 8318780 | Jan 08, 2030 | | | | |
| | 8318780 | Jan 08, 2030 | | | | |
| | 8383150 | Sep 22, 2026 | DP | | | |
| | 8420674 | Dec 18, 2027 | | | | |
| | 8592462 | Apr 22, 2029 | | | | |
| | 8592462 | Apr 22, 2029 | | | | |
| | 8592462 | Apr 22, 2029 | | | | |
| | 8592462 | Apr 22, 2029 | | | | |
| | 8592462 | Apr 22, 2029 | | | | |
| | 8592462 | Apr 22, 2029 | | | | |
| | 8592462 | Apr 22, 2029 | | | | |
| | 8592462 | Apr 22, 2029 | | | | |
| | 8592462 | Apr 22, 2029 | | | | |
| | 8592462 | Apr 22, 2029 | | | | |
| | 8592462 | Apr 22, 2029 | | | | |
| | 8592462 | Apr 22, 2029 | | | | |
| | 8609701 | Apr 22, 2029 | | | | |
| | 8609701 | Apr 22, 2029 | | | | |
| | 8609701 | Apr 22, 2029 | | | | |
| | 8609701 | Apr 22, 2029 | | | | |
| | 8609701 | Apr 22, 2029 | | | | |
| | 8609701 | Apr 22, 2029 | | | | |
| | 8609701 | Apr 22, 2029 | | | | |
| | 8609701 | Apr 22, 2029 | | | | |
| | 8609701 | Apr 22, 2029 | | | | |
| | 8648098 | Jan 08, 2030 | | | | |
| | 8648098 | Jan 08, 2030 | | | | |
| | 8754109 | Jan 08, 2030 | | | | |
| | 8778947 | Aug 30, 2033 | | | | |
| | 8778947 | Aug 30, 2033 | | | | |
| | 9561217 | Jan 25, 2022 | DP | | | |
| <u>PITAVASTATIN CALCIUM - LIVALO</u> | | | | | | |
| N 022363 001 | 5856336 | Dec 25, 2020 | DS | | | |
| | 7022713 | Feb 19, 2024 | | | | |
| | 8557993 | Feb 02, 2024 | DP | | | |
| <u>PITAVASTATIN CALCIUM - LIVALO</u> | | | | | | |
| N 022363 002 | 5856336 | Dec 25, 2020 | DS | | | |
| | 7022713 | Feb 19, 2024 | | | | |
| | 8557993 | Feb 02, 2024 | DP | | | |
| <u>PITAVASTATIN CALCIUM - LIVALO</u> | | | | | | |
| N 022363 003 | 5856336 | Dec 25, 2020 | DS | | | |
| | 7022713 | Feb 19, 2024 | | | | |
| | 8557993 | Feb 02, 2024 | DS DP | | | |
| <u>PITAVASTATIN MAGNESIUM - ZYPITAMAG</u> | | | | | | |
| N 208379 001 | 8829186 | Jan 19, 2031 | DS DP | | | |
| <u>PITAVASTATIN MAGNESIUM - ZYPITAMAG</u> | | | | | | |
| N 208379 002 | 8829186 | Jan 19, 2031 | DS DP | | | |
| <u>PITAVASTATIN MAGNESIUM - ZYPITAMAG</u> | | | | | | |
| N 208379 003 | 8829186 | Jan 19, 2031 | DS DP | | | |
| <u>PLAZOMICIN SULFATE - ZEMDRI</u> | | | | | | |
| N 210303 001 | 8383596 | Jun 02, 2031 | DS | | | |
| | 8822424 | Nov 21, 2028 | DP | | | |
| | 9266919 | Nov 21, 2028 | | | | |
| | 9688711 | Nov 21, 2028 | DS | | | |
| | | | | | NCE GAIN | Jun 25, 2023 Jun 25, 2028 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PLECANATIDE - TRULANCE</u> | | | | | | |
| N 208745 | 001 | 10011637 | Jun 05, 2034 | DS | I-764 | Jan 24, 2021 |
| | | 7041786 | Mar 25, 2023 | DS | NCE | Jan 19, 2022 |
| | | 7799897 | Jun 09, 2022 | DS | | |
| | | 8637451 | Mar 28, 2022 | | U-1964 | |
| | | 9610321 | Sep 15, 2031 | | U-1999 | |
| | | 9610321 | Sep 15, 2031 | | U-2230 | |
| | | 9616097 | Jul 02, 2032 | DP | | |
| | | 9919024 | Sep 15, 2031 | | U-1999 | |
| | | 9919024 | Sep 15, 2031 | | U-2230 | |
| | | 9925231 | Sep 15, 2031 | DP | | |
| <u>PLERIXAFOR - MOZOBIL</u> | | | | | | |
| N 022311 | 001 | 6987102 | Jul 22, 2023 | | U-936 | |
| | | 7897590 | Jul 22, 2023 | | U-936 | |
| <u>POLIDOCANOL - VARITHENA</u> | | | | | | |
| N 205098 | 001 | 6572873 | May 26, 2020 | | U-1461 | |
| | | 6846412 | Jul 19, 2022 | DP | | |
| | | 6942165 | May 26, 2020 | DP | | |
| | | 7025290 | May 26, 2020 | DP | U-1461 | |
| | | 7357336 | May 26, 2020 | | U-1461 | |
| | | 7604185 | May 26, 2020 | DS DP | U-1462 | |
| | | 7731986 | Nov 17, 2024 | DS DP | U-1463 | |
| | | 7814943 | Nov 19, 2027 | DP | U-1461 | |
| | | 7842282 | May 26, 2020 | | U-1461 | |
| | | 7842283 | May 26, 2020 | DP | | |
| | | 8122917 | Sep 09, 2024 | DP | | |
| | | 8323677 | May 26, 2020 | DS | | |
| | | 8734833 | May 26, 2020 | DS DP | | |
| | | 9480652 | May 12, 2032 | DP | | |
| <u>POMALIDOMIDE - POMALYST</u> | | | | | | |
| N 204026 | 001 | 6315720 | Oct 23, 2020 | | ODE-43 | Feb 08, 2020 |
| | | 6561977 | Oct 23, 2020 | | U-1361 | |
| | | 6755784 | Oct 23, 2020 | | U-1361 | |
| | | 8198262 | Jun 17, 2025 | | U-1360 | |
| | | 8198262 | Jun 17, 2025 | | U-2254 | |
| | | 8315886 | Oct 23, 2020 | | U-1361 | |
| | | 8626531 | Oct 23, 2020 | | U-1361 | |
| | | 8673939 | May 15, 2023 | | U-1360 | |
| | | 8673939 | May 15, 2023 | | U-2254 | |
| | | 8735428 | May 15, 2023 | | U-1360 | |
| | | 8735428 | May 15, 2023 | | U-2254 | |
| | | 8828427 | Jun 21, 2031 | DS DP | | |
| | | 9993467 | May 19, 2030 | DP | | |
| <u>POMALIDOMIDE - POMALYST</u> | | | | | | |
| N 204026 | 002 | 6315720 | Oct 23, 2020 | | ODE-43 | Feb 08, 2020 |
| | | 6561977 | Oct 23, 2020 | | U-1361 | |
| | | 6755784 | Oct 23, 2020 | | U-1361 | |
| | | 8198262 | Jun 17, 2025 | | U-1360 | |
| | | 8198262 | Jun 17, 2025 | | U-2254 | |
| | | 8315886 | Oct 23, 2020 | | U-1361 | |
| | | 8626531 | Oct 23, 2020 | | U-1361 | |
| | | 8673939 | May 15, 2023 | | U-1360 | |
| | | 8673939 | May 15, 2023 | | U-2254 | |
| | | 8735428 | May 15, 2023 | | U-1360 | |
| | | 8735428 | May 15, 2023 | | U-2254 | |
| | | 8828427 | Jun 21, 2031 | DS DP | | |
| | | 9993467 | May 19, 2030 | DP | | |
| <u>POMALIDOMIDE - POMALYST</u> | | | | | | |
| N 204026 | 003 | 6315720 | Oct 23, 2020 | | ODE-43 | Feb 08, 2020 |
| | | 6561977 | Oct 23, 2020 | | U-1361 | |
| | | 6755784 | Oct 23, 2020 | | U-1361 | |
| | | 8198262 | Jun 17, 2025 | | U-1360 | |
| | | 8198262 | Jun 17, 2025 | | U-2254 | |
| | | 8315886 | Oct 23, 2020 | | U-1361 | |
| | | 8626531 | Oct 23, 2020 | | U-1361 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>POMALIDOMIDE - POMALYST</u> | | | | | | |
| N 204026 003 | 8673939 | May 15, 2023 | U-1360 | | | |
| | 8673939 | May 15, 2023 | U-2254 | | | |
| | 8735428 | May 15, 2023 | U-1360 | | | |
| | 8735428 | May 15, 2023 | U-2254 | | | |
| | 8828427 | Jun 21, 2031 | DS DP | | | |
| | 9993467 | May 19, 2030 | DP | | | |
| <u>POMALIDOMIDE - POMALYST</u> | | | | | | |
| N 204026 004 | 6315720 | Oct 23, 2020 | U-1361 | | ODE-43 | Feb 08, 2020 |
| | 6561977 | Oct 23, 2020 | U-1361 | | | |
| | 6755784 | Oct 23, 2020 | U-1361 | | | |
| | 8198262 | Jun 17, 2025 | U-1360 | | | |
| | 8198262 | Jun 17, 2025 | U-2254 | | | |
| | 8315886 | Oct 23, 2020 | U-1361 | | | |
| | 8626531 | Oct 23, 2020 | U-1361 | | | |
| | 8673939 | May 15, 2023 | U-1360 | | | |
| | 8673939 | May 15, 2023 | U-2254 | | | |
| | 8735428 | May 15, 2023 | U-1360 | | | |
| | 8735428 | May 15, 2023 | U-2254 | | | |
| | 8828427 | Jun 21, 2031 | DS DP | | | |
| | 9993467 | May 19, 2030 | DP | | | |
| <u>PONATINIB HYDROCHLORIDE - ICLUSIG</u> | | | | | | |
| N 203469 001 | 8114874 | Dec 22, 2026 | DS DP | | ODE-35 | Dec 14, 2019 |
| | 9029533 | Dec 22, 2026 | U-1283 | | | |
| | 9029533 | Dec 22, 2026 | U-1699 | | | |
| | 9029533 | Dec 22, 2026 | U-1700 | | | |
| | 9029533 | Dec 22, 2026 | U-1701 | | | |
| | 9029533 | Dec 22, 2026 | U-836 | | | |
| | 9493470 | Dec 12, 2033 | DS DP U-1700 | | | |
| | 9493470 | Dec 12, 2033 | DS DP U-1948 | | | |
| <u>PONATINIB HYDROCHLORIDE - ICLUSIG</u> | | | | | | |
| N 203469 002 | 8114874 | Dec 22, 2026 | DS DP | | ODE-35 | Dec 14, 2019 |
| | 9029533 | Dec 22, 2026 | U-1283 | | | |
| | 9029533 | Dec 22, 2026 | U-1699 | | | |
| | 9029533 | Dec 22, 2026 | U-1700 | | | |
| | 9029533 | Dec 22, 2026 | U-1701 | | | |
| | 9029533 | Dec 22, 2026 | U-836 | | | |
| | 9493470 | Dec 12, 2033 | DS DP U-1700 | | | |
| | 9493470 | Dec 12, 2033 | DS DP U-1948 | | | |
| <u>PONATINIB HYDROCHLORIDE - ICLUSIG</u> | | | | | | |
| N 203469 003 | 8114874 | Dec 22, 2026 | DS DP | | ODE-35 | Dec 14, 2019 |
| | 9029533 | Dec 22, 2026 | U-1283 | | | |
| | 9029533 | Dec 22, 2026 | U-1699 | | | |
| | 9029533 | Dec 22, 2026 | U-1700 | | | |
| | 9029533 | Dec 22, 2026 | U-1701 | | | |
| | 9029533 | Dec 22, 2026 | U-836 | | | |
| | 9493470 | Dec 12, 2033 | DS DP U-1700 | | | |
| | 9493470 | Dec 12, 2033 | DS DP U-1948 | | | |
| <u>POSACONAZOLE - NOXAFIL</u> | | | | | | |
| N 022003 001 | 5661151 | Jul 19, 2019 | DS DP U-760 | | | |
| | 8263600 | Apr 01, 2022 | DP | | | |
| <u>POSACONAZOLE - NOXAFIL</u> | | | | | | |
| N 205053 001 | 5661151 | Jul 19, 2019 | DS DP U-1454 | | | |
| <u>POSACONAZOLE - NOXAFIL</u> | | | | | | |
| N 205596 001 | 10117951 | Mar 13, 2029 | DP | | | |
| | 5661151 | Jul 19, 2019 | DS DP U-1454 | | | |
| | 8410077 | Mar 13, 2029 | DP | | | |
| | 9023790 | Jul 04, 2031 | DP U-1698 | | | |
| | 9358297 | Jun 24, 2031 | DP U-1454 | | | |
| | 9493582 | Feb 27, 2033 | DP | | | |
| | 9750822 | Mar 13, 2029 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>POTASSIUM CHLORIDE - POTASSIUM CHLORIDE</u> | | | | | | |
| A 211067 | 001 | | | | CGT | Feb 25, 2019 |
| <u>POTASSIUM CHLORIDE - POTASSIUM CHLORIDE</u> | | | | | | |
| A 211067 | 002 | | | | CGT | Mar 06, 2019 |
| <u>PRALATREXATE - FOLOTYN</u> | | | | | | |
| N 022468 | 001 | 6028071 | Jul 16, 2022 | DS DP U-1004 | | |
| | | 7622470 | May 31, 2025 | U-1015 | | |
| | | 8299078 | May 31, 2025 | U-1004 | | |
| <u>PRALATREXATE - FOLOTYN</u> | | | | | | |
| N 022468 | 002 | 6028071 | Jul 16, 2022 | DS DP U-1004 | | |
| | | 7622470 | May 31, 2025 | U-1015 | | |
| | | 8299078 | May 31, 2025 | U-1004 | | |
| <u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u> | | | | | | |
| N 022421 | 001 | 7695734 | Apr 26, 2028 | DP | | |
| | | 8679533 | Sep 08, 2029 | DP U-219 | | |
| <u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u> | | | | | | |
| N 022421 | 002 | 7695734 | Apr 26, 2028 | DP | | |
| | | 8679533 | Sep 08, 2029 | DP U-219 | | |
| <u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u> | | | | | | |
| N 022421 | 003 | 7695734 | Apr 26, 2028 | DP | | |
| | | 8679533 | Sep 08, 2029 | DP U-219 | | |
| <u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u> | | | | | | |
| N 022421 | 004 | 7695734 | Apr 26, 2028 | DP | | |
| | | 8679533 | Sep 08, 2029 | DP U-219 | | |
| <u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u> | | | | | | |
| N 022421 | 005 | 7695734 | Apr 26, 2028 | DP | | |
| | | 8679533 | Sep 08, 2029 | DP U-219 | | |
| <u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u> | | | | | | |
| N 022421 | 006 | 7695734 | Apr 26, 2028 | DP | | |
| | | 8679533 | Sep 08, 2029 | DP U-219 | | |
| <u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u> | | | | | | |
| N 022421 | 007 | 7695734 | Apr 26, 2028 | DP | | |
| | | 8679533 | Sep 08, 2029 | DP U-219 | | |
| <u>PRAMLINTIDE ACETATE - SYMLIN</u> | | | | | | |
| N 021332 | 001 | 5686411 | Mar 16, 2019 | DS DP U-638 | | |
| <u>PRAMLINTIDE ACETATE - SYMLIN</u> | | | | | | |
| N 021332 | 002 | 5686411 | Mar 16, 2019 | DS DP U-638 | | |
| <u>PRAMLINTIDE ACETATE - SYMLIN</u> | | | | | | |
| N 021332 | 003 | 5686411 | Mar 16, 2019 | DS DP U-638 | | |
| <u>PRASTERONE - INTRAROSA</u> | | | | | | |
| N 208470 | 001 | 8268806 | Mar 19, 2031 | DP | NCE | Nov 16, 2021 |
| | | 8629129 | Aug 07, 2028 | DP | | |
| | | 8957054 | Aug 07, 2028 | U-1922 | | |
| <u>PRASUGREL HYDROCHLORIDE - EFFIENT</u> | | | | | | |
| N 022307 | 001 | 8404703 | Jan 02, 2023 | U-1381 | M-182 | Jul 12, 2019 |
| | | 8404703*PED | Jul 02, 2023 | | PED | Jan 12, 2020 |
| | | 8569325 | Jan 02, 2023 | U-1381 | | |
| | | 8569325*PED | Jul 02, 2023 | | | |
| <u>PRASUGREL HYDROCHLORIDE - EFFIENT</u> | | | | | | |
| N 022307 | 002 | 8404703 | Jan 02, 2023 | U-1381 | M-182 | Jul 12, 2019 |
| | | 8404703*PED | Jul 02, 2023 | | PED | Jan 12, 2020 |
| | | 8569325 | Jan 02, 2023 | U-1381 | | |
| | | 8569325*PED | Jul 02, 2023 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PREDNISOLONE ACETATE - FLO-PRED</u> | | | | | | |
| N 022067 | 001 | 7799331 | Oct 11, 2028 | DP U-1068 | | |
| | | 7799331 | Oct 11, 2028 | DP U-139 | | |
| <u>PREDNISOLONE ACETATE - FLO-PRED</u> | | | | | | |
| N 022067 | 002 | 7799331 | Oct 11, 2028 | DP U-1068 | | |
| | | 7799331 | Oct 11, 2028 | DP U-139 | | |
| <u>PREDNISOLONE SODIUM PHOSPHATE - ORAPRED ODT</u> | | | | | | |
| N 021959 | 001 | 6740341 | Nov 24, 2019 | DP | | |
| <u>PREDNISOLONE SODIUM PHOSPHATE - ORAPRED ODT</u> | | | | | | |
| N 021959 | 002 | 6740341 | Nov 24, 2019 | DP | | |
| <u>PREDNISOLONE SODIUM PHOSPHATE - ORAPRED ODT</u> | | | | | | |
| N 021959 | 003 | 6740341 | Nov 24, 2019 | DP | | |
| <u>PREDNISONE - RAYOS</u> | | | | | | |
| N 202020 | 001 | 6488960 | Mar 14, 2020 | DP U-1267 | | |
| | | 6677326 | Mar 14, 2020 | DP U-1268 | | |
| | | 8309124 | Apr 23, 2024 | U-1292 | | |
| | | 8394407 | Apr 23, 2024 | DP U-1362 | | |
| | | 9040085 | Apr 23, 2024 | U-1362 | | |
| | | 9186332 | Apr 23, 2024 | U-1362 | | |
| | | 9504699 | Aug 03, 2027 | U-1362 | | |
| <u>PREDNISONE - RAYOS</u> | | | | | | |
| N 202020 | 002 | 6488960 | Mar 14, 2020 | DP U-1267 | | |
| | | 6677326 | Mar 14, 2020 | DP U-1268 | | |
| | | 8309124 | Apr 23, 2024 | | | |
| | | 8394407 | Apr 23, 2024 | DP U-1362 | | |
| | | 9040085 | Apr 23, 2024 | U-1362 | | |
| | | 9186332 | Apr 23, 2024 | U-1362 | | |
| | | 9504699 | Aug 03, 2027 | U-1362 | | |
| <u>PREDNISONE - RAYOS</u> | | | | | | |
| N 202020 | 003 | 8168218 | Jan 07, 2028 | DP U-1269 | | |
| | | 8309124 | Apr 23, 2024 | U-1292 | | |
| | | 8394407 | Apr 23, 2024 | DP U-1362 | | |
| | | 9040085 | Apr 23, 2024 | U-1362 | | |
| | | 9186332 | Apr 23, 2024 | U-1362 | | |
| | | 9504699 | Aug 03, 2027 | U-1362 | | |
| <u>PREGABALIN - LYRICA</u> | | | | | | |
| N 021446 | 001 | 6001876*PED | Jun 30, 2019 | | M-193 | Dec 22, 2019 |
| | | 6197819*PED | Jun 30, 2019 | | NPP | May 03, 2021 |
| | | RE41920*PED | Jun 30, 2019 | | PED | Jun 22, 2020 |
| | | | | | PED | Nov 03, 2021 |
| <u>PREGABALIN - LYRICA</u> | | | | | | |
| N 021446 | 002 | 6001876*PED | Jun 30, 2019 | | M-193 | Dec 22, 2019 |
| | | 6197819*PED | Jun 30, 2019 | | NPP | May 03, 2021 |
| | | RE41920*PED | Jun 30, 2019 | | PED | Jun 22, 2020 |
| | | | | | PED | Nov 03, 2021 |
| <u>PREGABALIN - LYRICA</u> | | | | | | |
| N 021446 | 003 | 6001876*PED | Jun 30, 2019 | | M-193 | Dec 22, 2019 |
| | | 6197819*PED | Jun 30, 2019 | | NPP | May 03, 2021 |
| | | RE41920*PED | Jun 30, 2019 | | PED | Jun 22, 2020 |
| | | | | | PED | Nov 03, 2021 |
| <u>PREGABALIN - LYRICA</u> | | | | | | |
| N 021446 | 004 | 6001876*PED | Jun 30, 2019 | | M-193 | Dec 22, 2019 |
| | | 6197819*PED | Jun 30, 2019 | | NPP | May 03, 2021 |
| | | RE41920*PED | Jun 30, 2019 | | PED | Jun 22, 2020 |
| | | | | | PED | Nov 03, 2021 |
| <u>PREGABALIN - LYRICA</u> | | | | | | |
| N 021446 | 005 | 6001876*PED | Jun 30, 2019 | | M-193 | Dec 22, 2019 |
| | | 6197819*PED | Jun 30, 2019 | | NPP | May 03, 2021 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PREGABALIN - LYRICA</u> | | | | | | |
| N 021446 | 005 | RE41920*PED | | | | |
| | | Jun 30, 2019 | | | PED | Jun 22, 2020 |
| | | | | | PED | Nov 03, 2021 |
| <u>PREGABALIN - LYRICA</u> | | | | | | |
| N 021446 | 006 | 6001876*PED | | | | |
| | | Jun 30, 2019 | | | M-193 | Dec 22, 2019 |
| | | 6197819*PED | | | NPP | May 03, 2021 |
| | | Jun 30, 2019 | | | PED | Jun 22, 2020 |
| | | RE41920*PED | | | PED | Nov 03, 2021 |
| | | Jun 30, 2019 | | | | |
| <u>PREGABALIN - LYRICA</u> | | | | | | |
| N 021446 | 007 | 6001876*PED | | | | |
| | | Jun 30, 2019 | | | M-193 | Dec 22, 2019 |
| | | 6197819*PED | | | NPP | May 03, 2021 |
| | | Jun 30, 2019 | | | PED | Jun 22, 2020 |
| | | RE41920*PED | | | PED | Nov 03, 2021 |
| | | Jun 30, 2019 | | | | |
| <u>PREGABALIN - LYRICA</u> | | | | | | |
| N 021446 | 008 | 6001876*PED | | | | |
| | | Jun 30, 2019 | | | M-193 | Dec 22, 2019 |
| | | 6197819*PED | | | NPP | May 03, 2021 |
| | | Jun 30, 2019 | | | PED | Jun 22, 2020 |
| | | RE41920*PED | | | PED | Nov 03, 2021 |
| | | Jun 30, 2019 | | | | |
| <u>PREGABALIN - LYRICA</u> | | | | | | |
| N 022488 | 001 | 6001876*PED | | | | |
| | | Jun 30, 2019 | | | M-193 | Dec 22, 2019 |
| | | 6197819*PED | | | NPP | May 03, 2021 |
| | | Jun 30, 2019 | | | PED | Jun 22, 2020 |
| | | RE41920*PED | | | PED | Nov 03, 2021 |
| | | Jun 30, 2019 | | | | |
| <u>PREGABALIN - LYRICA CR</u> | | | | | | |
| N 209501 | 001 | 10022447 | | U-2136 | | |
| | | Nov 02, 2026 | | | NP | Oct 11, 2020 |
| | | 10022447 | | U-2137 | PED | Apr 11, 2021 |
| | | Nov 02, 2026 | | | | |
| | | 10022447*PED | | | | |
| | | May 02, 2027 | | | | |
| | | 6197819*PED | | | | |
| | | Jun 30, 2019 | | | | |
| | | 8945620 | | DP U-2136 | | |
| | | Nov 02, 2026 | | | | |
| | | 8945620 | | DP U-2137 | | |
| | | Nov 02, 2026 | | | | |
| | | 8945620*PED | | | | |
| | | May 02, 2027 | | | | |
| | | 9144559 | | DP | | |
| | | Nov 02, 2026 | | | | |
| | | 9144559*PED | | | | |
| | | May 02, 2027 | | | | |
| | | RE41920*PED | | | | |
| | | Jun 30, 2019 | | | | |
| <u>PREGABALIN - LYRICA CR</u> | | | | | | |
| N 209501 | 002 | 10022447 | | U-2136 | | |
| | | Nov 02, 2026 | | | NP | Oct 11, 2020 |
| | | 10022447 | | U-2137 | PED | Apr 11, 2021 |
| | | Nov 02, 2026 | | | | |
| | | 10022447*PED | | | | |
| | | May 02, 2027 | | | | |
| | | 6197819*PED | | | | |
| | | Jun 30, 2019 | | | | |
| | | 8945620 | | DP U-2136 | | |
| | | Nov 02, 2026 | | | | |
| | | 8945620 | | DP U-2137 | | |
| | | Nov 02, 2026 | | | | |
| | | 8945620*PED | | | | |
| | | May 02, 2027 | | | | |
| | | 9144559 | | DP | | |
| | | Nov 02, 2026 | | | | |
| | | 9144559*PED | | | | |
| | | May 02, 2027 | | | | |
| | | RE41920*PED | | | | |
| | | Jun 30, 2019 | | | | |
| <u>PREGABALIN - LYRICA CR</u> | | | | | | |
| N 209501 | 003 | 10022447 | | U-2136 | | |
| | | Nov 02, 2026 | | | NP | Oct 11, 2020 |
| | | 10022447 | | U-2137 | PED | Apr 11, 2021 |
| | | Nov 02, 2026 | | | | |
| | | 10022447*PED | | | | |
| | | May 02, 2027 | | | | |
| | | 6197819*PED | | | | |
| | | Jun 30, 2019 | | | | |
| | | 8945620 | | DP U-2136 | | |
| | | Nov 02, 2026 | | | | |
| | | 8945620 | | DP U-2137 | | |
| | | Nov 02, 2026 | | | | |
| | | 8945620*PED | | | | |
| | | May 02, 2027 | | | | |
| | | 9144559 | | DP | | |
| | | Nov 02, 2026 | | | | |
| | | 9144559*PED | | | | |
| | | May 02, 2027 | | | | |
| | | RE41920*PED | | | | |
| | | Jun 30, 2019 | | | | |
| <u>PROGESTERONE - ENDOMETRIN</u> | | | | | | |
| N 022057 | 001 | 7300664 | | U-856 | | |
| | | Nov 17, 2019 | | | | |
| | | 7320800 | | U-856 | | |
| | | Nov 17, 2019 | | | | |
| | | 7393543 | | DP U-880 | | |
| | | Nov 17, 2019 | | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>PROPOFOL - DIPRIVAN</u> | | | | | | |
| N 019627 | 002 8476010 | Dec 01, 2024 | DS DP | | | |
| | 8476010*PED | Jun 01, 2025 | | | | |
| <u>PROPRANOLOL HYDROCHLORIDE - INNOPRAN XL</u> | | | | | | |
| N 021438 | 001 6500454 | Oct 04, 2021 | DP | | | |
| <u>PROPRANOLOL HYDROCHLORIDE - INNOPRAN XL</u> | | | | | | |
| N 021438 | 002 6500454 | Oct 04, 2021 | DP | | | |
| <u>PROPRANOLOL HYDROCHLORIDE - HEMANGEOL</u> | | | | | | |
| N 205410 | 001 8338489 | Oct 16, 2028 | U-1496 | | ODE-62 | Mar 14, 2021 |
| <u>PRUCALOPRIDE SUCCINATE - MOTEGRITY</u> | | | | | | |
| N 210166 | 001 | | | | NCE | Dec 14, 2023 |
| <u>PRUCALOPRIDE SUCCINATE - MOTEGRITY</u> | | | | | | |
| N 210166 | 002 | | | | NCE | Dec 14, 2023 |
| <u>QUAZEPAM - DORAL</u> | | | | | | |
| N 018708 | 001 7608616 | Jun 03, 2028 | U-1012 | | | |
| <u>QUAZEPAM - DORAL</u> | | | | | | |
| N 018708 | 003 7608616 | Jun 03, 2028 | U-1012 | | | |
| <u>RADIUM RA-223 DICHLORIDE - XOFIGO</u> | | | | | | |
| N 203971 | 001 6635234 | Nov 17, 2022 | U-2271 | | | |
| <u>RALTEGRAVIR POTASSIUM - ISENTRESS</u> | | | | | | |
| N 022145 | 001 7169780 | Oct 03, 2023 | DS DP | | D-167 | May 26, 2020 |
| | 7169780*PED | Apr 03, 2024 | | | NPP | Nov 22, 2020 |
| | 7217713 | Oct 21, 2022 | U-257 | | PED | Nov 26, 2020 |
| | 7217713*PED | Apr 21, 2023 | | | PED | May 22, 2021 |
| | 7435734 | Oct 21, 2022 | U-257 | | | |
| | 7435734 | Oct 21, 2022 | U-900 | | | |
| | 7435734*PED | Apr 21, 2023 | | | | |
| | 7754731 | Mar 11, 2029 | DS DP U-257 | | | |
| | 7754731*PED | Sep 11, 2029 | | | | |
| <u>RALTEGRAVIR POTASSIUM - ISENTRESS HD</u> | | | | | | |
| N 022145 | 002 7169780 | Oct 03, 2023 | DS DP | | NPP | Nov 22, 2020 |
| | 7169780*PED | Apr 03, 2024 | | | NS | May 26, 2020 |
| | 7217713 | Oct 21, 2022 | U-257 | | PED | Nov 26, 2020 |
| | 7217713*PED | Apr 21, 2023 | | | PED | May 22, 2021 |
| | 7435734 | Oct 21, 2022 | U-257 | | | |
| | 7435734 | Oct 21, 2022 | U-900 | | | |
| | 7435734*PED | Apr 21, 2023 | | | | |
| | 7754731 | Mar 11, 2029 | DS DP U-257 | | | |
| | 7754731*PED | Sep 11, 2029 | | | | |
| | 9649311 | Oct 21, 2030 | DP | | | |
| | 9649311*PED | Apr 21, 2031 | | | | |
| <u>RALTEGRAVIR POTASSIUM - ISENTRESS</u> | | | | | | |
| N 203045 | 001 7169780 | Oct 03, 2023 | DS DP | | NPP | Nov 22, 2020 |
| | 7169780*PED | Apr 03, 2024 | | | PED | May 22, 2021 |
| | 7217713 | Oct 21, 2022 | U-257 | | | |
| | 7217713*PED | Apr 21, 2023 | | | | |
| | 7435734 | Oct 21, 2022 | U-257 | | | |
| | 7435734*PED | Apr 21, 2023 | | | | |
| | 7754731 | Mar 11, 2029 | DS DP U-257 | | | |
| | 7754731*PED | Sep 11, 2029 | | | | |
| <u>RALTEGRAVIR POTASSIUM - ISENTRESS</u> | | | | | | |
| N 203045 | 002 7169780 | Oct 03, 2023 | DS DP | | NPP | Nov 22, 2020 |
| | 7169780*PED | Apr 03, 2024 | | | PED | May 22, 2021 |
| | 7217713 | Oct 21, 2022 | U-257 | | | |
| | 7217713*PED | Apr 21, 2023 | | | | |
| | 7435734 | Oct 21, 2022 | U-257 | | | |
| | 7435734*PED | Apr 21, 2023 | | | | |
| | 7754731 | Mar 11, 2029 | DS DP U-257 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------|-----------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>RALTEGRAVIR POTASSIUM - ISENTRESS</u> | | | | | | |
| N 203045 | 002 7754731*PED | Sep 11, 2029 | | | | |
| <u>RALTEGRAVIR POTASSIUM - ISENTRESS</u> | | | | | | |
| N 205786 | 001 7169780 | Oct 03, 2023 | DS DP | | NPP | Nov 22, 2020 |
| | 7169780*PED | Apr 03, 2024 | | | PED | May 22, 2021 |
| | 7217713 | Oct 21, 2022 | U-257 | | | |
| | 7217713*PED | Apr 21, 2023 | | | | |
| | 7435734 | Oct 21, 2022 | U-257 | | | |
| | 7435734*PED | Apr 21, 2023 | | | | |
| | 7754731 | Mar 11, 2029 | DS DP U-257 | | | |
| | 7754731*PED | Sep 11, 2029 | | | | |
| <u>RAMELTEON - ROZEREM</u> | | | | | | |
| N 021782 | 001 10098866 | Nov 16, 2021 | | DP U-2433 | | |
| | 6034239 | Jul 22, 2019 | DS DP | U-674 | | |
| <u>RAMIPRIL - ALTACE</u> | | | | | | |
| N 019901 | 001 7368469 | Aug 30, 2020 | | U-871 | | |
| <u>RAMIPRIL - ALTACE</u> | | | | | | |
| N 019901 | 002 7368469 | Aug 30, 2020 | | U-871 | | |
| <u>RAMIPRIL - ALTACE</u> | | | | | | |
| N 019901 | 003 7368469 | Aug 30, 2020 | | U-871 | | |
| <u>RAMIPRIL - ALTACE</u> | | | | | | |
| N 019901 | 004 7368469 | Aug 30, 2020 | | U-871 | | |
| <u>RAMIPRIL - ALTACE</u> | | | | | | |
| N 022021 | 001 7368469 | Aug 30, 2020 | | U-871 | | |
| <u>RAMIPRIL - ALTACE</u> | | | | | | |
| N 022021 | 002 7368469 | Aug 30, 2020 | | U-871 | | |
| <u>RAMIPRIL - ALTACE</u> | | | | | | |
| N 022021 | 003 7368469 | Aug 30, 2020 | | U-871 | | |
| <u>RAMIPRIL - ALTACE</u> | | | | | | |
| N 022021 | 004 7368469 | Aug 30, 2020 | | U-871 | | |
| <u>RANOLAZINE - RANEXA</u> | | | | | | |
| N 021526 | 001 6303607 | May 27, 2019 | | U-705 | | |
| | 6369062 | May 27, 2019 | DP | | Y | |
| | 6479496 | May 27, 2019 | U-705 | | | |
| | 6503911 | May 27, 2019 | DP | | | |
| | 6525057 | May 27, 2019 | U-705 | | | |
| | 6562826 | May 27, 2019 | U-705 | | | |
| | 6617328 | May 27, 2019 | DP | | | |
| | 6620814 | May 27, 2019 | U-705 | | | |
| | 6852724 | May 27, 2019 | U-705 | | | |
| | 6864258 | May 27, 2019 | U-705 | | | |
| <u>RANOLAZINE - RANEXA</u> | | | | | | |
| N 021526 | 002 6303607 | May 27, 2019 | | U-705 | | |
| | 6369062 | May 27, 2019 | DP | | | |
| | 6479496 | May 27, 2019 | U-705 | | | |
| | 6503911 | May 27, 2019 | DP | | | |
| | 6525057 | May 27, 2019 | U-705 | | | |
| | 6562826 | May 27, 2019 | U-705 | | | |
| | 6617328 | May 27, 2019 | DP | | | |
| | 6620814 | May 27, 2019 | U-705 | | | |
| | 6852724 | May 27, 2019 | U-705 | | | |
| | 6864258 | May 27, 2019 | U-705 | | | |
| <u>RASAGILINE MESYLATE - AZILECT</u> | | | | | | |
| N 021641 | 001 7572834 | Dec 05, 2026 | | DP | | |
| | 7815942 | Aug 27, 2027 | DS DP | U-219 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>RASAGILINE MESYLATE - AZILECT</u> | | | | | | |
| N 021641 002 | 7572834 | Dec 05, 2026 | DP | | | |
| | 7815942 | Aug 27, 2027 | DS DP U-219 | | | |
| <u>REGADENOSON - LEXISCAN</u> | | | | | | |
| N 022161 001 | 6403567 | Apr 10, 2022 | DS DP U-869 | | M-194 | Jan 17, 2020 |
| | 6642210 | Jun 22, 2019 | DS DP U-869 | | | |
| | 7144872 | Jun 22, 2019 | DS DP U-116 | | | |
| | 7144872 | Jun 22, 2019 | DS DP U-869 | | | |
| | 7144872 | Jun 22, 2019 | DS DP U-870 | | | |
| | 7183264 | Jun 22, 2019 | DP U-116 | | | |
| | 7183264 | Jun 22, 2019 | DP U-869 | | | |
| | 7183264 | Jun 22, 2019 | DP U-870 | | | |
| | 7582617 | Jun 22, 2019 | U-1003 | | | |
| | 7655636 | Jun 22, 2019 | U-869 | | | |
| | 7655637 | Jun 22, 2019 | DS DP U-869 | | | |
| | 7683037 | Jun 22, 2019 | U-1042 | | | |
| | 8106029 | Jun 22, 2019 | U-1042 | | | |
| | 8106183 | Feb 02, 2027 | DS | | | |
| | 8133879 | Jun 22, 2019 | DP | | | |
| | 8183226 | Jun 22, 2019 | U-116 | | | |
| | 8470801 | Jun 22, 2019 | U-116 | | | |
| | 8536150 | Jun 22, 2019 | U-116 | | | |
| | 9045519 | Jun 22, 2019 | DP | | | |
| | 9085601 | Feb 02, 2027 | DP | | | |
| | 9289446 | Jun 22, 2019 | DP U-116 | | | |
| <u>REGORAFENIB - STIVARGA</u> | | | | | | |
| N 203085 001 | 7351834 | Jun 28, 2022 | DS | | I-744 | Apr 27, 2020 |
| | 8637553 | Feb 16, 2031 | DS DP | | ODE-139 | Apr 27, 2024 |
| | 8680124 | Jun 02, 2030 | U-1506 | | ODE-44 | Feb 25, 2020 |
| | 9458107 | Apr 08, 2031 | DP | | | |
| | 9957232 | Jul 09, 2032 | DS | | | |
| <u>RETAPAMULIN - ALTABAX</u> | | | | | | |
| N 022055 001 | 7875630 | Feb 14, 2027 | DS | | | |
| | 8207191 | Aug 30, 2024 | U-805 | | | |
| | RE43390 | Apr 12, 2021 | DS DP U-805 | | | |
| <u>REVEFENACIN - YUPELRI</u> | | | | | | |
| N 210598 001 | 10106503 | Mar 10, 2025 | U-2440 | | NCE | Nov 09, 2023 |
| | 7288657 | Dec 23, 2025 | DS | | | |
| | 7491736 | Mar 10, 2025 | U-2440 | | | |
| | 7521041 | Mar 10, 2025 | U-2440 | | | |
| | 7550595 | Mar 10, 2025 | DP | | | |
| | 7585879 | Mar 10, 2025 | DS DP U-2440 | | | |
| | 7910608 | Mar 10, 2025 | DS DP | | | |
| | 8034946 | Mar 10, 2025 | DP | | | |
| | 8053448 | Mar 10, 2025 | U-2440 | | | |
| | 8273894 | Mar 10, 2025 | DP | | | |
| <u>RIBAVIRIN - REBETOL</u> | | | | | | |
| N 021546 001 | 6790837 | Apr 05, 2023 | DP | | | |
| <u>RIBOCICLIB SUCCINATE - KISOALI</u> | | | | | | |
| N 209092 001 | 8324225 | Jun 17, 2028 | DS DP | | I-783 | Jul 18, 2021 |
| | 8415355 | Feb 19, 2031 | DS DP | | I-784 | Jul 18, 2021 |
| | 8685980 | May 25, 2030 | DS DP | | NCE | Mar 13, 2022 |
| | 8962630 | Dec 09, 2029 | U-1981 | | | |
| | 8962630 | Dec 09, 2029 | U-2355 | | | |
| | 8962630 | Dec 09, 2029 | U-2356 | | | |
| | 9193732 | Nov 09, 2031 | DS DP | | | |
| | 9416136 | Aug 20, 2029 | U-1981 | | | |
| | 9416136 | Aug 20, 2029 | U-2355 | | | |
| | 9416136 | Aug 20, 2029 | U-2356 | | | |
| | 9868739 | Nov 09, 2031 | U-1981 | | | |
| | 9868739 | Nov 09, 2031 | U-2355 | | | |
| | 9868739 | Nov 09, 2031 | U-2356 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>RIBOFLAVIN 5'-PHOSPHATE SODIUM - PHOTREXA</u> | | | | | | |
| N 203324 | 001 | | | | NP | Apr 15, 2019 |
| | | | | | ODE-116 | Apr 15, 2023 |
| | | | | | ODE-121 | Jul 15, 2023 |
| <u>RIBOFLAVIN 5'-PHOSPHATE SODIUM - PHOTREXA VISCOUS IN DEXTRAN 20%</u> | | | | | | |
| N 203324 | 002 | | | | NP | Apr 15, 2019 |
| | | | | | ODE-116 | Apr 15, 2023 |
| | | | | | ODE-121 | Jul 15, 2023 |
| <u>RIFAMYCIN - AEMCOLO</u> | | | | | | |
| N 210910 | 001 | 8263120 | May 03, 2025 | DP | | |
| | | 8486446 | May 03, 2025 | DP | | |
| | | 8529945 | May 03, 2025 | DP | | |
| | | 8741948 | May 03, 2025 | DP | U-2448 | |
| <u>RIFAXIMIN - XIFAXAN</u> | | | | | | |
| N 021361 | 001 | 7045620 | Jun 19, 2024 | DS DP | | |
| | | 7612199 | Jun 19, 2024 | DS DP | | |
| | | 7902206 | Jun 19, 2024 | DS DP | | |
| | | 7906542 | Jun 01, 2025 | DS DP | | |
| | | 7928115 | Jul 24, 2029 | | U-1121 | |
| | | 8158644 | Jun 19, 2024 | DP | | |
| | | 8158781 | Jun 19, 2024 | DS | | |
| | | 8193196 | Sep 02, 2027 | DS DP | | |
| | | 8518949 | Feb 27, 2026 | DP | | |
| | | 8741904 | Feb 27, 2026 | DS | U-1526 | |
| | | 8835452 | Jun 19, 2024 | DS DP | | |
| | | 8853231 | Jun 19, 2024 | DP | | |
| | | 9271968 | Feb 27, 2026 | DP | | |
| <u>RIFAXIMIN - XIFAXAN</u> | | | | | | |
| N 022554 | 001 | 6861053 | Aug 11, 2019 | | U-1707 | |
| | | 6861053 | Aug 11, 2019 | | U-1708 | |
| | | 7045620 | Jun 19, 2024 | DS | | |
| | | 7452857 | Aug 11, 2019 | | U-1707 | |
| | | 7452857 | Aug 11, 2019 | | U-1708 | |
| | | 7605240 | Aug 11, 2019 | | U-1707 | |
| | | 7605240 | Aug 11, 2019 | | U-1708 | |
| | | 7612199 | Jun 19, 2024 | DS DP | | |
| | | 7718608 | Aug 11, 2019 | | U-1707 | |
| | | 7718608 | Aug 11, 2019 | | U-1708 | |
| | | 7902206 | Jun 19, 2024 | DS DP | | |
| | | 7906542 | Jun 01, 2025 | DS DP | | |
| | | 7915275 | Feb 23, 2025 | | U-1707 | |
| | | 7915275 | Feb 23, 2025 | | U-1708 | |
| | | 7935799 | Aug 11, 2019 | | U-1707 | |
| | | 7935799 | Aug 11, 2019 | | U-1708 | |
| | | 8158644 | Jun 19, 2024 | DP | | |
| | | 8158781 | Jun 19, 2024 | DS | | |
| | | 8193196 | Sep 02, 2027 | DS DP | U-1707 | |
| | | 8193196 | Sep 02, 2027 | DS DP | U-1708 | |
| | | 8309569 | Jul 18, 2029 | | U-1707 | |
| | | 8309569 | Jul 18, 2029 | | U-1708 | |
| | | 8518949 | Feb 27, 2026 | DP | | |
| | | 8642573 | Oct 02, 2029 | | U-1481 | |
| | | 8741904 | Feb 27, 2026 | DS | U-1526 | |
| | | 8741904 | Feb 27, 2026 | DS | U-1707 | |
| | | 8741904 | Feb 27, 2026 | DS | U-1708 | |
| | | 8829017 | Jul 24, 2029 | | U-1562 | |
| | | 8835452 | Jun 19, 2024 | DS DP | | |
| | | 8853231 | Jun 19, 2024 | DP | | |
| | | 8946252 | Jul 24, 2029 | | U-1481 | |
| | | 8969398 | Oct 02, 2029 | | U-1481 | |
| | | 9271968 | Feb 27, 2026 | DP | | |
| | | 9421195 | Mar 10, 2030 | | U-1481 | |
| | | 9629828 | Jul 24, 2029 | | U-1994 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>RILPIVIRINE HYDROCHLORIDE - EDURANT</u> | | | | | | |
| N 202022 001 | 6838464 | Feb 26, 2021 | DS DP | | M-223 | Feb 01, 2021 |
| | 7067522 | Dec 20, 2019 | DS DP | | | |
| | 7125879 | Apr 21, 2025 | DS DP U-1153 | | | |
| | 7125879 | Apr 21, 2025 | DS DP U-1307 | | | |
| | 7125879 | Apr 21, 2025 | DS DP U-1740 | | | |
| | 7638522 | Apr 14, 2023 | DP | | | |
| | 8080551 | Apr 11, 2023 | DS DP | | | |
| | 8101629 | Aug 09, 2022 | DP | | | |
| <u>RILUZOLE - TIGLUTIK KIT</u> | | | | | | |
| N 209080 001 | 8765150 | Mar 12, 2029 | DP U-2401 | | | |
| <u>RIOCIGUAT - ADEMPAS</u> | | | | | | |
| N 204819 001 | 6743798 | Jul 16, 2019 | DS DP | | ODE-53 | Oct 08, 2020 |
| | 7173037 | Dec 04, 2026 | DS DP | | | |
| <u>RIOCIGUAT - ADEMPAS</u> | | | | | | |
| N 204819 002 | 6743798 | Jul 16, 2019 | DS DP | | ODE-53 | Oct 08, 2020 |
| | 7173037 | Dec 04, 2026 | DS DP | | | |
| <u>RIOCIGUAT - ADEMPAS</u> | | | | | | |
| N 204819 003 | 6743798 | Jul 16, 2019 | DS DP | | ODE-53 | Oct 08, 2020 |
| | 7173037 | Dec 04, 2026 | DS DP | | | |
| <u>RIOCIGUAT - ADEMPAS</u> | | | | | | |
| N 204819 004 | 6743798 | Jul 16, 2019 | DS DP | | ODE-53 | Oct 08, 2020 |
| | 7173037 | Dec 04, 2026 | DS DP | | | |
| <u>RIOCIGUAT - ADEMPAS</u> | | | | | | |
| N 204819 005 | 6743798 | Jul 16, 2019 | DS DP | | ODE-53 | Oct 08, 2020 |
| | 7173037 | Dec 04, 2026 | DS DP | | | |
| <u>RISEDRONATE SODIUM - ACTONEL</u> | | | | | | |
| N 020835 005 | 7192938 | May 06, 2023 | | U-353 | | |
| | 7718634 | May 06, 2023 | | U-662 | | |
| <u>RISEDRONATE SODIUM - ATELVIA</u> | | | | | | |
| N 022560 001 | 7645459 | Jan 09, 2028 | DP U-662 | | | |
| | 7645460 | Jan 09, 2028 | DP U-662 | | | |
| | 8246989 | Jan 16, 2026 | DP | | | |
| <u>RISPERIDONE - RISPERDAL CONSTA</u> | | | | | | |
| N 021346 001 | 6667061 | May 25, 2020 | DP | | | |
| <u>RISPERIDONE - RISPERDAL CONSTA</u> | | | | | | |
| N 021346 002 | 6667061 | May 25, 2020 | DP | | | |
| <u>RISPERIDONE - RISPERDAL CONSTA</u> | | | | | | |
| N 021346 003 | 6667061 | May 25, 2020 | DP | | | |
| <u>RISPERIDONE - PERSERIS KIT</u> | | | | | | |
| N 210655 001 | 10010612 | Feb 13, 2028 | DP | | NP | Jul 27, 2021 |
| | 10058554 | Sep 26, 2026 | U-2363 | | | |
| | 9180197 | Feb 13, 2028 | DP | | | |
| | 9186413 | Feb 13, 2028 | U-543 | | | |
| | 9597402 | Sep 26, 2026 | DP | | | |
| <u>RISPERIDONE - PERSERIS KIT</u> | | | | | | |
| N 210655 002 | 10010612 | Feb 13, 2028 | DP | | NP | Jul 27, 2021 |
| | 10058554 | Sep 26, 2026 | U-2363 | | | |
| | 9180197 | Feb 13, 2028 | DP | | | |
| | 9186413 | Feb 13, 2028 | U-543 | | | |
| | 9597402 | Sep 26, 2026 | DP | | | |
| <u>RITONAVIR - NORVIR</u> | | | | | | |
| N 020945 001 | 7141593 | May 22, 2020 | DP | | | |
| | 7432294 | May 22, 2020 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>RITONAVIR - NORVIR</u> | | | | | | |
| N 022417 001 | 7148359 | Jul 19, 2019 | DP | | | |
| | 7364752 | Nov 10, 2020 | DP U-688 | | | |
| | 8268349 | Aug 25, 2024 | DP | | | |
| | 8399015 | Aug 25, 2024 | DP | | | |
| | 8399015*PED | Feb 25, 2025 | | | | |
| | 8470347 | Sep 17, 2026 | DP | | | |
| | 8470347*PED | Mar 17, 2027 | | | | |
| | 8691878 | Aug 25, 2024 | U-688 | | | |
| | 8691878*PED | Feb 25, 2025 | | | | |
| <u>RITONAVIR - NORVIR</u> | | | | | | |
| N 209512 001 | | | | | ODE-184 | Jun 07, 2024 |
| <u>RIVAROXABAN - XARELTO</u> | | | | | | |
| N 022406 001 | 7157456 | Aug 28, 2024 | DS DP U-1301 | | D-168 | Oct 27, 2020 |
| | 7157456 | Aug 28, 2024 | DS DP U-1302 | | | |
| | 7585860 | Dec 11, 2020 | DS | | | |
| | 7592339 | Dec 11, 2020 | U-1167 | | | |
| | 7592339 | Dec 11, 2020 | U-1200 | | | |
| | 7592339 | Dec 11, 2020 | U-1301 | | | |
| | 7592339 | Dec 11, 2020 | U-1302 | | | |
| | 7592339 | Dec 11, 2020 | U-1303 | | | |
| | 7592339 | Dec 11, 2020 | U-2142 | | | |
| | 9415053 | Nov 13, 2024 | DP U-1167 | | | |
| | 9415053 | Nov 13, 2024 | DP U-1200 | | | |
| | 9415053 | Nov 13, 2024 | DP U-1301 | | | |
| | 9415053 | Nov 13, 2024 | DP U-1302 | | | |
| | 9415053 | Nov 13, 2024 | DP U-2142 | | | |
| | 9539218 | Feb 17, 2034 | U-1953 | | | |
| | 9539218 | Feb 17, 2034 | U-1954 | | | |
| | 9539218 | Feb 17, 2034 | U-1955 | | | |
| | 9539218 | Feb 17, 2034 | U-1957 | | | |
| | 9539218 | Feb 17, 2034 | U-2143 | | | |
| <u>RIVAROXABAN - XARELTO</u> | | | | | | |
| N 022406 002 | 7157456 | Aug 28, 2024 | DS DP U-1301 | | | |
| | 7157456 | Aug 28, 2024 | DS DP U-1302 | | | |
| | 7585860 | Dec 11, 2020 | DS | | | |
| | 7592339 | Dec 11, 2020 | U-1167 | | | |
| | 7592339 | Dec 11, 2020 | U-1200 | | | |
| | 7592339 | Dec 11, 2020 | U-1301 | | | |
| | 7592339 | Dec 11, 2020 | U-1302 | | | |
| | 7592339 | Dec 11, 2020 | U-1303 | | | |
| | 9415053 | Nov 13, 2024 | DP U-1167 | | | |
| | 9415053 | Nov 13, 2024 | DP U-1200 | | | |
| | 9415053 | Nov 13, 2024 | DP U-1301 | | | |
| | 9415053 | Nov 13, 2024 | DP U-1302 | | | |
| | 9415053 | Nov 13, 2024 | DP U-1303 | | | |
| | 9539218 | Feb 17, 2034 | U-1953 | | | |
| | 9539218 | Feb 17, 2034 | U-1954 | | | |
| | 9539218 | Feb 17, 2034 | U-1955 | | | |
| | 9539218 | Feb 17, 2034 | U-1956 | | | |
| | 9539218 | Feb 17, 2034 | U-1957 | | | |
| <u>RIVAROXABAN - XARELTO</u> | | | | | | |
| N 022406 003 | 7157456 | Aug 28, 2024 | DS DP U-1301 | | | |
| | 7157456 | Aug 28, 2024 | DS DP U-1302 | | | |
| | 7585860 | Dec 11, 2020 | DS | | | |
| | 7592339 | Dec 11, 2020 | U-1167 | | | |
| | 7592339 | Dec 11, 2020 | U-1200 | | | |
| | 7592339 | Dec 11, 2020 | U-1301 | | | |
| | 7592339 | Dec 11, 2020 | U-1302 | | | |
| | 7592339 | Dec 11, 2020 | U-1303 | | | |
| | 9415053 | Nov 13, 2024 | DP U-1167 | | | |
| | 9415053 | Nov 13, 2024 | DP U-1200 | | | |
| | 9415053 | Nov 13, 2024 | DP U-1301 | | | |
| | 9415053 | Nov 13, 2024 | DP U-1302 | | | |
| | 9539218 | Feb 17, 2034 | U-1953 | | | |
| | 9539218 | Feb 17, 2034 | U-1954 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>RIVAROXABAN - XARELTO</u> | | | | | | |
| N 022406 | 003 | 9539218 | Feb 17, 2034 | U-1955 | | |
| | | 9539218 | Feb 17, 2034 | U-1957 | | |
| <u>RIVAROXABAN - XARELTO</u> | | | | | | |
| N 022406 | 004 | 7157456 | Aug 28, 2024 | DS DP | | |
| | | 7585860 | Dec 11, 2020 | DS | | |
| | | 7592339 | Dec 11, 2020 | U-2435 | | |
| | | 9415053 | Nov 13, 2024 | DP U-2435 | | |
| <u>RIVASTIGMINE - EXELON</u> | | | | | | |
| N 022083 | 001 | 6316023 | Jan 08, 2019 | DP | | |
| | | 6335031 | Jan 08, 2019 | DP | | |
| <u>RIVASTIGMINE - EXELON</u> | | | | | | |
| N 022083 | 002 | 6316023 | Jan 08, 2019 | DP | | |
| | | 6335031 | Jan 08, 2019 | DP | | |
| <u>RIVASTIGMINE - EXELON</u> | | | | | | |
| N 022083 | 005 | 6316023 | Jan 08, 2019 | DP | | |
| | | 6335031 | Jan 08, 2019 | DP | | |
| <u>ROFLUMILAST - DALIRESP</u> | | | | | | |
| N 022522 | 001 | 5712298 | Jan 27, 2020 | DS DP U-1115 | D-171 | Jan 23, 2021 |
| | | 8431154 | Feb 19, 2023 | DP | M-208 | Aug 31, 2020 |
| | | 8536206 | Mar 08, 2024 | U-1115 | | |
| | | 8604064 | Mar 08, 2024 | U-1115 | | |
| | | 8618142 | Mar 08, 2024 | DP | | |
| | | 9468598 | Feb 19, 2023 | DP | | |
| <u>ROFLUMILAST - DALIRESP</u> | | | | | | |
| N 022522 | 002 | 5712298 | Jan 27, 2020 | DS DP U-1115 | NS | Jan 23, 2021 |
| | | 8431154 | Feb 19, 2023 | DP | | |
| | | 8536206 | Mar 08, 2024 | U-1115 | | |
| | | 8604064 | Mar 08, 2024 | U-1115 | | |
| | | 8618142 | Mar 08, 2024 | DP | | |
| | | 9468598 | Feb 19, 2023 | DP | | |
| <u>ROLAPITANT HYDROCHLORIDE - VARUBI</u> | | | | | | |
| N 206500 | 001 | 7049320 | Dec 08, 2023 | DS DP U-1741 | NCE | Sep 01, 2020 |
| | | 7563801 | Apr 04, 2027 | DP | | |
| | | 7981905 | Apr 04, 2027 | U-1741 | | |
| | | 8178550 | Apr 04, 2027 | DS DP | | |
| | | 8361500 | Oct 09, 2029 | DP | | |
| | | 8404702 | Apr 04, 2027 | U-1741 | | |
| | | 8470842 | Jan 18, 2029 | U-1741 | | |
| | | 8796299 | Dec 17, 2022 | U-1741 | | |
| <u>ROLAPITANT HYDROCHLORIDE - VARUBI</u> | | | | | | |
| N 208399 | 001 | 7049320 | Dec 08, 2023 | DS DP U-1741 | NCE | Sep 01, 2020 |
| | | 7981905 | Apr 04, 2027 | U-1741 | | |
| | | 8178550 | Apr 04, 2027 | DS DP | | |
| | | 8404702 | Apr 04, 2027 | U-1741 | | |
| | | 8470842 | Jan 18, 2029 | U-1741 | | |
| | | 8796299 | Dec 17, 2022 | U-1741 | | |
| | | 9101615 | Jul 14, 2032 | U-1741 | | |
| <u>ROMIDEPSIN - ISTODAX</u> | | | | | | |
| N 022393 | 001 | 7608280 | Aug 22, 2021 | DS | | |
| | | 7611724 | Aug 22, 2021 | DS | | |
| <u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u> | | | | | | |
| N 022008 | 001 | 7927624 | Dec 02, 2021 | DP U-20 | M-203 | Mar 23, 2020 |
| | | 8303986 | Apr 12, 2021 | DP | | |
| <u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u> | | | | | | |
| N 022008 | 002 | 7927624 | Dec 02, 2021 | DP U-20 | M-203 | Mar 23, 2020 |
| | | 8303986 | Apr 12, 2021 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u> | | | | | | |
| N 022008 003 | 7927624 | Dec 02, 2021 | DP U-20 | | M-203 | Mar 23, 2020 |
| | 8303986 | Apr 12, 2021 | DP | | | |
| <u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u> | | | | | | |
| N 022008 004 | 7927624 | Dec 02, 2021 | DP U-20 | | M-203 | Mar 23, 2020 |
| | 8303986 | Apr 12, 2021 | DP | | | |
| <u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u> | | | | | | |
| N 022008 005 | 7927624 | Dec 02, 2021 | DP U-20 | | M-203 | Mar 23, 2020 |
| | 8303986 | Apr 12, 2021 | DP | | | |
| <u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u> | | | | | | |
| N 022008 006 | 7927624 | Dec 02, 2021 | DP U-20 | | M-203 | Mar 23, 2020 |
| | 8303986 | Apr 12, 2021 | DP | | | |
| <u>ROPIVACAINE HYDROCHLORIDE - NAROPIN</u> | | | | | | |
| N 020533 006 | 7828787 | Oct 18, 2025 | DP | | | |
| | 7857802 | Nov 28, 2026 | DP | | | |
| | 8118802 | May 18, 2023 | DP | | | |
| | 8162915 | May 23, 2024 | DP | | | |
| <u>ROPIVACAINE HYDROCHLORIDE - NAROPIN</u> | | | | | | |
| N 020533 007 | 7828787 | Oct 18, 2025 | DP | | | |
| | 7857802 | Nov 28, 2026 | DP | | | |
| | 8118802 | May 18, 2023 | DP | | | |
| | 8162915 | May 23, 2024 | DP | | | |
| <u>ROSIGLITAZONE MALEATE - AVANDIA</u> | | | | | | |
| N 021071 002 | 7358366 | Apr 19, 2020 | DS | | Y | |
| | 7358366*PED | Oct 19, 2020 | | | | |
| <u>ROSIGLITAZONE MALEATE - AVANDIA</u> | | | | | | |
| N 021071 003 | 7358366 | Apr 19, 2020 | DS | | Y | |
| | 7358366*PED | Oct 19, 2020 | | | | |
| <u>ROSIGLITAZONE MALEATE - AVANDIA</u> | | | | | | |
| N 021071 004 | 7358366 | Apr 19, 2020 | DS | | Y | |
| | 7358366*PED | Oct 19, 2020 | | | | |
| <u>ROSUVASTATIN CALCIUM - CRESTOR</u> | | | | | | |
| N 021366 002 | 6316460 | Aug 04, 2020 | DP | | ODE-118 | May 27, 2023 |
| | 6858618 | Dec 17, 2021 | U-1032 | | | |
| | 6858618 | Dec 17, 2021 | U-1807 | | | |
| | 6858618 | Dec 17, 2021 | U-618 | | | |
| | 6858618*PED | Jun 17, 2022 | | | | |
| <u>ROSUVASTATIN CALCIUM - CRESTOR</u> | | | | | | |
| N 021366 003 | 6316460 | Aug 04, 2020 | DP | | ODE-118 | May 27, 2023 |
| | 6858618 | Dec 17, 2021 | U-1032 | | | |
| | 6858618 | Dec 17, 2021 | U-1807 | | | |
| | 6858618 | Dec 17, 2021 | U-618 | | | |
| | 6858618*PED | Jun 17, 2022 | | | | |
| <u>ROSUVASTATIN CALCIUM - CRESTOR</u> | | | | | | |
| N 021366 004 | 6316460 | Aug 04, 2020 | DP | | I-732 | May 27, 2019 |
| | 6858618 | Dec 17, 2021 | U-1032 | | ODE-118 | May 27, 2023 |
| | 6858618 | Dec 17, 2021 | U-1807 | | | |
| | 6858618 | Dec 17, 2021 | U-618 | | | |
| | 6858618*PED | Jun 17, 2022 | | | | |
| <u>ROSUVASTATIN CALCIUM - CRESTOR</u> | | | | | | |
| N 021366 005 | 6316460 | Aug 04, 2020 | DP | | ODE-118 | May 27, 2023 |
| | 6858618 | Dec 17, 2021 | U-618 | | | |
| <u>ROTIGOTINE - NEUPRO</u> | | | | | | |
| N 021829 001 | 10130589 | Dec 22, 2030 | DP | | | |
| | 6699498 | Nov 27, 2020 | DP | | | |
| | 6884434 | Mar 30, 2021 | DP | | | |
| | 7413747 | Mar 18, 2019 | DP | | | |
| | 8246979 | Sep 01, 2027 | DP U-1272 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------|-----------|------------------------|--------------|-------------------------|---------------------|-----------------------------|
| <u>ROTIGOTINE - NEUPRO</u> | | | | | | |
| N 021829 001 | 8246979 | Sep 01, 2027 | DP U-1273 | | | |
| | 8246980 | Nov 27, 2025 | DP | | | |
| | 8617591 | Jul 22, 2023 | DP U-1474 | | | |
| | 9925150 | Mar 01, 2032 | DP | | | |
| <u>ROTIGOTINE - NEUPRO</u> | | | | | | |
| N 021829 002 | 10130589 | Dec 22, 2030 | DP | | | |
| | 6699498 | Nov 27, 2020 | DP | | | |
| | 6884434 | Mar 30, 2021 | DP | | | |
| | 7413747 | Mar 18, 2019 | DP | | | |
| | 8246979 | Sep 01, 2027 | DP U-1272 | | | |
| | 8246979 | Sep 01, 2027 | DP U-1273 | | | |
| | 8246980 | Nov 27, 2025 | DP | | | |
| | 8617591 | Jul 22, 2023 | DP U-1474 | | | |
| | 9925150 | Mar 01, 2032 | DP | | | |
| <u>ROTIGOTINE - NEUPRO</u> | | | | | | |
| N 021829 003 | 10130589 | Dec 22, 2030 | DP | | | |
| | 6699498 | Nov 27, 2020 | DP | | | |
| | 6884434 | Mar 30, 2021 | DP | | | |
| | 7413747 | Mar 18, 2019 | DP | | | |
| | 8246979 | Sep 01, 2027 | DP U-1272 | | | |
| | 8246979 | Sep 01, 2027 | DP U-1273 | | | |
| | 8246980 | Nov 27, 2025 | DP | | | |
| | 8617591 | Jul 22, 2023 | DP U-1474 | | | |
| | 9925150 | Mar 01, 2032 | DP | | | |
| <u>ROTIGOTINE - NEUPRO</u> | | | | | | |
| N 021829 004 | 10130589 | Dec 22, 2030 | DP | | | |
| | 6699498 | Nov 27, 2020 | DP | | | |
| | 6884434 | Mar 30, 2021 | DP | | | |
| | 7413747 | Mar 18, 2019 | DP | | | |
| | 8246979 | Sep 01, 2027 | DP U-1272 | | | |
| | 8246979 | Sep 01, 2027 | DP U-1273 | | | |
| | 8246980 | Nov 27, 2025 | DP | | | |
| | 8617591 | Jul 22, 2023 | DP U-1474 | | | |
| | 9925150 | Mar 01, 2032 | DP | | | |
| <u>ROTIGOTINE - NEUPRO</u> | | | | | | |
| N 021829 005 | 10130589 | Dec 22, 2030 | DP | | | |
| | 6699498 | Nov 27, 2020 | DP | | | |
| | 6884434 | Mar 30, 2021 | DP | | | |
| | 7413747 | Mar 18, 2019 | DP | | | |
| | 8246979 | Sep 01, 2027 | DP U-1272 | | | |
| | 8246979 | Sep 01, 2027 | DP U-1273 | | | |
| | 8246980 | Nov 27, 2025 | DP | | | |
| | 8617591 | Jul 22, 2023 | DP U-1474 | | | |
| | 9925150 | Mar 01, 2032 | DP | | | |
| <u>ROTIGOTINE - NEUPRO</u> | | | | | | |
| N 021829 006 | 10130589 | Dec 22, 2030 | DP | | | |
| | 6699498 | Nov 27, 2020 | DP | | | |
| | 6884434 | Mar 30, 2021 | DP | | | |
| | 7413747 | Mar 18, 2019 | DP | | | |
| | 8246979 | Sep 01, 2027 | DP U-1272 | | | |
| | 8246979 | Sep 01, 2027 | DP U-1273 | | | |
| | 8246980 | Nov 27, 2025 | DP | | | |
| | 8617591 | Jul 22, 2023 | DP U-1474 | | | |
| | 9925150 | Mar 01, 2032 | DP | | | |
| <u>RUCAPARIB CAMSYLATE - RUBRACA</u> | | | | | | |
| N 209115 001 | 10130636 | Aug 17, 2035 | U-2012 | | I-772 | Apr 06, 2021 |
| | 10130636 | Aug 17, 2035 | U-2101 | | NCE | Dec 19, 2021 |
| | 10130636 | Aug 17, 2035 | U-2273 | | ODE-126 | Dec 19, 2023 |
| | 6495541 | Jan 10, 2020 | DS DP | | ODE-168 | Apr 06, 2025 |
| | 7351701 | Jul 23, 2024 | U-2012 | | | |
| | 7351701 | Jul 23, 2024 | U-2101 | | | |
| | 7351701 | Jul 23, 2024 | U-2273 | | | |
| | 7531530 | Jul 23, 2024 | U-2012 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>RUCAPARIB CAMSYLATE - RUBRACA</u> | | | | | | |
| N 209115 001 | 7531530 | Jul 23, 2024 | U-2101 | | | |
| | 7531530 | Jul 23, 2024 | U-2273 | | | |
| | 8071579 | Aug 12, 2027 | U-2012 | | | |
| | 8071579 | Aug 12, 2027 | U-2101 | | | |
| | 8071579 | Aug 12, 2027 | U-2273 | | | |
| | 8143241 | Aug 12, 2027 | U-2012 | | | |
| | 8143241 | Aug 12, 2027 | U-2101 | | | |
| | 8143241 | Aug 12, 2027 | U-2273 | | | |
| | 8754072 | Feb 10, 2031 | DS DP | | | |
| | 8859562 | Aug 04, 2031 | U-2012 | | | |
| | 8859562 | Aug 04, 2031 | U-2101 | | | |
| | 8859562 | Aug 04, 2031 | U-2273 | | | |
| | 9045487 | Feb 10, 2031 | DS DP | | | |
| | 9861638 | Feb 10, 2031 | U-2012 | | | |
| | 9861638 | Feb 10, 2031 | U-2101 | | | |
| | 9861638 | Feb 10, 2031 | U-2273 | | | |
| | 9987285 | Aug 17, 2035 | DP | | | |
| <u>RUCAPARIB CAMSYLATE - RUBRACA</u> | | | | | | |
| N 209115 002 | 10130636 | Aug 17, 2035 | U-2012 | | I-772 | Apr 06, 2021 |
| | 10130636 | Aug 17, 2035 | U-2101 | | NCE | Dec 19, 2021 |
| | 10130636 | Aug 17, 2035 | U-2273 | | ODE-126 | Dec 19, 2023 |
| | 6495541 | Jan 10, 2020 | DS DP | | ODE-168 | Apr 06, 2025 |
| | 7351701 | Jul 23, 2024 | U-2012 | | | |
| | 7351701 | Jul 23, 2024 | U-2101 | | | |
| | 7351701 | Jul 23, 2024 | U-2273 | | | |
| | 7531530 | Jul 23, 2024 | U-2012 | | | |
| | 7531530 | Jul 23, 2024 | U-2101 | | | |
| | 7531530 | Jul 23, 2024 | U-2273 | | | |
| | 8071579 | Aug 12, 2027 | U-2012 | | | |
| | 8071579 | Aug 12, 2027 | U-2101 | | | |
| | 8071579 | Aug 12, 2027 | U-2273 | | | |
| | 8143241 | Aug 12, 2027 | U-2012 | | | |
| | 8143241 | Aug 12, 2027 | U-2101 | | | |
| | 8143241 | Aug 12, 2027 | U-2273 | | | |
| | 8754072 | Feb 10, 2031 | DS DP | | | |
| | 8859562 | Aug 04, 2031 | U-2012 | | | |
| | 8859562 | Aug 04, 2031 | U-2101 | | | |
| | 8859562 | Aug 04, 2031 | U-2273 | | | |
| | 9045487 | Feb 10, 2031 | DS DP | | | |
| | 9861638 | Feb 10, 2031 | U-2012 | | | |
| | 9861638 | Feb 10, 2031 | U-2101 | | | |
| | 9861638 | Feb 10, 2031 | U-2273 | | | |
| | 9987285 | Aug 17, 2035 | DP | | | |
| <u>RUCAPARIB CAMSYLATE - RUBRACA</u> | | | | | | |
| N 209115 003 | 10130636 | Aug 17, 2035 | U-2012 | | I-772 | Apr 06, 2021 |
| | 10130636 | Aug 17, 2035 | U-2101 | | NCE | Dec 19, 2021 |
| | 10130636 | Aug 17, 2035 | U-2273 | | ODE-126 | Dec 19, 2023 |
| | 6495541 | Jan 10, 2020 | DS DP | | ODE-168 | Apr 06, 2025 |
| | 7351701 | Jul 23, 2024 | U-2012 | | | |
| | 7351701 | Jul 23, 2024 | U-2101 | | | |
| | 7351701 | Jul 23, 2024 | U-2273 | | | |
| | 7531530 | Jul 23, 2024 | U-2012 | | | |
| | 7531530 | Jul 23, 2024 | U-2101 | | | |
| | 7531530 | Jul 23, 2024 | U-2273 | | | |
| | 8071579 | Aug 12, 2027 | U-2012 | | | |
| | 8071579 | Aug 12, 2027 | U-2101 | | | |
| | 8071579 | Aug 12, 2027 | U-2273 | | | |
| | 8143241 | Aug 12, 2027 | U-2012 | | | |
| | 8143241 | Aug 12, 2027 | U-2101 | | | |
| | 8143241 | Aug 12, 2027 | U-2273 | | | |
| | 8754072 | Feb 10, 2031 | DS DP | | | |
| | 8859562 | Aug 04, 2031 | U-2012 | | | |
| | 8859562 | Aug 04, 2031 | U-2101 | | | |
| | 8859562 | Aug 04, 2031 | U-2273 | | | |
| | 9045487 | Feb 10, 2031 | DS DP | | | |
| | 9861638 | Feb 10, 2031 | U-2012 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>RUCAPARIB CAMSYLATE - RUBRACA</u> | | | | | | |
| N 209115 | 003 | 9861638 | Feb 10, 2031 | U-2101 | | |
| | | 9861638 | Feb 10, 2031 | U-2273 | | |
| | | 9987285 | Aug 17, 2035 | DP | | |
| <u>RUFINAMIDE - BANZEL</u> | | | | | | |
| N 021911 | 001 | 6740669 | Nov 14, 2022 | DS DP | | |
| | | 6740669*PED | May 14, 2023 | | | |
| | | 7750028*PED | Apr 19, 2019 | | | |
| <u>RUFINAMIDE - BANZEL</u> | | | | | | |
| N 021911 | 002 | 6740669 | Nov 14, 2022 | DS DP | | |
| | | 6740669*PED | May 14, 2023 | | | |
| | | 7750028*PED | Apr 19, 2019 | | | |
| <u>RUFINAMIDE - BANZEL</u> | | | | | | |
| N 021911 | 003 | 6740669 | Nov 14, 2022 | DS DP | | |
| | | 6740669*PED | May 14, 2023 | | | |
| | | 7750028*PED | Apr 19, 2019 | | | |
| <u>RUFINAMIDE - BANZEL</u> | | | | | | |
| N 201367 | 001 | 6740669 | Nov 14, 2022 | DS DP | | |
| | | 6740669*PED | May 14, 2023 | | | |
| | | 7750028*PED | Apr 19, 2019 | | | |
| <u>RUXOLITINIB PHOSPHATE - JAKAFI</u> | | | | | | |
| N 202192 | 001 | 7598257 | Dec 24, 2027 | DS DP U-1201 | ODE-79 | Dec 04, 2021 |
| | | 7598257 | Dec 24, 2027 | DS DP U-1622 | | |
| | | 8415362 | Dec 24, 2027 | DS DP | | |
| | | 8722693 | Jun 12, 2028 | DS DP | | |
| | | 8822481 | Jun 12, 2028 | | U-1573 | |
| | | 8829013 | Jun 12, 2028 | | U-1201 | |
| | | 8829013 | Jun 12, 2028 | | U-1622 | |
| | | 9079912 | Dec 12, 2026 | | U-1573 | |
| | | 9079912 | Dec 12, 2026 | | U-1721 | |
| <u>RUXOLITINIB PHOSPHATE - JAKAFI</u> | | | | | | |
| N 202192 | 002 | 7598257 | Dec 24, 2027 | DS DP U-1201 | ODE-79 | Dec 04, 2021 |
| | | 7598257 | Dec 24, 2027 | DS DP U-1622 | | |
| | | 8415362 | Dec 24, 2027 | DS DP | | |
| | | 8722693 | Jun 12, 2028 | DS DP | | |
| | | 8822481 | Jun 12, 2028 | | U-1573 | |
| | | 8829013 | Jun 12, 2028 | | U-1201 | |
| | | 8829013 | Jun 12, 2028 | | U-1622 | |
| | | 9079912 | Dec 12, 2026 | | U-1573 | |
| | | 9079912 | Dec 12, 2026 | | U-1721 | |
| <u>RUXOLITINIB PHOSPHATE - JAKAFI</u> | | | | | | |
| N 202192 | 003 | 7598257 | Dec 24, 2027 | DS DP U-1201 | ODE-79 | Dec 04, 2021 |
| | | 7598257 | Dec 24, 2027 | DS DP U-1622 | | |
| | | 8415362 | Dec 24, 2027 | DS DP | | |
| | | 8722693 | Jun 12, 2028 | DS DP | | |
| | | 8822481 | Jun 12, 2028 | | U-1573 | |
| | | 8829013 | Jun 12, 2028 | | U-1201 | |
| | | 8829013 | Jun 12, 2028 | | U-1622 | |
| | | 9079912 | Dec 12, 2026 | | U-1573 | |
| | | 9079912 | Dec 12, 2026 | | U-1721 | |
| <u>RUXOLITINIB PHOSPHATE - JAKAFI</u> | | | | | | |
| N 202192 | 004 | 7598257 | Dec 24, 2027 | DS DP U-1201 | ODE-79 | Dec 04, 2021 |
| | | 7598257 | Dec 24, 2027 | DS DP U-1622 | | |
| | | 8415362 | Dec 24, 2027 | DS DP | | |
| | | 8722693 | Jun 12, 2028 | DS DP | | |
| | | 8822481 | Jun 12, 2028 | | U-1573 | |
| | | 8829013 | Jun 12, 2028 | | U-1201 | |
| | | 8829013 | Jun 12, 2028 | | U-1622 | |
| | | 9079912 | Dec 12, 2026 | | U-1573 | |
| | | 9079912 | Dec 12, 2026 | | U-1721 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>RUXOLITINIB PHOSPHATE - JAKAFI</u> | | | | | | |
| N 202192 005 | 7598257 | Dec 24, 2027 | DS DP U-1201 | | ODE-79 | Dec 04, 2021 |
| | 7598257 | Dec 24, 2027 | DS DP U-1622 | | | |
| | 8415362 | Dec 24, 2027 | DS DP | | | |
| | 8722693 | Jun 12, 2028 | DS DP | | | |
| | 8822481 | Jun 12, 2028 | | U-1573 | | |
| | 8829013 | Jun 12, 2028 | | U-1201 | | |
| | 8829013 | Jun 12, 2028 | | U-1622 | | |
| | 9079912 | Dec 12, 2026 | | U-1573 | | |
| | 9079912 | Dec 12, 2026 | | U-1721 | | |
| <u>SACROSIDASE - SUCRAID</u> | | | | | | |
| N 020772 001 | 9255261 | Feb 07, 2034 | DS DP | | | |
| | 9469847 | Feb 07, 2034 | DS DP | | | |
| | 9849161 | Feb 07, 2034 | DS DP | | | |
| <u>SACUBITRIL; VALSARTAN - ENTRESTO</u> | | | | | | |
| N 207620 001 | 7468390 | Nov 27, 2023 | | DP | NCE | Jul 07, 2020 |
| | 8101659 | Jan 14, 2023 | | DP | | |
| | 8404744 | Jan 14, 2023 | | DP | | |
| | 8796331 | Jan 14, 2023 | | U-1723 | | |
| | 8877938 | May 27, 2027 | DS DP | | | |
| | 9388134 | Nov 08, 2026 | | U-1723 | | |
| <u>SACUBITRIL; VALSARTAN - ENTRESTO</u> | | | | | | |
| N 207620 002 | 7468390 | Nov 27, 2023 | | DP | NCE | Jul 07, 2020 |
| | 8101659 | Jan 14, 2023 | | DP | | |
| | 8404744 | Jan 14, 2023 | | DP | | |
| | 8796331 | Jan 14, 2023 | | U-1723 | | |
| | 8877938 | May 27, 2027 | DS DP | | | |
| | 9388134 | Nov 08, 2026 | | U-1723 | | |
| <u>SACUBITRIL; VALSARTAN - ENTRESTO</u> | | | | | | |
| N 207620 003 | 7468390 | Nov 27, 2023 | | DP | NCE | Jul 07, 2020 |
| | 8101659 | Jan 14, 2023 | | DP | | |
| | 8404744 | Jan 14, 2023 | | DP | | |
| | 8796331 | Jan 14, 2023 | | U-1723 | | |
| | 8877938 | May 27, 2027 | DS DP | | | |
| | 9388134 | Nov 08, 2026 | | U-1723 | | |
| <u>SAFINAMIDE MESYLATE - XADAGO</u> | | | | | | |
| N 207145 001 | 8076515 | Dec 10, 2028 | DS DP U-1993 | | NCE | Mar 21, 2022 |
| | 8278485 | Jun 08, 2027 | DS U-1993 | | | |
| | 8283380 | Sep 01, 2027 | | U-1993 | | |
| <u>SAFINAMIDE MESYLATE - XADAGO</u> | | | | | | |
| N 207145 002 | 8076515 | Dec 10, 2028 | DS DP U-1993 | | NCE | Mar 21, 2022 |
| | 8278485 | Jun 08, 2027 | DS U-1993 | | | |
| | 8283380 | Sep 01, 2027 | | U-1993 | | |
| <u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u> | | | | | | |
| N 022181 001 | 7566462 | Nov 16, 2025 | | DP | | |
| | 7566462*PED | May 16, 2026 | | | | |
| | 7566714 | Nov 17, 2024 | | U-989 | | |
| | 7566714*PED | May 17, 2025 | | | | |
| | 7612073 | Nov 17, 2024 | | U-1010 | | |
| | 7612073*PED | May 17, 2025 | | | | |
| | 7727987 | Nov 17, 2024 | | DP | | |
| | 7727987*PED | May 17, 2025 | | | | |
| | 7947681 | Nov 17, 2024 | | U-1156 | Y | |
| | 7947681*PED | May 17, 2025 | | | | |
| | 8003126 | Nov 16, 2025 | | | | |
| | 8003126*PED | May 16, 2026 | | | | |
| | 8067416 | Nov 17, 2024 | | U-989 | | |
| | 8067416*PED | May 17, 2025 | | | | |
| | 8318745 | Nov 17, 2024 | | DP | | |
| | 8318745*PED | May 17, 2025 | | | | |
| | 9433624 | Nov 17, 2024 | | U-1589 | | |
| | RE43797 | Nov 17, 2024 | | U-1156 | | |
| | RE43797*PED | May 17, 2025 | | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u> | | | | | | |
| N 022181 | 001 | 7566462 | Nov 16, 2025 | DP | | |
| | | 7566462*PED | May 16, 2026 | | | |
| | | 7566714 | Nov 17, 2024 | U-989 | | |
| | | 7566714*PED | May 17, 2025 | | | |
| | | 7612073 | Nov 17, 2024 | U-1010 | | |
| | | 7612073*PED | May 17, 2025 | | | |
| | | 7727987 | Nov 17, 2024 | DP | | |
| | | 7727987*PED | May 17, 2025 | | | |
| | | 7947681 | Nov 17, 2024 | U-1156 | Y | |
| | | 7947681*PED | May 17, 2025 | | | |
| | | 8003126 | Nov 16, 2025 | | | |
| | | 8003126*PED | May 16, 2026 | | | |
| | | 8067416 | Nov 17, 2024 | U-989 | | |
| | | 8067416*PED | May 17, 2025 | | | |
| | | 8318745 | Nov 17, 2024 | DP | | |
| | | 8318745*PED | May 17, 2025 | | | |
| | | 9433624 | Nov 17, 2024 | U-1589 | | |
| | | RE43797 | Nov 17, 2024 | U-1156 | | |
| | | RE43797*PED | May 17, 2025 | | | |
| <u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u> | | | | | | |
| N 205065 | 001 | 7566714 | Nov 17, 2024 | U-1589 | | |
| | | 7566714*PED | May 17, 2025 | | | |
| | | 7612073 | Nov 17, 2024 | U-1010 | | |
| | | 7612073*PED | May 17, 2025 | | | |
| | | 8067416 | Nov 17, 2024 | U-1589 | | |
| | | 8067416*PED | May 17, 2025 | | | |
| | | 9216178 | Nov 01, 2032 | DP | | |
| | | 9433624 | Nov 17, 2024 | U-1589 | | |
| | | RE43797 | Nov 17, 2024 | U-1590 | | |
| | | RE43797*PED | May 17, 2025 | | | |
| <u>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</u> | | | | | | |
| N 205065 | 002 | 7566714 | Nov 17, 2024 | U-1589 | | |
| | | 7566714*PED | May 17, 2025 | | | |
| | | 7612073 | Nov 17, 2024 | U-1010 | | |
| | | 7612073*PED | May 17, 2025 | | | |
| | | 8067416 | Nov 17, 2024 | U-1589 | | |
| | | 8067416*PED | May 17, 2025 | | | |
| | | 9216178 | Nov 01, 2032 | DP | | |
| | | 9433624 | Nov 17, 2024 | U-1589 | | |
| | | RE43797 | Nov 17, 2024 | U-1590 | | |
| | | RE43797*PED | May 17, 2025 | | | |
| <u>SAQUINAVIR - FORTOVASE</u> | | | | | | |
| N 020828 | 001 | 6352717 | Nov 16, 2019 | | | |
| <u>SARECYCLINE HYDROCHLORIDE - SEYSARA</u> | | | | | | |
| N 209521 | 001 | 8318706 | May 01, 2031 | DS DP U-2405 | NCE | Oct 01, 2023 |
| | | 8513223 | Dec 07, 2029 | U-2406 | | |
| | | 9255068 | Feb 09, 2033 | DS DP U-2407 | | |
| | | 9255068 | Feb 09, 2033 | DS DP U-2408 | | |
| | | 9481639 | Aug 10, 2028 | U-2409 | | |
| <u>SARECYCLINE HYDROCHLORIDE - SEYSARA</u> | | | | | | |
| N 209521 | 002 | 8318706 | May 01, 2031 | DS DP U-2405 | NCE | Oct 01, 2023 |
| | | 8513223 | Dec 07, 2029 | U-2406 | | |
| | | 9255068 | Feb 09, 2033 | DS DP U-2407 | | |
| | | 9255068 | Feb 09, 2033 | DS DP U-2408 | | |
| | | 9481639 | Aug 10, 2028 | U-2409 | | |
| <u>SARECYCLINE HYDROCHLORIDE - SEYSARA</u> | | | | | | |
| N 209521 | 003 | 8318706 | May 01, 2031 | DS DP U-2405 | NCE | Oct 01, 2023 |
| | | 8513223 | Dec 07, 2029 | U-2406 | | |
| | | 9255068 | Feb 09, 2033 | DS DP U-2407 | | |
| | | 9255068 | Feb 09, 2033 | DS DP U-2408 | | |
| | | 9481639 | Aug 10, 2028 | U-2409 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA</u> | | | | | | |
| N 022350 001 | 7951400 | Nov 30, 2028 | DP | | M-175 | Apr 05, 2019 |
| | RE44186 | Jul 31, 2023 | DS DP U-1837 | | M-198 | Feb 27, 2020 |
| | RE44186 | Jul 31, 2023 | DS DP U-995 | | | |
| <u>SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA</u> | | | | | | |
| N 022350 002 | 7951400 | Nov 30, 2028 | DP | | M-175 | Apr 05, 2019 |
| | RE44186 | Jul 31, 2023 | DS DP U-1837 | | M-198 | Feb 27, 2020 |
| | RE44186 | Jul 31, 2023 | DS DP U-995 | | | |
| <u>SECNIDAZOLE - SOLOSEC</u> | | | | | | |
| N 209363 001 | | | | | NCE GAIN | Sep 15, 2022 Sep 15, 2027 |
| <u>SELEXIPAG - UPTRAVI</u> | | | | | | |
| N 207947 001 | 7205302 | Apr 04, 2023 | DS DP U-1797 | | NCE | Dec 21, 2020 |
| | 8791122 | Aug 01, 2030 | DS DP | | ODE-106 | Dec 21, 2022 |
| | 9173881 | Aug 12, 2029 | U-1798 | | | |
| | 9284280 | Jun 25, 2030 | U-1831 | | | |
| <u>SELEXIPAG - UPTRAVI</u> | | | | | | |
| N 207947 002 | 7205302 | Apr 04, 2023 | DS DP U-1797 | | NCE | Dec 21, 2020 |
| | 8791122 | Aug 01, 2030 | DS DP | | ODE-106 | Dec 21, 2022 |
| | 9173881 | Aug 12, 2029 | U-1798 | | | |
| | 9284280 | Jun 25, 2030 | U-1831 | | | |
| <u>SELEXIPAG - UPTRAVI</u> | | | | | | |
| N 207947 003 | 7205302 | Apr 04, 2023 | DS DP U-1797 | | NCE | Dec 21, 2020 |
| | 8791122 | Aug 01, 2030 | DS DP | | ODE-106 | Dec 21, 2022 |
| | 9173881 | Aug 12, 2029 | U-1798 | | | |
| | 9284280 | Jun 25, 2030 | U-1831 | | | |
| <u>SELEXIPAG - UPTRAVI</u> | | | | | | |
| N 207947 004 | 7205302 | Apr 04, 2023 | DS DP U-1797 | | NCE | Dec 21, 2020 |
| | 8791122 | Aug 01, 2030 | DS DP | | ODE-106 | Dec 21, 2022 |
| | 9173881 | Aug 12, 2029 | U-1798 | | | |
| | 9284280 | Jun 25, 2030 | U-1831 | | | |
| <u>SELEXIPAG - UPTRAVI</u> | | | | | | |
| N 207947 005 | 7205302 | Apr 04, 2023 | DS DP U-1797 | | NCE | Dec 21, 2020 |
| | 8791122 | Aug 01, 2030 | DS DP | | ODE-106 | Dec 21, 2022 |
| | 9173881 | Aug 12, 2029 | U-1798 | | | |
| | 9284280 | Jun 25, 2030 | U-1831 | | | |
| <u>SELEXIPAG - UPTRAVI</u> | | | | | | |
| N 207947 006 | 7205302 | Apr 04, 2023 | DS DP U-1797 | | NCE | Dec 21, 2020 |
| | 8791122 | Aug 01, 2030 | DS DP | | ODE-106 | Dec 21, 2022 |
| | 9173881 | Aug 12, 2029 | U-1798 | | | |
| | 9284280 | Jun 25, 2030 | U-1831 | | | |
| <u>SELEXIPAG - UPTRAVI</u> | | | | | | |
| N 207947 007 | 7205302 | Apr 04, 2023 | DS DP U-1797 | | NCE | Dec 21, 2020 |
| | 8791122 | Aug 01, 2030 | DS DP | | ODE-106 | Dec 21, 2022 |
| | 9173881 | Aug 12, 2029 | U-1798 | | | |
| | 9284280 | Jun 25, 2030 | U-1831 | | | |
| <u>SELEXIPAG - UPTRAVI</u> | | | | | | |
| N 207947 008 | 7205302 | Apr 04, 2023 | DS DP U-1797 | | NCE | Dec 21, 2020 |
| | 8791122 | Aug 01, 2030 | DS DP | | ODE-106 | Dec 21, 2022 |
| | 9173881 | Aug 12, 2029 | U-1798 | | | |
| | 9284280 | Jun 25, 2030 | U-1831 | | | |
| <u>SEMAGLUTIDE - OZEMPIC</u> | | | | | | |
| N 209637 001 | 6899699 | Jan 02, 2022 | DP | | NCE | Dec 05, 2022 |
| | 7762994 | May 23, 2024 | DP | | | |
| | 8114833 | Aug 13, 2025 | DP | | | |
| | 8129343 | Jan 29, 2029 | DS DP U-2202 | | | |
| | 8536122 | Mar 20, 2026 | DS DP U-2202 | | | |
| | 8579869 | Jun 30, 2023 | DP | | | |
| | 8672898 | Jan 02, 2022 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>SEMAGLUTIDE - OZEMPIC</u> | | | | | | |
| N 209637 | 001 | 8684969 | | | | |
| | | Oct 20, 2025 | DP | | | |
| | | 8920383 | | | | |
| | | Jul 17, 2026 | DP | | | |
| | | 9108002 | | | | |
| | | Jan 20, 2026 | DP | | | |
| | | 9132239 | | | | |
| | | Feb 01, 2032 | DP | | | |
| | | 9457154 | | | | |
| | | Sep 27, 2027 | DP | | | |
| | | 9486588 | | | | |
| | | Jan 02, 2022 | DP | | | |
| | | 9616180 | | | | |
| | | Jan 20, 2026 | DP | | | |
| | | 9687611 | | | | |
| | | Feb 27, 2027 | DP | | | |
| | | 9775953 | | | | |
| | | Jul 17, 2026 | DP | | | |
| | | 9861757 | | | | |
| | | Jan 20, 2026 | DP | | | |
| | | RE46363 | | | | |
| | | Aug 03, 2026 | DP | | | |
| <u>SERTRALINE HYDROCHLORIDE - ZOLOFT</u> | | | | | | |
| N 020990 | 001 | 6727283 | | | | |
| | | Oct 11, 2019 | DP | U-580 | | |
| | | 7067555 | | | | |
| | | Oct 11, 2019 | DP | | | |
| | | 7067555*PED | | | | |
| | | Apr 11, 2020 | | | | |
| <u>SEVELAMER CARBONATE - RENVELA</u> | | | | | | |
| N 022127 | 001 | 7985418 | | | | |
| | | Oct 27, 2025 | DP | | NPP | Nov 25, 2019 |
| <u>SEVELAMER CARBONATE - RENVELA</u> | | | | | | |
| N 022318 | 001 | 9095509 | | | | |
| | | Dec 06, 2030 | DP | | NPP | Nov 25, 2019 |
| <u>SEVELAMER CARBONATE - RENVELA</u> | | | | | | |
| N 022318 | 002 | 9095509 | | | | |
| | | Dec 06, 2030 | DP | | NPP | Nov 25, 2019 |
| <u>SEVELAMER HYDROCHLORIDE - RENAGEL</u> | | | | | | |
| N 021179 | 001 | 6733780 | | | | |
| | | Oct 18, 2020 | DP | | | |
| <u>SEVELAMER HYDROCHLORIDE - RENAGEL</u> | | | | | | |
| N 021179 | 002 | 6733780 | | | | |
| | | Oct 18, 2020 | DP | | | |
| <u>SILDENAFIL CITRATE - VIAGRA</u> | | | | | | |
| N 020895 | 001 | 6469012 | | | | |
| | | Oct 22, 2019 | | U-155 | | |
| <u>SILDENAFIL CITRATE - VIAGRA</u> | | | | | | |
| N 020895 | 002 | 6469012 | | | | |
| | | Oct 22, 2019 | | U-155 | | |
| <u>SILDENAFIL CITRATE - VIAGRA</u> | | | | | | |
| N 020895 | 003 | 6469012 | | | | |
| | | Oct 22, 2019 | | U-155 | | |
| <u>SIMEPREVIR SODIUM - OLYSIO</u> | | | | | | |
| N 205123 | 001 | 7671032 | | | | |
| | | May 19, 2025 | DS DP | | M-171 | Feb 26, 2019 |
| | | 8148399 | | | | |
| | | Sep 05, 2029 | DS DP | U-1467 | M-179 | May 20, 2019 |
| | | 8349869 | | | | |
| | | Jul 28, 2026 | DS DP | U-1467 | | |
| | | 8741926 | | | | |
| | | Jul 28, 2026 | DS | U-1467 | | |
| | | 8754106 | | | | |
| | | Jul 28, 2026 | DS | U-1467 | | |
| | | 9040562 | | | | |
| | | Jul 28, 2026 | DS DP | U-1467 | | |
| | | 9353103 | | | | |
| | | Jul 28, 2026 | | U-1467 | | |
| | | 9623022 | | | | |
| | | Jul 28, 2026 | | U-1467 | | |
| | | 9856265 | | | | |
| | | Jul 28, 2026 | DS DP | U-1467 | | |
| <u>SIMVASTATIN - FLOLIPID</u> | | | | | | |
| N 206679 | 001 | 9597289 | | | | |
| | | Feb 23, 2030 | DP | | | |
| <u>SIMVASTATIN - FLOLIPID</u> | | | | | | |
| N 206679 | 002 | 9597289 | | | | |
| | | Feb 23, 2030 | DP | | | |
| <u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u> | | | | | | |
| N 202343 | 001 | 6699871 | | | | |
| | | Jul 26, 2022 | DS DP | U-1188 | | |
| | | 6890898 | | | | |
| | | Feb 02, 2019 | | U-1189 | | |
| | | 6890898 | | | | |
| | | Feb 02, 2019 | | U-1190 | | |
| | | 6890898 | | | | |
| | | Feb 02, 2019 | | U-1191 | | |
| | | 7078381 | | | | |
| | | Feb 02, 2019 | | U-1188 | | |
| | | 7125873 | | | | |
| | | Jul 26, 2022 | DP | U-1189 | | |
| | | 7125873 | | | | |
| | | Jul 26, 2022 | DP | U-1190 | | |
| | | 7125873 | | | | |
| | | Jul 26, 2022 | DP | U-1192 | | |
| | | 7125873 | | | | |
| | | Jul 26, 2022 | DP | U-1193 | | |
| | | 7326708 | | | | |
| | | Apr 11, 2026 | DS DP | U-1188 | | |
| | | 7459428 | | | | |
| | | Feb 02, 2019 | | U-1189 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u> | | | | | | |
| N 202343 001 | 8168637 | Jun 26, 2022 | DP U-1188 | | | |
| <u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u> | | | | | | |
| N 202343 002 | 6699871 | Jul 26, 2022 | DS DP U-1188 | | | |
| | 6890898 | Feb 02, 2019 | U-1189 | | | |
| | 6890898 | Feb 02, 2019 | U-1190 | | | |
| | 6890898 | Feb 02, 2019 | U-1191 | | | |
| | 7078381 | Feb 02, 2019 | U-1188 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1189 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1190 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1192 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1193 | | | |
| | 7326708 | Apr 11, 2026 | DS DP U-1188 | | | |
| | 7459428 | Feb 02, 2019 | U-1189 | | | |
| | 8168637 | Jun 26, 2022 | DP U-1188 | | | |
| <u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u> | | | | | | |
| N 202343 003 | 6699871 | Jul 26, 2022 | DS DP U-1188 | | | |
| | 6890898 | Feb 02, 2019 | U-1189 | | | |
| | 6890898 | Feb 02, 2019 | U-1190 | | | |
| | 6890898 | Feb 02, 2019 | U-1191 | | | |
| | 7078381 | Feb 02, 2019 | U-1188 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1189 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1190 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1192 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1193 | | | |
| | 7326708 | Apr 11, 2026 | DS DP U-1188 | | | |
| | 7459428 | Feb 02, 2019 | U-1189 | | | |
| | 8168637 | Jun 26, 2022 | DP U-1188 | | | |
| <u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u> | | | | | | |
| N 202343 004 | 6699871 | Jul 26, 2022 | DS DP U-1188 | | | |
| | 6890898 | Feb 02, 2019 | U-1189 | | | |
| | 6890898 | Feb 02, 2019 | U-1190 | | | |
| | 6890898 | Feb 02, 2019 | U-1191 | | | |
| | 7078381 | Feb 02, 2019 | U-1188 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1189 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1190 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1192 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1193 | | | |
| | 7326708 | Apr 11, 2026 | DS DP U-1188 | | | |
| | 7459428 | Feb 02, 2019 | U-1189 | | | |
| | 8168637 | Jun 26, 2022 | DP U-1188 | | | |
| <u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u> | | | | | | |
| N 202343 005 | 6699871 | Jul 26, 2022 | DS DP U-1188 | | | |
| | 6890898 | Feb 02, 2019 | U-1189 | | | |
| | 6890898 | Feb 02, 2019 | U-1190 | | | |
| | 6890898 | Feb 02, 2019 | U-1191 | | | |
| | 7078381 | Feb 02, 2019 | U-1188 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1189 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1190 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1192 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1193 | | | |
| | 7326708 | Apr 11, 2026 | DS DP U-1188 | | | |
| | 7459428 | Feb 02, 2019 | U-1189 | | | |
| | 8168637 | Jun 26, 2022 | DP U-1188 | | | |
| <u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u> | | | | | | |
| N 202343 006 | 6699871 | Jul 26, 2022 | DS DP U-1188 | | | |
| | 6890898 | Feb 02, 2019 | U-1189 | | | |
| | 6890898 | Feb 02, 2019 | U-1190 | | | |
| | 6890898 | Feb 02, 2019 | U-1191 | | | |
| | 7078381 | Feb 02, 2019 | U-1188 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1189 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1190 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1192 | | | |
| | 7125873 | Jul 26, 2022 | DP U-1193 | | | |
| | 7326708 | Apr 11, 2026 | DS DP U-1188 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u> | | | | | | |
| N 202343 006 | 7459428 | Feb 02, 2019 | | U-1189 | | |
| | 8168637 | Jun 26, 2022 | DP | U-1188 | | |
| <u>SINCALIDE - KINEVAC</u> | | | | | | |
| N 017697 001 | 6803046 | Aug 16, 2022 | DP | | | |
| <u>SINECATECHINS - VEREGEN</u> | | | | | | |
| N 021902 001 | 5795911 | Oct 31, 2020 | | U-172 | | |
| | 7858662 | Oct 02, 2026 | DP | U-172 | | |
| | 9770406 | Jul 12, 2025 | DP | U-172 | | |
| <u>SIROLIMUS - RAPAMUNE</u> | | | | | | |
| N 021083 001 | | | | | ODE-92 | May 28, 2022 |
| <u>SIROLIMUS - RAPAMUNE</u> | | | | | | |
| N 021110 001 | | | | | ODE-92 | May 28, 2022 |
| <u>SIROLIMUS - RAPAMUNE</u> | | | | | | |
| N 021110 002 | | | | | ODE-92 | May 28, 2022 |
| <u>SIROLIMUS - RAPAMUNE</u> | | | | | | |
| N 021110 003 | | | | | ODE-92 | May 28, 2022 |
| <u>SIROLIMUS - RAPAMUNE</u> | | | | | | |
| N 021110 004 | | | | | ODE-92 | May 28, 2022 |
| <u>SITAGLIPTIN PHOSPHATE - JANUVIA</u> | | | | | | |
| N 021995 001 | 6699871 | Jul 26, 2022 | DS DP | U-774 | | |
| | 6890898 | Feb 02, 2019 | | U-1997 | | |
| | 7078381 | Feb 02, 2019 | | U-1997 | | |
| | 7125873 | Jul 26, 2022 | | U-1036 | | |
| | 7125873 | Jul 26, 2022 | | U-1037 | | |
| | 7125873 | Jul 26, 2022 | | U-1038 | | |
| | 7125873 | Jul 26, 2022 | | U-775 | | |
| | 7326708 | Nov 24, 2026 | DS DP | U-802 | | |
| | 7459428 | Feb 02, 2019 | | U-1945 | | |
| <u>SITAGLIPTIN PHOSPHATE - JANUVIA</u> | | | | | | |
| N 021995 002 | 6699871 | Jul 26, 2022 | DS DP | U-774 | | |
| | 6890898 | Feb 02, 2019 | | U-1997 | | |
| | 7078381 | Feb 02, 2019 | | U-1997 | | |
| | 7125873 | Jul 26, 2022 | | U-1036 | | |
| | 7125873 | Jul 26, 2022 | | U-1037 | | |
| | 7125873 | Jul 26, 2022 | | U-1038 | | |
| | 7125873 | Jul 26, 2022 | | U-775 | | |
| | 7326708 | Nov 24, 2026 | DS DP | U-802 | | |
| | 7459428 | Feb 02, 2019 | | U-1945 | | |
| <u>SITAGLIPTIN PHOSPHATE - JANUVIA</u> | | | | | | |
| N 021995 003 | 6699871 | Jul 26, 2022 | DS DP | U-774 | | |
| | 6890898 | Feb 02, 2019 | | U-1997 | | |
| | 7078381 | Feb 02, 2019 | | U-1997 | | |
| | 7125873 | Jul 26, 2022 | | U-1036 | | |
| | 7125873 | Jul 26, 2022 | | U-1037 | | |
| | 7125873 | Jul 26, 2022 | | U-1038 | | |
| | 7125873 | Jul 26, 2022 | | U-775 | | |
| | 7326708 | Nov 24, 2026 | DS DP | U-802 | | |
| | 7459428 | Feb 02, 2019 | | U-1945 | | |
| <u>SODIUM NITRITE - SODIUM NITRITE</u> | | | | | | |
| N 203922 001 | 8568793 | Dec 24, 2031 | DS DP | | | |
| <u>SODIUM NITRITE; SODIUM THIOSULFATE - NITHIODOTE</u> | | | | | | |
| N 201444 001 | 8496973 | Mar 29, 2031 | DS DP | U-1419 | | |
| | 8568793 | Dec 24, 2031 | DS DP | | | |
| | 9345724 | Mar 29, 2031 | DS DP | U-2015 | | |
| | 9585912 | Mar 29, 2031 | DS DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>SODIUM OXYBATE - XYREM</u> | | | | | | |
| N 021196 001 | 6780889 | Jul 04, 2020 | DP | | NPP | Oct 26, 2021 |
| | 6780889*PED | Jan 04, 2021 | | | PED | Apr 26, 2022 |
| | 7262219 | Jul 04, 2020 | DP | | | |
| | 7262219*PED | Jan 04, 2021 | | | | |
| | 7668730 | Jun 16, 2024 | U-1110 | | | |
| | 7668730*PED | Dec 16, 2024 | | | | |
| | 7765106 | Jun 16, 2024 | U-1069 | | | |
| | 7765106*PED | Dec 16, 2024 | | | | |
| | 7765107 | Jun 16, 2024 | U-1070 | | | |
| | 7765107*PED | Dec 16, 2024 | | | | |
| | 7851506 | Dec 22, 2019 | U-1101 | | | |
| | 7851506 | Dec 22, 2019 | U-1102 | | | |
| | 7851506*PED | Jun 22, 2020 | | | | |
| | 7895059 | Dec 17, 2022 | U-1110 | | | |
| | 7895059*PED | Jun 17, 2023 | | | | |
| | 8263650 | Dec 22, 2019 | DP U-1101 | | | |
| | 8263650 | Dec 22, 2019 | DP U-1102 | | | |
| | 8263650*PED | Jun 22, 2020 | | | | |
| | 8324275 | Dec 22, 2019 | U-1101 | | | |
| | 8324275 | Dec 22, 2019 | U-1102 | | | |
| | 8324275*PED | Jun 22, 2020 | | | | |
| | 8457988 | Dec 17, 2022 | U-1110 | | | |
| | 8457988*PED | Jun 17, 2023 | | | | |
| | 8589182 | Dec 17, 2022 | U-1110 | | | |
| | 8589182*PED | Jun 17, 2023 | | | | |
| | 8731963 | Dec 17, 2022 | U-1110 | | | |
| | 8731963*PED | Jun 17, 2023 | | | | |
| | 8772306 | Mar 15, 2033 | U-1532 | | | |
| | 8772306*PED | Sep 15, 2033 | | | | |
| | 8859619 | Dec 22, 2019 | DP | | | |
| | 8859619*PED | Jun 22, 2020 | | | | |
| | 8952062 | Dec 22, 2019 | U-1101 | | | |
| | 8952062 | Dec 22, 2019 | U-1102 | | | |
| | 8952062*PED | Jun 22, 2020 | | | | |
| | 9050302 | Mar 15, 2033 | U-1532 | | | |
| | 9050302*PED | Sep 15, 2033 | | | | |
| | 9486426 | Mar 15, 2033 | U-1532 | | | |
| | 9486426*PED | Sep 15, 2033 | | | | |
| | 9539330 | Dec 22, 2019 | DP | | | |
| | 9539330*PED | Jun 22, 2020 | | | | |
| <u>SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE - OSMOPREP</u> | | | | | | |
| N 021892 001 | 7687075 | Jun 22, 2028 | DS DP | | | |
| <u>SODIUM THIOSULFATE - SODIUM THIOSULFATE</u> | | | | | | |
| N 203923 001 | 8496973 | Mar 29, 2031 | DS DP | U-1419 | | |
| | 9345724 | Mar 29, 2031 | DS DP | U-2015 | | |
| | 9585912 | Mar 29, 2031 | DS DP | | | |
| <u>SODIUM ZIRCONIUM CYCLOSILICATE - LOKELMA</u> | | | | | | |
| N 207078 001 | 6332985 | Mar 29, 2019 | | U-2312 | NCE | May 18, 2023 |
| | 8802152 | Apr 19, 2032 | DS | | | |
| | 8808750 | Feb 10, 2032 | | U-2312 | | |
| | 8877255 | Oct 22, 2033 | DS | | | |
| | 9592253 | Oct 14, 2035 | DS | U-2312 | | |
| | 9844567 | Feb 10, 2032 | | U-2312 | | |
| | 9861658 | Feb 10, 2032 | | U-2312 | | |
| <u>SODIUM ZIRCONIUM CYCLOSILICATE - LOKELMA</u> | | | | | | |
| N 207078 002 | 6332985 | Mar 29, 2019 | | U-2312 | NCE | May 18, 2023 |
| | 8802152 | Apr 19, 2032 | DS | | | |
| | 8808750 | Feb 10, 2032 | | U-2312 | | |
| | 8877255 | Oct 22, 2033 | DS | | | |
| | 9592253 | Oct 14, 2035 | DS | U-2312 | | |
| | 9844567 | Feb 10, 2032 | | U-2312 | | |
| | 9861658 | Feb 10, 2032 | | U-2312 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>SOFOSBUVIR - SOVALDI</u> | | | | | | |
| N 204671 | 001 | 7964580 | Mar 26, 2029 | DS DP U-1470 | NPP | Apr 07, 2020 |
| | | 8334270 | Mar 21, 2028 | DS DP U-1470 | ODE-135 | Apr 07, 2024 |
| | | 8580765 | Mar 21, 2028 | DS DP U-1470 | | |
| | | 8618076 | Dec 11, 2030 | DS DP U-1470 | | |
| | | 8633309 | Mar 26, 2029 | DS DP U-1470 | | |
| | | 8889159 | Mar 26, 2029 | DP U-1470 | | |
| | | 9085573 | Mar 21, 2028 | DS DP U-1470 | | |
| | | 9284342 | Sep 13, 2030 | DS DP U-1470 | | |
| | | 9549941 | Mar 26, 2029 | U-1958 | | |
| <u>SOFOSBUVIR; VELPATASVIR - EPCLUSA</u> | | | | | | |
| N 208341 | 001 | 10086011 | Jan 30, 2034 | U-1470 | NCE | Jun 28, 2021 |
| | | 7964580 | Mar 26, 2029 | DS DP U-1470 | NPP | Aug 01, 2020 |
| | | 8334270 | Mar 21, 2028 | DS DP U-1470 | | |
| | | 8575135 | Nov 16, 2032 | DS DP U-1470 | | |
| | | 8580765 | Mar 21, 2028 | DS DP U-1470 | | |
| | | 8618076 | Dec 11, 2030 | DS DP U-1470 | | |
| | | 8633309 | Mar 26, 2029 | DS DP U-1470 | | |
| | | 8735372 | Mar 21, 2028 | U-1470 | | |
| | | 8889159 | Mar 26, 2029 | DP U-1470 | | |
| | | 8921341 | Nov 16, 2032 | DS DP U-1470 | | |
| | | 8940718 | Nov 16, 2032 | DS DP U-1470 | | |
| | | 9085573 | Mar 21, 2028 | DS DP U-1470 | | |
| | | 9284342 | Sep 13, 2030 | DS DP U-1470 | | |
| | | 9757406 | Jan 30, 2034 | DP | | |
| <u>SOFOSBUVIR; VELPATASVIR; VOXILAPREVIR - VOSEVI</u> | | | | | | |
| N 209195 | 001 | 7964580 | Mar 26, 2029 | DS DP U-2039 | NCE | Jul 18, 2022 |
| | | 7964580 | Mar 26, 2029 | DS DP U-2040 | | |
| | | 8334270 | Mar 21, 2028 | DS DP U-2039 | | |
| | | 8334270 | Mar 21, 2028 | DS DP U-2040 | | |
| | | 8575135 | Nov 05, 2033 | DS DP U-2039 | | |
| | | 8575135 | Nov 05, 2033 | DS DP U-2040 | | |
| | | 8580765 | Mar 21, 2028 | DS DP U-2039 | | |
| | | 8580765 | Mar 21, 2028 | DS DP U-2040 | | |
| | | 8618076 | Dec 11, 2030 | DS DP U-2039 | | |
| | | 8618076 | Dec 11, 2030 | DS DP U-2040 | | |
| | | 8633309 | Mar 26, 2029 | DS DP U-2039 | | |
| | | 8633309 | Mar 26, 2029 | DS DP U-2040 | | |
| | | 8735372 | Mar 21, 2028 | DS DP U-2039 | | |
| | | 8735372 | Mar 21, 2028 | DS DP U-2040 | | |
| | | 8889159 | Mar 26, 2029 | DS DP U-2039 | | |
| | | 8889159 | Mar 26, 2029 | DS DP U-2040 | | |
| | | 8921341 | Nov 16, 2032 | DS DP U-2039 | | |
| | | 8921341 | Nov 16, 2032 | DS DP U-2040 | | |
| | | 8940718 | Nov 16, 2032 | DS DP U-2039 | | |
| | | 8940718 | Nov 16, 2032 | DS DP U-2040 | | |
| | | 9085573 | Mar 21, 2028 | DS DP U-2039 | | |
| | | 9085573 | Mar 21, 2028 | DS DP U-2040 | | |
| | | 9284342 | Sep 13, 2030 | DS DP U-2039 | | |
| | | 9284342 | Sep 13, 2030 | DS DP U-2040 | | |
| | | 9296782 | Jul 17, 2034 | DS DP | | |
| | | 9585906 | Mar 21, 2028 | DS DP U-2039 | | |
| | | 9585906 | Mar 21, 2028 | DS DP U-2040 | | |
| | | 9868745 | Nov 16, 2032 | DS DP | | |
| <u>SOLIFENACIN SUCCINATE - VESICARE</u> | | | | | | |
| N 021518 | 001 | 6017927*PED | May 19, 2019 | | | |
| <u>SOLIFENACIN SUCCINATE - VESICARE</u> | | | | | | |
| N 021518 | 002 | 6017927*PED | May 19, 2019 | | | |
| <u>SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX</u> | | | | | | |
| N 021148 | 004 | 6004297 | Jan 28, 2019 | DP | | |
| | | 6235004 | Jan 28, 2019 | DP | | |
| | | RE41956 | Jan 21, 2021 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX</u> | | | | | | |
| N 021148 | 005 | 6004297 | | | | |
| | | Jan 28, 2019 | DP | | | |
| | | 6235004 | | | | |
| | | Jan 28, 2019 | DP | | | |
| | | RE41956 | | | | |
| | | Jan 21, 2021 | DP | | | |
| <u>SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX</u> | | | | | | |
| N 021148 | 006 | 6004297 | | | | |
| | | Jan 28, 2019 | DP | | | |
| | | 6235004 | | | | |
| | | Jan 28, 2019 | DP | | | |
| | | RE41956 | | | | |
| | | Jan 21, 2021 | DP | | | |
| <u>SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX</u> | | | | | | |
| N 021148 | 007 | 6004297 | | | | |
| | | Jan 28, 2019 | DP | | | |
| | | RE41956 | | | | |
| | | Jan 21, 2021 | DP | | | |
| | | RE43834 | | | | |
| | | Jan 28, 2019 | DP | | | |
| <u>SOMATROPIN RECOMBINANT - NORDITROPIN FLEXPRO</u> | | | | | | |
| N 021148 | 008 | 6899699 | | | | |
| | | Jan 02, 2022 | DP | | | |
| | | 8672898 | | | | |
| | | Jan 02, 2022 | DP | | | |
| | | 8684969 | | | | |
| | | Oct 20, 2025 | DP | | | |
| | | 8920383 | | | | |
| | | Jul 17, 2026 | DP | | | |
| | | 9108002 | | | | |
| | | Jan 20, 2026 | DP | | | |
| | | 9132239 | | | | |
| | | Feb 01, 2032 | DP | | | |
| | | 9457154 | | | | |
| | | Sep 27, 2027 | DP | | | |
| | | 9486588 | | | | |
| | | Jan 02, 2022 | DP | | | |
| | | 9616180 | | | | |
| | | Jan 20, 2026 | DP | | | |
| | | 9687611 | | | | |
| | | Feb 27, 2027 | DP | | | |
| | | 9775953 | | | | |
| | | Jul 17, 2026 | DP | | | |
| | | 9861757 | | | | |
| | | Jan 20, 2026 | DP | | | |
| | | RE46363 | | | | |
| | | Aug 03, 2026 | DP | | | |
| <u>SOMATROPIN RECOMBINANT - NORDITROPIN FLEXPRO</u> | | | | | | |
| N 021148 | 009 | 6899699 | | | | |
| | | Jan 02, 2022 | DP | | | |
| | | 8672898 | | | | |
| | | Jan 02, 2022 | DP | | | |
| | | 8684969 | | | | |
| | | Oct 20, 2025 | DP | | | |
| | | 8920383 | | | | |
| | | Jul 17, 2026 | DP | | | |
| | | 9108002 | | | | |
| | | Jan 20, 2026 | DP | | | |
| | | 9132239 | | | | |
| | | Feb 01, 2032 | DP | | | |
| | | 9457154 | | | | |
| | | Sep 27, 2027 | DP | | | |
| | | 9486588 | | | | |
| | | Jan 02, 2022 | DP | | | |
| | | 9616180 | | | | |
| | | Jan 20, 2026 | DP | | | |
| | | 9687611 | | | | |
| | | Feb 27, 2027 | DP | | | |
| | | 9775953 | | | | |
| | | Jul 17, 2026 | DP | | | |
| | | 9861757 | | | | |
| | | Jan 20, 2026 | DP | | | |
| | | RE46363 | | | | |
| | | Aug 03, 2026 | DP | | | |
| <u>SOMATROPIN RECOMBINANT - NORDITROPIN FLEXPRO</u> | | | | | | |
| N 021148 | 010 | 6899699 | | | | |
| | | Jan 02, 2022 | DP | | | |
| | | 8672898 | | | | |
| | | Jan 02, 2022 | DP | | | |
| | | 8684969 | | | | |
| | | Oct 20, 2025 | DP | | | |
| | | 8920383 | | | | |
| | | Jul 17, 2026 | DP | | | |
| | | 9108002 | | | | |
| | | Jan 20, 2026 | DP | | | |
| | | 9132239 | | | | |
| | | Feb 01, 2032 | DP | | | |
| | | 9457154 | | | | |
| | | Sep 27, 2027 | DP | | | |
| | | 9486588 | | | | |
| | | Jan 02, 2022 | DP | | | |
| | | 9616180 | | | | |
| | | Jan 20, 2026 | DP | | | |
| | | 9687611 | | | | |
| | | Feb 27, 2027 | DP | | | |
| | | 9775953 | | | | |
| | | Jul 17, 2026 | DP | | | |
| | | 9861757 | | | | |
| | | Jan 20, 2026 | DP | | | |
| | | RE46363 | | | | |
| | | Aug 03, 2026 | DP | | | |
| <u>SOMATROPIN RECOMBINANT - NORDITROPIN FLEXPRO</u> | | | | | | |
| N 021148 | 011 | 6899699 | | | | |
| | | Jan 02, 2022 | DP | | | |
| | | 8672898 | | | | |
| | | Jan 02, 2022 | DP | | | |
| | | 8684969 | | | | |
| | | Oct 20, 2025 | DP | | | |
| | | 8920383 | | | | |
| | | Jul 17, 2026 | DP | | | |
| | | 9108002 | | | | |
| | | Jan 20, 2026 | DP | | | |
| | | 9132239 | | | | |
| | | Feb 01, 2032 | DP | | | |
| | | 9457154 | | | | |
| | | Sep 27, 2027 | DP | | | |
| | | 9486588 | | | | |
| | | Jan 02, 2022 | DP | | | |
| | | 9616180 | | | | |
| | | Jan 20, 2026 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>SOMATROPIN RECOMBINANT - NORDITROPIN FLEXPRO</u> | | | | | | |
| N 021148 | 011 | 9687611 | Feb 27, 2027 | DP | | |
| | | 9775953 | Jul 17, 2026 | DP | | |
| | | 9861757 | Jan 20, 2026 | DP | | |
| | | RE46363 | Aug 03, 2026 | DP | | |
| <u>SONIDEGIB PHOSPHATE - ODOMZO</u> | | | | | | |
| N 205266 | 001 | 8063043 | Sep 15, 2029 | DS DP | NCE | Jul 24, 2020 |
| | | 8178563 | Feb 06, 2029 | DS | U-1722 | |
| <u>SORAFENIB TOSYLATE - NEXAVAR</u> | | | | | | |
| N 021923 | 001 | 7235576 | Jan 12, 2020 | DS DP | ODE-56 | Nov 22, 2020 |
| | | 7351834 | Jan 12, 2020 | DS | | |
| | | 7897623 | Jan 12, 2020 | DP | | |
| | | 8124630 | Jan 12, 2020 | U-1459 | | |
| | | 8618141 | Feb 11, 2023 | U-1480 | | |
| | | 8841330 | Jan 12, 2020 | U-1696 | | |
| | | 8877933 | Dec 24, 2027 | DS DP U-1624 | | |
| | | 9737488 | Sep 10, 2028 | DP U-1480 | | |
| | | 9737488 | Sep 10, 2028 | DP U-1696 | | |
| | | 9737488 | Sep 10, 2028 | DP U-2107 | | |
| <u>SOTALOL HYDROCHLORIDE - SOTYLIZE</u> | | | | | | |
| N 205108 | 001 | 9724297 | Aug 31, 2035 | DP | U-2096 | |
| <u>SPINOSAD - NATROBA</u> | | | | | | |
| N 022408 | 001 | 6063771 | Jul 25, 2023 | DP | U-1670 | |
| | | 6342482 | Jun 22, 2019 | DP | U-1105 | |
| | | 7030095 | Jul 02, 2021 | DP | U-1105 | |
| <u>SPIRONOLACTONE - CAROSPIR</u> | | | | | | |
| N 209478 | 001 | 9757394 | Oct 28, 2036 | DP | U-2109 | |
| <u>STAVUDINE - ZERIT XR</u> | | | | | | |
| N 021453 | 001 | 7135465 | Feb 18, 2023 | DP | U-167 | |
| <u>STAVUDINE - ZERIT XR</u> | | | | | | |
| N 021453 | 002 | 7135465 | Feb 18, 2023 | DP | U-167 | |
| <u>STAVUDINE - ZERIT XR</u> | | | | | | |
| N 021453 | 003 | 7135465 | Feb 18, 2023 | DP | U-167 | |
| <u>STAVUDINE - ZERIT XR</u> | | | | | | |
| N 021453 | 004 | 7135465 | Feb 18, 2023 | DP | U-167 | |
| <u>STIRIPENTOL - DIACOMIT</u> | | | | | | |
| N 206709 | 001 | | | | NCE ODE-198 | Aug 20, 2023 Aug 20, 2025 |
| <u>STIRIPENTOL - DIACOMIT</u> | | | | | | |
| N 206709 | 002 | | | | NCE ODE-198 | Aug 20, 2023 Aug 20, 2025 |
| <u>STIRIPENTOL - DIACOMIT</u> | | | | | | |
| N 207223 | 001 | | | | NCE ODE-198 | Aug 20, 2023 Aug 20, 2025 |
| <u>STIRIPENTOL - DIACOMIT</u> | | | | | | |
| N 207223 | 002 | | | | NCE ODE-198 | Aug 20, 2023 Aug 20, 2025 |
| <u>SUCROFERRIC OXYHYDROXIDE - VELPHORO</u> | | | | | | |
| N 205109 | 001 | 6174442 | Dec 19, 2019 | DS | U-1468 | |
| | | 9561251 | Jan 23, 2030 | DP | U-1468 | |
| <u>SUFENTANIL CITRATE - DSUVIA</u> | | | | | | |
| N 209128 | 001 | 8202535 | Oct 22, 2030 | | U-1351 | NP Nov 02, 2021 |
| | | 8226978 | Jan 05, 2027 | DP | U-1351 | |
| | | 8231900 | Jan 05, 2027 | DP | | |
| | | 8252328 | Jan 05, 2027 | DP | | |
| | | 8252329 | Jan 05, 2027 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------------------------|--------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>SUFENTANIL CITRATE - DSUVIA</u> | | | | | | |
| N 209128 | 001 8535714 | Jan 05, 2027 | DP U-1351 | | | |
| | 8574189 | Mar 16, 2030 | DP | | | |
| | 8778393 | Jan 05, 2027 | U-1351 | | | |
| | 8778394 | Jan 05, 2027 | U-1351 | | | |
| | 8865211 | Jan 05, 2027 | U-1351 | | | |
| | 8865743 | Oct 22, 2030 | U-1351 | | | |
| | 8945592 | Jul 29, 2031 | DP | | | |
| | 9320710 | Jan 05, 2027 | U-1351 | | | |
| | 9744129 | Jan 05, 2027 | DP | | | |
| <u>SUGAMMADEX SODIUM - BRIDION</u> | | | | | | |
| N 022225 | 001 6949527 | Jan 27, 2021 | U-1795 | | NCE | Dec 15, 2020 |
| | 7265099 | Aug 07, 2020 | U-1795 | | | |
| | RE44733 | Jan 27, 2021 | DS DP U-1794 | | | |
| <u>SUGAMMADEX SODIUM - BRIDION</u> | | | | | | |
| N 022225 | 002 6949527 | Jan 27, 2021 | U-1795 | | NCE | Dec 15, 2020 |
| | 7265099 | Aug 07, 2020 | U-1795 | | | |
| | RE44733 | Jan 27, 2021 | DS DP U-1794 | | | |
| <u>SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES - LUMASON</u> | | | | | | |
| N 203684 | 001 5686060 | Nov 11, 2019 | DS DP | | I-728 NCE | Mar 31, 2019 Oct 10, 2019 |
| <u>SUMATRIPTAN SUCCINATE - SUMAVEL DOSEPRO</u> | | | | | | |
| N 022239 | 001 7776007 | Nov 22, 2026 | DP | | | |
| | 7901385 | Jul 31, 2026 | DP | | | |
| | 8118771 | Aug 10, 2023 | DP | | | |
| | 8241243 | Aug 10, 2023 | DP | | | |
| | 8241244 | Nov 21, 2022 | DP | | | |
| | 8267903 | Mar 18, 2023 | DP | | | |
| | 8287489 | Dec 06, 2024 | DP | | | |
| | 8343130 | Oct 18, 2022 | DP | | | |
| | 8491524 | Nov 21, 2022 | DP | | | |
| <u>SUMATRIPTAN SUCCINATE - ALSUMA</u> | | | | | | |
| N 022377 | 001 7811254 | Aug 26, 2027 | DP U-1083 | | | |
| <u>SUMATRIPTAN SUCCINATE - ZECURITY</u> | | | | | | |
| N 202278 | 001 6745071 | Feb 21, 2023 | DP | | | |
| | 7973058 | Apr 12, 2027 | U-1328 | | | |
| | 8155737 | Apr 12, 2027 | U-1328 | | | |
| | 8366600 | Apr 21, 2029 | U-1327 | | | |
| | 8470853 | Apr 12, 2027 | U-1328 | | | |
| | 8597272 | Apr 12, 2027 | DP | | | |
| | 8983594 | Nov 19, 2030 | DP U-1328 | | | |
| | 9272137 | Sep 07, 2027 | DP | | | |
| | 9327114 | Oct 08, 2032 | DP U-1328 | | | |
| | 9427578 | Apr 12, 2027 | DP U-1328 | | | |
| <u>SUMATRIPTAN SUCCINATE - ONZETRA XSAIL</u> | | | | | | |
| N 206099 | 001 10076614 | Oct 20, 2034 | DP | | | |
| | 10076615 | Jul 30, 2029 | U-2010 | | | |
| | 10076615 | Jul 30, 2029 | U-2011 | | | |
| | 10076615 | Jul 30, 2029 | U-2404 | | | |
| | 10124132 | Mar 06, 2027 | DP U-1719 | | | |
| | 10124132 | Mar 06, 2027 | DP U-2010 | | | |
| | 10124132 | Mar 06, 2027 | DP U-2011 | | | |
| | 6715485 | Mar 03, 2020 | DP | | | |
| | 7975690 | Aug 18, 2025 | DP U-1809 | | | |
| | 8047202 | Jul 02, 2023 | DP | | | |
| | 8327844 | Oct 03, 2023 | U-1809 | | | |
| | 8550073 | Oct 22, 2029 | DP | | | |
| | 8555877 | Mar 03, 2020 | DP | | | |
| | 8590530 | Sep 15, 2025 | DP U-1809 | | | |
| | 8875704 | Apr 07, 2028 | DP U-1809 | | | |
| | 8899229 | Aug 18, 2030 | DP | | | |
| | 8978647 | Dec 06, 2030 | DP | | | |
| | 9108015 | Sep 15, 2025 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>SUMATRIPTAN SUCCINATE - ONZETRA XSAIL</u> | | | | | | |
| N 206099 | 001 | 9119932 | Apr 23, 2024 | DP | | |
| | | 9649456 | Oct 21, 2030 | DP U-1719 | | |
| | | 9649456 | Oct 21, 2030 | DP U-2010 | | |
| | | 9649456 | Oct 21, 2030 | DP U-2011 | | |
| <u>SUNITINIB MALATE - SUTENT</u> | | | | | | |
| N 021938 | 001 | 6573293 | Feb 15, 2021 | DS DP U-1154 | I-755 | Nov 16, 2020 |
| | | 6573293 | Feb 15, 2021 | DS DP U-2171 | | |
| | | 7125905 | Feb 15, 2021 | DS DP | | |
| | | 7211600 | Dec 22, 2020 | U-883 | | |
| <u>SUNITINIB MALATE - SUTENT</u> | | | | | | |
| N 021938 | 002 | 6573293 | Feb 15, 2021 | DS DP U-1154 | I-755 | Nov 16, 2020 |
| | | 6573293 | Feb 15, 2021 | DS DP U-2171 | | |
| | | 7125905 | Feb 15, 2021 | DS DP | | |
| | | 7211600 | Dec 22, 2020 | U-883 | | |
| <u>SUNITINIB MALATE - SUTENT</u> | | | | | | |
| N 021938 | 003 | 6573293 | Feb 15, 2021 | DS DP U-1154 | I-755 | Nov 16, 2020 |
| | | 6573293 | Feb 15, 2021 | DS DP U-2171 | | |
| | | 7125905 | Feb 15, 2021 | DS DP | | |
| | | 7211600 | Dec 22, 2020 | U-883 | | |
| <u>SUNITINIB MALATE - SUTENT</u> | | | | | | |
| N 021938 | 004 | 6573293 | Feb 15, 2021 | DS DP U-1154 | I-755 | Nov 16, 2020 |
| | | 6573293 | Feb 15, 2021 | DS DP U-2171 | | |
| | | 7125905 | Feb 15, 2021 | DS DP | | |
| | | 7211600 | Dec 22, 2020 | U-883 | | |
| <u>SUVOREXANT - BELSOMRA</u> | | | | | | |
| N 204569 | 001 | 10098892 | May 29, 2033 | DP | NCE | Aug 13, 2019 |
| | | 7951797 | Nov 20, 2029 | DS DP U-620 | | |
| <u>SUVOREXANT - BELSOMRA</u> | | | | | | |
| N 204569 | 002 | 10098892 | May 29, 2033 | DP | NCE | Aug 13, 2019 |
| | | 7951797 | Nov 20, 2029 | DS DP U-620 | | |
| <u>SUVOREXANT - BELSOMRA</u> | | | | | | |
| N 204569 | 003 | 10098892 | May 29, 2033 | DP | NCE | Aug 13, 2019 |
| | | 7951797 | Nov 20, 2029 | DS DP U-620 | | |
| <u>SUVOREXANT - BELSOMRA</u> | | | | | | |
| N 204569 | 004 | 10098892 | May 29, 2033 | DP | NCE | Aug 13, 2019 |
| | | 7951797 | Nov 20, 2029 | DS DP U-620 | | |
| <u>TACROLIMUS - ASTAGRAF XL</u> | | | | | | |
| N 204096 | 001 | 6440458 | Mar 25, 2019 | DP | | |
| | | 6576259 | Mar 25, 2019 | DP U-1420 | | |
| | | 6884433 | Mar 25, 2019 | DP U-1420 | | |
| | | 8551522 | Mar 25, 2019 | DP | | |
| <u>TACROLIMUS - ASTAGRAF XL</u> | | | | | | |
| N 204096 | 002 | 6440458 | Mar 25, 2019 | DP | | |
| | | 6576259 | Mar 25, 2019 | DP U-1420 | | |
| | | 6884433 | Mar 25, 2019 | DP U-1420 | | |
| | | 8551522 | Mar 25, 2019 | DP | | |
| <u>TACROLIMUS - ASTAGRAF XL</u> | | | | | | |
| N 204096 | 003 | 6440458 | Mar 25, 2019 | DP | | |
| | | 6576259 | Mar 25, 2019 | DP U-1420 | | |
| | | 6884433 | Mar 25, 2019 | DP U-1420 | | |
| | | 8551522 | Mar 25, 2019 | DP | | |
| <u>TACROLIMUS - ENVARUS XR</u> | | | | | | |
| N 206406 | 001 | 7994214 | Aug 30, 2024 | DP | ODE-94 | Jul 10, 2022 |
| | | 8486993 | Aug 30, 2024 | DP U-1752 | | |
| | | 8586084 | Aug 30, 2024 | U-1752 | | |
| | | 8591946 | Aug 30, 2024 | DP | | |
| | | 8617599 | Aug 30, 2024 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TACROLIMUS - ENVARUSUS XR</u> | | | | | | |
| N 206406 001 | 8623410 | Aug 30, 2024 | DP | | | |
| | 8623411 | Aug 30, 2024 | U-1752 | | | |
| | 8664239 | May 30, 2028 | U-1752 | | | |
| | 8685998 | May 30, 2028 | DP U-1752 | | | |
| | 8889185 | Aug 30, 2024 | U-1752 | | | |
| | 8889186 | Aug 30, 2024 | U-1752 | | | |
| | 9161907 | Aug 30, 2024 | DP U-1752 | | | |
| | 9549918 | May 30, 2028 | DP | | | |
| | 9757362 | Aug 30, 2024 | DP | | | |
| | 9763920 | Aug 30, 2024 | DP | | | |
| <u>TACROLIMUS - ENVARUSUS XR</u> | | | | | | |
| N 206406 002 | 7994214 | Aug 30, 2024 | DP | | ODE-94 | Jul 10, 2022 |
| | 8486993 | Aug 30, 2024 | DP U-1752 | | | |
| | 8586084 | Aug 30, 2024 | U-1752 | | | |
| | 8591946 | Aug 30, 2024 | DP | | | |
| | 8617599 | Aug 30, 2024 | DP | | | |
| | 8623410 | Aug 30, 2024 | DP | | | |
| | 8623411 | Aug 30, 2024 | U-1752 | | | |
| | 8664239 | May 30, 2028 | U-1752 | | | |
| | 8685998 | May 30, 2028 | DP U-1752 | | | |
| | 8889185 | Aug 30, 2024 | U-1752 | | | |
| | 8889186 | Aug 30, 2024 | U-1752 | | | |
| | 9161907 | Aug 30, 2024 | DP U-1752 | | | |
| | 9549918 | May 30, 2028 | DP | | | |
| | 9757362 | Aug 30, 2024 | DP | | | |
| | 9763920 | Aug 30, 2024 | DP | | | |
| <u>TACROLIMUS - ENVARUSUS XR</u> | | | | | | |
| N 206406 003 | 7994214 | Aug 30, 2024 | DP | | ODE-94 | Jul 10, 2022 |
| | 8486993 | Aug 30, 2024 | DP U-1752 | | | |
| | 8586084 | Aug 30, 2024 | U-1752 | | | |
| | 8591946 | Aug 30, 2024 | DP | | | |
| | 8617599 | Aug 30, 2024 | DP | | | |
| | 8623410 | Aug 30, 2024 | DP | | | |
| | 8623411 | Aug 30, 2024 | U-1752 | | | |
| | 8664239 | May 30, 2028 | U-1752 | | | |
| | 8685998 | May 30, 2028 | DP U-1752 | | | |
| | 8889185 | Aug 30, 2024 | U-1752 | | | |
| | 8889186 | Aug 30, 2024 | U-1752 | | | |
| | 9161907 | Aug 30, 2024 | DP U-1752 | | | |
| | 9549918 | May 30, 2028 | DP | | | |
| | 9757362 | Aug 30, 2024 | DP | | | |
| | 9763920 | Aug 30, 2024 | DP | | | |
| <u>TADALAFIL - TADALAFIL</u> | | | | | | |
| A 090141 001 | | | | | PC | Mar 26, 2019 |
| <u>TADALAFIL - TADALAFIL</u> | | | | | | |
| A 090141 002 | | | | | PC | Mar 26, 2019 |
| <u>TADALAFIL - TADALAFIL</u> | | | | | | |
| A 090141 003 | | | | | PC | Mar 26, 2019 |
| <u>TADALAFIL - TADALAFIL</u> | | | | | | |
| A 090141 004 | | | | | PC | Mar 26, 2019 |
| <u>TADALAFIL - TADALAFIL</u> | | | | | | |
| A 200630 001 | | | | | PC | Feb 05, 2019 |
| <u>TADALAFIL - CIALIS</u> | | | | | | |
| N 021368 001 | 6821975 | Nov 19, 2020 | DS DP U-1184 | Y | M-219 | Feb 15, 2021 |
| | 6821975 | Nov 19, 2020 | DS DP U-533 | Y | PED | Aug 15, 2021 |
| | 6821975 | Nov 19, 2020 | DS DP U-614 | Y | | |
| | 6821975*PED | May 19, 2021 | | | | |
| | 6943166 | Apr 26, 2020 | U-1184 | | | |
| | 6943166 | Apr 26, 2020 | U-155 | | | |
| | 6943166 | Apr 26, 2020 | U-614 | | | |
| | 6943166*PED | Oct 26, 2020 | | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TADALAFIL - CIALIS</u> | | | | | | |
| N 021368 001 | 7182958 | Apr 26, 2020 | DP U-1184 | Y | | |
| | 7182958 | Apr 26, 2020 | DP U-155 | Y | | |
| | 7182958*PED | Oct 26, 2020 | | | | |
| <u>TADALAFIL - CIALIS</u> | | | | | | |
| N 021368 002 | 6821975 | Nov 19, 2020 | DS DP U-533 | Y | M-219 | Feb 15, 2021 |
| | 6821975 | Nov 19, 2020 | DS DP U-614 | Y | PED | Aug 15, 2021 |
| | 6821975*PED | May 19, 2021 | | | | |
| | 6943166 | Apr 26, 2020 | U-155 | | | |
| | 6943166 | Apr 26, 2020 | U-614 | | | |
| | 6943166*PED | Oct 26, 2020 | | | | |
| | 7182958 | Apr 26, 2020 | DP U-155 | Y | | |
| | 7182958*PED | Oct 26, 2020 | | | | |
| <u>TADALAFIL - CIALIS</u> | | | | | | |
| N 021368 003 | 6821975 | Nov 19, 2020 | DS DP U-533 | Y | M-219 | Feb 15, 2021 |
| | 6821975 | Nov 19, 2020 | DS DP U-614 | Y | PED | Aug 15, 2021 |
| | 6821975*PED | May 19, 2021 | | | | |
| | 6943166 | Apr 26, 2020 | U-614 | | | |
| | 6943166*PED | Oct 26, 2020 | | | | |
| | 7182958 | Apr 26, 2020 | DP U-155 | Y | | |
| | 7182958*PED | Oct 26, 2020 | | | | |
| <u>TADALAFIL - CIALIS</u> | | | | | | |
| N 021368 004 | 6821975 | Nov 19, 2020 | DS DP U-533 | Y | M-219 | Feb 15, 2021 |
| | 6821975 | Nov 19, 2020 | DS DP U-614 | Y | PED | Aug 15, 2021 |
| | 6821975*PED | May 19, 2021 | | | | |
| | 6943166 | Apr 26, 2020 | U-155 | | | |
| | 6943166*PED | Oct 26, 2020 | | | | |
| | 7182958 | Apr 26, 2020 | DP U-155 | Y | | |
| | 7182958*PED | Oct 26, 2020 | | | | |
| <u>TADALAFIL - ADCIRCA</u> | | | | | | |
| N 022332 001 | 6821975 | Nov 19, 2020 | DS DP | Y | | |
| | 6821975*PED | May 19, 2021 | | | | |
| | 7182958 | Apr 26, 2020 | DP | Y | | |
| | 7182958*PED | Oct 26, 2020 | | | | |
| <u>TAFENOQUINE SUCCINATE - ARAKODA</u> | | | | | | |
| N 210607 001 | | | | | NCE NP | Jul 20, 2023 Aug 08, 2021 |
| <u>TAFENOQUINE SUCCINATE - KRINTAFEL</u> | | | | | | |
| N 210795 001 | | | | | NCE ODE-201 | Jul 20, 2023 Jul 20, 2025 |
| <u>TAFLUPROST - ZIOPTAN</u> | | | | | | |
| N 202514 001 | 5886035 | Dec 18, 2022 | DS DP U-778 | | | |
| | 9999593 | May 28, 2029 | DP | | | |
| <u>TALAZOPARIB TOSYLATE - TALZENNA</u> | | | | | | |
| N 211651 001 | 8012976 | Oct 19, 2029 | DS DP | | | |
| | 8420650 | Jul 27, 2029 | DS DP | | | |
| | 8735392 | Oct 20, 2031 | DS DP | | | |
| | 9820985 | Jul 27, 2029 | U-2437 | | | |
| <u>TALAZOPARIB TOSYLATE - TALZENNA</u> | | | | | | |
| N 211651 002 | 8012976 | Oct 19, 2029 | DS DP | | | |
| | 8420650 | Jul 27, 2029 | DS DP | | | |
| | 8735392 | Oct 20, 2031 | DS DP | | | |
| | 9820985 | Jul 27, 2029 | U-2437 | | | |
| <u>TALC - STERITALC</u> | | | | | | |
| N 205555 001 | | | | | ODE-143 ODE-191 | May 01, 2024 May 01, 2024 |
| <u>TALC - STERITALC</u> | | | | | | |
| N 205555 002 | | | | | ODE-143 ODE-191 | May 01, 2024 May 01, 2024 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|----------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TALC - STERITALC</u> | | | | | | |
| N 205555 | 003 | | | | ODE-143 ODE-191 | May 01, 2024 May 01, 2024 |
| <u>TALIGLUCERASE ALFA - ELELYSO</u> | | | | | | |
| N 022458 | 001 | 8227230 | Feb 24, 2024 | DS DP | | |
| | | 8741620 | Feb 24, 2024 | DS DP | | |
| | | 8790641 | Oct 18, 2025 | U-1564 | | |
| | | 8790641 | Oct 18, 2025 | U-1574 | | |
| <u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u> | | | | | | |
| N 022304 | 001 | 7994364 | Jun 27, 2025 | DS DP U-931 | | |
| | | RE39593 | Aug 05, 2022 | DS DP U-931 | | |
| <u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u> | | | | | | |
| N 022304 | 002 | 7994364 | Jun 27, 2025 | DS DP U-931 | | |
| | | RE39593 | Aug 05, 2022 | DS DP U-931 | | |
| <u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u> | | | | | | |
| N 022304 | 003 | 7994364 | Jun 27, 2025 | DS DP U-931 | | |
| | | RE39593 | Aug 05, 2022 | DS DP U-931 | | |
| <u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u> | | | | | | |
| N 200533 | 001 | 7994364 | Jun 27, 2025 | DS DP U-1178 | | |
| | | 7994364 | Jun 27, 2025 | DS DP U-1276 | | |
| | | 8075872 | Nov 20, 2023 | DP | | |
| | | 8114383 | Oct 10, 2024 | DP | Y | |
| | | 8309060 | Nov 20, 2023 | DP U-1178 | | |
| | | 8309060 | Nov 20, 2023 | DP U-1276 | | |
| | | 8420056 | Nov 20, 2023 | DP | | |
| | | 8536130 | Sep 22, 2028 | U-1276 | | |
| | | RE39593 | Aug 05, 2022 | DS DP U-1178 | | |
| | | RE39593 | Aug 05, 2022 | DS DP U-1276 | | |
| <u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u> | | | | | | |
| N 200533 | 002 | 7994364 | Jun 27, 2025 | DS DP U-1178 | | |
| | | 7994364 | Jun 27, 2025 | DS DP U-1276 | | |
| | | 8075872 | Nov 20, 2023 | DP | | |
| | | 8114383 | Oct 10, 2024 | DP | Y | |
| | | 8309060 | Nov 20, 2023 | DP U-1178 | | |
| | | 8309060 | Nov 20, 2023 | DP U-1276 | | |
| | | 8420056 | Nov 20, 2023 | DP | | |
| | | 8536130 | Sep 22, 2028 | U-1276 | | |
| | | RE39593 | Aug 05, 2022 | DS DP U-1178 | | |
| | | RE39593 | Aug 05, 2022 | DS DP U-1276 | | |
| <u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u> | | | | | | |
| N 200533 | 003 | 7994364 | Jun 27, 2025 | DS DP U-1178 | | |
| | | 7994364 | Jun 27, 2025 | DS DP U-1276 | | |
| | | 8075872 | Nov 20, 2023 | DP | | |
| | | 8114383 | Oct 10, 2024 | DP | Y | |
| | | 8309060 | Nov 20, 2023 | DP U-1178 | | |
| | | 8309060 | Nov 20, 2023 | DP U-1276 | | |
| | | 8420056 | Nov 20, 2023 | DP | | |
| | | 8536130 | Sep 22, 2028 | U-1276 | | |
| | | RE39593 | Aug 05, 2022 | DS DP U-1178 | | |
| | | RE39593 | Aug 05, 2022 | DS DP U-1276 | | |
| <u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u> | | | | | | |
| N 200533 | 004 | 7994364 | Jun 27, 2025 | DS DP U-1178 | | |
| | | 7994364 | Jun 27, 2025 | DS DP U-1276 | | |
| | | 8075872 | Nov 20, 2023 | DP | | |
| | | 8114383 | Oct 10, 2024 | DP | Y | |
| | | 8309060 | Nov 20, 2023 | DP U-1178 | | |
| | | 8309060 | Nov 20, 2023 | DP U-1276 | | |
| | | 8420056 | Nov 20, 2023 | DP | | |
| | | 8536130 | Sep 22, 2028 | U-1276 | | |
| | | RE39593 | Aug 05, 2022 | DS DP U-1178 | | |
| | | RE39593 | Aug 05, 2022 | DS DP U-1276 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u> | | | | | | |
| N 200533 | 005 | 7994364 | Jun 27, 2025 | DS DP U-1178 | | |
| | | 7994364 | Jun 27, 2025 | DS DP U-1276 | | |
| | | 8075872 | Nov 20, 2023 | DP | | |
| | | 8114383 | Oct 10, 2024 | DP | Y | |
| | | 8309060 | Nov 20, 2023 | DP U-1178 | | |
| | | 8309060 | Nov 20, 2023 | DP U-1276 | | |
| | | 8420056 | Nov 20, 2023 | DP | | |
| | | 8536130 | Sep 22, 2028 | U-1276 | | |
| | | RE39593 | Aug 05, 2022 | DS DP U-1178 | | |
| | | RE39593 | Aug 05, 2022 | DS DP U-1276 | | |
| <u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u> | | | | | | |
| N 203794 | 001 | 7994364 | Jun 27, 2025 | DS DP U-1289 | | |
| | | RE39593 | Aug 05, 2022 | DS DP U-1289 | | |
| <u>TASIMELTEON - HETLIOZ</u> | | | | | | |
| N 205677 | 001 | 10071977 | Feb 12, 2035 | DS DP | NCE | Jan 31, 2019 |
| | | 10149829 | Jan 25, 2033 | U-2477 | ODE-59 | Jan 31, 2021 |
| | | 5856529 | Dec 09, 2022 | DS DP U-2149 | | |
| | | 9060995 | Jan 25, 2033 | U-1710 | | |
| | | 9539234 | Jan 25, 2033 | U-1934 | | |
| | | 9549913 | Jan 25, 2033 | U-1486 | | |
| | | 9730910 | May 17, 2034 | U-2085 | | |
| | | 9855241 | Jan 25, 2033 | U-2149 | | |
| | | RE46604 | Jan 25, 2033 | U-2147 | | |
| <u>TAVABOROLE - KERYDIN</u> | | | | | | |
| N 204427 | 001 | 7582621 | May 26, 2027 | U-2016 | Y NCE | Jul 07, 2019 |
| | | 7582621*PED | Nov 26, 2027 | | | |
| | | 9549938 | Feb 16, 2026 | U-1951 | | |
| | | 9549938*PED | Aug 16, 2026 | | | |
| | | 9566289 | Feb 16, 2026 | DP | | |
| | | 9566289*PED | Aug 16, 2026 | | | |
| | | 9566290 | Feb 16, 2026 | U-1970 | | |
| | | 9566290*PED | Aug 16, 2026 | | | |
| | | 9572823 | Feb 16, 2026 | U-1970 | | |
| | | 9572823*PED | Aug 16, 2026 | | | |
| <u>TAZAROTENE - FABIOR</u> | | | | | | |
| N 202428 | 001 | 8808716 | Feb 24, 2030 | DP | | |
| <u>TECHNETIUM TC-99M SULFUR COLLOID KIT - AN-SULFUR COLLOID</u> | | | | | | |
| N 017858 | 001 | | | | ODE-29 | Aug 13, 2019 |
| <u>TECHNETIUM TC-99M TEBOROXIME KIT - CARDIOTEC</u> | | | | | | |
| N 019928 | 001 | 6056941 | Jul 28, 2019 | DP | | |
| <u>TECHNETIUM TC-99M TETROFOSMIN KIT - MYOVIEW 30ML</u> | | | | | | |
| N 020372 | 002 | 9549999 | Mar 10, 2030 | DP | | |
| <u>TECHNETIUM TC-99M TILMANOCEPT - LYMPHOSEEK KIT</u> | | | | | | |
| N 202207 | 001 | 6409990 | May 12, 2020 | DS | ODE-67 | Jun 13, 2021 |
| | | 9439985 | Sep 27, 2033 | DS DP | | |
| <u>TECOVIRIMAT - TPOXX</u> | | | | | | |
| N 208627 | 001 | 7737168 | May 03, 2027 | U-2346 | NCE | Jul 13, 2023 |
| | | 8039504 | Jul 23, 2027 | DP | ODE-200 | Jul 13, 2025 |
| | | 8124643 | Jun 18, 2024 | DS DP | | |
| | | 8530509 | Jun 18, 2024 | DP | | |
| | | 8802714 | Jun 18, 2024 | U-2346 | | |
| | | 9339466 | Mar 23, 2031 | DS DP | | |
| <u>TEDIZOLID PHOSPHATE - SIVEXTRO</u> | | | | | | |
| N 205435 | 001 | 7816379 | Feb 23, 2028 | DS DP U-282 | NCE | Jun 20, 2019 |
| | | 8420676 | Feb 23, 2028 | DS DP U-282 | GAIN | Jun 20, 2024 |
| | | 8426389 | Dec 31, 2030 | DS DP U-282 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TEDIZOLID PHOSPHATE - SIVEXTRO</u> | | | | | | |
| N 205436 001 | 7816379 | Feb 23, 2028 | DS DP U-282 | | NCE | Jun 20, 2019 |
| | 8420676 | Feb 23, 2028 | DS DP U-282 | | GAIN | Jun 20, 2024 |
| | 8426389 | Dec 31, 2030 | DS DP U-282 | | | |
| <u>TEDUGLUTIDE RECOMBINANT - GATTEX KIT</u> | | | | | | |
| N 203441 001 | 5789379 | Apr 14, 2020 | DS DP U-1320 | | ODE-37 | Dec 21, 2019 |
| | 7056886 | Sep 18, 2022 | DP U-1320 | | | |
| | 7847061 | Nov 01, 2025 | U-1320 | | | |
| | 9060992 | Nov 01, 2025 | U-1320 | | | |
| | 9539310 | Nov 01, 2025 | U-1320 | | | |
| | 9545434 | Nov 01, 2025 | U-1320 | | | |
| | 9545435 | Nov 01, 2025 | U-1320 | | | |
| | 9555079 | Nov 01, 2025 | U-1320 | | | |
| | 9572867 | Nov 01, 2025 | U-1320 | | | |
| | 9592273 | Nov 01, 2025 | U-1320 | | | |
| | 9592274 | Nov 01, 2025 | U-1320 | | | |
| | 9968655 | Nov 01, 2025 | U-2308 | | | |
| | 9968656 | Nov 01, 2025 | U-2308 | | | |
| | 9968658 | Nov 01, 2025 | U-1320 | | | |
| | 9974835 | Nov 01, 2025 | U-1320 | | | |
| | 9974837 | Nov 01, 2025 | U-1320 | | | |
| | 9981014 | Nov 01, 2025 | U-1320 | | | |
| | 9981016 | Nov 01, 2025 | U-1320 | | | |
| | 9987334 | Nov 01, 2025 | U-1320 | | | |
| | 9987335 | Nov 01, 2025 | U-1320 | | | |
| | 9993528 | Nov 01, 2025 | U-1320 | | | |
| <u>TELAPREVIR - INCIVEK</u> | | | | | | |
| N 201917 001 | 7820671 | Feb 25, 2025 | DS DP | | | |
| | 8431615 | May 30, 2028 | U-1398 | | | |
| | 8529882 | Aug 31, 2021 | U-1398 | | | |
| <u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u> | | | | | | |
| N 022110 001 | 6635618 | Sep 11, 2023 | DS DP U-728 | | | |
| | 6858584 | Aug 24, 2022 | DP | | | |
| | 6872701 | Jun 05, 2021 | DP | | | |
| | 7008923 | May 06, 2021 | U-1005 | | | |
| | 7208471 | May 01, 2021 | DS DP | | | |
| | 7351691 | May 01, 2021 | DS DP U-728 | | | |
| | 7531623 | Jan 01, 2027 | DS | | | |
| | 7544364 | May 01, 2021 | DP | | | |
| | 7700550 | May 01, 2021 | U-282 | | | |
| | 8101575 | May 01, 2021 | DP | | | |
| | 8158580 | May 01, 2021 | DP | | | |
| <u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u> | | | | | | |
| N 022110 002 | 6635618 | Sep 11, 2023 | DS DP U-728 | | | |
| | 6858584 | Aug 24, 2022 | DP | | | |
| | 6872701 | Jun 05, 2021 | DP | | | |
| | 7008923 | May 06, 2021 | U-1005 | | | |
| | 7208471 | May 01, 2021 | DS DP | | | |
| | 7351691 | May 01, 2021 | DS DP U-728 | | | |
| | 7531623 | Jan 01, 2027 | DS | | | |
| | 7544364 | May 01, 2021 | DP | | | |
| | 7700550 | May 01, 2021 | U-282 | | | |
| | 8101575 | May 01, 2021 | DP | | | |
| | 8158580 | May 01, 2021 | DP | | | |
| <u>TELBIVUDINE - TYZEKA</u> | | | | | | |
| N 022011 001 | 6395716 | Aug 10, 2019 | U-782 | | | |
| | 6444652 | Aug 10, 2019 | U-782 | | | |
| | 6566344 | Aug 10, 2019 | U-782 | | | |
| | 6569837 | Oct 25, 2020 | U-782 | | | |
| | 6569837 | Oct 25, 2020 | U-999 | | | |
| | 7589079 | Sep 11, 2023 | DS DP U-999 | | | |
| | 7795238 | Aug 10, 2019 | U-999 | | | |
| | 7858594 | Sep 11, 2023 | DS DP U-999 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TELBIVUDINE - TYZEKA</u> | | | | | | |
| N 022154 | 001 | 6395716 | Aug 10, 2019 | U-999 | | |
| | | 6444652 | Aug 10, 2019 | U-999 | | |
| | | 6566344 | Aug 10, 2019 | U-999 | | |
| | | 6569837 | Oct 25, 2020 | U-999 | | |
| | | 7795238 | Aug 10, 2019 | U-999 | | |
| | | 7858594 | Sep 11, 2023 | DS DP U-999 | | |
| <u>TELMISARTAN - MICARDIS</u> | | | | | | |
| N 020850 | 001 | 6358986 | Jan 10, 2020 | | | |
| <u>TELMISARTAN - MICARDIS</u> | | | | | | |
| N 020850 | 002 | 6358986 | Jan 10, 2020 | | | |
| | | 7998953 | Jun 06, 2020 | U-1177 | | |
| | | 8003679 | Oct 06, 2022 | U-1176 | | |
| <u>TELMISARTAN - MICARDIS</u> | | | | | | |
| N 020850 | 003 | 6358986 | Jan 10, 2020 | | | |
| <u>TELOTTRISTAT ETIPRATE - XERMELO</u> | | | | | | |
| N 208794 | 001 | 7553840 | Dec 11, 2027 | DS | NCE | Feb 28, 2022 |
| | | 7709493 | Dec 11, 2027 | DS U-1979 | ODE-132 | Feb 28, 2024 |
| | | 7968559 | Dec 11, 2027 | U-1979 | | |
| | | 8193204 | Feb 27, 2031 | DS | | |
| | | 8653094 | Dec 19, 2028 | U-1979 | | |
| <u>TEMOZOLOMIDE - TEMODAR</u> | | | | | | |
| N 022277 | 001 | 6987108 | Sep 08, 2023 | DP | | |
| | | 7786118 | Feb 21, 2023 | DP | | |
| | | 8623868 | Feb 21, 2023 | DP | | |
| <u>TEMSIROLIMUS - TORISEL</u> | | | | | | |
| N 022088 | 001 | 5362718 | Feb 15, 2019 | DS DP | Y | |
| | | 5362718*PED | Aug 15, 2019 | | | |
| | | 8026276 | Jan 20, 2026 | DP | | |
| | | 8299116 | Jul 25, 2023 | DP | | |
| | | 8455539 | Jul 25, 2023 | DP | | |
| | | 8455539*PED | Jan 25, 2024 | | | |
| | | 8722700 | Jul 25, 2023 | DP | | |
| | | 8722700*PED | Jan 25, 2024 | | | |
| | | 8791097 | May 10, 2032 | U-1550 | | |
| | | 8791097 | May 10, 2032 | U-1551 | | |
| | | 8791097*PED | Nov 10, 2032 | | | |
| | | RE44768 | Feb 15, 2019 | DS DP | | |
| | | RE44768*PED | Aug 15, 2019 | | | |
| <u>TENOFOVIR ALAFENAMIDE FUMARATE - VEMLIDY</u> | | | | | | |
| N 208464 | 001 | 7390791 | May 07, 2022 | DS DP | NCE | Nov 05, 2020 |
| | | 7803788 | Feb 02, 2022 | U-999 | NP | Nov 11, 2019 |
| | | 8754065 | Aug 15, 2032 | DS DP U-999 | | |
| | | 9296769 | Aug 15, 2032 | DS DP U-999 | | |
| <u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u> | | | | | | |
| N 021356 | 001 | | | | NPP | Dec 11, 2021 |
| | | | | | PED | Jun 11, 2022 |
| <u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u> | | | | | | |
| N 021356 | 002 | | | | NPP | Dec 11, 2021 |
| | | | | | PED | Jun 11, 2022 |
| <u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u> | | | | | | |
| N 021356 | 003 | | | | NPP | Dec 11, 2021 |
| | | | | | PED | Jun 11, 2022 |
| <u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u> | | | | | | |
| N 021356 | 004 | | | | NPP | Dec 11, 2021 |
| | | | | | PED | Jun 11, 2022 |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u> | | | | | | |
| N 022577 | 001 | | | | NPP PED | Dec 11, 2021 Jun 11, 2022 |
| <u>TERIFLUNOMIDE - AUBAGIO</u> | | | | | | |
| N 202992 | 001 | 6794410 | Sep 12, 2026 | U-1285 | | |
| | | 8802735 | Sep 14, 2030 | DP | | |
| | | 9186346 | Feb 04, 2034 | U-1786 | | |
| <u>TERIFLUNOMIDE - AUBAGIO</u> | | | | | | |
| N 202992 | 002 | 6794410 | Sep 12, 2026 | U-1285 | | |
| | | 8802735 | Sep 14, 2030 | DP | | |
| | | 9186346 | Feb 04, 2034 | U-1786 | | |
| <u>TERIPARATIDE RECOMBINANT HUMAN - FORTEO</u> | | | | | | |
| N 021318 | 001 | 6977077 | Aug 19, 2019 | U-597 | | |
| | | 7163684 | Aug 19, 2019 | U-790 | | |
| | | 7351414 | Aug 19, 2019 | U-865 | | |
| | | 7517334 | Mar 25, 2025 | DP | | |
| <u>TERIPARATIDE RECOMBINANT HUMAN - FORTEO</u> | | | | | | |
| N 021318 | 002 | 6977077 | Aug 19, 2019 | U-982 | | |
| | | 6977077 | Aug 19, 2019 | U-994 | | |
| | | 7163684 | Aug 19, 2019 | U-983 | | |
| | | 7163684 | Aug 19, 2019 | U-994 | | |
| | | 7351414 | Aug 19, 2019 | U-984 | | |
| | | 7351414 | Aug 19, 2019 | U-994 | | |
| | | 7517334 | Mar 25, 2025 | DP | | |
| <u>TESAMORELIN ACETATE - EGRIFTA</u> | | | | | | |
| N 022505 | 001 | 5861379 | May 26, 2020 | DS DP U-1100 | | |
| | | 7144577 | Jul 14, 2020 | U-1100 | | |
| | | 7316997 | Aug 14, 2023 | U-1100 | | |
| | | 8314066 | Aug 14, 2023 | U-1100 | | |
| | | 8435945 | Aug 14, 2023 | U-1100 | | |
| <u>TESAMORELIN ACETATE - EGRIFTA</u> | | | | | | |
| N 022505 | 002 | 5861379 | May 26, 2020 | DS DP U-1100 | | |
| | | 7144577 | Jul 14, 2020 | U-1100 | | |
| | | 7316997 | Aug 14, 2023 | U-1100 | | |
| | | 8314066 | Aug 14, 2023 | U-1100 | | |
| | | 8435945 | Aug 14, 2023 | U-1100 | | |
| <u>TESTOSTERONE - TESTOSTERONE</u> | | | | | | |
| A 204268 | 001 | | | | PC | Apr 10, 2019 |
| <u>TESTOSTERONE - TESTODERM TTS</u> | | | | | | |
| N 020791 | 001 | 6348210 | Nov 10, 2019 | U-440 | | |
| <u>TESTOSTERONE - ANDROGEL</u> | | | | | | |
| N 021015 | 001 | 6503894 | Aug 30, 2020 | U-490 | | |
| | | 9125816 | Aug 30, 2020 | U-490 | | |
| | | 9125816*PED | Mar 02, 2021 | | | |
| | | 9132089 | Aug 30, 2020 | U-490 | | |
| | | 9132089*PED | Mar 02, 2021 | | | |
| <u>TESTOSTERONE - ANDROGEL</u> | | | | | | |
| N 021015 | 002 | 6503894 | Aug 30, 2020 | U-490 | | |
| | | 9125816 | Aug 30, 2020 | U-490 | | |
| | | 9125816*PED | Mar 02, 2021 | | | |
| | | 9132089 | Aug 30, 2020 | U-490 | | |
| | | 9132089*PED | Mar 02, 2021 | | | |
| <u>TESTOSTERONE - ANDROGEL</u> | | | | | | |
| N 021015 | 003 | 6503894 | Aug 30, 2020 | U-490 | | |
| | | 9125816 | Aug 30, 2020 | U-490 | | |
| | | 9125816*PED | Mar 02, 2021 | | | |
| | | 9132089 | Aug 30, 2020 | U-490 | | |
| | | 9132089*PED | Mar 02, 2021 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------|-------------|------------------------|--------------|-------------------------|---------------------|-----------------------------|
| <u>TESTOSTERONE - TESTIM</u> | | | | | | |
| N 021454 001 | 7320968 | Jan 18, 2025 | | | | |
| | 7608605 | Apr 21, 2023 | | | U-843 | |
| | 7608606 | Apr 21, 2023 | | | U-1009 | |
| | 7608607 | Apr 21, 2023 | | | U-1009 | |
| | 7608608 | Apr 21, 2023 | | | U-1009 | |
| | 7608609 | Apr 21, 2023 | | | U-1009 | |
| | 7608610 | Apr 21, 2023 | | | U-1009 | |
| | 7935690 | Apr 21, 2023 | | | U-1009 | |
| | 8063029 | Apr 21, 2023 | | | U-843 | |
| | 8178518 | Apr 21, 2023 | DP | | | |
| <u>TESTOSTERONE - STRIANT</u> | | | | | | |
| N 021543 001 | 6248358 | Aug 23, 2019 | | | U-527 | |
| <u>TESTOSTERONE - ANDROGEL</u> | | | | | | |
| N 022309 001 | 6503894 | Aug 30, 2020 | | | U-1103 | |
| | 6503894*PED | Mar 02, 2021 | | | | |
| | 8466136 | Oct 12, 2026 | DP | | | |
| | 8466137 | Oct 12, 2026 | | | U-1103 | |
| | 8466138 | Oct 12, 2026 | | | U-1103 | |
| | 8486925 | Oct 12, 2026 | DP | | | |
| | 8729057 | Oct 12, 2026 | DP | | | |
| | 8741881 | Oct 12, 2026 | | | U-1103 | |
| | 8754070 | Oct 12, 2026 | DP | | | |
| | 8759329 | Oct 12, 2026 | DP | | | |
| | 9125816 | Aug 30, 2020 | | | U-1103 | |
| | 9125816*PED | Mar 02, 2021 | | | | |
| | 9132089 | Aug 30, 2020 | | | U-1103 | |
| | 9132089*PED | Mar 02, 2021 | | | | |
| <u>TESTOSTERONE - ANDROGEL</u> | | | | | | |
| N 022309 002 | 6503894 | Aug 30, 2020 | | | U-1103 | |
| | 6503894*PED | Mar 02, 2021 | | | | |
| | 8466136 | Oct 12, 2026 | DP | | | |
| | 8466137 | Oct 12, 2026 | | | U-1103 | |
| | 8466138 | Oct 12, 2026 | | | U-1103 | |
| | 8486925 | Oct 12, 2026 | DP | | | |
| | 8729057 | Oct 12, 2026 | DP | | | |
| | 8741881 | Oct 12, 2026 | | | U-1103 | |
| | 8754070 | Oct 12, 2026 | DP | | | |
| | 8759329 | Oct 12, 2026 | DP | | | |
| | 9125816 | Aug 30, 2020 | | | U-1103 | |
| | 9125816*PED | Mar 02, 2021 | | | | |
| | 9132089 | Aug 30, 2020 | | | U-1103 | |
| | 9132089*PED | Mar 02, 2021 | | | | |
| <u>TESTOSTERONE - ANDROGEL</u> | | | | | | |
| N 022309 003 | 6503894 | Aug 30, 2020 | | | U-1103 | |
| | 6503894*PED | Mar 02, 2021 | | | | |
| | 8466136 | Oct 12, 2026 | DP | | | |
| | 8466137 | Oct 12, 2026 | | | U-1103 | |
| | 8466138 | Oct 12, 2026 | | | U-1103 | |
| | 8486925 | Oct 12, 2026 | DP | | | |
| | 8729057 | Oct 12, 2026 | DP | | | |
| | 8741881 | Oct 12, 2026 | | | U-1103 | |
| | 8754070 | Oct 12, 2026 | DP | | | |
| | 8759329 | Oct 12, 2026 | DP | | | |
| | 9125816 | Aug 30, 2020 | | | U-1103 | |
| | 9125816*PED | Mar 02, 2021 | | | | |
| | 9132089 | Aug 30, 2020 | | | U-1103 | |
| | 9132089*PED | Mar 02, 2021 | | | | |
| <u>TESTOSTERONE - AXIRON</u> | | | | | | |
| N 022504 001 | 8419307 | Feb 26, 2027 | | | U-1386 | |
| | 8435944 | Sep 27, 2027 | | | U-1390 | |
| | 8784878 | Jul 13, 2023 | DP | | U-1545 | |
| | 8807861 | Feb 26, 2027 | DP | | U-1563 | |
| | 8993520 | Jun 02, 2026 | | | U-1390 | |
| | 9180194 | Jun 02, 2026 | | | U-1390 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------|-----------|------------------------|--------------|-------------------------|---------------------|-----------------------------|
| <u>TESTOSTERONE - AXIRON</u> | | | | | | |
| N 022504 001 | 9289586 | Feb 26, 2027 | U-1390 | | | |
| <u>TESTOSTERONE - VOGELXO</u> | | | | | | |
| N 204399 002 | 8785426 | Feb 11, 2034 | DP U-1531 | | | |
| | 9295675 | Feb 11, 2034 | DP U-1531 | | | |
| | 9662340 | Feb 11, 2034 | DP U-1531 | | | |
| <u>TESTOSTERONE - VOGELXO</u> | | | | | | |
| N 204399 003 | 8785426 | Feb 11, 2034 | DP U-1531 | | | |
| | 9295675 | Feb 11, 2034 | DP U-1531 | | | |
| | 9662340 | Feb 11, 2034 | DP U-1531 | | | |
| <u>TESTOSTERONE - NATESTO</u> | | | | | | |
| N 205488 001 | 8574622 | Feb 04, 2024 | DP | | | |
| | 8784869 | Feb 04, 2024 | DP | | | |
| | 8784882 | Feb 04, 2024 | DP U-1557 | | | |
| | 8877230 | Feb 04, 2024 | U-1616 | | | |
| <u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u> | | | | | | |
| N 209863 001 | 8021335 | Oct 04, 2026 | DP | | NP | Sep 28, 2021 |
| | 8562564 | Jan 24, 2026 | DP | | | |
| | 9180259 | Jan 24, 2026 | DP | | | |
| | 9533102 | Jan 24, 2026 | DP | | | |
| | 9629959 | Jan 24, 2026 | DP | | | |
| | 9744302 | Nov 19, 2035 | DP | | | |
| | 9950125 | Sep 04, 2036 | DP U-2418 | | | |
| | RE44846 | Aug 10, 2019 | DP | | | |
| | RE44847 | Aug 10, 2019 | DP U-2419 | | | |
| <u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u> | | | | | | |
| N 209863 002 | 8021335 | Oct 04, 2026 | DP | | NP | Sep 28, 2021 |
| | 8562564 | Jan 24, 2026 | DP | | | |
| | 9180259 | Jan 24, 2026 | DP | | | |
| | 9533102 | Jan 24, 2026 | DP | | | |
| | 9629959 | Jan 24, 2026 | DP | | | |
| | 9744302 | Nov 19, 2035 | DP | | | |
| | 9950125 | Sep 04, 2036 | DP U-2418 | | | |
| | RE44846 | Aug 10, 2019 | DP | | | |
| | RE44847 | Aug 10, 2019 | DP U-2419 | | | |
| <u>TESTOSTERONE ENANTHATE - XYOSTED (AUTOINJECTOR)</u> | | | | | | |
| N 209863 003 | 8021335 | Oct 04, 2026 | DP | | NP | Sep 28, 2021 |
| | 8562564 | Jan 24, 2026 | DP | | | |
| | 9180259 | Jan 24, 2026 | DP | | | |
| | 9533102 | Jan 24, 2026 | DP | | | |
| | 9629959 | Jan 24, 2026 | DP | | | |
| | 9744302 | Nov 19, 2035 | DP | | | |
| | 9950125 | Sep 04, 2036 | DP U-2418 | | | |
| | RE44846 | Aug 10, 2019 | DP | | | |
| | RE44847 | Aug 10, 2019 | DP U-2419 | | | |
| <u>TESTOSTERONE UNDECANOATE - AVEED</u> | | | | | | |
| N 022219 001 | 7718640 | Mar 14, 2027 | DP | | | |
| | 8338395 | Feb 27, 2026 | U-1500 | | | |
| <u>THALIDOMIDE - THALOMID</u> | | | | | | |
| N 020785 001 | 6315720 | Oct 23, 2020 | U-442 | | | |
| | 6315720 | Oct 23, 2020 | U-731 | | | |
| | 6561977 | Oct 23, 2020 | U-371 | | | |
| | 6561977 | Oct 23, 2020 | U-731 | | | |
| | 6755784 | Oct 23, 2020 | U-371 | | | |
| | 6755784 | Oct 23, 2020 | U-731 | | | |
| | 6869399 | Oct 23, 2020 | U-371 | | | |
| | 6869399 | Oct 23, 2020 | U-731 | | | |
| | 6869399 | Oct 23, 2020 | U-732 | | | |
| | 6869399 | Oct 23, 2020 | U-733 | | | |
| | 7141018 | Oct 23, 2020 | U-371 | | | |
| | 7141018 | Oct 23, 2020 | U-731 | | | |
| | 7141018 | Oct 23, 2020 | U-732 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>THALIDOMIDE - THALOMID</u> | | | | | | |
| N 020785 001 | 7141018 | Oct 23, 2020 | U-733 | | | |
| | 7230012 | Dec 09, 2023 | DP | | | |
| | 7959566 | Oct 23, 2020 | U-1155 | | | |
| | 8315886 | Oct 23, 2020 | U-1249 | | | |
| | 8626531 | Oct 23, 2020 | U-1465 | | | |
| <u>THALIDOMIDE - THALOMID</u> | | | | | | |
| N 020785 002 | 6315720 | Oct 23, 2020 | U-442 | | | |
| | 6315720 | Oct 23, 2020 | U-731 | | | |
| | 6561977 | Oct 23, 2020 | U-371 | | | |
| | 6561977 | Oct 23, 2020 | U-731 | | | |
| | 6755784 | Oct 23, 2020 | U-371 | | | |
| | 6755784 | Oct 23, 2020 | U-731 | | | |
| | 6869399 | Oct 23, 2020 | U-371 | | | |
| | 6869399 | Oct 23, 2020 | U-731 | | | |
| | 6869399 | Oct 23, 2020 | U-732 | | | |
| | 6869399 | Oct 23, 2020 | U-733 | | | |
| | 7141018 | Oct 23, 2020 | U-371 | | | |
| | 7141018 | Oct 23, 2020 | U-731 | | | |
| | 7141018 | Oct 23, 2020 | U-732 | | | |
| | 7141018 | Oct 23, 2020 | U-733 | | | |
| | 7230012 | Dec 09, 2023 | DP | | | |
| | 7959566 | Oct 23, 2020 | U-1155 | | | |
| | 8315886 | Oct 23, 2020 | U-1249 | | | |
| | 8626531 | Oct 23, 2020 | U-1465 | | | |
| <u>THALIDOMIDE - THALOMID</u> | | | | | | |
| N 020785 003 | 6315720 | Oct 23, 2020 | U-442 | | | |
| | 6315720 | Oct 23, 2020 | U-731 | | | |
| | 6561977 | Oct 23, 2020 | U-371 | | | |
| | 6561977 | Oct 23, 2020 | U-731 | | | |
| | 6755784 | Oct 23, 2020 | U-371 | | | |
| | 6755784 | Oct 23, 2020 | U-731 | | | |
| | 6869399 | Oct 23, 2020 | U-371 | | | |
| | 6869399 | Oct 23, 2020 | U-731 | | | |
| | 6869399 | Oct 23, 2020 | U-732 | | | |
| | 6869399 | Oct 23, 2020 | U-733 | | | |
| | 7141018 | Oct 23, 2020 | U-371 | | | |
| | 7141018 | Oct 23, 2020 | U-731 | | | |
| | 7141018 | Oct 23, 2020 | U-732 | | | |
| | 7141018 | Oct 23, 2020 | U-733 | | | |
| | 7230012 | Dec 09, 2023 | DP | | | |
| | 7959566 | Oct 23, 2020 | U-1155 | | | |
| | 8315886 | Oct 23, 2020 | U-1249 | | | |
| | 8626531 | Oct 23, 2020 | U-1465 | | | |
| <u>THALIDOMIDE - THALOMID</u> | | | | | | |
| N 020785 004 | 6315720 | Oct 23, 2020 | U-731 | | | |
| | 6561977 | Oct 23, 2020 | U-731 | | | |
| | 6755784 | Oct 23, 2020 | U-731 | | | |
| | 6869399 | Oct 23, 2020 | U-731 | | | |
| | 7141018 | Oct 23, 2020 | U-731 | | | |
| | 7959566 | Oct 23, 2020 | U-1155 | | | |
| | 8315886 | Oct 23, 2020 | U-1249 | | | |
| | 8626531 | Oct 23, 2020 | U-1465 | | | |
| <u>THIOTEPA - TEPADINA</u> | | | | | | |
| N 208264 001 | | | | | I-747 ODE-129 | Jan 26, 2020 Jan 26, 2024 |
| <u>THIOTEPA - TEPADINA</u> | | | | | | |
| N 208264 002 | | | | | I-747 ODE-129 | Jan 26, 2020 Jan 26, 2024 |
| <u>TICAGRELOR - BRILINTA</u> | | | | | | |
| N 022433 001 | 6525060 | Dec 02, 2019 | DS DP U-1171 | | | Y |
| | 6525060 | Dec 02, 2019 | DS DP U-1860 | | | Y |
| | 6525060 | Dec 02, 2019 | DS DP U-1862 | | | Y |
| | 6525060 | Dec 02, 2019 | DS DP U-1863 | | | Y |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TICAGRELOR - BRILINTA</u> | | | | | | |
| N 022433 001 | 7250419 | Dec 02, 2019 | DS DP U-1171 | | | |
| | 7250419 | Dec 02, 2019 | DS DP U-1860 | | | |
| | 7250419 | Dec 02, 2019 | DS DP U-1864 | | | |
| | 7250419 | Dec 02, 2019 | DS DP U-1865 | | | |
| | 7250419 | Dec 02, 2019 | DS DP U-1866 | | | |
| | 7250419 | Dec 02, 2019 | DS DP U-1867 | | | |
| | 7265124 | Jul 09, 2021 | DS DP U-1171 | | | |
| | 7265124 | Jul 09, 2021 | DS DP U-1860 | | | |
| | 7265124 | Jul 09, 2021 | DS DP U-1868 | | | |
| | 7265124 | Jul 09, 2021 | DS DP U-1869 | | | |
| | 8425934 | Apr 17, 2030 | DP | | | |
| | RE46276 | Oct 30, 2024 | DS DP U-1935 | | | |
| | RE46276 | Oct 30, 2024 | DS DP U-1936 | | | |
| | RE46276 | Oct 30, 2024 | DS DP U-1937 | | | |
| | RE46276 | Oct 30, 2024 | DS DP U-1938 | | | |
| <u>TICAGRELOR - BRILINTA</u> | | | | | | |
| N 022433 002 | 6525060 | Dec 02, 2019 | DS DP U-1171 | | | Y |
| | 6525060 | Dec 02, 2019 | DS DP U-1860 | | | Y |
| | 6525060 | Dec 02, 2019 | DS DP U-1862 | | | Y |
| | 6525060 | Dec 02, 2019 | DS DP U-1863 | | | Y |
| | 7250419 | Dec 02, 2019 | DS DP U-1171 | | | |
| | 7250419 | Dec 02, 2019 | DS DP U-1860 | | | |
| | 7250419 | Dec 02, 2019 | DS DP U-1864 | | | |
| | 7250419 | Dec 02, 2019 | DS DP U-1865 | | | |
| | 7250419 | Dec 02, 2019 | DS DP U-1866 | | | |
| | 7250419 | Dec 02, 2019 | DS DP U-1867 | | | |
| | 7265124 | Jul 09, 2021 | DS DP U-1171 | | | |
| | 7265124 | Jul 09, 2021 | DS DP U-1860 | | | |
| | 7265124 | Jul 09, 2021 | DS DP U-1868 | | | |
| | 7265124 | Jul 09, 2021 | DS DP U-1869 | | | |
| | 8425934 | Apr 17, 2030 | DP | | | |
| | RE46276 | Oct 30, 2024 | DS DP U-1935 | | | |
| | RE46276 | Oct 30, 2024 | DS DP U-1936 | | | |
| | RE46276 | Oct 30, 2024 | DS DP U-1937 | | | |
| | RE46276 | Oct 30, 2024 | DS DP U-1938 | | | |
| <u>TIGECYCLINE - TYGACIL</u> | | | | | | |
| N 021821 001 | 7879828 | Feb 05, 2029 | DP | | | |
| | 8372995 | Oct 08, 2030 | DP | | | |
| | 8975242 | Oct 24, 2028 | DP | | | |
| | 9254328 | Mar 13, 2026 | DP | | | |
| | 9694078 | Mar 13, 2026 | DP | | | |
| <u>TIGECYCLINE - TIGECYCLINE</u> | | | | | | |
| N 211158 001 | 9855335 | Apr 07, 2033 | DP | | | |
| <u>TIMOLOL MALEATE - TIMOLOL MALEATE</u> | | | | | | |
| N 020963 001 | 6174524 | Mar 26, 2019 | DP | | | |
| <u>TIMOLOL MALEATE - TIMOLOL MALEATE</u> | | | | | | |
| N 020963 002 | 6174524 | Mar 26, 2019 | DP | | | |
| <u>TIOTROPIUM BROMIDE - SPIRIVA</u> | | | | | | |
| N 021395 001 | 6777423 | Sep 24, 2021 | DS DP | | | |
| | 6777423*PED | Mar 24, 2022 | | | | |
| | 6908928 | Sep 24, 2021 | DS DP U-566 | | | |
| | 6908928 | Sep 24, 2021 | DS DP U-762 | | | |
| | 6908928*PED | Mar 24, 2022 | | | | |
| | 7070800 | Jan 22, 2022 | DP U-566 | | | |
| | 7070800*PED | Jul 22, 2022 | | | | |
| | 7309707 | Sep 24, 2021 | DS DP | | | |
| | 7309707*PED | Mar 24, 2022 | | | | |
| | 7642268 | Sep 24, 2021 | DS DP | | | |
| | 7642268*PED | Mar 24, 2022 | | | | |
| | 7694676 | Mar 12, 2027 | DP | | | |
| | 7694676*PED | Sep 12, 2027 | | | | |
| | 8022082 | Jan 19, 2026 | DP U-1186 | | | |
| | 8022082*PED | Jul 19, 2026 | | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TIOTROPIUM BROMIDE - SPIRIVA</u> | | | | | | |
| N 021395 | 001 | 9010323 | | | | |
| | | RE38912 | | | | |
| | | RE38912*PED | | | | |
| | | Apr 19, 2030 | DP | | | |
| | | Oct 11, 2021 | DP | | | |
| | | Apr 11, 2022 | | | | |
| <u>TIOTROPIUM BROMIDE - SPIRIVA RESPIMAT</u> | | | | | | |
| N 021936 | 001 | 6846413*PED | | | NPP | Feb 15, 2020 |
| | | 6977042*PED | | | PED | Aug 15, 2020 |
| | | 6988496 | | DP | | |
| | | 6988496*PED | | | | |
| | | 7284474 | | DP | | |
| | | 7284474*PED | | | | |
| | | 7396341 | | DP | | |
| | | 7396341*PED | | | | |
| | | 7802568 | | DP | | |
| | | 7802568*PED | | | | |
| | | 7837235 | | DP | | |
| | | 7837235*PED | | | | |
| | | 7896264 | | DP | | |
| | | 7896264*PED | | | | |
| | | 7988001 | | DP | | |
| | | 7988001*PED | | | | |
| | | 8733341 | | DP | | |
| | | 8733341*PED | | | | |
| | | 9027967 | | DP | | |
| | | 9027967*PED | | | | |
| <u>TIOTROPIUM BROMIDE - SPIRIVA RESPIMAT</u> | | | | | | |
| N 021936 | 002 | 6846413*PED | | | NPP | Feb 15, 2020 |
| | | 6977042*PED | | | PED | Aug 15, 2020 |
| | | 6988496 | | DP | | |
| | | 6988496*PED | | | | |
| | | 7284474 | | DP | | |
| | | 7284474*PED | | | | |
| | | 7396341 | | DP | | |
| | | 7396341*PED | | | | |
| | | 7802568 | | DP | | |
| | | 7802568*PED | | | | |
| | | 7837235 | | DP | | |
| | | 7837235*PED | | | | |
| | | 7896264 | | DP | | |
| | | 7896264*PED | | | | |
| | | 7988001 | | DP | | |
| | | 7988001*PED | | | | |
| | | 8733341 | | DP | | |
| | | 8733341*PED | | | | |
| | | 9027967 | | DP | | |
| | | 9027967*PED | | | | |
| <u>TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE - LONSURE</u> | | | | | | |
| N 207981 | 001 | 6479500 | | | | |
| | | 9527833 | | DS DP | | |
| | | RE46284 | | | | |
| | | Mar 16, 2020 | | U-1751 | NCE | Sep 22, 2020 |
| | | Jun 17, 2034 | | | | |
| | | Dec 16, 2026 | | U-1751 | | |
| <u>TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE - LONSURE</u> | | | | | | |
| N 207981 | 002 | 6479500 | | | | |
| | | 9527833 | | DS DP | | |
| | | RE46284 | | | | |
| | | Mar 16, 2020 | | U-1751 | NCE | Sep 22, 2020 |
| | | Jun 17, 2034 | | | | |
| | | Dec 16, 2026 | | U-1751 | | |
| <u>TIPRANAVIR - APTIVUS</u> | | | | | | |
| N 021814 | 001 | 5852195 | | | | |
| | | 6147095 | | DS | | |
| | | Jun 22, 2019 | | | | |
| | | Oct 29, 2019 | | U-670 | | |
| <u>TIPRANAVIR - APTIVUS</u> | | | | | | |
| N 022292 | 001 | 5852195 | | | | |
| | | 6147095 | | DS | | |
| | | Jun 22, 2019 | | | | |
| | | Oct 29, 2019 | | U-670 | | |
| <u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u> | | | | | | |
| N 020912 | 001 | 6136794 | | | | |
| | | 6770660 | | | | |
| | | Jan 29, 2019 | | | | |
| | | May 01, 2023 | | U-1444 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|--------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u> | | | | | | |
| N 020912 001 | 6136794 | Jan 29, 2019 | | | | |
| | 6770660 | May 01, 2023 | U-1444 | | | |
| <u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u> | | | | | | |
| N 020912 002 | 6136794 | Jan 29, 2019 | | U-1898 | | |
| | 6770660 | May 01, 2023 | U-1444 | | | |
| <u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u> | | | | | | |
| N 020913 001 | 6136794 | Jan 29, 2019 | | | | |
| | 6770660 | May 01, 2023 | U-1444 | | | |
| <u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u> | | | | | | |
| N 020913 002 | 6136794 | Jan 29, 2019 | | | | |
| | 6770660 | May 01, 2023 | U-1444 | | | |
| <u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u> | | | | | | |
| N 020913 003 | 6136794 | Jan 29, 2019 | | | | |
| | 6770660 | May 01, 2023 | U-1444 | | | |
| <u>TOBRAMYCIN - TOBI PODHALER</u> | | | | | | |
| N 201688 001 | 7368102 | Dec 19, 2022 | DP U-909 | | | |
| | 7442388 | May 10, 2020 | DP | | | |
| | 7516741 | Jan 11, 2024 | DP | | | |
| | 7559325 | Oct 27, 2025 | DP | | | |
| | 8069851 | Sep 24, 2024 | DP | | | |
| | 8349294 | May 10, 2020 | DP | | | |
| | 8715623 | Dec 19, 2022 | DP U-909 | | | |
| <u>TOBRAMYCIN - BETHKIS</u> | | | | | | |
| N 201820 001 | 6987094 | Sep 22, 2022 | DP | | | |
| | 7696178 | Sep 22, 2022 | DP | | | |
| | 7939502 | Jun 14, 2022 | U-1324 | | | |
| <u>TOFACITINIB CITRATE - XELJANZ</u> | | | | | | |
| N 203214 001 | 6956041 | Dec 08, 2020 | DP | | I-761 | Dec 14, 2020 |
| | 6965027 | Mar 25, 2023 | DS | | I-780 | May 30, 2021 |
| | 7091208 | Dec 08, 2020 | | U-247 | | |
| | 7265221 | Dec 08, 2020 | DS | | | |
| | 7301023 | May 23, 2022 | DS | | | |
| | 7842699 | Dec 08, 2020 | | U-2322 | | |
| | RE41783 | Dec 08, 2025 | DS | | | |
| <u>TOFACITINIB CITRATE - XELJANZ</u> | | | | | | |
| N 203214 002 | 6956041 | Dec 08, 2020 | DP | | I-780 | May 30, 2021 |
| | 6965027 | Mar 25, 2023 | DS | | | |
| | 7265221 | Dec 08, 2020 | DS | | | |
| | 7301023 | May 23, 2022 | DS | | | |
| | 7842699 | Dec 08, 2020 | | U-2322 | | |
| | RE41783 | Dec 08, 2025 | DS | | | |
| <u>TOFACITINIB CITRATE - XELJANZ XR</u> | | | | | | |
| N 208246 001 | 6956041 | Dec 08, 2020 | DP | | I-761 | Dec 14, 2020 |
| | 6965027 | Mar 25, 2023 | DS | | | |
| | 7091208 | Dec 08, 2020 | | U-247 | | |
| | 7265221 | Dec 08, 2020 | DS | | | |
| | 7301023 | May 23, 2022 | DS | | | |
| | 9937181 | Mar 14, 2034 | DP | | | |
| | RE41783 | Dec 08, 2025 | DS | | | |
| <u>TOLTERODINE TARTRATE - DETROL LA</u> | | | | | | |
| N 021228 001 | 6630162 | Nov 11, 2019 | DP U-544 | | | |
| | 6770295 | Aug 26, 2019 | DP U-544 | | | |
| | 6911217 | Nov 11, 2019 | DP U-544 | | | |
| | 6911217*PED | May 11, 2020 | | | | |
| <u>TOLTERODINE TARTRATE - DETROL LA</u> | | | | | | |
| N 021228 002 | 6630162 | Nov 11, 2019 | DP U-544 | | | |
| | 6770295 | Aug 26, 2019 | DP U-544 | | | |
| | 6911217 | Nov 11, 2019 | DP U-544 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------|------------------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TOLTERODINE TARTRATE - DETROL LA</u> | | | | | | |
| N 021228 | 002 6911217*PED | May 11, 2020 | | | | |
| <u>TOLVAPTAN - SAMSCA</u> | | | | | | |
| N 022275 | 001 5753677 8501730 | May 19, 2020 Sep 01, 2026 | DS | U-978 | | |
| <u>TOLVAPTAN - SAMSCA</u> | | | | | | |
| N 022275 | 002 5753677 8501730 | May 19, 2020 Sep 01, 2026 | DS | U-978 | | |
| <u>TOLVAPTAN - SAMSCA</u> | | | | | | |
| N 022275 | 003 5753677 8501730 | May 19, 2020 Sep 01, 2026 | DS | U-978 | | |
| <u>TOLVAPTAN - JYNARQUE</u> | | | | | | |
| N 204441 | 001 5753677 8501730 | May 19, 2020 Sep 01, 2026 | DS | U-2307 | I-779 ODE-178 | Apr 23, 2021 Apr 23, 2025 |
| <u>TOLVAPTAN - JYNARQUE</u> | | | | | | |
| N 204441 | 002 5753677 8501730 | May 19, 2020 Sep 01, 2026 | DS | U-2307 | I-779 ODE-178 | Apr 23, 2021 Apr 23, 2025 |
| <u>TOLVAPTAN - JYNARQUE</u> | | | | | | |
| N 204441 | 003 5753677 8501730 | May 19, 2020 Sep 01, 2026 | DS | U-2307 | I-779 ODE-178 | Apr 23, 2021 Apr 23, 2025 |
| <u>TOLVAPTAN - JYNARQUE</u> | | | | | | |
| N 204441 | 004 5753677 8501730 | May 19, 2020 Sep 01, 2026 | DS | U-2307 | I-779 ODE-178 | Apr 23, 2021 Apr 23, 2025 |
| <u>TOLVAPTAN - JYNARQUE</u> | | | | | | |
| N 204441 | 005 5753677 8501730 | May 19, 2020 Sep 01, 2026 | DS | U-2307 | I-779 ODE-178 | Apr 23, 2021 Apr 23, 2025 |
| <u>TOPIRAMATE - TOPAMAX</u> | | | | | | |
| N 020844 | 001 7125560 | Mar 01, 2019 | | U-766 | | |
| <u>TOPIRAMATE - TOPAMAX</u> | | | | | | |
| N 020844 | 002 7125560 | Mar 01, 2019 | | U-766 | | |
| <u>TOPIRAMATE - TOPAMAX SPRINKLE</u> | | | | | | |
| N 020844 | 003 7125560 | Mar 01, 2019 | | U-766 | | |
| <u>TOPIRAMATE - TROKENDI XR</u> | | | | | | |
| N 201635 | 001 8298576 | Apr 04, 2028 | DP | U-106 | | |
| | 8298576 | Apr 04, 2028 | DP | U-1992 | | |
| | 8298580 | Nov 16, 2027 | DP | U-106 | | |
| | 8298580 | Nov 16, 2027 | DP | U-1992 | | |
| | 8663683 | Nov 16, 2027 | DP | U-106 | | |
| | 8663683 | Nov 16, 2027 | DP | U-1992 | | |
| | 8877248 | Nov 16, 2027 | DP | U-106 | | |
| | 8877248 | Nov 16, 2027 | DP | U-1992 | | |
| | 8889191 | Nov 16, 2027 | | U-106 | | |
| | 8889191 | Nov 16, 2027 | | U-1992 | | |
| | 8992989 | Nov 16, 2027 | DP | U-1675 | | |
| | 8992989 | Nov 16, 2027 | DP | U-1992 | | |
| | 9549940 | Nov 16, 2027 | DP | U-1675 | | |
| | 9549940 | Nov 16, 2027 | DP | U-1992 | | |
| | 9555004 | Nov 16, 2027 | DP | U-1675 | | |
| | 9555004 | Nov 16, 2027 | DP | U-1992 | | |
| | 9622983 | Nov 16, 2027 | DP | U-1675 | | |
| | 9622983 | Nov 16, 2027 | DP | U-1992 | | |
| <u>TOPIRAMATE - TROKENDI XR</u> | | | | | | |
| N 201635 | 002 8298576 | Apr 04, 2028 | DP | U-106 | | |
| | 8298576 | Apr 04, 2028 | DP | U-1992 | | |
| | 8298580 | Nov 16, 2027 | DP | U-106 | | |
| | 8298580 | Nov 16, 2027 | DP | U-1992 | | |
| | 8663683 | Nov 16, 2027 | DP | U-106 | | |
| | 8663683 | Nov 16, 2027 | DP | U-1992 | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TOPIRAMATE - TROKENDI XR</u> | | | | | | |
| N 201635 | 002 | 8877248 | Nov 16, 2027 | DP U-106 | | |
| | | 8877248 | Nov 16, 2027 | DP U-1992 | | |
| | | 8889191 | Nov 16, 2027 | U-106 | | |
| | | 8889191 | Nov 16, 2027 | U-1992 | | |
| | | 8992989 | Nov 16, 2027 | DP U-1675 | | |
| | | 8992989 | Nov 16, 2027 | DP U-1992 | | |
| | | 9549940 | Nov 16, 2027 | DP U-1675 | | |
| | | 9549940 | Nov 16, 2027 | DP U-1992 | | |
| | | 9555004 | Nov 16, 2027 | DP U-1675 | | |
| | | 9555004 | Nov 16, 2027 | DP U-1992 | | |
| | | 9622983 | Nov 16, 2027 | DP U-1675 | | |
| | | 9622983 | Nov 16, 2027 | DP U-1992 | | |
| <u>TOPIRAMATE - TROKENDI XR</u> | | | | | | |
| N 201635 | 003 | 8298576 | Apr 04, 2028 | DP U-106 | | |
| | | 8298576 | Apr 04, 2028 | DP U-1992 | | |
| | | 8298580 | Nov 16, 2027 | DP U-106 | | |
| | | 8298580 | Nov 16, 2027 | DP U-1992 | | |
| | | 8663683 | Nov 16, 2027 | DP U-106 | | |
| | | 8663683 | Nov 16, 2027 | DP U-1992 | | |
| | | 8877248 | Nov 16, 2027 | DP U-106 | | |
| | | 8877248 | Nov 16, 2027 | DP U-1992 | | |
| | | 8889191 | Nov 16, 2027 | U-106 | | |
| | | 8889191 | Nov 16, 2027 | U-1992 | | |
| | | 8992989 | Nov 16, 2027 | DP U-1675 | | |
| | | 8992989 | Nov 16, 2027 | DP U-1992 | | |
| | | 9549940 | Nov 16, 2027 | DP U-1675 | | |
| | | 9549940 | Nov 16, 2027 | DP U-1992 | | |
| | | 9555004 | Nov 16, 2027 | DP U-1675 | | |
| | | 9555004 | Nov 16, 2027 | DP U-1992 | | |
| | | 9622983 | Nov 16, 2027 | DP U-1675 | | |
| | | 9622983 | Nov 16, 2027 | DP U-1992 | | |
| <u>TOPIRAMATE - TROKENDI XR</u> | | | | | | |
| N 201635 | 004 | 8298576 | Apr 04, 2028 | DP U-106 | | |
| | | 8298576 | Apr 04, 2028 | DP U-1992 | | |
| | | 8298580 | Nov 16, 2027 | DP U-106 | | |
| | | 8298580 | Nov 16, 2027 | DP U-1992 | | |
| | | 8663683 | Nov 16, 2027 | DP U-106 | | |
| | | 8663683 | Nov 16, 2027 | DP U-1992 | | |
| | | 8877248 | Nov 16, 2027 | DP U-106 | | |
| | | 8877248 | Nov 16, 2027 | DP U-1992 | | |
| | | 8889191 | Nov 16, 2027 | U-106 | | |
| | | 8889191 | Nov 16, 2027 | U-1992 | | |
| | | 8992989 | Nov 16, 2027 | DP U-1675 | | |
| | | 8992989 | Nov 16, 2027 | DP U-1992 | | |
| | | 9549940 | Nov 16, 2027 | DP U-1675 | | |
| | | 9549940 | Nov 16, 2027 | DP U-1992 | | |
| | | 9555004 | Nov 16, 2027 | DP U-1675 | | |
| | | 9555004 | Nov 16, 2027 | DP U-1992 | | |
| | | 9622983 | Nov 16, 2027 | DP U-1675 | | |
| | | 9622983 | Nov 16, 2027 | DP U-1992 | | |
| <u>TOPIRAMATE - OUDEXY XR</u> | | | | | | |
| N 205122 | 001 | 8652527 | Mar 19, 2033 | DP | | |
| | | 8889190 | Mar 19, 2033 | DP | | |
| | | 9101545 | Mar 19, 2033 | DP | | |
| | | 9555005 | Mar 19, 2033 | DP | | |
| <u>TOPIRAMATE - OUDEXY XR</u> | | | | | | |
| N 205122 | 002 | 8652527 | Mar 19, 2033 | DP | | |
| | | 8889190 | Mar 19, 2033 | DP | | |
| | | 9101545 | Mar 19, 2033 | DP | | |
| | | 9555005 | Mar 19, 2033 | DP | | |
| <u>TOPIRAMATE - OUDEXY XR</u> | | | | | | |
| N 205122 | 003 | 8652527 | Mar 19, 2033 | DP | | |
| | | 8889190 | Mar 19, 2033 | DP | | |
| | | 9101545 | Mar 19, 2033 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TOPIRAMATE - OUDEXY XR</u> | | | | | | |
| N 205122 | 003 | 9555005 | Mar 19, 2033 | DP | | |
| <u>TOPIRAMATE - OUDEXY XR</u> | | | | | | |
| N 205122 | 004 | 8652527 | Mar 19, 2033 | DP | | |
| | | 8889190 | Mar 19, 2033 | DP | | |
| | | 9101545 | Mar 19, 2033 | DP | | |
| | | 9555005 | Mar 19, 2033 | DP | | |
| <u>TOPIRAMATE - OUDEXY XR</u> | | | | | | |
| N 205122 | 005 | 8652527 | Mar 19, 2033 | DP | | |
| | | 8889190 | Mar 19, 2033 | DP | | |
| | | 9101545 | Mar 19, 2033 | DP | | |
| | | 9555005 | Mar 19, 2033 | DP | | |
| <u>TOPOTECAN HYDROCHLORIDE - HYCAMTIN</u> | | | | | | |
| N 020981 | 001 | 8158645 | Dec 10, 2024 | DP | | |
| <u>TOPOTECAN HYDROCHLORIDE - HYCAMTIN</u> | | | | | | |
| N 020981 | 002 | 8158645 | Dec 10, 2024 | DP | | |
| <u>TRABECTEDIN - YONDELIS</u> | | | | | | |
| N 207953 | 001 | 8895557 | Jan 07, 2028 | DP | M-232 | Jun 29, 2021 |
| | | 8895557*PED | Jul 07, 2028 | | NCE | Oct 23, 2020 |
| | | | | | ODE-100 | Oct 23, 2022 |
| | | | | | PED | Apr 23, 2021 |
| | | | | | PED | Dec 29, 2021 |
| | | | | | PED | Apr 23, 2023 |
| <u>TRAMADOL HYDROCHLORIDE - ULTRAM</u> | | | | | | |
| N 020281 | 001 | 6339105 | Oct 12, 2019 | U-435 | | |
| <u>TRAMADOL HYDROCHLORIDE - ULTRAM</u> | | | | | | |
| N 020281 | 002 | 6339105 | Oct 12, 2019 | U-435 | | |
| <u>TRAMADOL HYDROCHLORIDE - RYZOLT</u> | | | | | | |
| N 021745 | 001 | 6607748 | Jun 29, 2020 | DP | | |
| | | 7988998 | Oct 27, 2023 | DP | | |
| <u>TRAMADOL HYDROCHLORIDE - RYZOLT</u> | | | | | | |
| N 021745 | 002 | 6607748 | Jun 29, 2020 | DP | | |
| | | 7988998 | Oct 27, 2023 | DP | | |
| <u>TRAMADOL HYDROCHLORIDE - RYZOLT</u> | | | | | | |
| N 021745 | 003 | 6607748 | Jun 29, 2020 | DP | | |
| | | 7988998 | Oct 27, 2023 | DP | | |
| <u>TRAMADOL HYDROCHLORIDE - CONZIP</u> | | | | | | |
| N 022370 | 001 | 7858118 | Apr 11, 2022 | DP | U-1104 | |
| <u>TRAMADOL HYDROCHLORIDE - CONZIP</u> | | | | | | |
| N 022370 | 002 | 7858118 | Apr 11, 2022 | DP | U-1104 | |
| <u>TRAMADOL HYDROCHLORIDE - CONZIP</u> | | | | | | |
| N 022370 | 003 | 7858118 | Apr 11, 2022 | DP | U-1104 | |
| <u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u> | | | | | | |
| N 204114 | 001 | 7378423 | May 29, 2027 | DS DP | I-745 | Jun 22, 2020 |
| | | 8580304 | Jan 28, 2032 | DP | I-778 | Apr 30, 2021 |
| | | 8703781 | Oct 15, 2030 | DS DP | U-1712 | May 04, 2021 |
| | | 8703781 | Oct 15, 2030 | DS DP | U-2033 | Jun 22, 2024 |
| | | 8835443 | Jun 10, 2025 | | U-1581 | Apr 30, 2025 |
| | | 8835443 | Jun 10, 2025 | | U-1582 | May 04, 2025 |
| | | 8835443 | Jun 10, 2025 | | U-2020 | May 29, 2020 |
| | | 8835443 | Jun 10, 2025 | | U-2037 | Jan 08, 2021 |
| | | 8835443 | Jun 10, 2025 | | U-2302 | |
| | | 8835443 | Jun 10, 2025 | | U-2305 | |
| | | 8952018 | Oct 15, 2030 | | U-2020 | |
| | | 9155706 | Jan 28, 2032 | DP | | |
| | | 9271941 | Jan 28, 2032 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u> | | | | | | |
| N 204114 002 | 7378423 | May 29, 2027 | DS DP | | I-745 | Jun 22, 2020 |
| | 8580304 | Jan 28, 2032 | DP | | I-778 | Apr 30, 2021 |
| | 8703781 | Oct 15, 2030 | DS DP U-1712 | | I-781 | May 04, 2021 |
| | 8703781 | Oct 15, 2030 | DS DP U-2033 | | ODE-148 | Jun 22, 2024 |
| | 8835443 | Jun 10, 2025 | U-1581 | | ODE-182 | Apr 30, 2025 |
| | 8835443 | Jun 10, 2025 | U-1582 | | ODE-183 | May 04, 2025 |
| | 8835443 | Jun 10, 2025 | U-2020 | | ODE-48 | May 29, 2020 |
| | 8835443 | Jun 10, 2025 | U-2037 | | ODE-57 | Jan 08, 2021 |
| | 8835443 | Jun 10, 2025 | U-2302 | | | |
| | 8835443 | Jun 10, 2025 | U-2305 | | | |
| | 8952018 | Oct 15, 2030 | U-2020 | | | |
| | 9155706 | Jan 28, 2032 | DP | | | |
| | 9271941 | Jan 28, 2032 | DP | | | |
| <u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u> | | | | | | |
| N 204114 003 | 7378423 | May 29, 2027 | DS DP | | I-745 | Jun 22, 2020 |
| | 8580304 | Jan 28, 2032 | DP | | I-778 | Apr 30, 2021 |
| | 8703781 | Oct 15, 2030 | DS DP U-1712 | | I-781 | May 04, 2021 |
| | 8703781 | Oct 15, 2030 | DS DP U-2033 | | ODE-148 | Jun 22, 2024 |
| | 8835443 | Jun 10, 2025 | U-1581 | | ODE-182 | Apr 30, 2025 |
| | 8835443 | Jun 10, 2025 | U-1582 | | ODE-183 | May 04, 2025 |
| | 8835443 | Jun 10, 2025 | U-2020 | | ODE-48 | May 29, 2020 |
| | 8835443 | Jun 10, 2025 | U-2037 | | ODE-57 | Jan 08, 2021 |
| | 8835443 | Jun 10, 2025 | U-2302 | | | |
| | 8835443 | Jun 10, 2025 | U-2305 | | | |
| | 8952018 | Oct 15, 2030 | U-2020 | | | |
| | 9155706 | Jan 28, 2032 | DP | | | |
| | 9271941 | Jan 28, 2032 | DP | | | |
| <u>TRANEXAMIC ACID - LYSTEDA</u> | | | | | | |
| N 022430 001 | 7947739 | Mar 04, 2025 | DP | | | |
| | 8022106 | Mar 04, 2025 | U-1182 | | | |
| | 8273795 | Mar 04, 2025 | U-1182 | | | |
| | 8487005 | Mar 04, 2025 | DP U-1182 | | | |
| | 8791160 | Mar 04, 2025 | DP U-1182 | | | |
| | 8809394 | Mar 04, 2025 | DP U-1182 | | | |
| | 8957113 | Mar 04, 2025 | DP U-1182 | | | |
| | 9060939 | Mar 04, 2025 | DP | | | |
| <u>TRAVOPROST - TRAVATAN Z</u> | | | | | | |
| N 021994 001 | 8268299 | Oct 13, 2029 | DP | | | |
| | 8323630 | Sep 20, 2027 | DP | | | |
| | 8388941 | Sep 20, 2027 | DP | | | |
| <u>TRAVOPROST - IZBA</u> | | | | | | |
| N 204822 001 | 8178582 | Oct 10, 2029 | DP | | | |
| | 8722735 | Oct 10, 2029 | DP | | | |
| | 8754123 | May 19, 2029 | DP | | | |
| | 9144561 | Mar 13, 2029 | DP | | | |
| <u>TRAZODONE HYDROCHLORIDE - DESYREL</u> | | | | | | |
| N 018207 001 | 8133893 | Mar 13, 2029 | DS DP | | | |
| <u>TRAZODONE HYDROCHLORIDE - DESYREL</u> | | | | | | |
| N 018207 002 | 8133893 | Mar 13, 2029 | DS DP | | | |
| <u>TRAZODONE HYDROCHLORIDE - DESYREL</u> | | | | | | |
| N 018207 003 | 8133893 | Mar 13, 2029 | DS DP | | | |
| <u>TRAZODONE HYDROCHLORIDE - DESYREL</u> | | | | | | |
| N 018207 004 | 8133893 | Mar 13, 2029 | DS DP | | | |
| <u>TRAZODONE HYDROCHLORIDE - OLEPTRO</u> | | | | | | |
| N 022411 001 | 6607748 | Jun 29, 2020 | DP | | | |
| | 7829120 | Mar 27, 2027 | DP U-796 | | | |
| | 8133893 | Mar 13, 2029 | DS DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TRAZODONE HYDROCHLORIDE - OLEPTRO</u> | | | | | | |
| N 022411 | 002 | 6607748 | Jun 29, 2020 | DP | | |
| | | 7829120 | Mar 27, 2027 | DP | U-796 | |
| | | 8133893 | Mar 13, 2029 | DS | DP | |
| <u>TREPROSTINIL - REMODULIN</u> | | | | | | |
| N 021272 | 001 | 10076505 | Dec 16, 2024 | DP | | |
| | | 7999007 | Mar 29, 2029 | DP | U-1437 | |
| | | 8653137 | Sep 05, 2028 | | U-1437 | |
| | | 8658694 | Sep 05, 2028 | | U-1437 | |
| | | 9199908 | May 24, 2024 | | U-1771 | |
| | | 9593066 | Dec 15, 2028 | DS | | |
| | | 9604901 | Dec 15, 2028 | DS | | |
| | | 9713599 | Dec 16, 2024 | | U-2036 | |
| <u>TREPROSTINIL - REMODULIN</u> | | | | | | |
| N 021272 | 002 | 10076505 | Dec 16, 2024 | DP | | |
| | | 7999007 | Mar 29, 2029 | DP | U-1437 | |
| | | 8653137 | Sep 05, 2028 | | U-1437 | |
| | | 8658694 | Sep 05, 2028 | | U-1437 | |
| | | 9199908 | May 24, 2024 | | U-1771 | |
| | | 9593066 | Dec 15, 2028 | DS | | |
| | | 9604901 | Dec 15, 2028 | DS | | |
| | | 9713599 | Dec 16, 2024 | | U-2036 | |
| <u>TREPROSTINIL - REMODULIN</u> | | | | | | |
| N 021272 | 003 | 10076505 | Dec 16, 2024 | DP | | |
| | | 7999007 | Mar 29, 2029 | DP | U-1437 | |
| | | 8653137 | Sep 05, 2028 | | U-1437 | |
| | | 8658694 | Sep 05, 2028 | | U-1437 | |
| | | 9199908 | May 24, 2024 | | U-1771 | |
| | | 9593066 | Dec 15, 2028 | DS | | |
| | | 9604901 | Dec 15, 2028 | DS | | |
| | | 9713599 | Dec 16, 2024 | | U-2036 | |
| <u>TREPROSTINIL - REMODULIN</u> | | | | | | |
| N 021272 | 004 | 10076505 | Dec 16, 2024 | DP | | |
| | | 7999007 | Mar 29, 2029 | DP | U-1437 | |
| | | 8653137 | Sep 05, 2028 | | U-1437 | |
| | | 8658694 | Sep 05, 2028 | | U-1437 | |
| | | 9199908 | May 24, 2024 | | U-1771 | |
| | | 9593066 | Dec 15, 2028 | DS | | |
| | | 9604901 | Dec 15, 2028 | DS | | |
| | | 9713599 | Dec 16, 2024 | | U-2036 | |
| <u>TREPROSTINIL - TYVASO</u> | | | | | | |
| N 022387 | 001 | 8497393 | Dec 15, 2028 | DS | | Y |
| | | 9339507 | Mar 10, 2028 | DP | | |
| | | 9358240 | May 05, 2028 | | U-1849 | |
| | | 9593066 | Dec 15, 2028 | DS | | |
| | | 9604901 | Dec 15, 2028 | DS | | |
| <u>TREPROSTINIL - REMODULIN</u> | | | | | | |
| N 208276 | 001 | 10076505 | Dec 16, 2024 | DP | | |
| | | 9593066 | Dec 15, 2028 | DS | | |
| | | 9604901 | Dec 15, 2028 | DS | | |
| | | 9713599 | Dec 16, 2024 | | U-2036 | |
| <u>TREPROSTINIL - REMODULIN</u> | | | | | | |
| N 208276 | 002 | 10076505 | Dec 16, 2024 | DP | | |
| | | 9593066 | Dec 15, 2028 | DS | | |
| | | 9604901 | Dec 15, 2028 | DS | | |
| | | 9713599 | Dec 16, 2024 | | U-2036 | |
| <u>TREPROSTINIL - REMODULIN</u> | | | | | | |
| N 208276 | 003 | 10076505 | Dec 16, 2024 | DP | | |
| | | 9593066 | Dec 15, 2028 | DS | | |
| | | 9604901 | Dec 15, 2028 | DS | | |
| | | 9713599 | Dec 16, 2024 | | U-2036 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TREPROSTINIL - REMODULIN</u> | | | | | | |
| N 208276 | 004 | 10076505 | Dec 16, 2024 | DP | | |
| | | 9593066 | Dec 15, 2028 | DS | | |
| | | 9604901 | Dec 15, 2028 | DS | | |
| | | 9713599 | Dec 16, 2024 | | U-2036 | |
| <u>TREPROSTINIL DIOLAMINE - ORENITRAM</u> | | | | | | |
| N 203496 | 001 | 7417070 | Jul 30, 2026 | DS | | D-156 Jan 28, 2019 |
| | | 7544713 | Jul 14, 2024 | | U-1475 | D-157 Jan 28, 2019 |
| | | 8252839 | May 24, 2024 | DP | | |
| | | 8349892 | Jan 22, 2031 | DP | | |
| | | 8410169 | Feb 13, 2030 | DP | | |
| | | 8747897 | Oct 08, 2029 | DP | | |
| | | 9050311 | May 24, 2024 | DS DP | | |
| | | 9278901 | May 24, 2024 | | U-1475 | |
| | | 9393203 | Apr 27, 2026 | DP | U-1877 | |
| | | 9422223 | May 24, 2024 | DP | | |
| | | 9593066 | Dec 15, 2028 | DS | | |
| | | 9604901 | Dec 15, 2028 | DS | | |
| <u>TREPROSTINIL DIOLAMINE - ORENITRAM</u> | | | | | | |
| N 203496 | 002 | 7417070 | Jul 30, 2026 | DS | | D-156 Jan 28, 2019 |
| | | 7544713 | Jul 14, 2024 | | U-1475 | D-157 Jan 28, 2019 |
| | | 8252839 | May 24, 2024 | DP | | |
| | | 8349892 | Jan 22, 2031 | DP | | |
| | | 8410169 | Feb 13, 2030 | DP | | |
| | | 8497393 | Dec 15, 2028 | DS | | Y |
| | | 8747897 | Oct 08, 2029 | DP | | |
| | | 9050311 | May 24, 2024 | DS DP | | |
| | | 9278901 | May 24, 2024 | | U-1475 | |
| | | 9393203 | Apr 27, 2026 | DP | U-1877 | |
| | | 9422223 | May 24, 2024 | DP | | |
| | | 9593066 | Dec 15, 2028 | DS | | |
| | | 9604901 | Dec 15, 2028 | DS | | |
| <u>TREPROSTINIL DIOLAMINE - ORENITRAM</u> | | | | | | |
| N 203496 | 003 | 7417070 | Jul 30, 2026 | DS | | D-156 Jan 28, 2019 |
| | | 7544713 | Jul 14, 2024 | | U-1475 | D-157 Jan 28, 2019 |
| | | 8252839 | May 24, 2024 | DP | | |
| | | 8349892 | Jan 22, 2031 | DP | | |
| | | 8410169 | Feb 13, 2030 | DP | | |
| | | 8497393 | Dec 15, 2028 | DS | | Y |
| | | 8747897 | Oct 08, 2029 | DP | | |
| | | 9050311 | May 24, 2024 | DS DP | | |
| | | 9278901 | May 24, 2024 | | U-1475 | |
| | | 9393203 | Apr 27, 2026 | DP | U-1877 | |
| | | 9422223 | May 24, 2024 | DP | | |
| | | 9593066 | Dec 15, 2028 | DS | | |
| | | 9604901 | Dec 15, 2028 | DS | | |
| <u>TREPROSTINIL DIOLAMINE - ORENITRAM</u> | | | | | | |
| N 203496 | 004 | 7417070 | Jul 30, 2026 | DS | | D-156 Jan 28, 2019 |
| | | 7544713 | Jul 14, 2024 | | U-1475 | D-157 Jan 28, 2019 |
| | | 8252839 | May 24, 2024 | DP | | |
| | | 8349892 | Jan 22, 2031 | DP | | |
| | | 8410169 | Feb 13, 2030 | DP | | |
| | | 8497393 | Dec 15, 2028 | DS | | Y |
| | | 8747897 | Oct 08, 2029 | DP | | |
| | | 9050311 | May 24, 2024 | DS DP | | |
| | | 9278901 | May 24, 2024 | | U-1475 | |
| | | 9393203 | Apr 27, 2026 | DP | U-1877 | |
| | | 9422223 | May 24, 2024 | DP | | |
| | | 9593066 | Dec 15, 2028 | DS | | |
| | | 9604901 | Dec 15, 2028 | DS | | |
| <u>TREPROSTINIL DIOLAMINE - ORENITRAM</u> | | | | | | |
| N 203496 | 005 | 7417070 | Jul 30, 2026 | DS | | D-156 Jan 28, 2019 |
| | | 7544713 | Jul 14, 2024 | | U-1475 | D-157 Jan 28, 2019 |
| | | 8252839 | May 24, 2024 | DP | | |
| | | 8349892 | Jan 22, 2031 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>TREPROSTINIL DIOLAMINE - ORENITRAM</u> | | | | | | |
| N 203496 005 | 8410169 | Feb 13, 2030 | DP | | | |
| | 8747897 | Oct 08, 2029 | DP | | | |
| | 9050311 | May 24, 2024 | DS DP | | | |
| | 9278901 | May 24, 2024 | | U-1475 | | |
| | 9393203 | Apr 27, 2026 | DP | U-1877 | | |
| | 9422223 | May 24, 2024 | DP | | | |
| | 9593066 | Dec 15, 2028 | DS | | | |
| | 9604901 | Dec 15, 2028 | DS | | | |
| <u>TRETINOIN - RENOVA</u> | | | | | | |
| N 021108 001 | 6531141 | Mar 07, 2020 | | | | |
| <u>TRETINOIN - ALTRENO</u> | | | | | | |
| N 209353 001 | | | | | NDF | Aug 23, 2021 |
| <u>TRIAMCINOLONE ACETONIDE - TRIESENCE</u> | | | | | | |
| N 022048 001 | 6395294 | Jan 13, 2020 | DP | U-846 | | |
| | 8128960 | Dec 17, 2029 | DP | | | |
| | 8211880 | Mar 10, 2029 | | U-1257 | | |
| | 8211880 | Mar 10, 2029 | | U-1258 | | |
| <u>TRIAMCINOLONE ACETONIDE - ZILRETTA</u> | | | | | | |
| N 208845 001 | 8828440 | Aug 04, 2031 | DP | | NP | Oct 06, 2020 |
| | 9555048 | Aug 04, 2031 | | U-2151 | | |
| <u>TRIPTORELIN PAMOATE - TRIPTODUR KIT</u> | | | | | | |
| N 208956 001 | | | | | NP ODE-149 | Jun 29, 2020 Jun 29, 2024 |
| <u>TROSPIDIUM CHLORIDE - SANCTURA XR</u> | | | | | | |
| N 022103 001 | 7410978 | Feb 01, 2025 | DP | | Y | |
| | 7759359 | Nov 04, 2024 | | U-1071 | Y | |
| | 7763635 | Nov 04, 2024 | | U-1071 | Y | |
| | 7781448 | Nov 04, 2024 | | U-1071 | Y | |
| | 7781449 | Nov 04, 2024 | | U-1071 | Y | |
| <u>TRYPAN BLUE - VISIONBLUE</u> | | | | | | |
| N 021670 001 | 6367480 | Oct 26, 2019 | | U-2321 | | |
| | 6720314 | May 07, 2019 | | U-2321 | | |
| <u>TRYPAN BLUE - MEMBRANEBLUE</u> | | | | | | |
| N 022278 001 | 6372449 | Nov 12, 2019 | | U-2379 | | |
| | 6696430 | May 07, 2019 | | U-2377 | | |
| <u>ULIPRISTAL ACETATE - ELLA</u> | | | | | | |
| N 022474 001 | 8426392 | Jun 12, 2030 | | U-1389 | | |
| | 8512745 | Jun 02, 2030 | DP | | | |
| | 8735380 | Feb 20, 2029 | DP | | | |
| | 8962603 | Jun 12, 2030 | | U-1657 | | |
| | 9283233 | Apr 13, 2030 | | U-1821 | | |
| <u>UMECLIDINIUM BROMIDE - INCRUSE ELLIPTA</u> | | | | | | |
| N 205382 001 | 7488827 | Dec 18, 2027 | DS DP | | M-172 | Feb 24, 2019 |
| | 7498440 | Apr 27, 2025 | DS DP | | | |
| | 8113199 | Oct 23, 2027 | DP | | | |
| | 8161968 | Feb 05, 2028 | DP | | | |
| | 8183257 | Jul 27, 2025 | | U-1476 | | |
| | 8201556 | Feb 05, 2029 | DP | | | |
| | 8309572 | Apr 27, 2025 | | U-1476 | | |
| | 8534281 | Mar 08, 2030 | DP | | | |
| | 8746242 | Oct 11, 2030 | DP | | | |
| | 9333310 | Oct 02, 2027 | DP | | | |
| <u>UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - ANORO ELLIPTA</u> | | | | | | |
| N 203975 001 | 7439393 | May 21, 2025 | DS DP | U-1476 | | |
| | 7488827 | Dec 18, 2027 | DS DP | | | |
| | 7498440 | Apr 27, 2025 | DS DP | | | |
| | 7776895 | Sep 11, 2022 | DP | | | |
| | 8113199 | Oct 23, 2027 | DP | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - ANORO ELLIPTA</u> | | | | | | |
| N 203975 | 001 | 8161968 | Feb 05, 2028 | DP | | |
| | | 8183257 | Jul 27, 2025 | | U-1476 | |
| | | 8309572 | Apr 27, 2025 | | U-1476 | |
| | | 8511304 | Jun 14, 2027 | DP | U-1476 | |
| | | 8534281 | Mar 08, 2030 | DP | | |
| | | 8746242 | Oct 11, 2030 | DP | | |
| | | 9333310 | Oct 02, 2027 | DP | | |
| | | 9750726 | Nov 29, 2030 | DP | | |
| | | RE44874 | Mar 23, 2023 | DS DP | U-1476 | |
| <u>UNOPROSTONE ISOPROPYL - RESCULA</u> | | | | | | |
| N 021214 | 001 | 6458836 | Jul 09, 2021 | | U-1315 | |
| | | 6458836 | Jul 09, 2021 | | U-333 | |
| <u>URIDINE TRIACETATE - VISTOGARD</u> | | | | | | |
| N 208159 | 001 | 6258795 | Jul 10, 2019 | DP | | NCE Sep 04, 2020 |
| | | 7776838 | Aug 17, 2027 | | U-1791 | ODE-104 Dec 11, 2022 |
| <u>URIDINE TRIACETATE - XURIDEN</u> | | | | | | |
| N 208169 | 001 | 6258795 | Jul 10, 2019 | DP | | NCE Sep 04, 2020 |
| | | | | | | ODE-98 Sep 04, 2022 |
| <u>VALBENZAZINE TOSYLATE - INGREZZA</u> | | | | | | |
| N 209241 | 001 | 10065952 | Oct 28, 2036 | DS DP | U-1995 | NCE Apr 11, 2022 |
| | | 8039627 | Oct 06, 2029 | DS DP | | |
| | | 8357697 | Nov 08, 2027 | | U-1995 | |
| <u>VALBENZAZINE TOSYLATE - INGREZZA</u> | | | | | | |
| N 209241 | 002 | 10065952 | Oct 28, 2036 | DS DP | U-1995 | NCE Apr 11, 2022 |
| | | 8039627 | Oct 06, 2029 | DS DP | | |
| | | 8357697 | Nov 08, 2027 | | U-1995 | |
| <u>VALGANCICLOVIR HYDROCHLORIDE - VALCYTE</u> | | | | | | |
| N 022257 | 001 | 8889109 | Dec 11, 2027 | DP | | |
| | | 9642911 | Dec 11, 2027 | DP | | |
| <u>VANDETANIB - CAPRELSA</u> | | | | | | |
| N 022405 | 001 | 7173038 | Aug 14, 2021 | DS DP | | |
| | | 8067427 | Aug 08, 2028 | | DP | |
| | | 8642608 | Feb 06, 2022 | | U-1490 | |
| | | RE42353 | Jun 27, 2022 | DS DP | | |
| <u>VANDETANIB - CAPRELSA</u> | | | | | | |
| N 022405 | 002 | 7173038 | Aug 14, 2021 | DS DP | | |
| | | 8067427 | Aug 08, 2028 | | DP | |
| | | 8642608 | Feb 06, 2022 | | U-1490 | |
| | | RE42353 | Jun 27, 2022 | DS DP | | |
| <u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u> | | | | | | |
| N 021400 | 001 | 8273876 | Jul 23, 2027 | | U-1288 | |
| | | 8841446 | Jul 03, 2023 | DP | | |
| <u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u> | | | | | | |
| N 021400 | 002 | 8273876 | Jul 23, 2027 | | U-1288 | |
| | | 8841446 | Jul 03, 2023 | DP | | |
| <u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u> | | | | | | |
| N 021400 | 003 | 8273876 | Jul 23, 2027 | | U-1288 | |
| | | 8841446 | Jul 03, 2023 | DP | | |
| <u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u> | | | | | | |
| N 021400 | 004 | 8273876 | Jul 23, 2027 | | U-1288 | |
| | | 8841446 | Jul 03, 2023 | DP | | |
| <u>VARDENAFIL HYDROCHLORIDE - STAXYN</u> | | | | | | |
| N 200179 | 001 | 8613950 | Dec 23, 2028 | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-------------------------------------------|-------------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>VARENICLINE TARTRATE - CHANTIX</u> | | | | | | |
| N 021928 001 | 6410550 | May 10, 2020 | DS DP U-56 | | M-183 | Aug 12, 2019 |
| | 6410550*PED | Nov 10, 2020 | | | M-192 | Dec 16, 2019 |
| | 6890927 | May 06, 2022 | DS DP U-56 | | PED | Feb 12, 2020 |
| | 6890927*PED | Nov 06, 2022 | | | PED | Jun 16, 2020 |
| | 7265119 | Aug 03, 2022 | DS DP U-56 | | | |
| | 7265119*PED | Feb 03, 2023 | | | | |
| <u>VARENICLINE TARTRATE - CHANTIX</u> | | | | | | |
| N 021928 002 | 6410550 | May 10, 2020 | DS DP U-56 | | M-183 | Aug 12, 2019 |
| | 6410550*PED | Nov 10, 2020 | | | M-192 | Dec 16, 2019 |
| | 6890927 | May 06, 2022 | DS DP U-56 | | PED | Feb 12, 2020 |
| | 6890927*PED | Nov 06, 2022 | | | PED | Jun 16, 2020 |
| | 7265119 | Aug 03, 2022 | DS DP U-56 | | | |
| | 7265119*PED | Feb 03, 2023 | | | | |
| <u>VASOPRESSIN - VASOSTRICT</u> | | | | | | |
| N 204485 001 | 9375478 | Jan 30, 2035 | | U-1857 | | |
| | 9687526 | Jan 30, 2035 | | U-1857 | | |
| | 9744209 | Jan 30, 2035 | | U-1857 | | |
| | 9744239 | Jan 30, 2035 | | U-1857 | | |
| | 9750785 | Jan 30, 2035 | | DP | | |
| <u>VASOPRESSIN - VASOSTRICT</u> | | | | | | |
| N 204485 002 | 9375478 | Jan 30, 2035 | | U-1857 | | |
| | 9687526 | Jan 30, 2035 | | U-1857 | | |
| | 9744209 | Jan 30, 2035 | | U-1857 | | |
| | 9744239 | Jan 30, 2035 | | U-1857 | | |
| | 9750785 | Jan 30, 2035 | | DP | | |
| | 9937223 | Jan 30, 2035 | | U-1857 | | |
| <u>VEMURAFENIB - ZELBORAF</u> | | | | | | |
| N 202429 001 | 7504509 | Oct 22, 2026 | DS DP | | I-757 | Nov 06, 2020 |
| | 7863288 | Jun 20, 2029 | DS DP | | M-184 | Aug 31, 2019 |
| | 8143271 | Jun 21, 2026 | DS DP | | ODE-158 | Nov 06, 2024 |
| | 8470818 | Aug 02, 2026 | | U-1418 | | |
| | 8470818 | Aug 02, 2026 | | U-2164 | | |
| | 8741920 | Jul 27, 2030 | DS DP | | | |
| | 9447089 | Jun 06, 2032 | | DP | | |
| <u>VENETOCLAX - VENCLEXTA</u> | | | | | | |
| N 208573 001 | 8546399 | Jun 27, 2031 | DS DP | | I-782 | Jun 08, 2021 |
| | 9174982 | May 26, 2030 | | U-2323 | I-789 | Nov 21, 2021 |
| | 9174982 | May 26, 2030 | | U-2445 | M-228 | Jun 08, 2021 |
| | 9174982 | May 26, 2030 | | U-2446 | NCE | Apr 11, 2021 |
| | | | | | ODE-114 | Apr 11, 2023 |
| | | | | | ODE-185 | Jun 08, 2025 |
| | | | | | ODE-211 | Nov 21, 2025 |
| <u>VENETOCLAX - VENCLEXTA</u> | | | | | | |
| N 208573 002 | 8546399 | Jun 27, 2031 | DS DP | | I-782 | Jun 08, 2021 |
| | 9174982 | May 26, 2030 | | U-2323 | I-789 | Nov 21, 2021 |
| | 9174982 | May 26, 2030 | | U-2445 | M-228 | Jun 08, 2021 |
| | 9174982 | May 26, 2030 | | U-2446 | NCE | Apr 11, 2021 |
| | | | | | ODE-114 | Apr 11, 2023 |
| | | | | | ODE-185 | Jun 08, 2025 |
| | | | | | ODE-211 | Nov 21, 2025 |
| <u>VENETOCLAX - VENCLEXTA</u> | | | | | | |
| N 208573 003 | 8546399 | Jun 27, 2031 | DS DP | | I-782 | Jun 08, 2021 |
| | 9174982 | May 26, 2030 | | U-2323 | I-789 | Nov 21, 2021 |
| | 9174982 | May 26, 2030 | | U-2445 | M-228 | Jun 08, 2021 |
| | 9174982 | May 26, 2030 | | U-2446 | NCE | Apr 11, 2021 |
| | | | | | ODE-114 | Apr 11, 2023 |
| | | | | | ODE-185 | Jun 08, 2025 |
| | | | | | ODE-211 | Nov 21, 2025 |
| <u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u> | | | | | | |
| N 022567 001 | 5532241 | Sep 29, 2019 | DS DP | | | |
| | 7834020 | Jun 05, 2022 | DS DP U-839 | | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u> | | | | | | |
| N 022567 | 001 | 8193195 | Jun 05, 2022 | U-839 | | |
| | | 8236804 | Jun 05, 2022 | U-839 | | |
| | | 8673921 | Jun 05, 2022 | DS DP | | |
| <u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u> | | | | | | |
| N 022567 | 002 | 5532241 | Sep 29, 2019 | DS DP | | |
| | | 7834020 | Jun 05, 2022 | DS DP | U-839 | |
| | | 8193195 | Jun 05, 2022 | | U-839 | |
| | | 8236804 | Jun 05, 2022 | | U-839 | |
| | | 8673921 | Jun 05, 2022 | DS DP | | |
| <u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u> | | | | | | |
| N 022567 | 003 | 5532241 | Sep 29, 2019 | DS DP | | |
| | | 7834020 | Jun 05, 2022 | DS DP | U-839 | |
| | | 8193195 | Jun 05, 2022 | | U-839 | |
| | | 8236804 | Jun 05, 2022 | | U-839 | |
| | | 8673921 | Jun 05, 2022 | DS DP | | |
| <u>VINCRIStINE SULFATE - MARQIBO KIT</u> | | | | | | |
| N 202497 | 001 | 6723338 | Mar 31, 2020 | U-1271 | | |
| | | 7247316 | Sep 25, 2020 | DP | | ODE-28 |
| | | 7887836 | Mar 31, 2020 | U-1271 | | Aug 09, 2019 |
| <u>VISMODEGIB - ERIVEDGE</u> | | | | | | |
| N 203388 | 001 | 7888364 | Nov 11, 2028 | DS DP | | |
| | | 9278961 | Dec 15, 2028 | | U-1825 | |
| <u>VORAPAXAR SULFATE - ZONTIVITY</u> | | | | | | |
| N 204886 | 001 | 7235567 | Jun 13, 2021 | DS DP | | |
| | | 7304078 | Apr 06, 2024 | DS DP | U-1512 | |
| | | 7713999 | May 30, 2024 | DS DP | U-2291 | |
| <u>VORINOSTAT - ZOLINZA</u> | | | | | | |
| N 021991 | 001 | 7399787 | Feb 09, 2025 | | U-892 | |
| | | 7456219 | Mar 11, 2027 | DS | | |
| | | 7652069 | Mar 04, 2023 | | DP | |
| | | 7732490 | Mar 04, 2023 | | U-892 | |
| | | 7851509 | Feb 21, 2024 | | DP | U-892 |
| | | 8067472 | Mar 04, 2023 | | U-892 | |
| | | 8093295 | May 16, 2026 | | DP | |
| | | 8101663 | Mar 04, 2023 | | U-892 | |
| | | 8450372 | Mar 18, 2028 | | U-892 | |
| <u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u> | | | | | | |
| N 204447 | 001 | 7144884 | Jun 17, 2026 | DS DP | U-1439 | |
| | | 8476279 | Oct 02, 2022 | | DP | U-1439 |
| | | 8722684 | Jun 30, 2031 | DS DP | | |
| | | 8969355 | Jun 15, 2027 | | U-1668 | |
| | | 9125908 | Jun 15, 2027 | | U-2309 | |
| | | 9125909 | Jun 15, 2027 | | U-2309 | |
| | | 9125910 | Jun 15, 2027 | | U-2309 | |
| | | 9227946 | Jun 15, 2027 | | U-1668 | |
| | | 9278096 | Mar 21, 2032 | | U-2436 | |
| | | 9861630 | Jun 15, 2027 | | U-1668 | |
| <u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u> | | | | | | |
| N 204447 | 002 | 7144884 | Jun 17, 2026 | DS DP | U-1439 | |
| | | 8476279 | Oct 02, 2022 | | DP | U-1439 |
| | | 8722684 | Jun 30, 2031 | DS DP | | |
| | | 8969355 | Jun 15, 2027 | | U-1668 | |
| | | 9125908 | Jun 15, 2027 | | U-2309 | |
| | | 9125909 | Jun 15, 2027 | | U-2309 | |
| | | 9125910 | Jun 15, 2027 | | U-2309 | |
| | | 9227946 | Jun 15, 2027 | | U-1668 | |
| | | 9278096 | Mar 21, 2032 | | U-2436 | |
| | | 9861630 | Jun 15, 2027 | | U-1668 | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|-----------------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u> | | | | | | |
| N 204447 003 | 7144884 | Jun 17, 2026 | DS DP U-1439 | | M-227 | May 02, 2021 |
| | 8476279 | Oct 02, 2022 | DP U-1439 | | M-234 | Oct 19, 2021 |
| | 8722684 | Jun 30, 2031 | DS DP | | | |
| | 8969355 | Jun 15, 2027 | | U-1668 | | |
| | 9125908 | Jun 15, 2027 | | U-2309 | | |
| | 9125909 | Jun 15, 2027 | | U-2309 | | |
| | 9125910 | Jun 15, 2027 | | U-2309 | | |
| | 9227946 | Jun 15, 2027 | | U-1668 | | |
| | 9278096 | Mar 21, 2032 | | U-2436 | | |
| | 9861630 | Jun 15, 2027 | | U-1668 | | |
| <u>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</u> | | | | | | |
| N 204447 004 | 7144884 | Jun 17, 2026 | DS DP U-1439 | | M-227 | May 02, 2021 |
| | 8476279 | Oct 02, 2022 | DP U-1439 | | M-234 | Oct 19, 2021 |
| | 8722684 | Jun 30, 2031 | DS DP | | | |
| | 8969355 | Jun 15, 2027 | | U-1668 | | |
| | 9125908 | Jun 15, 2027 | | U-2309 | | |
| | 9125909 | Jun 15, 2027 | | U-2309 | | |
| | 9125910 | Jun 15, 2027 | | U-2309 | | |
| | 9227946 | Jun 15, 2027 | | U-1668 | | |
| | 9278096 | Mar 21, 2032 | | U-2436 | | |
| | 9861630 | Jun 15, 2027 | | U-1668 | | |
| <u>ZICONOTIDE ACETATE - PRIALT</u> | | | | | | |
| N 021060 001 | 8653033 | Oct 01, 2024 | | U-48 | | |
| | 8653033 | Oct 01, 2024 | | U-55 | | |
| | 8765680 | Oct 01, 2024 | | U-48 | | |
| | 8765680 | Oct 01, 2024 | | U-55 | | |
| | 9707270 | Oct 01, 2024 | | U-2084 | | |
| <u>ZICONOTIDE ACETATE - PRIALT</u> | | | | | | |
| N 021060 002 | 8653033 | Oct 01, 2024 | | U-48 | | |
| | 8653033 | Oct 01, 2024 | | U-55 | | |
| | 8765680 | Oct 01, 2024 | | U-48 | | |
| | 8765680 | Oct 01, 2024 | | U-55 | | |
| | 9707270 | Oct 01, 2024 | | U-2084 | | |
| <u>ZICONOTIDE ACETATE - PRIALT</u> | | | | | | |
| N 021060 003 | 8653033 | Oct 01, 2024 | | U-48 | | |
| | 8653033 | Oct 01, 2024 | | U-55 | | |
| | 8765680 | Oct 01, 2024 | | U-48 | | |
| | 8765680 | Oct 01, 2024 | | U-55 | | |
| | 9707270 | Oct 01, 2024 | | U-2084 | | |
| <u>ZICONOTIDE ACETATE - PRIALT</u> | | | | | | |
| N 021060 004 | 8653033 | Oct 01, 2024 | | U-48 | | |
| | 8653033 | Oct 01, 2024 | | U-55 | | |
| | 8765680 | Oct 01, 2024 | | U-48 | | |
| | 8765680 | Oct 01, 2024 | | U-55 | | |
| | 9707270 | Oct 01, 2024 | | U-2084 | | |
| <u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u> | | | | | | |
| N 020825 001 | 6150366 | May 27, 2019 | | DP | | |
| <u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u> | | | | | | |
| N 020825 002 | 6150366 | May 27, 2019 | | DP | | |
| <u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u> | | | | | | |
| N 020825 003 | 6150366 | May 27, 2019 | | DP | | |
| <u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u> | | | | | | |
| N 020825 004 | 6150366 | May 27, 2019 | | DP | | |
| <u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u> | | | | | | |
| N 021483 001 | 6150366 | May 27, 2019 | | DP U-719 | | |
| | 7175855 | May 18, 2020 | | DP | | |

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

| APPL/PROD NO | PATENT NO | PATENT EXPIRATION DATE | PATENT CODES | PATENT DELIST REQUESTED | EXCLUSIVITY CODE(S) | EXCLUSIVITY EXPIRATION DATE |
|---------------------------------------|-----------|------------------------------|-----------------|-------------------------------|------------------------|-----------------------------------|
| <u>ZOLEDRONIC ACID - ZOMETA</u> | | | | | | |
| N 021223 | 002 | 8324189 | May 29, 2025 | U-1308 | | |
| | | 8324189 | May 29, 2025 | U-1309 | | |
| | | 8324189 | May 29, 2025 | U-53 | | |
| <u>ZOLEDRONIC ACID - ZOMETA</u> | | | | | | |
| N 021223 | 003 | 7932241 | Feb 05, 2028 | DP | | |
| | | 8324189 | May 29, 2025 | U-1308 | | |
| | | 8324189 | May 29, 2025 | U-1309 | | |
| | | 8324189 | May 29, 2025 | U-53 | | |
| <u>ZOLEDRONIC ACID - RECLAST</u> | | | | | | |
| N 021817 | 001 | 7932241 | Feb 05, 2028 | DP | | |
| | | 8052987 | Oct 27, 2023 | U-1199 | | |
| <u>ZOLMITRIPTAN - ZOMIG</u> | | | | | | |
| N 021450 | 003 | 6750237 | Nov 28, 2020 | DP | | |
| | | 6750237*PED | May 28, 2021 | | | |
| | | 7220767 | Nov 28, 2020 | DP | | |
| | | 7220767*PED | May 28, 2021 | | | |
| <u>ZOLMITRIPTAN - ZOMIG</u> | | | | | | |
| N 021450 | 004 | 6750237 | Nov 28, 2020 | DP | | |
| | | 7220767 | Nov 28, 2020 | DP | | |
| <u>ZOLPIDEM TARTRATE - AMBIEN CR</u> | | | | | | |
| N 021774 | 001 | 6514531 | Dec 01, 2019 | DP | | |
| <u>ZOLPIDEM TARTRATE - AMBIEN CR</u> | | | | | | |
| N 021774 | 002 | 6514531 | Dec 01, 2019 | DP | | |
| <u>ZOLPIDEM TARTRATE - EDLUAR</u> | | | | | | |
| N 021997 | 001 | 6761910 | Sep 24, 2019 | DP | U-674 | |
| | | 8512747 | Sep 24, 2019 | | U-674 | |
| | | 9265720 | Feb 25, 2031 | | U-674 | |
| | | 9597281 | Apr 06, 2027 | | U-674 | |
| <u>ZOLPIDEM TARTRATE - EDLUAR</u> | | | | | | |
| N 021997 | 002 | 6761910 | Sep 24, 2019 | DP | U-674 | |
| | | 8512747 | Sep 24, 2019 | | U-674 | |
| | | 9265720 | Feb 25, 2031 | | U-674 | |
| | | 9597281 | Apr 06, 2027 | | U-674 | |
| <u>ZOLPIDEM TARTRATE - ZOLPIMIST</u> | | | | | | |
| N 022196 | 001 | 8236285 | Aug 07, 2032 | DS DP | U-70 | |
| <u>ZOLPIDEM TARTRATE - INTERMEZZO</u> | | | | | | |
| N 022328 | 001 | 7658945 | Apr 15, 2027 | DP | U-1194 | |
| | | 7682628 | Feb 16, 2025 | | U-1194 | |
| | | 8242131 | Aug 20, 2029 | | U-1266 | |
| | | 8252809 | Feb 16, 2025 | DP | | |
| <u>ZOLPIDEM TARTRATE - INTERMEZZO</u> | | | | | | |
| N 022328 | 002 | 7658945 | Apr 15, 2027 | DP | U-1194 | |
| | | 7682628 | Feb 16, 2025 | | U-1194 | |
| | | 8242131 | Aug 20, 2029 | | U-1266 | |
| | | 8252809 | Feb 16, 2025 | DP | | |

Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons

PATENT AND EXCLUSIVITY TERMS

ADB 1 of 133

PATENT & EXCLUSIVITY ABBREVIATIONS

| | |
|-------|----------------------------------------------------------------------------------------------------------------------------------------------|
| CGT | COMPETITIVE GENERIC THERAPY |
| D | NEW DOSING SCHEDULE (SEE INDIVIDUAL REFERENCES) |
| GAIN | GAIN EXCLUSIVITY |
| I | NEW INDICATION (SEE INDIVIDUAL REFERENCES) |
| M | MISCELLANEOUS EXCLUSIVITY CODES (SEE INDIVIDUAL REFERENCES) |
| NC | NEW COMBINATION |
| NCE | NEW CHEMICAL ENTITY |
| NCE* | NEW CHEMICAL ENTITY (AN ENANTIOMER OF PREVIOUSLY APPROVED RACEMIC MIXTURE. SEE SECTION 505(U) OF THE FEDERAL FOOD AND DRUG COSMETIC ACT). |
| NDF | NEW DOSAGE FORM |
| NE | NEW ESTER OR SALT OF AN ACTIVE INGREDIENT |
| NP | NEW PRODUCT |
| NP* | NEW PRODUCT (MINT FLAVORED) |
| NPP | NEW PATIENT POPULATION |
| NR | NEW ROUTE |
| NS | NEW STRENGTH |
| ODE | ORPHAN DRUG EXCLUSIVITY (SEE INDIVIDUAL REFERENCES) |
| PC | PATENT CHALLENGE |
| PED | PEDIATRIC EXCLUSIVITY |
| RTO | RX TO OTC SWITCH OR OTC USE |
| RTO* | OTC USE FOR WOMEN AGES 15 AND 16 |
| RTO** | OTC USE FOR WOMEN 14 AND BELOW |
| U | PATENT USE CODE (SEE INDIVIDUAL REFERENCES) |
| W | EXCLUSIVITY ON THIS APPLICATION EXPIRING ON THIS DATE HAS BEEN WAIVED BY SPONSOR - SEE SECTION 1.8 OF ORANGE BOOK PREFACE WAIVED EXCLUSIVITY |

EXCLUSIVITY DOSING SCHEDULE

| | |
|------|---------------------------------------------------------------------------------------------------------------------------------------------|
| D-1 | ONCE A DAY APPLICATION |
| D-2 | ONCE DAILY DOSING |
| D-3 | SEVEN DAYS/SEVEN DAYS/SEVEN DAYS DOSING SCHEDULE |
| D-4 | SEVEN DAYS/FOURTEEN DAYS DOSING SCHEDULE |
| D-5 | TEN DAYS/ELEVEN DAYS DOSING SCHEDULE |
| D-6 | SEVEN DAYS/NINE DAYS/FIVE DAYS DOSING SCHEDULE |
| D-7 | BID DOSING |
| D-8 | INTRAVENOUS, EPIDURAL AND INTRATHECAL DOSING |
| D-9 | NARCOTIC OVERDOSE IN ADULTS |
| D-10 | NARCOTIC OVERDOSE IN CHILDREN |
| D-11 | POSTOPERATIVE NARCOTIC DEPRESSION IN CHILDREN |
| D-12 | BEDTIME DOSING OF 800MG FOR TREATMENT OF ACTIVE DUODENAL ULCER |
| D-13 | INCREASED MAXIMUM DAILY DOSAGE RECOMMENDATION |
| D-14 | BEDTIME DOSING OF 800MG FOR TREATMENT OF ACTIVE BENIGN GASTRIC ULCER |
| D-15 | SINGLE DAILY DOSE OF 25MG/37.5MG |
| D-16 | CONTINUOUS INTRAVENOUS INFUSION |
| D-17 | 400MG EVERY 12 HOURS FOR THREE DAYS FOR UNCOMPLICATED URINARY TRACT INFECTIONS |
| D-18 | LOWER RECOMMENDED STARTING DOSE GUIDELINES |
| D-19 | BOLUS DOSING GUIDELINES |
| D-20 | SINGLE 32MG DOSE |
| D-21 | ALTERNATIVE DOSAGE OF 300MG ONCE DAILY AFTER THE EVENING MEAL |
| D-22 | REDUCTION IN INFUSION TIME FROM 24 TO 4 HOURS FOR THE 60MG DOSE |
| D-23 | INCREASE MAXIMUM DOSE AND VARIATIONS IN THE DOSING REGIMEN |
| D-24 | FOR OVARIAN CANCER THE RECOMMENDED REGIMEN IS 135MG/M ² OR 175MG/M ² INTRAVENOUSLY OVER THREE HOURS EVERY THREE WEEKS |
| D-25 | ADDITIONAL DOSAGE REGIMEN EQUAL TO HALF THE ORIGINAL DOSING REGIMEN |

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

- D-26 ONCE WEEKLY APPLICATION
- D-27 BID DOSING IN PATIENTS 12 YEARS OF AGE AND OLDER FOR PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH MODERATE EMETOGENIC CANCER CHEMOTHERAPY
- D-28 USE OF ISOVUE-370 IN EXCRETORY UROGRAPHY AT EQUIVALENT GRAMS OF IODINE TO THE CURRENTLY APPROVED ISOVUE-250 AND ISOVUE-300
- D-29 INCREASE OF CUMULATIVE DOSE TO 0.3MMOL/KG FOR MRI OF CNS IN ADULTS
- D-30 5000 IU DOSE FOR PROPHYLAXIS AGAINST DEEP VEIN THROMBOSIS
- D-31 CHANGE IN RECOMMENDED TOTAL DAILY DOSE TO 80MG (40MG BID)
- D-32 REMOVAL OF THE RESTRICTIONS LIMITING TREATMENT TO TWO CONSECUTIVE WEEKS AND TO SMALL AREAS
- D-33 ONCE DAILY DOSING FOR PLAQUE PSORIASIS
- D-34 EVERY FOUR MONTHS DOSAGE REGIMEN
- D-35 FOR A ONE WEEK DOSING OF INTERDIGITAL TINEA PEDIS
- D-36 FOR A SINGLE 2MG DOSE AS AN ALTERNATIVE TO THE 1MG DOSE GIVEN TWICE DAILY
- D-37 DOSING REGIMEN FOR ADMINISTRATION EITHER ONCE DAILY (QD) OR TWICE DAILY (BID)
- D-38 CONTINUOUS INFUSION AS AN ALTERNATE METHOD OF ADMINISTRATION
- D-39 CHANGE IN TIME TO TAKE THE DRUG PRIOR TO A MEAL TO PREVENT MEAL-INDUCED HEARTBURN SYMPTOMS FROM ".1/2 TO 1 HOUR BEFORE EATING" TO ".. RIGHT BEFORE EATING OR UP TO 60MIN BEFORE CONSUMING..."
- D-40 ONCE-A-DAY DOSING REGIMEN
- D-41 DRUG MAY BE DOSED RIGHT BEFORE A MEAL OR ANY TIME UP TO 30MIN BEFORE EATING OR DRINKING FOOD AND BEVERAGES THAT WOULD BE EXPECTED TO CAUSE SYMPTOMS
- D-42 TEN DAY DOSING REGIMEN FOR TRIPLE THERAPY, PREVACID IN COMBINATION WITH CLARITHROMYCIN AND AMOXICILLIN, FOR THE ERADICATION OF H.PYLORI IN PATIENTS WITH DUODENAL ULCER DISEASE
- D-43 INITIATION OF TREATMENT WITH 900MG/DAY BY DELETION OF THE REQUIREMENT TO TITRATE TO 900MG/DAY OVER A 3-DAY PERIOD
- D-44 IN A CLINICAL TRIAL, FEWER DISCONTINUATIONS DUE TO ADVERSE EVENTS, ESPECIALLY DIZZINESS AND VERTIGO, WERE OBSERVED WHEN TITRATING THE DOSE IN INCREMENTS OF 50MG/DAY EVERY 3 DAYS UNTIL AN EFFECTIVE DOSE (NOT EXCEEDING 400MG/DAY) WAS REACHED
- D-45 ONCE DAILY DOSING FOR MAINTENANCE ONLY
- D-46 NEW DOSING REGIMEN OF 80MG DAILY
- D-47 PREVENTION OF HEARTBURN SYMPTOMS WHEN ADMINISTERED FROM 15 MINUTES UP TO, BUT NOT INCLUDING, 1 HOUR PRIOR TO A PROVOCATIVE MEAL
- D-48 ADMINISTRATION OF CISATRACURIUM A NEUROMUSCULAR BLOCKING AGENT AT DOSES OF 3 AND 4X THE ED95 OF CISATRACURIUM FOLLOWING INDUCTION WITH THIOPIENTAL
- D-49 PEDIATRIC DOSING GUIDELINES
- D-50 INFORMATION FOR USE OF CORVERT IN POST-CARDIAC SURGERY PATIENTS
- D-51 OPTIONAL STARTING DOSE OF 40MG/DAY
- D-52 ALTERNATE DOSING REGIMEN OF 1250MG TWICE DAILY
- D-53 USE IN PEDIATRIC PATIENTS FROM 1 MONTH TO 16 YEARS OF AGE
- D-54 USE OF ZYBAN FOR MAINTENANCE THERAPY. TREATMENT UP TO 6 MONTHS WAS SHOWN EFFICACIOUS
- D-55 ADDITION OF A HIGHER DOSE OF NUTROPIN FOR PUBERTAL PATIENTS (PUBERTAL DOSE LESS THAN OR EQUAL TO 0.7MG/KG/WEEK)
- D-56 ADDITION OF POSTPRANDIAL DOSING
- D-57 3-HOUR INFUSION OF TAXOL GIVEN EVERY THREE WEEKS AT A DOSE OF 175MG/M2 FOLLOWED BY CISPLATIN AT A DOSE OF 75MG/M2 FOR THE FIRST-LINE TREATMENT OF ADVANCED OVARIAN CANCER
- D-58 CHANGE IN DOSING INTERVAL TO ONCE-DAILY ADMINISTRATION
- D-59 REDUCTION OF ELEVATED LDL-C IN A NEW, HIGHER STRENGTH TABLET, 0.8MG, AND FOR EXTENSION OF THE DOSAGE RANGE TO 0.8MG DAILY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

- D-60 ADDITION OF A POST-OPERATIVE DOSING REGIMEN
- D-61 ONCE WEEKLY DOSING FOR THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- D-62 ONCE WEEKLY DOSING FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-63 TO ALLOW A TITRATION DOSING REGIMEN USING A 25MG DOSE
- D-64 INCREASING DOSAGE FOR NERVE BLOCK ANESTHESIA USING NAROPIN 7.5MG/ML AND FOR EXTENDING THE DURATION OF TREATMENT FOR POSTOPERATIVE ANALGESIA USING NAROPIN 2MG/ML
- D-65 CHANGE DOSING AND ADMINISTRATION TO INDICATE MAINTENANCE OF WEIGHT LOSS OVER AN 18 MONTH PERIOD THUS EXTENDING THE USE OF THIS DRUG FROM ONE TO TWO YEARS
- D-66 DOSING RECOMMENDATIONS FOR PATIENTS UNDERGOING PCI
- D-67 SHORTER TREATMENT COURSE OF THREE DAYS IN THE TREATMENT OF RECURRENT EPISODES OF GENITAL HERPES
- D-68 CHANGE OF ADMIN RATE FOR INFUSION OF AREDIA FOR TREATMENT OF MODERATE AND SEVERE HYPERCALCEMIA OF MALIGNANCY FROM 24 HOURS TO 2 HOURS UP TO BUT NOT INCLUDING 24 HOURS
- D-69 SHORTENED DOSING REGIMEN TO 5 DAYS FOR THE TREATMENT OF ACUTE EXACERBATION OF CHRONIC BRONCHITIS
- D-70 80MG ONCE DAILY DOSING REGIMEN
- D-71 EIGHT WEEK DOSING REGIMEN
- D-72 INFORMATION REGARDING INCREASED RATE OF INFUSION FOR DEPACON
- D-73 ONCE A WEEK DOSING FOR THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- D-74 ONCE A WEEK DOSING FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-75 INTERMITTENT DOSING REGIMEN, STARTING DAILY DOSE 14 DAYS PRIOR TO THE ANTICIPATED ONSET OF MENSTRUATION THROUGH THE FIRST FULL DAY OF MENSES AND REPEATING WITH EACH NEW CYCLE
- D-76 FOR USE ON AN "AS NEEDED" OR PRN BASIS FOR THE MANAGEMENT OF NASAL SYMPTOMS IN PATIENTS FOR WHOM THE DRUG IS INDICATED
- D-77 ADDITION OF 20MG AND 40MG DAILY AS OPTIONAL STARTING DOSES WITH 40MG INTENDED FOR PATIENTS WHO REQUIRE A LARGE REDUCTION IN LDL-C (MORE THAN 45%)
- D-78 USE OF FLEXERIL 5MG FOR THE RELIEF OF MUSCLE SPASM ASSOCIATED WITH ACUTE, PAINFUL, MUSCULOSKELETAL CONDITIONS
- D-79 NEW LOWER STARTING DOSE FOR TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS AND/OR MODERATE TO SEVERE SYMPTOMS OF VULVAR AND VAGINAL ATROPHY ASSOCIATED W/ THE MENOPAUSE
- D-80 CHANGE OF DOSING SCHEDULE FOR LANTUS FROM ONCE DAILY AT BEDTIME TO FLEXIBLE DAILY DOSING
- D-81 NEW LOWER STARTING DOSE FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPORSIS
- D-82 USE OF PREMARIN 0.3 MG AND 0.45 MG FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-83 750 MG, ONCE DAILY FOR 5 DAYS FOR COMMUNITY ACQUIRED PNEUMONIA (CAP)
- D-84 ONCE-A-DAY DOSING OF FLOXACIN OTIC FOR THE TREATMENT OF ADULTS AND PEDIATRIC PATIENTS (AGES 6 MO & OLDER) W/ OTITIS EXTERNA CAUSED BY SUSCEPTIBLE STRAINS OF E.COLI, P.AERUGINOSA AND S.AUREUS
- D-85 LOWER RECOMMENDED STARTING DOSE GUIDELINES FOR TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH THE MENOPAUSE
- D-86 FOR USE IN SELECT EXTERNAL INSULIN PUMPS
- D-87 ADDITION OF ONCE-WEEKLY DOSING FOR THE TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
- D-88 NEW DOSING RANGE OF 200-400MG PER DAY IN TWO DIVIDED DOSES FOR ADULTS WITH PARTIAL SEIZURES
- D-89 USE OF REYATAZ 300 MG/RITONAVIR 100 MG ONCE DAILY FOR TREATMENT IN HIV-INFECTED ANTIRETROVIRAL-EXPERIENCED PATIENTS
- D-90 ADDITION OF DAYTIME ADMINISTRATION TO TREAT VULVOVAGINAL CANDIDIASIS
- D-91 ALTERNATE INTERMITTENT DOSING REGIMEN

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

| | |
|-------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| D-92 | ALTERNATIVE DOSAGE OF 1000MG ONCE DAILY AT BEDTIME |
| D-93 | ALTERNATE TWO OR THREE TIMES DAILY DOSING REGIMENS |
| D-94 | NEW MAXIMUM DOSAGE OF 72 MG/DAY IN ADOLESCENTS 13-17 YEARS OF AGE WITH ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) |
| D-95 | BROADENED INITIAL STARTING DOSE FOR HYPERTENSION FROM 50 MG TO 100 MG TO 25 MG TO 100 MG DOSE RANGE |
| D-96 | ONCE-MONTHLY TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS WITH BONIVA (IBANDRONATE SODIUM) 150 MG TABLETS |
| D-97 | PED CANCER PT POPULATION EXPANDED TO INCLUDE PTS 6 MOS UP TO BUT NOT INCLUDING 4 YRS AND DOSING INSTRUCTIONS TO ADMIN 30 MIN BEFORE CHEMO WITH SECOND AND THIRD DOSES 4 & 8 HOURS AFTER FIRST DOSE |
| D-98 | DOSING FOR PED SURGICAL PTS EXPANDED TO INCLUDE PTS 1 MONTH UP TO BUT NOT INCLUDING 2 YEARS OF AGE |
| D-99 | ONCE DAILY ADMINISTRATION FOR THE TREATMENT OF HIV INFECTION IN THERAPY NAIVE ADULT PATIENTS |
| D-100 | 750 MG ONCE DAILY FOR FIVE DAYS FOR THE TREATMENT OF ACUTE BACTERIAL SINUSITIS |
| D-101 | ONCE DAILY IN CHRONIC IDIOPATHIC URTICARIA FOR ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER |
| D-102 | NEW DOSING REGIMEN OF ONE SPRAY TWICE DAILY FOR SEASONAL ALLERIC RHINITIS IN PATIENTS 12 YRS OF AGE AND OLDER |
| D-103 | NEW DOSING RECOMMENDATION FOR THE TREATMENT OF RECURRENT GENITAL HERPES IN IMMUNOCOMPETENT PATIENTS, SPECIFICALLY A REDUCTION IN COURSE OF THERAPY FROM FAMCICLOVIR 125 MG TWICE-A-DAY FOR 5 DAYS TO 1000 MG TWICE-A-DAY FOR 1 DAY. |
| D-104 | 0.5MG/0.1MG FOR THE TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE IN WOMEN WHO HAVE A UTERUS |
| D-105 | USE OF ACTONEL 75MG TWO CONSECUTIVE DAYS PER MONTH FOR THE PREVENTION AND TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS |
| D-106 | FIVE DAY TREATMENT OF SELECTED SUSCEPTIBLE STRAINS OF STREPTOCOCCUS PNEUMONIAE, HAEMOPHILUS INFLUENZA, MYCOPLASMA PNEUMONIAE, AND CHLAMYDIA PNEUMONIAE FOR COMMUNITY-ACQUIRED PNEUMONIA |
| D-107 | PROVIDES FOR THE COMBINATION TABLET OF 70MG ALENDRONATE AND 5600 IU OF VITAMIN D3 FOR THE TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN AND TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS |
| D-108 | TREATMENT OF COMPLICATED URINARY TRACT INFECTION AND ACUTE PYELONEPHRITIS WITH LEVAQUIN 750MG ONCE DAILY FOR FIVE DAYS |
| D-109 | PROVIDE FOR THE USE OF A LOWER DOSE FOR THE TREATMENT OF ADULTS WITH CHRONIC PHASE CHRONIC MYELOID LEUKEMIA (CML) WITH RESISTANCE OR INTOLERANCE TO PRIOR THERAPY INCLUDING IMATINIB MESYLATE |
| D-110 | TREATMENT OF SCHIZOPHRENIA IN ADOLESCENTS AGED 13-17 |
| D-111 | PROVIDES FOR ONCE DAILY USE OF CIALIS, 2.5 MG AND 5 MG, FOR THE TREATMENT OF ERECTILE DYSFUNCTION |
| D-112 | PROVIDES FOR PEDIATRIC PUMP USE |
| D-113 | ONCE DAILY DOSING REGIMEN FOR PATIENTS WHO BECOME CONSTIPATED ON TWICE DAILY REGIMEN |
| D-114 | NEW DOSING RECOMMENDATIONS FOR USE OF SIROLIMUS IN COMBINATION WITH CYCLOSPORINE FOR THE PROPHYLAXIS OF REJECTION IN HIGH-RISK RENAL TRANSPLANT RECIPIENTS |
| D-115 | STARTING DOSE OF 15MG/DAY FOR MONOTHERAPY IN ACUTE TREATMENT OF BIPOLAR DISORDER, MANIC OR MIXED |
| D-116 | ALTERNATIVE DOSING REGIMEN ATAZANAVIR SULATE CO-ADMINISTERED WITH RITONAVIR FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT NAIVE PATIENTS |
| D-117 | 50 MG TABLET FOR INITIATION OF DOSE TITRATION FOR BIPOLAR DISORDER |
| D-118 | TWO 400MG TABLETS ONCE DAILY, CO-ADMINISTERED WITH 100MG RITONAVIR |
| D-119 | DOSING RECOMMENDATIONS FOR HIV INFECTED PEDIATRIC PATIENTS 6 TO LESS THAN 18 YEARS OF AGE |
| D-120 | DOSING REGIMEN ADJUSTMENTS |
| D-121 | CHANGE TO REMOVE 20 MG MAXIMUM DOSAGE RESTRICTION |

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

| | |
|-------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| D-122 | USE OF VAGIFEM 10 MCG FOR THE TREATMENT OF ATROPHIC VAGINITIS DUE TO MENOPAUSE |
| D-123 | ALTERNATIVE DOSING REGIMEN DOSE OF 20 MG/METER SQUARE BY CONTINUOUS INTRAVENOUS INFUSION OVER 1 HOUR REPEATED DAILY FOR 5 DAYS |
| D-124 | ONCE DAILY DOSING REGIMEN IN ADULT PATIENTS WITH LESS THAN THREE LOPINAVIR RESISTANCE-ASSOCIATED SUBSTITUTIONS |
| D-125 | EXTEND CURRENT DOSING REGIMEN TO 900MG (2-450MG TABLETS) ONCE A DAY WITHIN 10 DAYS OF TRANSPLANTATION UNTIL 200 DAYS POST-TRANSPLANTATION FOR THE PREVENTION OF CYTOMEGALOVIRUS (CMV) DISEASE IN ADULT KIDNEY TRANSPLANT PATIENTS AT HIGH RISK. |
| D-126 | CHANGE DOSAGE REGIMEN FROM 250MG TO 500MG |
| D-127 | DOSING REGIMEN FOR ADULT PATIENTS WITH CHRONIC HEPATITIS B (CHB) AND DECOMPENSATED LIVER DISEASE |
| D-128 | SINGLE IV DOSE OF FOSAPREPITANT 150MG, DOSED CONCOMITANTLY WITH 5HT3 RECEPTOR ANTAGONIST & CORTICOSTEROID, FOR PREVENTION OF ACUTE & DELAYED NAUSEA & VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF HIGHLY EMETOGENIC CANCER CHEMO |
| D-129 | 800/100 MG DARUNAVIR/RITONAVIR, ONCE DAILY, IN TREATMENT-EXPERIENCED HIV-1 INFECTED PATIENTS WITH NO DARUNAVIR RESISTANCE ASSOCIATED SUBSTITUTIONS |
| D-130 | DOSING RECOMMENDATIONS FOR TREATMENT OF HIV-1 INFECTION DURING PREGNANCY BASED ON DATA FROM STUDY AI424-182, A STUDY OF ATAZANAVIR/RITONAVIR IN COMBINATION WITH ZIDOVUDINE/LAMIVUDINE IN HIV INFECTED PREGNANT WOMEN |
| D-131 | EVERY 6 TO 8 WEEKS FOR THE 120MG STRENGTH FOR PATIENTS WHO ARE CONTROLLED ON SOMATULINE DEPOT 60MG OR 90MG |
| D-132 | 45MG FOR 6 MONTH ADMINISTRATION |
| D-133 | NEW EFFICACY DATA AND DOSING REGIMEN FOR PREGNANCY IN NORMAL OVULATORY WOMEN UNDERGOING CONTROLLED OVARIAN STIMULATION AS PART OF AN IVF OR INTRACYTOPLASMIC SPERM INJECTION (ICSI) CYCLE |
| D-134 | INCREASING MAXIMUM DOSING OF PATIENTS WITH SCHIZOPHRENIA TO 160 MG/DAY |
| D-135 | UPDATE LABELING WITH ONCE DAILY DOSING IN HIV-1 INFECTED, TREATMENT-NAIVE PEDIATRIC PATIENTS 12 TO LESS THAN 18 YEARS OF AGE |
| D-136 | ALTERNATE DOSING REGIMEN FOR UNCOMPLICATED URETHRAL OR ENDOCERVICAL INFECTION CAUSED BY CHLAMYDIA TRACHOMATIS, ADMINISTER 200 MG BY MOUTH ONCE-A-DAY FOR 7 DAYS |
| D-137 | NEW LOWER DOSING REGIMEN FOR REVATIO IN THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (WHO GROUP 1) IN ADULTS |
| D-138 | 80 MG DOSING REGIMEN FOR THE RISK REDUCTION OF REBLEEDING OF GASTRIC AND DUODENAL ULCERS IN THE FIRST 72 HOURS FOLLOWING THERAPEUTIC ENDOSCOPY IN ADULTS |
| D-139 | ADDITIONAL INFORMATION ADDED TO THE DOSING AND ADMINISTRATION SECTION OF THE LABELING REGARDING THE ADMINISTRATION OF BRAVELLE AND MENOPUR IN THE SAME SYRINGE TO OVULATORY WOMEN AS PART OF AN ART CYCLE |
| D-140 | REVISED DOSING SCHEDULE TO ADMINISTER AVANAFIL 15 MINUTES PRIOR TO SEXUAL ACTIVITY |
| D-141 | DOSING INFORMATION IN PREVIOUSLY UNTREATED MANTLE CELL LYMPHOMA |
| D-142 | DOSE MODIFICATION GUIDELINES FOR BORTEZOMIB WHEN GIVEN IN COMBINATION WITH RITUXIMAB, CYCLOPHOSPHAMIDE, DOXORUBICIN, AND PREDNISONE |
| D-143 | INITIATION OF VIMPAT THERAPY WITH A LOADING DOSE OF 200MG |
| D-144 | LOWER LIMIT OF 15 MINUTES FOR THE INFUSION DURATION |
| D-145 | UPDATES TO THE DOSAGE AND ADMINISTRATION SECTION OF THE LABELING TO REFLECT THE RESULTS OF TWO SHORT TERM STUDIES EVALUATING THE SAFETY AND EFFICACY OF INTUNIV IN CHILDREN AND ADOLESCENTS AGES 6 TO 17 WITH ADHD. |
| D-146 | CHANGE IN TARGET DOSING TO 20MG TO 40MG ORALLY ONCE DAILY |
| D-147 | ONCE DAILY DOSING IN PEDIATRIC PATIENTS 3 MONTHS OF AGE AND OLDER IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION |
| D-148 | EXTENDED THE DURATION OF THE DOSING REGIMEN FROM 100 DAYS TO 200 DAYS POST-TRANSPLANTATION FOR THE PREVENTION OF CMV DISEASE IN PEDIATRIC KIDNEY TRANSPLANT |
| D-149 | DOSING INFORMATION ADDED TO THE LABELING REGARDING PEDIATRIC PATIENTS 6 YEARS AND OLDER WITH ITP |

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

- D-150 1600MG DAILY FOR PATIENTS ON ADJUNCTIVE THERAPY WHO DID NOT ACHIEVE A SATISFACTORY RESPONSE ON 1200MG DAILY DOSE
- D-151 DOSING RECOMMENDATIONS FOR THE TREATMENT OF CHRONIC HEPATITIS C IN PATIENTS CO-INFECTED WITH HIV-1
- D-152 DOSING RECOMMENDATIONS AS NECESSARY FOR FEVER AND PAIN FOR AGES 6MO TO LESS THAN 12 YEARS AND 12 TO 17 YEARS.
- D-153 IN COMBINATION WITH RIBAVIRIN FOR 12 WEEKS, FOR THE TREATMENT OF GENOTYPE 1, CHRONIC HEPATITIS C TREATMENT EXPERIENCED PATIENTS WITH COMPENSATED CIRRHOSIS BASED UPON THE RESULTS OF THE SIRIUS STUDY
- D-154 ADDITION OF A 1500MG-SINGLE-DOSE REGIMEN FOR THE TREATMENT OF ADULT PATIENTS WITH ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS (ABSSI)
- D-155 SINGLE IV DOSE OF FOSAPREPITANT 150MG DOSED CONCOMITANTLY WITH 5HT3 RECEPTOR ANTAGONIST & CORTICOSTEROID FOR PREVENTION OF DELAYED NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF MODERATELY EMETOGENIC CANCER CHEMOTHERAPY
- D-156 DOSING INFORMATION ADDED TO THE LABELING PROVIDING INFORMATION ON TRANSITIONING FROM SUBCUTANEOUS OR INTRAVENOUS ROUTES OF ADMINISTRATION OF TREPROSTINIL
- D-157 UPDATED INFORMATION ADDED TO THE DOSAGE AND ADMINISTRATION SECTION OF THE LABELING PROVIDING DOSAGE RECOMMENDATIONS FOR INTERRUPTIONS AND DISCONTINUATION OF THERAPY
- D-158 REVISED DOSING TO EXPAND PATIENT POPULATION TO INCLUDE LIVER TRANSPLANT RECIPIENTS WITH GENOTYPE 1 HCV INFECTION
- D-159 REVISED DOSING TO EXPAND PATIENT POPULATION TO INCLUDE LIVER TRANSPLANT RECIPIENTS WITH GENOTYPE 4 HCV INFECTION
- D-160 REVISED DOSING TO EXPAND PATIENT POPULATION TO INCLUDE PATIENTS WITH DECOMPENSATED CIRRHOSIS WITH GENOTYPE 1 HCV INFECTION
- D-161 DOSAGE RECOMMENDATIONS ADDED TO INCLUDE TREATMENT OF HCV GENOTYPE 3 SUBJECTS CO-INFECTED WITH HIV-1
- D-162 DOSING TO INCLUDE PATIENTS WITH CHRONIC HCV GENOTYPE 1 INFECTION WITH COMPENSATED (CHILD-PUGH A) OR DECOMPENSATED (CHILD-PUGH B OR C) CIRRHOSIS AND TREATMENT OF CHRONIC HCV GENOTYPE 3 INFECTION IN SUBJECTS WITH DECOMPENSATED (CHILD-PUGH B OR C) CIRRHOSIS
- D-163 DOSING TO INCLUDE PATIENTS WITH CHRONIC HCV GENOTYPE 1A INFECTION WITH COMPENSATED (CHILD-PUGH A) CIRRHOSIS AND GENOTYPE 1B WITH OR WITHOUT COMPENSATED (CHILD-PUGH A) CIRRHOSIS
- D-164 UPDATES TO THE DOSAGE AND ADMINISTRATION, DOSE MODIFICATIONS SECTION OF THE LABELING
- D-165 DOSING RECOMMENDATION ADDED TO THE LABELING FOR IMBRUVICA USE IN COMBINATION WITH BENDAMUSTINE AND RITUXIMAB FOR THE TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)/SMALL LYMPHOCYTIC LEUKEMIA (SLL)
- D-166 BROADEN INITIAL STARTING DOSE FOR BIPOLAR I DISORDER TO 5-10MG TWICE DAILY
- D-167 ADDITION OF 1200 MG ONCE DAILY DOSING FOR TREATMENT-NAIVE PATIENTS OR PATIENTS WHO ARE VIROLOGICALLY SUPPRESSED ON AN INITIAL REGIMEN OF RALTEGRAVIR FILM-COATED TABLETS 400 MG TWICE DAILY
- D-168 NEW DOSING REGIMEN OF 10 MG ONCE DAILY FOR THE REDUCTION IN THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS (DVT) AND/OR PULMONARY EMBOLISM (PE) IN PATIENTS AT CONTINUED RISK FOR DVT AND/OR PE AFTER COMPLETION OF INITIAL TREATMENT LASTING AT LEAST 6 MONTHS
- D-169 ONCE-DAILY DOSING FOR PATIENTS 5 YEARS OF AGE AND OLDER WHO HAVE UNDETECTABLE SERUM AND URINE SUCCINYLACETONE CONCENTRATIONS AFTER A MINIMUM OF 4 WEEKS ON A STABLE DOSAGE OF NITISINONE
- D-170 TO ALLOW WITHDRAWAL THERAPY OF PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA IN CHRONIC PHASE WHO HAVE ACHIEVED A SUSTAINED MOLECULAR RESPONSE ON NILOTINIB THERAPY FOR A MINIMUM OF ONE YEAR PRIOR TO DISCONTINUATION
- D-171 REVISED DOSING TO INCLUDE UP-TITRATION AS A STRATEGY TO IMPROVE TOLERABILITY AND THEREBY REDUCE TREATMENT DISCONTINUATION FOR ROFLUMILAST MAINTENANCE DOSAGE OF 500 MCG DAILY
- D-172 ADDITION OF A ONCE WEEKLY DOSING REGIMEN FOR CARFILZOMIB IN COMBINATION WITH DEXAMETHASONE FOR PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY DOSING SCHEDULE**

RECEIVED ONE TO THREE LINES OF THERAPY

D-173 DOSING RECOMMENDATION FOR THE USE OF
ELVITEGRAVIR/COBICISTAT/EMTRICITABINE/TENOFOVIR ALAFENAMIDE FIXED DOSE
COMBINATION IN HIV-1 INFECTED ADULT PATIENTS WITH END-STAGE-RENAL DISEASE WHO
ARE RECEIVING CHRONIC HEMODIALYSIS

EXCLUSIVITY INDICATION

I-1 DYSMENORRHEA
I-2 CHOLANGIOPANCREATOGRAPHY
I-3 INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY
I-4 PERIPHERAL VENOGRAPHY (PHLEBOGRAPHY)
I-5 HYSTEROSALPINGOGRAPHY
I-6 TREATMENT OF JUVENILE ARTHRITIS
I-7 BIOPSY PROVEN MINIMAL CHANGE NEPHROTIC SYNDROME IN CHILDREN
I-8 ADULT INTRAVENOUS CONTRAST-ENHANCED COMPUTED TOMOGRAPHY OF THE HEAD AND BODY
I-9 PREVENTION OF POSTOPERATIVE NAUSEA AND VOMITING
I-10 PREVENTION OF POSTOPERATIVE DEEP VEIN THROMBOSIS AND PULMONARY EMBOLISM IN
TOTAL HIP REPLACEMENT SURGERY
I-11 RELIEF OF MILD TO MODERATE PAIN
I-12 TREATMENT OF CUTANEOUS CANDIDIASIS
I-13 URINARY TRACT INFECTION (UTI) PREVENTION FOR PERIODS UP TO FIVE MONTHS IN WOMEN
WITH A HISTORY OF RECURRENT UTI
I-14 SEBORRHEIC DERMATITIS
I-15 PHOTOPHERESIS IN THE PALLIATIVE TREATMENT OF SKIN MANIFESTATIONS OF CUTANEOUS T-
CELL LYMPHOMA IN PERSONS NOT RESPONSIVE TO OTHER TREATMENT
I-16 STIMULATE THE DEVELOPMENT OF MULTIPLE FOLLICLES/OOCYTES IN OVULATORY PATIENTS
PARTICIPATING IN AN IN VITRO FERTILIZATION PROGRAM
I-17 MANAGEMENT OF CONGESTIVE HEART FAILURE
I-18 ENDOSCOPIC RETROGRADE PANCREATOGRAPHY
I-19 HERNIOGRAPHY
I-20 KNEE ARTHROGRAPHY
I-21 HIGH DOSE METHOTREXATE WITH LEUCOVORIN RESCUE IN COMBINATION WITH OTHER
CHEMOTHERAPEUTIC AGENTS TO DELAY RECURRENCE IN PATIENTS WITH NONMETASTATIC
OSTEOSARCOMA WHO HAVE UNDERGONE SURGICAL RESECTION OR AMPUTATION FOR THE PRIMARY
TUMOR
I-22 RESCUE AFTER HIGH-DOSE METHOTREXATE THERAPY IN OSTEOSARCOMA
I-23 SHORT-TERM TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
I-24 TREATMENT OF RHEUMATOID ARTHRITIS
I-25 ADULT INTRA-ARTERIAL DIGITAL SUBTRACTION ANGIOGRAPHY OF THE HEAD, NECK,
ABDOMINAL, RENAL AND PERIPHERAL VESSELS
I-26 TREATMENT OF LIVER FLUKES
I-27 ADJUNCTIVE THERAPY TO DIET TO REDUCE THE RISK OF CORONARY ARTERY DISEASE
I-28 SELECTIVE ADULT VISCERAL ARTERIOGRAPHY
I-29 METASTATIC BREAST CANCER IN PREMENOPAUSAL WOMEN AS AN ALTERNATIVE TO
OOPHORECTOMY OR OVARIAN IRRADIATION
I-30 TREATMENT OF TINEA PEDIS
I-31 CONTRAST ENHANCEMENT AGENT TO FACILITATE VISUALIZATION OF LESIONS IN THE SPINE
AND ASSOCIATED TISSUES
I-32 PEDIATRIC MYELOGRAPHY
I-33 ORAL USE OF DILUTED OMNIPAQUE INJECTION IN ADULTS FOR CONTRAST ENHANCED
COMPUTED TOMOGRAPHY OF THE ABDOMEN

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-34 ORAL USE IN ADULTS FOR PASS-THROUGH EXAMINATION OF THE GASTROINTESTINAL TRACT

I-35 PEDIATRIC CONTRAST ENHANCEMENT OF COMPUTED TOMOGRAPHIC HEAD IMAGING

I-36 ARTHROGRAPHY OF THE SHOULDER JOINTS IN ADULTS

I-37 RADIOGRAPHY OF THE TEMPOROMANDIBULAR JOINT IN ADULTS

I-38 CONTRAST ENHANCEMENT AGENT TO FACILITATE VISUALIZATION OF LESIONS OF THE CENTRAL NERVOUS SYSTEM IN CHILDREN (2 YEARS OF AGE AND OLDER)

I-39 TREATMENT OF ACUTE MYOCARDIAL INFARCTION

I-40 PRIMARY NOCTURNAL ENURESIS

I-41 MIGRAINE HEADACHE PROPHYLAXIS

I-42 HERPES ZOSTER

I-43 HERPES SIMPLEX ENCEPHALITIS

I-44 MAINTENANCE THERAPY IN HEALED DUODENAL ULCER PATIENTS AT DOSE OF 1 GRAM TWICE DAILY

I-45 ACUTE TREATMENT OF VARICELLA ZOSTER VIRUS

I-46 USE IN PEDIATRIC COMPUTED TOMOGRAPHIC HEAD AND BODY IMAGING

I-47 TREATMENT OF PEDIATRIC PATIENTS WITH SYMPTOMATIC HUMAN IMMUNODEFICIENCY VIRUS (HIV) DISEASE

I-48 PEDIATRIC ANGIOCARDIOGRAPHY

I-49 TREATMENT OF TRAVELERS' DIARRHEA DUE TO SUSCEPTIBLE STRAINS OF ENTEROTOXIGENIC ESCHERICHIA COLI

I-50 FOR USE IN WOMEN WITH AXILLARY NODE-NEGATIVE BREAST CANCER

I-51 TREATMENT OF PRIMARY DYSMENORRHEA AND FOR THE TREATMENT OF IDIOPATHIC HEAVY MENSTRUAL BLOOD LOSS

I-52 PEDIATRIC EXCRETORY UROGRAPHY

I-53 TREATMENT OF PANIC DISORDER, WITH OR WITHOUT AGORAPHOBIA

I-54 RENAL CONCENTRATION CAPACITY TEST

I-55 HYPERTENSION

I-56 EROSIIVE GASTROESOPHAGEAL REFLUX DISEASE

I-57 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER

I-58 INITIAL TREATMENT OF ADVANCED OVARIAN CARCINOMA IN COMBINATION WITH OTHER APPROVED CHEMOTHERAPEUTIC AGENTS

I-59 ENDOSCOPICALLY DIAGNOSED ESOPHAGITIS, INCLUDING EROSIIVE AND ULCERATIVE ESOPHAGITIS, AND ASSOCIATED HEARTBURN DUE TO GASTROESOPHAGEAL REFLUX DISEASE

I-60 SINGLE APPLICATION TREATMENT OF HEAD LICE IN CHILDREN TWO MONTHS TO TWO YEARS IN AGE

I-61 FEMALE ANDROGENETIC ALOPECIA

I-62 PREVENTION AND TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS

I-63 ONCE DAILY TREATMENT AS INITIAL THERAPY IN THE TREATMENT OF HYPERTENSION

I-64 PREVENTION OF SUPRAVENTRICULAR TACHYCARDIAS

I-65 PREVENTION OF UPPER GASTROINTESTINAL BLEEDING IN CRITICALLY ILL PATIENTS

I-66 UNCOMPLICATED GONORRHEA

I-67 TREATMENT OF ACUTE ASTHMATIC ATTACKS IN CHILDREN SIX YEARS OF AGE AND OLDER

I-68 CENTRAL PRECOCIOUS PUBERTY

I-69 SHORT TERM TREATMENT OF PATIENTS WITH SYMPTOMS OF GASTROESOPHAGEAL REFLUX DISEASE (GERD), AND FOR THE SHORT TERM TREATMENT OF ESOPHAGITIS DUE TO GERD INCLUDING ULCERATIVE DISEASE DIAGNOSED BY ENDOSCOPY

I-70 USE IN COMBINATION WITH 5-FLUOROURACIL TO PROLONG SURVIVAL IN THE PALLIATIVE TREATMENT OF PATIENTS WITH ADVANCED COLORECTAL CANCER

I-71 VARICELLA INFECTIONS (CHICKENPOX)

I-72 PREVENTION OF CMV DISEASE IN TRANSPLANT PATIENTS AT RISK FOR CMV DISEASE

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

| | |
|-------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| I-73 | INITIATE AND MAINTAIN MONITORED ANESTHESIA CARE (MAC) SEDATION DURING DIAGNOSTIC PROCEDURES |
| I-74 | INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY |
| I-75 | TREATMENT OF ENDOSCOPICALLY DIAGNOSED EROSIVE ESOPHAGITIS |
| I-76 | PREVENTION OF OSTEOPOROSIS |
| I-77 | DERMAL INFECTIONS-TINEA PEDIS, TINEA CORPORIS, TINEA CRURIS DUE TO EPIDERMOPHYTON FLOCCOSUM |
| I-78 | CONTRAST ENHANCED COMPUTED TOMOGRAPHIC IMAGING OF THE HEAD AND BODY AND INTRAVENOUS EXCRETORY UROGRAPHY |
| I-79 | MANAGEMENT OF CHRONIC STABLE ANGINA AND ANGINA DUE TO CORONARY ARTERY SPASM |
| I-80 | DIAGNOSIS AND LOCALIZATION OF ISCHEMIA AND CORONARY HEART DISEASE |
| I-81 | PROPHYLAXIS IN DESIGNATED IMMUNOCOMPROMISED CONDITIONS TO REDUCE THE INCIDENCE OF OROPHARYNGEAL CANDIDIASIS |
| I-82 | TREATMENT OF TRAVELERS' DIARRHEA |
| I-83 | ANGIOCARDIOGRAPHY, CONTRAST ENHANCED COMPUTED TOMOGRAPHIC IMAGING OF THE HEAD AND BODY, AND INTRAVENOUS EXCRETORY UROGRAPHY IN CHILDREN |
| I-84 | INTRAOPERATIVE AND POSTOPERATIVE TACHYCARDIA AND/OR HYPERTENSION |
| I-85 | TREATMENT OF ANOREXIA ASSOCIATED WITH WEIGHT LOSS IN PATIENTS WITH AIDS |
| I-86 | TREATMENT OF SECONDARY CARNITINE DEFICIENCY |
| I-87 | RENAL IMAGING AGENT FOR USE IN CHILDREN |
| I-88 | MANAGEMENT OF ENDOMETRIOSIS |
| I-89 | EPIDURAL USE IN LABOR AND DELIVERY AS AN ANALGESIC ADJUNCT TO BUPIVACAINE |
| I-90 | INTENSIVE CARE UNIT SEDATION |
| I-91 | MONOTHERAPY USE FOR HYPERTENSION |
| I-92 | ADJUNCTIVE THERAPY IN THE MANAGEMENT OF HEART FAILURE |
| I-93 | PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN CHILDREN AGES 4-11 YEARS |
| I-94 | USE WITH MRI IN ADULTS TO PROVIDE CONTRAST ENHANCEMENT AND FACILITATE VISUALIZATION OF LESIONS IN THE BODY [EXCLUDING THE HEART] |
| I-95 | TREATMENT OF LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION |
| I-96 | TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA |
| I-97 | ORAL OR RECTAL USE IN CHILDREN FOR THE EXAMINATION OF THE GASTROINTESTINAL TRACT |
| I-98 | TREATMENT OF CHILDREN WHO HAVE GROWTH FAILURE ASSOCIATED WITH CHRONIC RENAL INSUFFICIENCY |
| I-99 | PEDIATRIC ANESTHESIA IN CHILDREN 3 YEARS AND OLDER |
| I-100 | TO DECREASE THE INCIDENCE OF CANDIDIASIS IN PATIENTS UNDERGOING BONE MARROW TRANSPLANTATION WHO RECEIVE CYTOTOXIC CHEMOTHERAPY AND/OR RADIATION THERAPY |
| I-101 | TREATMENT OF DIABETIC NEPHROPATHY IN PATIENTS WITH TYPE I INSULIN-DEPENDENT DIABETES MELLITUS AND RETINOPATHY |
| I-102 | TREATMENT OF OBSESSIVE-COMPULSIVE DISORDER |
| I-103 | PROPHYLAXIS AGAINST PNEUMOCYSTIS CARINII PNEUMONIA IN INDIVIDUALS WHO ARE IMMUNOCOMPROMISED AND CONSIDERED TO BE AT RISK OF DEVELOPING PNEUMOCYSTIS CARINII PNEUMONIA |
| I-104 | TREATMENT OF PULMONARY AND EXTRAPULMONARY ASPERGILLOSIS IN PATIENTS WHO ARE INTOLERANT OF OR WHO ARE REFRACTORY TO AMPHOTERICIN B THERAPY |
| I-105 | TREATMENT OF METASTATIC CARCINOMA OF THE BREAST AFTER FAILURE OF FIRST-LINE OR SUBSEQUENT CHEMOTHERAPY |
| I-106 | TREATMENT OF ACROMEGALY |
| I-107 | VAGINAL CANDIDIASIS |
| I-108 | EXPANDED USE-FOR ICU PATIENTS UNDERGOING LONG-TERM INFUSION DURING MECHANICAL VENTILATION |
| I-109 | TYPHOID FEVER |

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-110 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH RADIOTHERAPY

I-111 TREATMENT OF PAGET'S DISEASE OF BONE

I-112 MANAGEMENT OF MODERATE TO SEVERE PAIN

I-113 TREATMENT OF PROSTATITIS

I-114 USE IN CHILDREN TO VISUALIZE LESIONS WITH ABNORMAL VASCULARITY IN THE BRAIN (INTRACRANIAL LESIONS), SPINE, AND ASSOCIATED TISSUE

I-115 USE IN MRI IN ADULTS TO VISUALIZE LESIONS IN THE HEAD AND NECK

I-116 MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS

I-117 TO SLOW THE PROGRESSION FO CORONARY ATHEROSCLEROSIS IN PATIENTS WITH CORONARY HEART DISEASE

I-118 PREVENTION OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM FOLLOWING KNEE REPLACEMENT SURGERY

I-119 TREATMENT OF ANEMIA CAUSED BY UTERINE LEIOMYOMATA IN WOMEN WHO FAIL IRON THERAPY

I-120 MAINTENANCE THERAPY FOR GASTRIC ULCER PATIENTS AT REDUCED DOSAGE AFTER HEALING ACUTE ULCERS

I-121 EXPANDED PATIENT POPULATION -- USE IN ICU PATIENTS

I-122 PSORIASIS OF THE SCALP

I-123 RELIEF OF MILD TO MODERATE PAIN IN PATIENTS AGED 6 MONTHS AND OLDER

I-124 LEUKOCYTE LABELED SCINTIGRAPHY AS AN ADJUNCT IN THE LOCALIZATION OF INTRA-ABDOMINAL INFECTION AND INFLAMMATORY BOWEL DISEASE

I-125 EXPANSION OF CONSCIOUS SEDATION INDICATION TO INCLUDE SHORT THERAPEUTIC PROCEDURES

I-126 ADJUNCT TO THALLIUM- 201 MYOCARDIAL PERFUSION IN PATIENTS UNABLE TO EXERCISE ADEQUATELY

I-127 TREATMENT OF ACYCLOVIR-RESISTANT HERPES IN IMMUNOCOMPROMISED PATIENTS

I-128 IN PT W/ CH DISEASE AND HYPERCHOLESTEROLEMIA: REDUCE RISK TOTAL MORTALITY BY REDUCING CORONARY DEATH; REDUCE RISK NON-FATAL MI; REDUCE RISK UNDERGOING MYOCARDIAL REVASCULARIZATION PROCEDURES; REDUCTION ELEVATED TOTAL AND LDL CHOL LEVELS...

I-129 TREATMENT OF ALCOHOL DEPENDENCE

I-130 MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS

I-131 PERIPHERAL ARTERIOGRAPHY

I-132 TREATMENT OF MANIC PHASE OF BIPOLAR DISORDER

I-133 MANAGEMENT OF CHRONIC STABLE ANGINA

I-134 HEART FAILURE POST MYOCARDIAL INFARCTION

I-135 BONE METASTASES ASSOCIATED WITH MULTIPLE MYELOMA

I-136 IDIOPATHIC CHRONIC URTICARIA

I-137 PREVENTION OF METAL-INDUCED HEART BURN, ACID INDIGESTION, AND SOUR STOMACH WHEN TAKEN 30 MINUTES PRIOR TO CONSUMING FOOD OR BEVERAGES

I-138 TREATMENT OF ACUTE RECURRENT GENITAL HERPES

I-139 PALLIATIVE TREATMENT OF ADVANCED BREAST CANCER IN PRE- AND PERIMENOPAUSAL WOMEN

I-140 PREVENTION OF CYTOMEGALOVIRUS (CMV) DISEASE IN INDIVIDUALS WITH HIV INFECTION AT RISK FOR DEVELOPING CMV DISEASE

I-141 TREATMENT OF HEMODYNAMICALLY STABLE PATIENTS WITHIN 24 HOURS OF ACUTE MYOCARDIAL INFARCTION TO IMPROVE SURVIVAL

I-142 LOCALIZE MYOCARDIAL ISCHEMIA (REVERSIBLE DEFECT) AND INFARCTION (NON-REVERSIBLE DEFECTS) IN EVALUATING MYOCARDIAL FUNCTION

I-143 EPISODIC TREATMENT OF RECURRENT GENITAL HERPES IN IMMUNOCOMPETENT ADULTS

I-144 ENHANCEMENT OF MRI OF THE ADULT BODY INTERNAL ORGANS

I-145 0.1MMOL/KG AS A SINGLE INTRAVENOUS BOLUS FOR MRI OF THE CNS IN CHILDREN

I-146 CONTRAST ENHANCEMENT AND FACILITATION OF VISUALIZATION OF EXTRACRANIAL HEAD AND NECK LESIONS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

| | |
|-------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| I-147 | PREVENTION OF GALLSTONE FORMATION IN OBESE PATIENTS EXPERIENCING RAPID WEIGHT LOSS |
| I-148 | TREATMENT OF ACUTE PNEUMOCYSTIS CARINI PNEUMONIA (PCP) IN HIV-INFECTED PATIENTS WHOSE ALVEOLAR-ARTERIAL OXYGEN DIFFERENCE (AaDO ₂) IS LESS THAN OR EQUAL TO 55 TORR |
| I-149 | TREATMENT OF PATIENTS WITH NON-SMALL CELL LUNG CANCER |
| I-150 | TREATMENT OF OBSESSIVE COMPULSIVE DISORDER AND PANIC DISORDER |
| I-151 | PREVENTION OF AND PREVENTION OF FURTHER POSTOPERATIVE NAUSEA AND VOMITING IN PEDIATRIC PATIENTS RECEIVING GENERAL ANESTHESIA |
| I-152 | SLOWING THE PROGRESSION OF CORONARY ATHEROSCLEROSIS AND REDUCING THE RISK OF ACUTE CORONARY EVENTS |
| I-153 | MANAGEMENT OF SEVERE SPASTICITY [ENCOMPASSES SPINAL AND CEREBRAL ORIGIN] |
| I-154 | PATIENT POPULATION ALTERED TO INCLUDE PEDIATRIC USE |
| I-155 | TREATMENT OF ONYCHOMYCOSIS DUE TO DERMATOPHYTES (TINEA UNGUIUM) OF THE TOENAIL WITH OR WITHOUT FINGERNAIL INVOLVEMENT |
| I-156 | ADDITIONAL DATA REGARDING THE SAFE USE OF NORVASC IN PATIENTS WITH HEART FAILURE |
| I-157 | TREATMENT OF ACUTE UNCOMPLICATED CYSTITIS IN FEMALES |
| I-158 | TREATMENT OF OSTEOLYTIC BONE METASTASES OF BREAST CANCER |
| I-159 | FOR HYPERCHOLESTEROLEMIC PATIENTS WITHOUT CLINICALLY EVIDENT HEART DISEASE REDUCE THE RISK OF MYOCARDIAL INFARCTION, REVASCULARIZATION, AND DEATH DUE TO CARDIOVASCULAR CAUSES WITH NO INCREASE IN DEATH FROM NON-CARDIOVASCULAR CAUSES |
| I-160 | TREATMENT OF BACTERIAL CORNEAL ULCERS |
| I-161 | TREATMENT OF ADULT-ONSET OR CHILDHOOD-ONSET ADULT GROWTH HORMONE DEFICIENCY |
| I-162 | FOR USE IN PATIENTS 6-11 YEARS OF AGE |
| I-163 | TREATMENT OF PHOTOPHOBIA |
| I-164 | CHRONIC BACTERIAL PROSTATITIS |
| I-165 | MANAGEMENT OF ADULTS WITH ACTIVE, CLASSIC AND DEFINITIVE RHEUMATOID ARTHRITIS WHO HAVE HAD INSUFFICIENT THERAPEUTIC RESPONSE TO OR ARE INTOLERANT OF AN ADEQUATE TRIAL OF FULL DOSES OF ONE OR MORE NON-STEROIDAL ANTI-INFLAMMATORY DRUGS |
| I-166 | TREATMENT OF BULIMIA |
| I-167 | COMPLICATED INTRA-ABDOMINAL INFECTIONS (USED IN COMBINATION WITH METRONIDAZOLE) CAUSED BY MIXED AEROBIC/ANAEROBIC PATHOGENS |
| I-168 | MANAGEMENT OF LOCALLY CONFINED STAGE B2-C METASTATIC CARCINOMA OF THE PROSTATE (IN COMBINATION WITH LHRH AGONISTS) |
| I-169 | USE IN COMBINATION WITH CORTICOSTEROIDS AS INITIAL CHEMOTHERAPY FOR THE TREATMENT OF PATIENTS WITH PAIN RELATED TO ADVANCED HORMONE-REFRACTORY PROSTATE CANCER |
| I-170 | PROPHYLACTIC USE DURING HEAD LICE EPIDEMICS |
| I-171 | RELIEF OF SYMPTOMS OF THE COMMON COLD |
| I-172 | TREATMENT OF INITIAL EPISODE OF GENITAL HERPES |
| I-173 | PREOPERATIVELY FOR THE PREVENTION OF INFECTION IN TRANSRECTAL PROSTATE BIOPSY |
| I-174 | PELVIC INFLAMMATORY DISEASE |
| I-175 | TREATMENT OF TINEA CORPORIS AND TINEA CRURIS |
| I-176 | TREATMENT OF POSTOPERATIVE INFLAMMATION IN PATIENTS WHO HAVE UNDERGONE CATARACT EXTRACTION |
| I-177 | TX OF MODERATE ACNE VULGARIS IN FEMALES, GREATER OR EQUAL TO 15YRS OF AGE, WHO HAVE NO KNOWN CONTRAINDICATIONS TO ORAL CONTRACEPTIVE THERAPY, DESIRE CONTRACEPTION, HAVE ACHIEVED MENARCHE AND ARE UNRESPONSIVE TO TOPICAL ANTI-ACNE MEDICATIONS |
| I-178 | TREATMENT OF ONYCHOMYCOSIS OF THE FINGERNAIL WITHOUT CONCOMITANT ONCHOMYCOSIS OF THE TOENAIL WITH A PULSE DOSING REGIMEN |
| I-179 | NOSOCOMIAL PNEUMONIA-MILD TO MODERATE AND SEVERE CAUSED BY HAEMOPHILUS INFLUENZAE OR KLEBSIELLA PNEUMONIAE |

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-180 TREATMENT OF PLANTAR TINEA PEDIS (MOCCASIN TYPE)

I-181 TREATMENT OF PATIENTS WITH COMPLEX PARTIAL SEIZURES WITH AND WITHOUT SECONDARY GENERALIZATION

I-182 TREATMENT OF GROWTH FAILURE ASSOCIATED WITH TURNER SYNDROME

I-183 MAINTENANCE THERAPY IN THE MANAGEMENT OF MILD TO MODERATE ASTHMA IN PEDIATRIC PATIENTS AGES 6-11

I-184 TREATMENT OF PANIC DISORDER AT A RECOMMENDED DOSE RANGE OF 1 TO 2MG/DAY (MAXIMUM OF 4MG)

I-185 PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN

I-186 TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR CAUSED BY OR PRESUMED TO BE CAUSED BY PITYROSPORUM ORBICULARE (ALSO KNOWN AS MALASSEZIA FURFUR OR M. ORBICULARE)

I-187 PREVENTION OF FRACTURES IN THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS

I-188 TREATMENT OF ACUTE SINUSITIS AND ACUTE EXACERBATION OF CHRONIC SINUSITIS

I-189 TREATMENT OF ACUTE OTITIS MEDIA IN PEDIATRIC PATIENTS

I-190 PLANAR IMAGING AS A SECOND LINE DIAGNOSTIC DRUG AFTER MAMMOGRAPHY TO ASSIST IN THE EVALUATION OF BREAST LESIONS IN PATIENTS WITH AN ABNORMAL MAMMOGRAM OR A PALPABLE BREAST MASS

I-191 ENDOMETRIAL THINNING AGENT PRIOR TO ENDOMETRIAL ABLATION FOR DYSFUNCTIONAL UTERINE BLEEDING

I-192 THE PREVENTION OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING ABDOMINAL SURGERY WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS AND A NEW DOSAGE REGIMEN, 40MG ONCE DAILY, FOR THIS INDICATION

I-193 TREATMENT OF PANIC DISORDER IN A RECOMMENDED DOSE RANGE OF 50 TO 200MG/DAY

I-194 CONGESTIVE HEART FAILURE

I-195 FOR USE OF LANSOPRAZOLE IN COMBINATION WITH CLARITHROMYCIN AND AMOXICILLIN FOR THE ERADICATION OF HELICOBACTER PYLORI IN PATIENTS WITH ACTIVE DUODENAL ULCER DISEASE OR A ONE-YEAR HISTORY OF DUODENAL ULCER

I-196 ACUTE TREATMENT OF ACTIVE BENIGN GASTRIC ULCER

I-197 MAINTENANCE OF HEALING OF DUODENAL ULCER

I-198 FOR THE USE OF LANSOPRAZOLE IN COMBINATION WITH AMOXICILLIN FOR THE ERADICATION OF HELICOBACTER PYLORI IN PATIENTS WITH ACTIVE DUODENAL ULCER DISEASE OR A ONE-YEAR HISTORY OF A DUODENAL ULCER

I-199 MONOTHERAPY AND COMBINATION THERAPY WITH SULFONYLUREA IN THE TREATMENT OF TYPE II DIABETES

I-200 TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR

I-201 EMPIRICAL THERAPY FOR FEBRILE NEUTROPENIC PATIENTS

I-202 SECOND-LINE TREATMENT OF AIDS-RELATED KAPOSI'S SARCOMA

I-203 MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS

I-204 USE IN PEDIATRIC PATIENTS BETWEEN THE AGES OF 6 AND 11 FOR THE TREATMENT OF THE NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS

I-205 INITIAL ANTICONVULSANT TREATMENT OF STATUS EPILEPTICUS

I-206 TREATMENT OF EDEMA ASSOCIATED WITH CHRONIC RENAL FAILURE

I-207 FOR THE SUPPRESSION OF RECURRENT EPISODES OF GENITAL HERPES IN IMMUNOCOMPETENT ADULTS

I-208 TREATMENT OF OBSESSIVE COMPULSIVE DISORDER IN THE PEDIATRIC POPULATION

I-209 PAROXYSMAL SUPRAVENTRICULAR TACHYCARDIA (PSVT)

I-210 TO SLOW THE PROGRESSION OF CORONARY ATHEROSCLEROSIS IN PATIENTS WITH CORONARY HEART DISEASE AS PART OF A TREATMENT STRATEGY TO LOWER TOTAL AND LDL CHOLESTEROL TO TARGET LEVELS

I-211 FOR USE IN PEDIATRIC POPULATION

I-212 TREATMENT OF SYMPTOMS OF DRY MOUTH IN PATIENTS WITH SJOGREN'S SYNDROME

I-213 TEMPORARY RELIEF OF PAIN AND PHOTOPHOBIA IN PATIENTS UNDERGOING CORNEAL REFRACTIVE SURGERY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

| | |
|-------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| I-214 | TREATMENT OF OSTEOPOROSIS |
| I-215 | PRE-PROCEDURAL APPLICATION TO ADULT MALE GENITAL SKIN PRIOR TO SITE-SPECIFIC SUBCUTANEOUS INFILTRATION WITH LIDOCAINE FOR THE REMOVAL OF GENITAL WARTS |
| I-216 | FOR THE LONG-TERM TWICE-DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD, INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA |
| I-217 | PREVENTION (DURING AND FOLLOWING HOSPITALIZATION) OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING HIP REPLACEMENT SURGERY |
| I-218 | USE OF LIPITOR AS AN ADJUNCTIVE THERAPY TO DIET FOR THE TREATMENT OF PATIENTS WITH ELEVATED SERUM TRIGLYCERIDE LEVELS (FREDERICKSON TYPE IV) |
| I-219 | USE OF LIPITOR BY PATIENTS WITH PRIMARY DYSBETALIPOPROTEINEMIA (FREDERICKSON TYPE III) WHO DO NOT RESPOND ADEQUATELY TO DIET |
| I-220 | TREATMENT OF EPISODIC- HEARTBURN, ACID INDIGESTION AND SOUR STOMACH |
| I-221 | TREATMENT OF BENIGN PROSTATIC HYPERPLASIA (BPH) IN MEN WITH AN ENLARGED PROSTATE TO IMPROVE SYMPTOMS, REDUCE THE RISK OF ACUTE URINARY RETENTION AND REDUCE THE RISK OF THE NEED OF SURGERY |
| I-222 | PREVENTION OF ISCHEMIC COMPLICATIONS OF UNSTABLE ANGINA AND NON-Q-WAVE MYOCARDIAL INFARCTION, WHEN CONCURRENTLY ADMINISTERED WITH ASPIRIN |
| I-223 | USE IN THE SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH ALLERGIC AND NONALLERGIC-PERENNIAL RHINITIS IN CHILDREN AGE 6-11 YEARS |
| I-224 | FOR THE USE IN PEDIATRIC PATIENTS 4 TO 11 YEARS OF AGE FOR THE MANAGEMENT OF THE NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS |
| I-225 | USE IN PATIENTS WITH PREVIOUS MI AND NORMAL CHOLESTEROL LEVELS, TO REDUCE RISK OF RECURRENT MI, MYOCARDIAL REVASCULARIZATION, AND CEREBROVASCULAR DISEASE EVENTS |
| I-226 | FIRST-LINE THERAPY FOR THE TREATMENT OF ADVANCED CARCINOMA OF THE OVARY IN COMBINATION WITH CISPLATIN |
| I-227 | SHORT-TERM TREATMENT OF SYMPTOMATIC GASTROESOPHAGEAL REFLUX DISEASE (GERD) |
| I-228 | PREVENTION OF MEAL INDUCED HEARTBURN AT A DOSE OF 75MG TAKEN 30-60MIN PRIOR TO A MEAL |
| I-229 | PRIOLOSEC (OMEPRAZOLE), AMOXICILLIN, AND CLARITHROMYCIN FOR THE ERADICATION OF H. PYLORI IN PATIENTS WITH DUODENAL ULCER DISEASE |
| I-230 | IN COMBINATION WITH CIS-PLATIN, FOR THE FIRST LINE TREATMENT OF NON-SMALL CELL LUNG CANCER IN PATIENTS WHO ARE NOT CANDIDATES FOR POTENTIALLY CURATIVE SURGERY AND/OR RADIATION |
| I-231 | TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC BREAST CANCER AFTER FAILURE OF PRIOR CHEMOTHERAPY |
| I-232 | TREATMENT OF RECURRENT MUCOCUTANEOUS HERPES SIMPLEX INFECTIONS IN HIV-AFFECTED PATIENTS AT A DOSE OF 500MG TWICE DAILY |
| I-233 | PROPHYLACTIC USE TO REDUCE PERIOPERATIVE BLOOD LOSS AND THE NEED FOR BLOOD TRANSFUSION IN PATIENTS UNDERGOING CARDIOPULMONARY BYPASS IN THE COURSE OF CORONARY ARTERY BYPASS GRAFT SURGERY |
| I-234 | FOR USE IN COMBINATION WITH CISPLATIN FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH INOPERABLE LOCALLY ADVANCED (STAGE IIIA OR IIIB) OR METASTATIC (STAGE IV) NON-SMALL CELL LUNG CANCER |
| I-235 | PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN PATIENTS 12 YEARS OF AGE AND OLDER |
| I-236 | PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN PATIENTS 4 YEARS OF AGE AND OLDER |
| I-237 | MAINTENANCE TREATMENT OF ASTHMA AND PREVENTION OF BRONCHOSPASM IN PATIENTS 4 YEARS OF AGE AND OLDER |
| I-238 | ADJUNCTIVE TREATMENT OF LENNOX-GASTAUT SYNDROME IN PEDIATRIC AND ADULT PATIENTS |
| I-239 | TREATMENT OF PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA |
| I-240 | MANAGEMENT OF SECONDARY HYPERPARATHYROIDISM AND RESULTANT METABOLIC BONE DISEASE IN PATIENTS WITH MODERATE TO SEVERE CHRONIC RENAL FAILURE (CCR 15 TO 55ML/MIN) NOT YET ON DIALYSIS |
| I-241 | USE IN PHOTODYNAMIC THERAPY (PDT) FOR REDUCTION OF OBSTRUCTION AND PALLIATION OF SYMPTOMS IN PATIENTS WITH COMPLETELY OR PARTIALLY OBSTRUCTING ENDOBRONCHIAL |

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

NONSMALL CELL LUNG CANCER

- I-242 TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH THE MENOPAUSE AND IN THE TREATMENT OF VULVAR AND VAGINAL ATROPHY IN WOMEN WITH AN INTACT UTERUS
- I-243 USE IN THE SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH THE COMMON COLD IN CHILDREN AGE 5 TO 11 YEARS
- I-244 REDUCE THE INCIDENCE OF BREAST CANCER IN WOMEN AT HIGH RISK FOR BREAST CANCER
- I-245 TREATMENT OF ACUTE SINUSITIS
- I-246 TREATMENT OF UNCOMPLICATED URINARY TRACT INFECTIONS
- I-247 USE IN CONVERSION TO MONOTHERAPY IN ADULTS WITH PARTIAL SEIZURES WHO ARE RECEIVING TREATMENT WITH A SINGLE ENZYME-INDUCING ANTIEPILEPTIC DRUG
- I-248 INPATIENT TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITH/WITHOUT PULMONARY EMBOLISM WHEN ADMIN WITH WARFARIN SODIUM AND OUTPATIENT TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITHOUT PULMONARY EMBOLISM WHEN ADMIN WITH WARFARIN SODIUM
- I-249 TREATMENT OF CHRONIC HEPATITIS C IN PATIENTS WITH COMPENSATED LIVER DISEASE PREVIOUSLY UNTREATED WITH ALPHA INTERFERON THERAPY
- I-250 PRIMARY PREVENTION OF CORONARY HEART DISEASE IN PATIENTS WITHOUT SYMPATOMATIC CARDIOVASCULAR DISEASE WHO HAVE AVERAGE TO MODERATELY ELEVATED TOTAL-C AND LDL-C AND BELOW AVERAGE HDL-C
- I-251 TREATMENT OF GENERALIZED ANXIETY DISORDER
- I-252 NEW COMBINATION USE OF PRECOSE FOR PATIENTS WITH TYPE 2 DIABETES TREATED WITH DIET PLUS METFORMIN
- I-253 COMBINATION USE OF PRECOSE FOR PATIENTS WITH TYPE 2 DIABETES TREATED WITH DIET PLUS INSULIN
- I-254 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS (LOSS OF BONE MASS)
- I-255 PREVENTION OF PNEUMOCYSTIS CARINII PNEUMONIA (PCP)
- I-256 USE IN TREATMENT OF SMALL CELL LUNG CANCER SENSITIVE DISEASE AFTER FAILURE OF FIRST-LINE CHEMOTHERAPY
- I-257 TREATMENT OF CHRONIC HEPATITIS B ASSOCIATED WITH EVIDENCE OF HEPATITIS B VIRAL REPLICATION AND ACTIVE LIVER INFLAMMATION
- I-258 FOR PERENNIAL NONALLERGIC RHINITIS FOR AGES 4 AND ABOVE
- I-259 PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT), WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING HIP REPLACEMENT SURGERY
- I-260 EXPANDED PEDIATRIC USE IN CHILDREN YOUNGER THAN ONE MONTH OF AGE TO BIRTH (WITH A GESTATIONAL AGE OF 37 WEEKS OR GREATER)
- I-261 TREATMENT OF SOCIAL ANXIETY DISORDER
- I-262 TREATMENT OR PREVENTION OF BRONCHOSPASM WITH REVERSIBLE OBSTRUCTIVE AIRWAY DISEASE AND FOR THE PREVENTION OF EXERCISE INDUCED BRONCHOSPASM IN CHILDREN AGES 4-12
- I-263 TREATMENT OF UNSTABLE ANGINA AND NON-Q-WAVE MYOCARDIAL INFARCTION FOR THE PREVENTION OF ISCHEMIC COMPLICATIONS IN PATIENTS ON CONCURRENT ASPIRIN THERAPY
- I-264 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH RADIATION, INCLUDING TOTAL BODY IRRADIATION (TBI) AND FRACTIONATED ABDOMINAL RADIATION
- I-265 TREATMENT OF ATOPIC DERMATITIS IN PEDIATRIC PATIENTS 6 YEARS AND OLDER
- I-266 USE OF TOPAMAX AS ADJUNCTIVE THERAPY IN PEDIATRIC PATIENTS AGES 2-16 YEARS WITH PARTIAL ONSET SEIZURES
- I-267 USE IN PEDIATRIC PATIENTS 3 MONTHS OLD AND OLDER - FOR CORTICOSTEROID-RESPONSIVE DERMATOSES
- I-268 PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA IN PATIENTS 7-11 YEARS OF AGE
- I-269 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH HIGHLY EMETOGENIC CANCER CHEMOTHERAPY, INCLUDING CISPLATIN
- I-270 ADJUVANT TREATMENT OF NODE-POSITIVE BREAST CANCER ADMINISTREDED SEQUENTIALLY TO STANDARD DOXORUBICIN-CONTAINING COMBINATION CHEMOTHERAPY
- I-271 TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- I-272 TREATMENT OF GLUCOCORTICOID-INDUCED OSTEOPOROSIS IN MEN AND WOMEN RECEIVING

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- GLUCOCORTICOIDS IN A DAILY DOSE EQUIVALENT TO 7.5MG OR GREATER OF PREDNISONE AND WHO HAVE LOW BONE MINERAL DENSITY
- I-273 ADJUNCT TO DIET TO INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA (HETEROZYGOUS FAMILIAL AND NON FAMILIAL) AND MIXED DYSLIPIDEMIA (FREDERICKSON TYPES IIA AND IIB)
- I-274 USE OF TOPAMAX AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES
- I-275 USE IN COMBINATION WITH METFORMIN AND SULFONYLUREA IN PATIENTS WITH TYPE 2 DIABETES
- I-276 USE OF REZULIN IN COMBINATION WITH METFORMIN AND SULFONYLUREAS IN PATIENTS WITH TYPE 2 DIABETES
- I-277 TREATMENT OF TYPE III HYPERLIPOPROTEINEMIA
- I-278 TREATMENT OF PATIENTS WITH ISOLATED HYPERTRIGLYCERIDEMIA (FREDERICKSON TYPE IV)
- I-279 TREATMENT OF POST-TRAUMATIC STRESS DISORDER
- I-280 USE OF CARNITOR INJECTION FOR THE PREVENTION AND TREATMENT OF CARNITINE DEFICIENCY IN PATIENTS WITH END STAGE RENAL DISEASE WHO ARE UNDERGOING DIALYSIS
- I-281 INCREASING HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA (HETEROZYGOUS FAMILIAL AND NONFAMILIAL) AND MIXED DYSLIPIDEMIA (FREDERICKSON TYPES IIA AND IIB)
- I-282 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER AFTER FAILURE OF PRIOR PLATINUM-BASED CHEMOTHERAPY
- I-283 TO REDUCE THE INCIDENCE OF MODERATE TO SEVERE XEROSTOMIA IN PATIENTS UNDERGOING POST-OPERATIVE RADIATION TREATMENT FOR HEAD AND NECK CANCER, WHERE THE RADIATION PORT INCLUDES A SUBSTANTIAL PORTION OF THE PAROTID GLANDS
- I-284 TO REDUCE THE NUMBER OF ADENOMATOUS COLORECTAL POLYPS IN FAMILIAL ADENOMATOUS POLYPOSIS PATIENTS AS AN ADJUNCT TO USUAL CARE
- I-285 TREATMENT OF NASAL SYMPTOMS OF SEASONAL AND PERENNIAL RHINITIS IN ADULTS AND CHILDREN 3 YEARS OF AGE AND OLDER
- I-286 TREATMENT OF PATIENTS WITH FREDERICKSON TYPE III
- I-287 USE OF PRAVASTATIN IN PATIENTS WITH EVIDENT CORONARY HEART DISEASE TO REDUCE THE RISK OF TOTAL MORTALITY BY REDUCING CORONARY DEATH
- I-288 CHANGES IN SEVERAL SECTIONS OF THE INSERT TO INCORPORATE STATEMENTS CONCERNING THE USE OF HIGH DOSES OF LISINAPRIL TO REDUCE THE RISK OF THE COMBINED OUTCOMES OF MORTALITY AND HOSPITALIZATION IN PATIENTS WITH CONGESTIVE HEART FAILURE
- I-289 USE OF AVANDIA IN COMBINATION WITH A SULFONYLUREA IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHEN DIET AND EXERCISE WITH EITHER SINGLE AGENT DOES NOT ACHIEVE ADEQUATE GLYCEMIC CONTROL
- I-290 PREVENTION OF CORTICOSTEROID-INDUCED OSTEOPOROSIS
- I-291 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- I-292 TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- I-293 TREATMENT OF CORTICOSTEROID-INDUCED OSTEOPOROSIS
- I-294 TREATMENT OF UNCOMPLICATED ACUTE ILLNESS DUE TO INFLUENZA A AND B IN PEDIATRIC PATIENTS 7 YEARS AND OLDER WHO HAVE BEEN SYMPTOMATIC FOR NO MORE THAN 2 DAYS
- I-295 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS FOR WOMEN WITH AN INTACT UTERUS
- I-296 LONG-TERM INTRAVENOUS TREATMENT OF PULMONARY HYPERTENSION ASSOCIATED WITH THE SCLERODERMA SPECTRUM OF DISEASE IN NYHA CLASS III AND CLASS IV PATIENTS WHO DO NOT RESPOND TO CONVENTIONAL THERAPY
- I-297 SHORT-TERM TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-298 TREATMENT OF PATIENTS WITH FREDERICKSON TYPE IIA AND IIB HYPERLIPOPROTEINEMIA
- I-299 USE OF CAMPTOSAR AS A COMPONENT OF FIRST-LINE THERAPY IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR PATIENTS WITH METASTATIC CARCINOMA OF THE COLON OR RECTUM
- I-300 PROPHYLAXIS FOR ASTHMA IN CHILDREN 2-5 YEARS OF AGE
- I-301 TREATMENT OF SIGNS AND SYMPTOMS OF ALLERGIC CONJUNCTIVITIS
- I-302 TREATMENT OF PEDIATRIC PATIENTS WITH PRADER-WILLI SYNDROME

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

| | |
|-------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| I-303 | INCREASING HDL-CHOLESTEROL IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA AND MIXED DYSLIPIDEMIAS |
| I-304 | TREATMENT OF PATIENTS WITH FREDERICKSON TYPE IV |
| I-305 | TREATMENT OF LEVOFLOXACIN SUSCEPTIBLE STRAINS OF PENICILLIN-RESISTANT STREPTOCOCCUS PNEUMONIAE IN PATIENTS WITH COMMUNITY ACQUIRED PNEUMONIA |
| I-306 | INDUCTION OF SPERMATOGENESIS IN MEN WITH PRIMARY AND SECONDARY HYPOGONADOTROPIC HYPOGONADISM IN WHOM THE CAUSE OF INFERTILITY IS NOT DUE TO PRIMARY TESTICULAR FAILURE |
| I-307 | NEW COMBINATION USE OF METFORMIN AND INSULIN IN TYPE 2 DIABETES |
| I-308 | TREATMENT OF PEDIATRIC PATIENTS WITH POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS WHO RESPONDED INADEQUATELY TO SALICYLATES OR OTHER NSAIDS |
| I-309 | USE OF ACTONEL 35MG ONCE A WEEK TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS |
| I-310 | REDUCTION IN RISK OF MYOCARDIAL INFARCTION, STROKE, AND DEATH FROM CARDIOVASCULAR CAUSES |
| I-311 | ADJUNCTIVE THERAPY IN THE TREATMENT OF PARTIAL SEIZURES IN PEDIATRIC PATIENTS AGE 3 TO 12 YEARS |
| I-312 | FIRST LINE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE OR HORMONE RECEPTOR UNKNOWN LOCALLY ADVANCED OR METASTATIC BREAST CANCER |
| I-313 | EXTENSION OF INDICATION TO PROVIDE FOR MAINTENANCE OF RESPONSE |
| I-314 | TOPICAL ANESTHETIC FOR SUPERFICIAL MINOR SURGERY OF GENITAL MUCOUS MEMBRANES AND AS AN ADJUNCT FOR LOCAL INFILTRATION ANESTHESIA IN GENITAL MUCOUS MEMBRANES |
| I-315 | THROMBOPROPHYLAXIS OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN MEDICAL PATIENTS WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS DUE TO SEVERELY RESTRICTED MOBILITY DURING ACUTE ILLNESS |
| I-316 | TREATMENT OF NSAID-ASSOCIATED GASTRIC ULCER PATIENTS WHO CONTINUE NSAID USE AND REDUCING RISK OF NSAID-ASSOCIATED GASTRIC ULCERS IN PATIENTS WITH HISTORY OF DOCUMENTED GASTRIC ULCER WHO REQUIRE USE OF AN NSAID |
| I-317 | PROPHYLAXIS OF INFLUENZA IN ADULTS AND ADOLESCENTS 13 YEARS AND OLDER |
| I-318 | FIRSTLINE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE OR HORMONE RECEPTOR UNKNOWN LOCALLY ADVANCED OR METASTATIC BREAST CANCER |
| I-319 | USE FOR SUSPECTED OR CONFIRMED METHANOL POISONING, EITHER ALONE OR IN COMBINATION WITH HEMODIALYSIS |
| I-320 | TREATMENT OF TYPE 2 DIABETES IN PEDIATRIC PATIENTS (AGES 10-16 YEARS) |
| I-321 | JUVENILE RHEUMATOID ARTHRITIS |
| I-322 | USE OF DIPRIVAN IN PATIENTS 3 MONTHS TO 16 YEARS |
| I-323 | COLORECTAL CANCER |
| I-324 | REDUCING NEUROLOGIC DISABILITY AND/OR FREQUENCY OF CLINICAL RELAPSES IN PATIENTS WITH SECONDARY (CHRONIC) PROGRESSIVE, PROGRESSIVE RELAPSING, OR WORSENING RELAPSING-REMITTING MULTIPLE SCLEROSIS |
| I-325 | PREVENTION OF RELAPSE AND RECURRENCE OF DEPRESSION |
| I-326 | GENERALIZED ANXIETY DISORDER |
| I-327 | SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN PATIENTS 5 YEARS AND OLDER |
| I-328 | PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA IN PATIENTS 5-6 YEARS OF AGE |
| I-329 | UNCOMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS |
| I-330 | MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS AND CONTROL OF DAYTIME AND NIGHTTIME HEARTBURN SYMPTOMS IN PATIENTS WITH GERD |
| I-331 | TREATMENT OF MODERATE ACNE VULGARIS |
| I-332 | EMPIRIC THERAPY IN FEBRILE NEUTROPENIC PATIENTS WITH SUSPECTED FUNGAL INFECTIONS (EFTN) |
| I-333 | TOPICAL TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR DUE TO MALASSEZIA FURFUR (FORMERLY PITYROSPORUM ORBICULARE) |
| I-334 | LONG-TERM TREATMENT OF GROWTH FAILURE IN CHILDREN BORN SMALL FOR GESTATIONAL AGE WHO FAIL TO MANIFEST CATCH-UP GROWTH BY TWO YEARS OF AGE |

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

| | |
|-------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| I-335 | ADJUNCTIVE THERAPY IN PATIENTS TWO YEARS AND OLDER WITH SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME |
| I-336 | EXPANSION OF INDICATION TO INCLUDE THE TREATMENT OF PATIENTS WITH PREDOMINATELY CLASSIC SUBFOVEAL CHOROIDAL NEOVASCULARIZATION DUE TO PATHOLOGIC MYOPIA OR PRESUMED OCULAR HISTOPLASMOSIS |
| I-337 | PATHOLOGICAL HYPERSECRETION ASSOCIATED WITH ZOLLINGER-ELLISON SYNDROME |
| I-338 | MANAGEMENT OF ACUTE PAIN IN ADULTS AND TREATMENT OF PRIMARY DYSMENORRHEA |
| I-339 | TREATMENT OF HEPATITIS B IN PEDIATRIC PATIENTS AGES 2-17 YEARS |
| I-340 | ATOPIC DERMATITIS IN PEDIATRIC PATIENTS AGES 2-5 |
| I-341 | BREAST CANCER COMBINATION THERAPY |
| I-342 | USE OF FORADIL FOR LONG-TERM, TWICE DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHO-CONSTRICTION IN PATIENTS WITH COPD INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA |
| I-343 | USE OF COREG FOR SEVERE HEART FAILURE |
| I-344 | ACNE VULGARIS |
| I-345 | TREATMENT OF POSTTRAUMATIC STRESS DISORDER |
| I-346 | TREATMENT OF SYMPTOMATIC GASTRO ESOPHAGEAL REFLUX DISEASE (GERD) |
| I-347 | TREATMENT OR PREVENTION OF BRONCHOSPASM IN CHILDREN 6 YEARS OF AGE AND OLDER WITH OBSTRUCTIVE AIRWAY DISEASE |
| I-348 | LONG-TERM, TWICE-DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD (INCLUDING EMPHYSEMA AND CHRONIC BRONCHITIS) |
| I-349 | ACUTE CORONARY SYNDROME |
| I-350 | TREATMENT OF HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA IN ADOLESCENT BOYS AND GIRLS AT LEAST ONE YEAR POSTMENARCHAL, AGES 10 TO 17 YEARS, WITH A RECOMMENDED DOSING RANGE OF 10 TO 40MG ONCE DAILY |
| I-351 | PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS FOR ALL STRENGTHS |
| I-352 | ANTICOAGULANT IN PATIENTS WITH OR AT RISK FOR HEPARIN-INDUCED THROMBOCYTOPENIA UNDERGOING PERCUTANEOUS CORONARY INTERVENTIONS (PCI) |
| I-353 | TREATMENT OF SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS |
| I-354 | MANAGEMENT OF POST HERPETIC NEURALGIA |
| I-355 | PREMENSTRUAL DYSPHORIC DISORDER |
| I-356 | TREATMENT OF PATHOLOGICAL HYPERSECRETORY CONDITIONS, INCLUDING ZOLLINGER-ELLISON SYNDROME |
| I-357 | TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS |
| I-358 | TREATMENT OF PANIC DISORDER |
| I-359 | TREATMENT OF VULVAR AND VAGINAL ATROPHY ASSOCIATED WITH MENOPAUSE |
| I-360 | TREATMENT OF NASAL SYMPTOMS OF SEASONAL AND PERENNIAL RHINITIS IN CHILDREN AGES TWO UP TO AGE THREE |
| I-361 | TREATMENT OF MULTIPLE MYELOMA AND DOCUMENTED BONE METASTASES FROM SOLID TUMORS, IN CONJUNCTION WITH STANDARD ANTINEOPLASTIC THERAPY. PROSTATE CANCER SHOULD HAVE PROGRESSED AFTER TREATMENT WITH AT LEAST ONE HORMONAL THERAPY |
| I-362 | TREATMENT OF PANIC DISORDER, WITH OR WITHOUT AGORAPHOBIA |
| I-363 | ADJUVANT TREATMENT OF POST MENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE EARLY BREAST CANCER |
| I-364 | TREATMENT OF COMMUNITY-ACQUIRED PNEUMONIA IN ADULTS |
| I-365 | TREATMENT OF HEART FAILURE (NYHA CLASS II-IV) IN PATIENTS WHO ARE INTOLERANT TO AN ACE INHIBITOR |
| I-366 | PREVENTION OF RELAPSE FOLLOWING LONG-TERM TREATMENT OF MAJOR DEPRESSIVE DISORDER |
| I-367 | COMBINATION THERAPY WITH THIAZOLIDINEDIONE TO LOWER BLOOD GLUCOSE IN PTS WHOSE HYPERGLYCEMIA CANNOT BE CONTROLLED BY DIET/EXERCISE PLUS MONOTHERAPY WITH ANY OF THE FOLLOWING AGENTS: METFORMIN, SULFONYLUREAS, REPAGLINIDE, OR THIAZOLIDINEDIONES |

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-368 USE OF GLUCOVANCE WITH A THIAZOLIDINEDIONE WHEN GLYCEMIC CONTROL IS NOT OBTAINED WITH GLUCOVANCE ALONE
- I-369 PREVENTION AND TREATMENT OF POSTOPERATIVE NAUSEA AND VOMITING
- I-370 TREATMENT OF HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA IN CHILDREN, AGES 8-13 YEARS, WITH RECOMMENDED DOSE OF 20MG ONCE DAILY AND IN ADOLESCENTS, AGES 14-18 WITH A RECOMMENDED DOSE OF 40MG ONCE DAILY
- I-371 HELICOBACTER PYLORI ERADICATION TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE
- I-372 NOSOCOMIAL PNEUMONIA
- I-373 TREATMENT OF TYPE 2 DIABETIC NEPHROPATHY
- I-374 SHORT TERM TOPICAL TREATMENT OF MILD TO MODERATE PLAQUE-TYPE PSORIASIS OF NON SCALP REGIONS
- I-375 FIRST LINE THERAPY FOR THE REDUCTION OF INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- I-376 TREATMENT OF NEWLY DIAGNOSED ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (CML)
- I-377 USE OF BRAVELLE FOR MULTIPLE FOLLICULAR DEVELOPMENT (CONTROLLED OVARIAN STIMULATION) DURING ASSISTED REPRODUCTIVE TECHNOLOGY CYCLES IN PATIENTS WHO HAVE PREVIOUSLY RECEIVED PITUITARY SUPPRESSION
- I-378 RELIEF OF SYMPTOMS OF SEASONAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- I-379 USE TAXOTERE IN COMBINATION WITH CISPLATIN FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER WHO HAVE NOT PREVIOUSLY RECEIVED CHEMOTHERAPY FOR THIS CONDITION
- I-380 TO TREAT PATIENTS WITH SCHIZOPHRENIA OR SCHIZOAFFECTIVE DISORDER AT RISK FOR EMERGENT SUICIDAL BEHAVIOR
- I-381 TREATMENT OF COLD SORES (HERPES LABIALIS) IN ADULT AND ADOLESCENT PATIENTS 12 YEARS OF AGE AND OLDER
- I-382 FOR NEWLY-DIAGNOSED HIGH GRADE MALIGNANT GLIOMA PATIENTS AS AN ADJUNCT TO SURGERY AND RADIATION
- I-383 TREATMENT OF TYPE 2 DIABETIC NEPHROPATHY
- I-384 USE IN COMBINATION WITH INSULIN FOR THE TREATMENT OF PATIENTS WITH TYPE 2 DIABETES MELLITUS
- I-385 MODIFICATION OF THE INDICATION FOR COMMUNITY ACQUIRED PNEUMONIA TO ADD "INCLUDING PENICILLIN-RESISTANT STRAINS, MIC PENICILLIN>=2MCG/ML" TO STREPTOCOCCUS PNEUMONIAE
- I-386 RAPAMUNE (SIROLIMUS) WITHIN AN IMMUNOSUPPRESSIVE REGIMEN THAT WOULD ALLOW FOR THE WITHDRAWAL OF CYCLOSPORINE 2 TO 4 MONTHS AFTER RENAL TRANSPLANTATION IN PATIENTS CONSIDERED AT LOW TO MODERATE IMMUNOLOGIC RISK FOR RENAL TRANSPLANT REJECTION
- I-387 ADJUNCTIVE THERAPY OF PARTIAL SEIZURES IN PEDIATRIC PATIENTS GREATER THAN OR EQUAL TO 2 YEARS OF AGE
- I-388 TREATMENT OF PATIENTS WITH LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION
- I-389 SUPPRESSION OF RECURRENT GENITAL HERPES IN HIV-INFECTED INDIVIDUALS
- I-390 USE IN PTS AT HIGH RISK CORONARY EVENTS DUE TO EXISTING CORONARY HEART DISEASE, DIABETES, PERIPHERAL VESSEL DISEASE, STROKE HISTORY, OTHER CV DISEASE TO REDUCE RISK TOTAL MORTALITY BY REDUCING CORONARY DEATH, REDUCE NONFATAL MI & STROKE.....
- I-391 ABLATION OF HIGH-GRADE DYSPLASIA IN BARRETT'S ESOPHAGUS PATIENTS WHO DO NOT UNDERGO ESOPHAGECTOMY
- I-392 TX OF PED PATIENTS W/PH+ CHRONIC PHASE CML DISEASE RECURRENCE AFTER STEM CELL TRANSPLANT OR RESISTANCE TO INTERFERON ALPHA THERAPY. NO CONTROLLED TRIALS DEMONSTRATING A CLINICAL BENEFIT SUCH AS IMPROVEMENT IN DISEASE RELATED SX OR INCREASED SURVIVAL
- I-393 CHRONIC BACTERIAL PROSTATITIS
- I-394 USE IN PATIENTS WITH CORONARY HEART DISEASE TO REDUCE THE RISK OF UNDERGOING CORONARY REVASCULARIZATION PROCEDURES

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-395 TO IMPROVE PHYSICAL FUNCTION

I-396 EXPANDED INDICATION TO INCLUDE THE ASSESSMENT OF VENTRICULAR FUNCTION IN SUBJECTS BEING EVALUATED FOR HEART DISEASE AND/OR VENTRICULAR FUNCTION

I-397 EXTENDED PROPHYLAXIS IN PATIENTS UNDERGOING HIP FRACTURE SURGERY

I-398 IDIOPATHIC SHORT STATURE

I-399 TREATMENT OF CANDIDEMIA AND THE FOLLOWING CANDIDA INFECTIONS: INTRA-ABDOMINAL ABSCESSSES, PERITONITIS AND PLEURAL SPACE INFECTIONS

I-400 USE OF OLANZAPINE IN COMBINATION WITH LITHIUM OR VALPROATE FOR THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR DISORDER

I-401 LONGER-TERM EFFICACY OF ARIPIRAZOLE IN THE TREATMENT OF SCHIZOPHRENIA

I-402 DIABETIC FOOT INFECTIONS WITHOUT CONCOMITANT OSTEOMYELITIS

I-403 USE OF VALTRESX IN COMBINATION WITH SAFER SEX PRACTICES FOR THE REDUCTION OF THE RISK OF TRANSMISSION OF GENITAL HERPES DURING SUPPRESSIVE THERAPY OF THE SOURCE PARTNER IN A HETEROSEXUAL COUPLE

I-404 MAINTENANCE TREATMENT OF BIPOLAR I DISORDER TO DELAY THE TIME TO OCCURRENCE OF MOOD EPISODES (DEPRESSION, MANIA, HYPOMANIA, MIXED EPISODES) IN PATIENTS TREATED FOR ACUTE MOOD EPISODES WITH STANDARD THERAPY

I-405 TREATMENT OF PREMENSTRUAL DYSPHORIC DISORDER (PMDD) USING AN INTERMITTENT DOSING REGIMEN

I-406 PREVENTION OF CYTOMEGALOVIRUS DISEASE IN KIDNEY, HEART, AND KIDNEY-PANCREAS TRANSPLANT PATIENTS AT HIGH RISK (DONOR CMV SEROPOSITIVE/RECIPIENT CMV SERONEGATIVE)

I-407 IMPROVE SURVIVAL OF STABLE PATIENTS WITH LEFT VENTRICULAR SYSTOLIC DYSFUNCTION (EJECTION FRACTION<=40%) AND CLINICAL EVIDENCE OF CONGESTIVE HEART FAILURE AFTER AN ACUTE MYOCARDIAL INFARCTION

I-408 STIMULATION OF PANCREATIC SECRETIONS TO FACILITATE THE IDENTIFICATION OF THE AMPULLA OF VATER AND ACCESSORY PAPILLA DURING ENDOSCOPIC RETROGRADE CHOLANGIO-PANCREATOGRAPHY (ERCP)

I-409 ESOPHAGEAL CANDIDIASIS

I-410 USE OF ADVAIR DISKUS 250/50 FOR CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) ASSOCIATED WITH CHRONIC BRONCHITIS

I-411 EXPANDED INDICATION FOR USE IN COMBINATION WITH ANTIDIABETIC DRUGS IN THE THIAZOLIDINEDIONE CLASS

I-412 MONOTHERAPY FOR THE SHORT TERM TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER

I-413 ADJUNCTIVE THERAPY FOR THE SHORT TERM TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER

I-414 PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT), WHICH MAY LEAD TO PULMONARY EMBOLISM (PE) IN MEDICAL PATIENTS WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS DUE TO SEVERELY RESTRICTED MOBILITY DURING ACUTE ILLNESS

I-415 SEVERE HYPERTENSION WHEN THE VALUE OF ACHIEVING PROMPT BLOOD PRESSURE CONTROL EXCEEDS THE RISK OF INITIATING COMBINATION THERAPY

I-416 THE USE OF CIPRO XR FOR COMPLICATED URINARY TRACT INFECTIONS AND ACUTE UNCOMPLICATED PYELONEPHRITIS

I-417 USE IN THE LONG TERM TREATMENT OF BIPOLAR I DISORDER

I-418 ADJUNCTIVE THERAPY W/ MOOD STABILIZERS (LITHIUM OR DIVALPROEX) IN THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDERS

I-419 MONOTHERAPY IN THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDER

I-420 TOPICAL TREATMENT OF CLINICALLY TYPICAL, NONHYPERKERATOTIC, NONHYPERTROPHIC ACTINIC KERATOSES ON THE FACE OR SCALP IN IMMUNOCOMPETENT ADULTS

I-421 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS AND PYELONEPHRITIS DUE TO E.COLI FOR PED PATIENTS (1-17) NOT AS FIRST CHOICE

I-422 INDICATED FOR THE IN-HOSPITAL SHORT-TERM (UP TO 4 HOURS) REDUCTION IN BLOOD PRESSURE IN PEDIATRIC PATIENTS

I-423 ACUTE TREATMENT OF MIGRAINE ATTACKS WITH OR WITHOUT AURA IN ADULTS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-424 MANAGEMENT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS WITH MODERATE TO SEVERE CHRONIC RENAL INSUFFICIENCY NOT YET ON DIALYSIS
- I-425 ELOXATIN IN COMBINATION WITH INFUSIONAL 5-FLUOROURACIL (5-FU) AND LEUCOVORIN (LV) FOR THE TREATMENT OF PATIENTS PREVIOUSLY UNTREATED FOR ADVANCED COLORECTAL CANCER
- I-426 TREATMENT OF ACUTE PULMONARY EMBOLISM WHEN ADMINISTERED IN CONJUNCTION WITH WARFARIN SODIUM
- I-427 TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITHOUT PULMONARY EMBOLISM WHEN ADMINISTERED IN CONJUNCTION WITH WARFARIN SODIUM
- I-428 FOR USE IN COMBINATION WITH PACLITAXEL FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC BREAST CANCER AFTER FAILURE OF PRIOR ANTHRACYCLINE CONTAINING ADJUVANT CHEMOTHERAPY UNLESS ANTHRACYCLINES WERE CLINICALLY CONTRAINDICATED
- I-429 FOR USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH ANDROGEN INDEPENDENT (HORMONE REFRACTORY) METASTATIC PROSTATE CANCER
- I-430 FOR USE IN THE RELIEF OF THE SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS IN ADULTS
- I-431 NOSOCOMIAL PNEUMONIA AND COMMUNITY-ACQUIRED PNEUMONIA CAUSED BY STREPTOCOCCUS PNEUMONIAE INDICATION EXPANDED TO INCLUDE MULTI-DRUG RESISTANT STRAINS
- I-432 TREATMENT OF COMMUNITY ACQUIRED PNEUMONIA CAUSED BY MULTI-DRUG RESISTANT STREPTOCOCCUS PNEUMONIAE
- I-433 TREATMENT OF BIOPSY-CONFIRMED, PRIMARY SUPERFICIAL BASAL CELL CARCINOMA IN IMMUNOCOMPETENT ADULTS, WITH A MAXIMUM TUMOR DIAMETER OF 2.0CM, LOCATED ON THE TRUNK (EXCLUDING ANOGENITAL SKIN), NECK, OR EXTREMITIES (EXCLUDING HANDS AND FEET)
- I-434 PREVENTION OF CARDIOVASCULAR DISEASE IN ADULT PATIENTS WITHOUT CLINICALLY EVIDENT HEART DISEASE, BUT WITH MULTIPLE RISK FACTORS FOR CORONARY HEART DISEASE TO REDUCE RISK OF MI AND RISK FOR REVASCULARIZATION PROCEDURES AND ANGINA
- I-435 CHRONIC IDIOPATHIC CONSTIPATION
- I-436 FOR USE IN COMBINATION WITH DOXORUBICIN AND CYCLOPHOSPHAMIDE FOR THE ADJUVANT TREATMENT OF PATIENTS WITH OPERABLE NODE-POSITIVE BREAST CANCER
- I-437 TREATMENT OF ACUTE MANIC AND MIXED EPISODES ASSOCIATED WITH BIPOLAR DISORDER
- I-438 EMPIRICAL THERAPY FOR PRESUMED FUNGAL INFECTIONS IN FEBRILE, NEUTROPENIC PATIENTS
- I-439 USED TO TREAT ADULTS WITH GROWTH HORMONE DEFICIENCY
- I-440 FOR THE REPLACEMENT OF ENDOGENOUS GROWTH HORMONE IN ADULTS WITH GROWTH HORMONE DEFICIENCY
- I-441 USE COMBINATION WITH INFUSIONAL 5-FU/LV FOR ADJUVANT TREATMENT STAGE III COLON CANCER PTS WHO HAVE UNDERGONE COMPLETE RESECTION PRIMARY TUMOR-BASED ON IMPROVEMENT IN DISEASE FREE SURVIVAL, NO DEMONSTRATED BENEFIT OVERALL SURVIVAL AFTER 4YRS
- I-442 USED FOR CANDIDEMIA IN NONNEUTROPENIC PATIENTS AND THE FOLLOWING CANDIDA INFECTIONS: DISSEMINATED INFECTIONS IN SKIN & INFECTIONS IN ABDOMEN, KIDNEY, BLADDER WALL, AND WOUNDS
- I-443 TREATMENT OF NASAL POLYPS IN PATIENTS 18 YEARS OF AGE AND OLDER
- I-444 USE OF PROTONIX IV FOR INJECTION AS STAND ALONE THERAPY FOR THE SHORT-TERM TREATMENT OF PATIENTS HAVING GASTROESOPHAGEAL REFLUX (GERD) WITH A HISTORY OF EROSIIVE ESOPHAGITIS
- I-445 TO IMPROVE (COMPARED TO 4.25% DEXTROSE) LONG-DWELL ULTRAFILTRATION AND CLEARANCE OF CREATININE AND UREA NITROGEN IN PATIENTS WITH HIGH AVERAGE OR GREATER TRANSPORT CHARACTERISTICS, AS DEFINED USING THE PERITONEAL EQUILIBRATION TEST (PET)
- I-446 EXTENDED ADJUVANT TREATMENT OF EARLY BREAST CANCER IN POSTMENOPAUSAL WOMEN WHO HAVE RECEIVED 5 YRS ADJUVANT TAMOXIFEN THERAPY-EFFECTIVENESS BASED ON AN ANALYSIS OF DISEASE FREE SURVIVAL IN PATIENTS TREATED FOR A MEDIAN 24 MONTHS
- I-447 USE OF COPEGUS (RIBAVIRIN) FOR TREATMENT OF CHRONIC HEPATITIS C IN ADULT PATIENTS COINFECTED WITH HIV IN COMBINATION WITH PEGASYS (PEGINTERFERON ALFA-2A)
- I-448 TREATMENT OF HEART FAILURE (NYHA CLASS II-IV AND EJECTION FRACTION <=40%) TO REDUCE THE RISK OF DEATH FROM CARDIOVASCULAR CAUSES AND TO REDUCE HOSPITALIZATIONS FOR HEART FAILURE

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-449 TO IMPROVE WAKEFULNESS IN TWO NEW PATIENT POPULATIONS WITH EXCESSIVE SLEEPINESS: THOSE WITH OBSTRUCTIVE SLEEP APNEA/HYPOPNEA SYNDROME AND THOSE WITH SHIFT WORK SLEEP DISORDER
- I-450 TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED HIGH GRADE GLIOMAS CONCOMITANTLY WITH RADIOTHERAPY AND THEN AS ADJUVANT TREATMENT
- I-451 MANAGEMENT OF ENDOMETRIOSIS ASSOCIATED PAIN
- I-452 EXPANDED INDICATION TO INCLUDE TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST 1 PRIOR THERAPY
- I-453 USE IN COMBINATION WITH A SULFONYLUREA PLUS METFORMIN WHEN DIET, EXERCISE AND BOTH AGENTS DO NOT RESULT IN ADEQUATE GLYCEMIC CONTROL (TRIPLE THERAPY)
- I-454 MAINTENANCE OF CLINICAL REMISSION OF MILD TO MODERATE CROHN'S DISEASE INVOLVING THE ILEUM AND/OR THE ASCENDING COLON FOR UP TO 3 MONTHS
- I-455 MODIFIED HEART FAILURE INDICATION TO INCLUDE TREATMENT OF HEART FAILURE IN PATIENTS WITH LEFT VENTRICULAR SYSTOLIC DYSFUNCTION (NYHA CLASS II-IV; EJECTION FRACTION LESS THAN OR EQUAL TO 40%)
- I-456 TO REDUCE CARDIOVASCULAR DEATH AND TO REDUCE HEART FAILURE HOSPITALIZATIONS. INCLUDES ADDITIONAL INFORMATION ON THE ADDED EFFECT ON THESE OUTCOMES WHEN USED WITH AN ACE INHIBITOR
- I-457 TREATMENT OF PATIENTS UNDERGOING ABDOMINAL SURGERY WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS
- I-458 USE OF BIVALIRUDIN FOR INJECTION WITH PROVISIONAL USE OF GLYCOPROTEIN IIB/IIIA INHIBITOR (GPI) AS LISTED IN THE CLINICAL TRIALS REPLACE-2 SECTION FOR USE AS AN ANTICOAGULANT IN PATIENTS UNDERGOING PERCUTANEOUS CORONARY INTERVENTION (PCI)
- I-459 NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE (NDD-CKD) PATIENTS RECEIVING OR NOT RECEIVING AN ERYTHROPOIETIN
- I-460 TREATMENT OF DIARRHEA CAUSED BY CRYPTOSPORIDIUM PARVUM IN NON-HIV INFECTED PATIENTS 12 YEARS OF AGE AND OLDER
- I-461 USE AS A SINGLE AGENT FOR ADJUVANT TREATMENT IN PATIENTS WITH DUKES' C COLON CANCER WHO HAVE UNDERGONE COMPLETE RESECTION OF THE PRIMARY TUMOR WHEN TREATMENT WITH FLUOROPYRIMIDINE THERAPY ALONE IS PREFERRED
- I-462 LONG TERM TREATMENT OF IDIOPATHIC SHORT STATURE
- I-463 TREATMENT OF PATIENTS POST MYOCARDIAL INFARCTION
- I-464 TREATMENT OF MODERATE TO SEVERE PRIMARY RESTLESS LEGS SYNDROME
- I-465 PERENNIAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 6 MONTHS OF AGE AND OLDER
- I-466 FOR RELIEF OF THE SIGNS AND SYMPTOMS OF ANKYLOSING SPONDYLITIS
- I-467 USE OF TOPIRAMATE AS INITIAL MONOTHERAPY IN PATIENTS 10 YEARS OF AGE AND OLDER WITH PARTIAL ONSET OR PRIMARY GENERALIZED TONIC CLONIC SEIZURES
- I-468 USE IN PATIENTS WITH STABLE CORONARY ARTERY DISEASE TO REDUCE THE RISK OF CARDIOVASCULAR MORTALITY OR NON-FATAL MYOCARDIAL INFARCTION
- I-469 RELIEF OF THE SIGNS AND SYMPTOMS OF PAUCIARTICULAR OR POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS IN PATIENTS 2 YEARS OF AGE AND OLDER
- I-470 DIABETIC PERIPHERAL NEUROPATHIC PAIN
- I-471 INDICATED TO REDUCE THE RISK OF MYOCARDIAL INFARCTION AND STROKE IN PATIENTS WITH TYPE 2 DIABETES AND WITHOUT CLINICALLY EVIDENT CORONARY HEART DISEASE BUT WITH MULTIPLE RISK FACTORS FOR CORONARY HEART DISEASE
- I-472 USE IN PATIENTS WITH ANGIOGRAPHICALLY DOCUMENTED CORONARY ARTERY DISEASE
- I-473 USE IN COMBINATION WITH GEMCITABINE FOR THE FIRST LINE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC PANCREATIC CANCER
- I-474 TREATMENT OF IRON DEFICIENCY ANEMIA IN PERITONEAL DIALYSIS DEPENDANT CHRONIC KIDNEY DISEASE IN PATIENTS RECEIVING AN ERYTHROPOIETIN
- I-475 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF MODERATELY EMETOGENIC CANCER CHEMOTHERAPY
- I-476 TREATMENT OF DIABETIC FOOT INFECTIONS WITHOUT OSTEOMYELITIS
- I-477 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS CAUSED BY METHICILLIN SUSCEPTIBLE STAPHYLOCOCCUS AUREUS, ESCHERICHIA COLI, KLEBSIELLA PNEUMONIAE, OR ENTEROBACTER CLOACAE

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- I-478 FOR USE AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PARTIAL SEIZURES IN CHILDREN WITH EPILEPSY AGED 2-4 YEARS
- I-479 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTIONS CAUSED BY E.COLI, B. FRAGILIS, S.ANGINOSUS, S.CONSTELLATUS, E. FAECALIS, P. MIRABILIS, C. PERFRINGENS, B. THETA IOTAOMICRON OR PEPTOSTREPTOCOCCUS SPECIES
- I-480 PROPHYLAXIS OF INFLUENZA FOR PATIENTS BETWEEN 1-12 YEARS OF AGE
- I-481 INDICATED FOR THE ADJUVANT TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE EARLY BREAST CANCER
- I-482 TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER WITH OR WITHOUT PSYCHOTIC FEATURES
- I-483 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- I-484 FOR THE RISK REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCERS
- I-485 TREATMENT OF POSTOPERATIVE INFLAMMATION AND REDUCTION OF OCULAR PAIN IN PATIENTS WHO HAVE UNDERGONE CATARACT EXTRACTION
- I-486 ANGIOMAX IS INDICATED FOR PATIENTS WITH, OR AT RISK OF, HIT/HITTS UNDERGOING PCI
- I-487 INDICATED FOR THE RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF CORTICOSTEROID RESPONSIVE DERMATOSES IN PATIENTS 12 YRS OF AGE OR OLDER
- I-488 MAINTENANCE THERAPY IN BIPOLAR I DISORDER
- I-489 FOR USE IN PEDIATRIC PATIENTS WITH TYPE I DIABETES
- I-490 FOR USE IN COMBINATION WITH CISPLATIN AND FLUOROURACIL FOR THE TREATMENT OF PATIENTS WITH ADVANCED GASTRIC ADENOCARCINOMA, INCLUDING ADENOCARCINOMA OF GASTROESOPHAGEAL JUNCTION, WHO HAVE NOT RECEIVED PRIOR CHEMOTHERAPY FOR ADVANCED DISEASE
- I-491 INFLUENZA PROPHYLAXIS
- I-492 MONOTHERAPY IN THE TREATMENT OF ACUTE MANIC OR MIXED EPISODES IN BIPOLAR I DISORDER, WITH OR WITHOUT PSYCHOTIC FEATURES
- I-493 ADMINISTERED IN COMBINATION WITH FENOFIBRATE, AS ADJUNCTIVE THERAPY TO DIET FOR THE REDUCTION OF ELEVATED TOTAL-C, LDL-C, APO B, AND NON-HDL-C IN PATIENTS WITH MIXED HYPERLIPIDEMIA
- I-494 CLINICAL DATA IN SUPPORT OF AVANDAMET AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHEN TREATMENT WITH DUAL ROSIGLITAZONE AND METFORMIN THERAPY IS APPROPRIATE
- I-495 ADJUVANT TX OF POSTMENOPAUSAL WOMEN WITH ESTROGEN-RECEPTOR POSITIVE EARLY BREAST CANCER WHO HAVE RECEIVED 2 TO 3 YRS OF TAMOXIFEN AND ARE SWITCHED TO AROMASIN FOR COMPLETION OF A TOTAL OF 5 CONSECUTIVE YRS OF ADJUVANT HORMONAL THERAPY
- I-496 LONG TERM TREATMENT OF GROWTH FAILURE ASSOCIATED WITH TURNER SYNDROME IN PATIENTS WHO HAVE OPEN EPIPHYSES
- I-497 PREVENTION OF SEASONAL MAJOR DEPRESSIVE EPISODES IN PATIENTS WITH SEASONAL AFFECTIVE DISORDER
- I-498 PREVENTION OF POSTOPERATIVE NAUSEA AND VOMITING
- I-499 USE OF GEMZAR IN COMBINATION WITH CARBOPLATIN FOR THE TREATMENT OF PATIENTS WITH ADVANCED OVARIAN CANCER THAT HAS RELAPSED AT LEAST 6 MONTHS AFTER COMPLETION OF PLATINUM-BASED THERAPY
- I-500 FOR USE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- I-501 TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) IN IMMUNOCOMPETENT PATIENTS WITH A SINGLE DOSE OF FAMCICLOVIR 1500 MG.
- I-502 FOR PTS WITH ST-SEGMENT ELEVATION ACUTE MYOCARDIAL INFARCTION, PLAVIX TO REDUCE RATE OF DEATH FROM ANY CAUSE AND THE RATE OF A COMBINED ENDPOINT OF DEATH, REINFARCTION OR STROKE. NOT KNOWN TO PERTAIN TO PTS WHO RECEIVE PRIMARY ANGIOPLASTY
- I-503 TREATMENT OF MAJOR DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR DISORDER
- I-504 TREATMENT OF PATHOLOGICAL HYPERSECRETORY CONDITIONS INCLUDING ZOLLINGER-ELLISON SYNDROME
- I-505 TREATMENT OF STAPHYLOCOCCUS AUREUS BLOODSTREAM INFECTIONS (BACTEREMIA), INCLUDING THOSE WITH RIGHT SIDED INFECTIVE ENDOCARDITIS, CAUSED BY METHICILLIN-SUSCEPTIBLE AND METHICILLIN-RESISTANT ISOLATES

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

| | |
|-------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| I-506 | ADJUNCTIVE THERAPY OF MYOCLONIC SEIZURES IN ADULTS AND ADOLESCENTS AGE 12 AND OVER WITH JUVENILE MYOCLONIC EPILEPSY |
| I-507 | ADJUNCT TO DIET TO REDUCE TOTAL-C, LDL-C AND APO B LEVELS IN ADOLESCENT BOYS AND GIRLS WHO ARE AT LEAST ONE YEAR POST-MENARCHE, 10-16 YEARS OF AGE, WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA |
| I-508 | PREMENSTRUAL DYSPHONIC DISORDER |
| I-509 | TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER |
| I-510 | ADULT DERMATOFIBROSARCOMA PROTUBERANS (DFSP) |
| I-511 | ADULT MYELODYSPLASTIC SYNDROME/MYELOPROLIFERATIVE DISEASES (MDS/MDP) |
| I-512 | ADULT PH+ ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) MONOTHERAPY |
| I-513 | ADULT AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM) |
| I-514 | ADULT HYPEREOSINOPHILIC SYNDROME/CHRONIC EOSINOPHILIC LEUKEMIA (HES/CEL) |
| I-515 | PROPHYLAXIS OF SURGICAL SITE INFECTION FOLLOWING ELECTIVE COLORECTAL SURGERY |
| I-516 | PRIMARY GENERALIZED TONIC CLONIC SEIZURES IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER |
| I-517 | TREATMENT OF MODERATE TO SEVERE PRIMARY RESTLESS LEG SYNDROME (RLS) |
| I-518 | TREATMENT OF SHORT STATURE OR GROWTH FAILURE IN CHILDREN WITH SHOX (SHORT STATURE HOMEBOX CONTAINING GENE) DEFICIENCY WHOSE EPIPHYSES ARE NOT CLOSED |
| I-519 | USE OF TAXOTERE (DOCETAXEL) INJECTION CONCENTRATE IN COMBINATION WITH CISPLATIN AND FLUOROURACIL FOR THE INDUCTION OF PATIENTS WITH INOPERABLE LOCALLY ADVANCED SQUAMOUS CELL CARCINOMA OF THE HEAD AND NECK (SCCHN) |
| I-520 | USE OF EXENATIDE IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHO ARE USING A THIAZOLIDINEDIONE ALONE OR IN COMBINATION WITH METFORMIN BUT HAVE NOT ACHIEVED ADEQUATE GLYCEMIC CONTROL |
| I-521 | TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST 1 YEAR PRIOR THERAPY |
| I-522 | TREATMENT OF MODERATE ACNE VULGARIS IN WOMEN AT LEAST 14 YRS OF AGE, WHO HAVE NO KNOWN CONTRAINDICATIONS TO ORAL CONTRACEPTIVE THERAPY, AND HAVE ACHIEVED MENARCHE, IF THE PATIENT DESIRES AN ORAL CONTRACEPTIVE FOR BIRTH CONTROL. |
| I-523 | USE IN ADULT PATIENTS WITH CLINICALLY EVIDENT CORONARY HEART DISEASE TO REDUCE THE RISK OF NONFATAL MYOCARDIAL INFARCTION, FATAL AND NONFATAL STROKE, ANGINA, REVASCULARIZATION PROCEDURES AND HOSPITALIZATION FOR CONGESTIVE HEART FAILURE |
| I-524 | GENERALIZED ANXIETY DISORDER (GAD) |
| I-525 | USE OF 0.5MG/0.1MG FOR PREVENTION OF POST-MENOPAUSAL OSTEOPOROSIS |
| I-526 | TREATMENT OF HYPONATREMIA IN HOSPITALIZED PATIENTS |
| I-527 | ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER WITH IDIOPATHIC GENERALIZED EPILEPSY |
| I-528 | TREATMENT OF MODERATE TO SEVERE VAGINAL DRYNESS AND PAIN WITH INTERCOURSE, SYMPTOMS OF VULVAR AND VAGINAL ATROPHY ASSOCIATED WITH MENOPAUSE |
| I-529 | TREATMENT OF DEMENTIA OF THE ALZHEIMER'S TYPE IN PATIENTS WITH SEVERE ALZHEIMER'S DISEASE |
| I-530 | PREVENTION OF EXERCISE-INDUCED BRONCHOCONSTRICTION IN PATIENTS 15 YEARS OF AGE AND OLDER |
| I-531 | MAINTENANCE TREATMENT OF SCHIZOPHRENIA |
| I-532 | TREATMENT OF BACTERIAL VAGINOSIS IN NON-PREGNANT FEMALES |
| I-533 | ACUTE ST-SEGMENT ELEVATION MYOCARDIAL INFARCTION (STEMI) |
| I-534 | EXTENDED TREATMENT OF SYMPTOMATIC VENOUS THROMBOEMBOLISM (VTE) AND/OR PULMONARY EMBOLISM TO REDUCE THE RECURRENCE OF VTE IN PATIENTS WITH CANCER |
| I-535 | MANAGEMENT OF FIBROMYALGIA |
| I-536 | FOR THE TREATMENT OF SHORT STATURE IN CHILDREN WITH NOONAN SYNDROME |
| I-537 | LONG TERM TREATMENT OF PANIC DISORDER |
| I-538 | SHORT TERM TREATMENT OF PANIC DISORDER |

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

I-539 REDUCTION IN RISK OF INVASIVE BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH OSTEOPOROSIS OR AT HIGH RISK FOR INVASIVE BREAST CANCER

I-540 TREATMENT OF SCHIZOPHRENIA IN ADOLESCENTS AGES 13-17

I-541 TREATMENT OF BIPOLAR I DISORDER IN CHILDREN AGES 10-12 AND ADOLESCENTS AGES 13-17

I-542 EXPANSION OF PATIENT POPULATION FOR HEAD AND NECK CANCER FROM "INOPERABLE" PATIENTS TO ALL PATIENTS

I-543 USE IN COMBINATION WITH CISPLATIN AND FLUOROURACIL FOR THE INDUCTION TREATMENT OF PATIENTS WITH LOCALLY ADVANCED SQUAMOUS CELL CARCINOMA OF THE HEAD AND NECK (SCCHN)

I-544 ADJUNCTIVE THERAPY OF MYOCLONIC SEIZURES IN ADULTS AND ADOLESCENTS AGE 16 AND OVER WITH JUVENILE MYOCLONIC EPILEPSY

I-545 ADJUNCTIVE TREATMENT TO TREAT PATIENTS WITH MAJOR DEPRESSIVE DISORDER

I-546 TREATMENT OF UNRESECTABLE HEPATOCELLULAR CARCINOMA

I-547 ADJUNCTIVE THERAPY TO DIET TO SLOW THE PROGRESSION OF ARTEROSCLEROSIS IN ADULT PATIENTS AS PART OF A TREATMENT STRATEGY TO LOWER TOTAL-C AND LDL-C TO TARGET LEVELS

I-548 SEASONAL ALLERGIC RHINITIS IN PATIENTS 6 THROUGH LESS THAN 12 YEARS OF AGE

I-549 USE OF AVALIDE TABLETS AS INITIAL THERAPY IN PATIENTS WHO ARE LIKELY TO NEED MULTIPLE DRUGS TO ACHIEVE THEIR BLOOD PRESSURE GOALS

I-550 TREATMENT OF HYPERTENSION IN PEDIATRIC PATIENTS 6-16 YEARS OF AGE

I-551 TREATMENT OF SHORT STATURE IN CHILDREN WITH TURNER'S SYNDROME

I-552 ADJUNCTIVE TREATMENT FOR RADIOIODINE ABLATION OF THYROID TISSUE REMNANTS IN PATIENTS WHO HAVE UNDERGONE THYROIDECTOMY FOR WELL-DIFFERENTIATED THYROID CANCER AND WHO DO NOT HAVE EVIDENCE OF METASTATIC THYROID CANCER

I-553 FOR USE AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

I-554 TREATMENT OF PATIENTS WITH CANDIDEMIA, ACUTE DISSEMINATED CANDIDIASIS, CANDIDA PERITONITIS AND ABSCESES

I-555 TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER IN PEDIATRIC PATIENTS AGED 10-17 YEARS

I-556 PREVENTION OF POST OPERATIVE NAUSEA AND VOMITING FOR UP TO 24 HOURS FOLLOWING SURGERY

I-557 USE OF AMITIZA (LUBIPROSTONE) 8 MCG TWICE DAILY FOR TREATMENT OF IRRITABLE BOWEL SYNDROME WITH CONSTIPATION IN WOMEN GREATER THAN OR EQUAL TO 18 YEARS OLD

I-558 MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION AND REDUCING EXACERBATIONS IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA

I-559 ADJUNCTIVE THERAPY ADDED TO LITHIUM OR VALPROATE IN SHORT TERM TREATMENT OF BIPOLAR DISORDER, MANIC OR MIXED

I-560 MAINTENANCE TREATMENT FOR BIPOLAR I DISORDER, AS ADJUNCTIVE THERAPY TO LITHIUM OR DIVALPROEX

I-561 LONG-TERM TREATMENT OF SOCIAL ANXIETY DISORDER

I-562 MAINTENANCE TREATMENT OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER (ADHD) IN CHILDREN AND ADOLESCENTS

I-563 ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN ADULTS AND CHILDREN 16 YEARS OF AGE AND OLDER WITH IDIOPATHIC GENERALIZED EPILEPSY

I-564 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA

I-565 USE OF DUTASTERIDE IN COMBINATION WITH TAMSULOSIN FOR THE TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA (BPH)

I-566 MANAGEMENT OF FIBROMYALGIA

I-567 INITIAL THERAPY IN PATIENTS LIKELY TO NEED MULTIPLE DRUGS TO ACHIEVE THEIR BLOOD PRESSURE GOALS

I-568 USE OF APTIVUS, CO-ADMINISTERED W/RITONAVIR, FOR COMBINATION ANTIRETROVIRAL TREATMENT OF HIV-1 INFECTED PED (AGE 2-18 YRS) PATIENTS WHO ARE TREATMENT-

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

EXPERIENCED AND INFECTED W/HIV-1 STRAINS RESISTANT TO MORE THAN ONE PROTEASE INHIBITOR

I-569 TREATMENT OF CHRONIC HEPATITIS B

I-570 TREATMENT OF CHICKEN POX IN IMMUNOCOMPETENT PEDIATRIC PATIENTS 2 TO <18 YEARS OF AGE

I-571 NON-SMALL CELL LUNG CANCER IN COMBINATION WITH CISPLATIN AND AS SINGLE AGENT FOR NONSQUAMOUS NON-SMALL CELL LUNG CANCER

I-572 TREATMENT OF GROWTH FAILURE IN CHILDREN BORN SMALL FOR GESTATIONAL AGE (SGA) WITH NO CATCH-UP BY AGE 2-4 YRS.

I-573 TO TREAT PATIENTS WITH PRIMARY DYSBETALIPOPROTEINEMIA (FREDRICKSON TYPE III HYPERLIPOPROTEINEMIA) AS AN ADJUNCT TO DIET

I-574 MONOTHERAPY IN THE TREATMENT OF BIPOLAR DEPRESSION

I-575 MONOTHERAPY IN THE TREATMENT OF BIPOLAR MANIA

I-576 ADJUNCTIVE THERAPY IN THE TREATMENT OF BIPOLAR MANIA

I-577 SEDATION OF NON-INTUBATED PATIENTS PRIOR TO AND/OR DURING SURGICAL AND OTHER PROCEDURES

I-578 EXPANSION OF INDICATION TO INCLUDE TREATMENT OF HIV IN TREATMENT NAIVE ADULTS

I-579 TREATMENT OF MODERATE TO SEVERE DYSpareunia, A SYMPTOM OF VULVAR AND VAGINAL ATROPHY, DUE TO MENOPAUSE AND NEW TWICE WEEKLY DOSING REGIMEN FOR THIS INDICATION

I-580 INDOLENT B-CELL NON-HODGKINS LYMPHOMA (NHL) THAT HAS PROGRESSED DURING OR WITHIN SIX MONTHS OF TREATMENT WITH RITUXIMAB OR A RITUXIMAB CONTAINING REGIMEN

I-581 TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS

I-582 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE

I-583 ADJUVANT TREATMENT OF ADULT PATIENTS FOLLOWING COMPLETE GROSS RESECTION OF KIT (CD117) POSITIVE GASTROINTESTINAL STROMAL TUMORS (GIST)

I-584 TREATMENT AND PREVENTION OF GLUCOCORTICOID-INDUCED OSTEOPOROSIS IN PATIENTS EXPECTED TO BE ON GLUCOCORTICIDS FOR AT LEAST 12 MONTHS

I-585 TREATMENT OF SHORT STATURE IN PEDIATRIC PATIENTS SMALL FOR GESTATIONAL AGE WHO DO NOT MANIFEST CATCH UP GROWTH BY AGE 2 TO 4 YEARS

I-586 COMMUNITY ACQUIRED BACTERIAL PNEUMONIA

I-587 ADDITIONAL PATHOGENS TO COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS INDICATION

I-588 ADDITIONAL PATHOGENS TO COMPLICATED INTRA-ABDOMINAL INFECTIONS INDICATION

I-589 TREATMENT OF TREATMENT RESISTANT DEPRESSION (TRD) IN COMBINATION WITH OLANZAPINE

I-590 ACUTE TREATMENT OF DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR DISORDER (IN COMBINATION WITH OLANZAPINE)

I-591 TREATMENT OF TREATMENT RESISTANT DEPRESSION (TRD) IN COMBINATION WITH FLUOXETINE

I-592 ACUTE TREATMENT OF DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR DISORDER (IN COMBINATION WITH FLUOXETINE)

I-593 TREATMENT OF TREATMENT RESISTANT DEPRESSION (TRD)

I-594 INDICATION EXPANDED TO INCLUDE PATIENTS WHO HAVE EXPERIENCED A FIRST CLINICAL EPISODE AND HAVE MRI FEATURES CONSISTENT WITH MULTIPLE SCLEROSIS

I-595 PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN

I-596 USE AS ADJUNCTIVE THERAPY WITH LITHIUM OR VALPROATE FOR THE MAINTENANCE TREATMENT OF BIPOLAR I DISORDER

I-597 MONOTHERAPY FOR THE MAINTENANCE TREATMENT OF BIPOLAR I DISORDER

I-598 TREATMENT OF PULMONARY ARTERIAL HYPERTENSION INDICATION EXPANDED TO INCLUDE DELAY IN CLINICAL WORSENING

I-599 PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGE 5 IN PATIENTS ON HEMODIALYSIS OR PERITONEAL DIALYSIS

I-600 FOR USE AS INITIAL THERAPY IN PATIENTS WHO ARE LIKELY TO NEED MULTIPLE DRUGS TO ACHIEVE THEIR BLOOD PRESSURE GOALS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

| | |
|-------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| I-601 | MAINTENANCE TREATMENT IN PATIENTS WITH ADVANCED OR METASTATIC NONSQUAMOUS NON-SMALL CELL LUNG CANCER WHOSE DISEASE HAS NOT PROGRESSED AFTER FOUR CYCLES OF PLATINUM-BASED FIRST LINE CHEMOTHERAPY |
| I-602 | TREATMENT OF MEN AND WOMEN WITH OSTEOPOROSIS ASSOCIATED WITH SUSTAINED SYSTEMIC GLUCOCORTICOID THERAPY AT HIGH RISK FOR FRACTURE |
| I-603 | GOUT FLARES |
| I-604 | PREVENTION OF CMV DISEASE IN KIDNEY AND HEART TRANSPLANT PATIENTS 4 MONTHS TO 16 YEARS AT HIGH RISK |
| I-605 | ADJUNCT TO MOOD STABILIZERS AND/OR ANTIDEPRESSANTS FOR SCHIZOAFFECTIVE DISORDER |
| I-606 | TREATMENT OF SCHIZOAFFECTIVE DISORDER AS MONOTHERAPY |
| I-607 | INDICATION EXPANDED TO INCLUDE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (WHO GROUP I) IN PATIENTS WITH CLASS II SYMPTOMS |
| I-608 | REDUCE LDL-C LEVELS IN BOYS AND POSTMENARCHAL GIRLS, 10 TO 17 YEARS OF AGE, WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA AS MONOTHERAPY OR IN COMBINATION WITH A STATIN AFTER FAILING AN ADEQUATE TRIAL OF DIET THERAPY |
| I-610 | TREATMENT OF HEAVY MENSTRUAL BLEEDING FOR WOMEN WHO CHOOSE TO USE INTRAUTERINE CONTRACEPTION AS THEIR METHOD OF CONTRACEPTION |
| I-611 | TREATMENT OF HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA IN ADOLESCENT BOYS AND POSTMENARCHAL GIRLS, AGES 10 TO 17 YEARS, WITH A RECOMMENDATION DOSING RANGE OF 5 TO 20 MG ONCE DAILY |
| I-612 | MICARDIS 80 MG FOR REDUCTION OF THE RISK OF MYOCARDIAL INFARCTION, STROKE, OR DEATH FROM CARDIOVASCULAR CAUSES IN PATIENTS 55 YEARS OF AGE OR OLDER AT HIGH RISK OF DEVELOPING MAJOR CARDIOVASCULAR EVENTS WHO ARE UNABLE TO TAKE ACE INHIBITORS |
| I-613 | MILD TO MODERATE ATOPIC DERMATITIS IN PATIENTS 3 MONTHS OF AGE TO LESS THAN 18 YEARS OF AGE |
| I-614 | SHORT TERM TREATMENT OF EROSIIVE ESOPHAGITIS ASSOCIATED WITH GERD IN PEDIATRIC PATIENTS AGES FIVE YEARS AND OLDER |
| I-615 | MAINTENANCE TREATMENT OF BIPOLAR DISORDER AS AN ADJUNCT TO LITHIUM OR VALPROATE |
| I-616 | TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER IN PEDIATRIC PATIENTS AGES 6-17 YEARS OF AGE |
| I-617 | MAINTENANCE OF GENERALIZED ANXIETY DISORDER (GAD) |
| I-618 | ADJUNCTIVE THERAPY IN THE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD) |
| I-619 | INTRAVENOUS CONTRAST ENHANCED COMPUTER TOMOGRAPHY OF THE HEAD AND BODY |
| I-620 | FOR USE IN COMBINATION WITH LETROZOLE FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE METASTATIC BREAST CANCER THAT OVEREXPRESSES THE HER2 RECEPTOR FOR WHOM HORMONAL THERAPY IS INDICATED |
| I-621 | PRIMARY PREVENTION OF CARDIOVASCULAR DISEASE, BASED ON THE RESULTS OF JUSTIFICATION FOR THE USE OF STATINS IN PRIMARY PREVENTION; AN INTERVENTION TRIAL EVALUATING ROSUVASTATIN (JUPITER) |
| I-622 | ADJUNCTIVE THERAPY FOR PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN PATIENTS THIRTEEN YEARS OF AGE AND OLDER |
| I-623 | TREATMENT OF SIGNS AND SYMPTOMS OF ADVANCED IDIOPATHIC PARKINSON'S DISEASE |
| I-624 | MAINTENANCE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER WHOSE DISEASE HAS NOT PROGRESSED AFTER FOUR CYCLES OF PLATINUM-BASED FIRST-LINE CHEMOTHERAPY |
| I-625 | PANCREATIC INSUFFICIENCY DUE TO CHRONIC PANCREATITIS AND PANCREATECTOMY |
| I-626 | RELIEF OF NASAL CONGESTION ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER |
| I-627 | TREATMENT OF NEWLY DIAGNOSED ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH & CML) IN CHRONIC PHASE. |
| I-628 | MAINTENANCE TREATMENT OF SCHIZOPHRENIA IN ADULTS |
| I-629 | ADJUNCTIVE THERAPY WITH EITHER LITHIUM OR VALPROATE FOR THE ACUTE TREATMENT OF MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER |
| I-630 | TREATMENT OF PATIENTS WITH SUBEPENDYMAL GIANT CELL ASTROCYTOMA (SEGA) ASSOCIATED WITH TUBEROUS SCLEROSIS (TS) WHO REQUIRE THERAPEUTIC INTERVENTION BUT |

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

ARE NOT CANDIDATES FOR CURATIVE SURGICAL RESECTION.

- I-631 PREVENTION OF RELAPSE TO OPIOID DEPENDENCE FOLLOWING OPIOID DETOXIFICATION
- I-632 MANAGEMENT OF CHRONIC MUSCULOSKELETAL PAIN
- I-633 MAINTENANCE TREATMENT OF BIPOLAR I DISORDER AS AN ADJUNCT TO LITHIUM OR VALPROATE
- I-634 TREATMENT OF SEVERE HYPERCALCEMIA IN PATIENTS WITH PRIMARY HYPERPARATHYROIDISM WHO ARE UNABLE TO UNDERGO PARATHYROIDECTOMY
- I-635 ADJUNCTIVE TREATMENT WITH LONG-ACTING ORAL PSYCHOSTIMULANTS FOR THE TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)
- I-636 TREATMENT OF EXTERNAL GENITAL AND PERIANAL WARTS/CONDYLOMA ACUMINATA IN PATIENTS 12 YEARS OR OLDER
- I-637 USE IN COMBINATION CHEMOTHERAPY WITH 5-FLUOROURACIL IN THE PALLIATIVE TREATMENT OF PATIENTS WITH ADVANCED METASTATIC COLORECTAL CANCER
- I-638 FOR PATIENTS WITH PROGRESSIVE NEUROENDOCRINE TUMORS OF PANCREATIC ORIGIN (PNET) THAT ARE UNRESECTABLE, LOCALLY ADVANCED, OR METASTATIC.
- I-639 TREATMENT OF PROGRESSIVE, WELL-DIFFERENTIATED PANCREATIC NEUROENDOCRINE TUMORS IN PATIENTS WITH UNRESECTABLE, LOCALLY ADVANCED, OR METASTATIC DISEASE
- I-640 MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS
- I-641 TREATMENT OF THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA (BPH)
- I-642 TREATMENT OF ERECTILE DYSFUNCTION (ED) AND THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA (BPH)
- I-643 REDUCE THE RISK OF STROKE AND SYSTEMIC EMBOLISM IN PATIENTS WITH NONVALVULAR ATRIAL FIBRILLATION.
- I-644 MONOTHERAPY IN PATIENTS 13 YEARS OF AGE AND OLDER WITH PARTIAL SEIZURES WHO ARE RECEIVING THERAPY WITH A SINGLE ANTIEPILEPTIC DRUG (AED)
- I-645 MAINTENANCE TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) IN ADULTS
- I-646 SIGNS AND SYMPTOMS OF ADVANCED PARKINSON'S DISEASE (APD)
- I-647 SIGNS AND SYMPTOMS OF MODERATE TO SEVERE PRIMARY RESTLESS LEGS SYNDROME (RLS)
- I-648 TREATMENT OF HEAVY MENSTRUAL BLEEDING IN WOMEN WITHOUT ORGANIC PATHOLOGY WHO CHOOSE TO USE AN ORAL CONTRACEPTIVE AS THEIR METHOD OF CONTRACEPTION
- I-649 TREATMENT OF PATIENTS WITH ADVANCED SOFT TISSUE SARCOMA (STS) WHO HAVE RECEIVED PRIOR CHEMOTHERAPY
- I-650 TREATMENT OF ADULTS WITH RENAL ANGIOMYOLIPOMA AND TUBEROUS SCLEROSIS COMPLEX (TSC), NOT REQUIRING IMMEDIATE SURGERY
- I-651 MANAGEMENT OF NEUROPATHIC PAIN ASSOCIATED WITH SPINAL CORD INJURY
- I-652 MANAGEMENT OF POSTHERPETIC NEURALGIA
- I-653 TREATMENT OF ENDOGENOUS ANTERIOR UVEITIS
- I-654 MAGNETIC RESONANCE ANGIOGRAPHY (MRA) TO EVALUATE ADULTS WITH KNOWN OR SUSPECTED RENAL OR AORTO-ILIO-FEMORAL OCCLUSIVE VASCULAR DISEASE
- I-655 TREATMENT OF POSTMENOPAUSAL WOMEN WITH ADVANCED HORMONE RECEPTOR-POSITIVE, HER2-NEGATIVE BREAST CANCER (ADVANCED HR+BC) IN COMBINATION WITH EXEMESTANE, AFTER FAILURE OF TREATMENT WITH LETROZOLE OR ANASTROZOLE
- I-656 MANAGEMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY (DPN) IN ADULTS WHEN A CONTINUOUS, AROUND-THE-CLOCK OPIOID ANALGESIC IS NEEDED FOR AN EXTENDED PERIOD OF TIME
- I-657 PLAQUE PSORIASIS OF THE SCALP
- I-658 FIRST-LINE TREATMENT OF LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER, IN COMBINATION WITH CARBOPLATIN, IN PATIENTS WHO ARE NOT CANDIDATES FOR CURATIVE SURGERY OR RADIATION THERAPY
- I-659 PLAQUE PSORIASIS OF THE BODY
- I-660 TREATMENT OF DEEP VEIN THROMBOSIS
- I-661 TREATMENT OF PULMONARY EMBOLISM
- I-662 REDUCTION IN RISK FOR DEEP VEIN THROMBOSIS AND THE REDUCTION IN RISK FOR

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

PULMONARY EMBOLISM

- I-663 IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER
- I-664 TREATMENT OF THROMBOCYTOPENIA IN PATIENTS WITH CHRONIC HEPATITIS C TO ALLOW THE INITIATION AND MAINTENANCE OF INTERFERON-BASED THERAPY
- I-665 TREATMENT OF CHRONIC IRON OVERLOAD IN PATIENTS 10 YRS OF AGE AND OLDER WITH (NTDT)SYNDROMES AND WITH A (LIC) OF AT LEAST 5 MG OF IRON PER GRAM OF LIVER DRY WEIGHT (MG FE/G DW) AND SERUM FERRITIN GREATER THAN 300MCG/L
- I-666 TREATMENT OF PEDIATRIC PATIENTS WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (PH+ALL) IN COMBINATION WITH CHEMOTHERAPY
- I-667 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED, UNRESECTABLE OR METASTATIC GASTROINTESTINAL STROMAL TUMOR (GIST) WHO HAVE BEEN PREVIOUSLY TREATED WITH IMATINIB MESYLATE AND SUNITINIB MALATE
- I-668 PROPHYLAXIS OF ALLOGRAFT REJECTION IN ADULT PATIENTS RECEIVING A LIVER TRANSPLANT
- I-669 SCINTIGRAPHIC ASSESSMENT OF SYMPATHETIC INNERVATION OF THE MYOCARDIUM BY MEASUREMENT OF THE HEART TO MEDIASTINUM (H/M) RATIO OF RADIOACTIVITY UPTAKE IN PATIENTS WITH NYHA CLASS II OR CLASS III HEART FAILURE AND LVEF LESS THAN 35%
- I-670 TREATMENT OF OPIOID-INDUCED CONSTIPATION (OIC) IN ADULTS WITH CHRONIC, NON-CANCER PAIN
- I-671 FIRSTLINE TREATMENT OF PATIENTS WITH METASTATIC NON- SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21(L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- I-672 USE IN PATIENTS WITH MANTLE CELL LYMPHOMA WHOSE DISEASE HAS RELAPSED OR PROGRESSED AFTER TWO PRIOR THERAPIES, ONE OF WHICH INCLUDED BORTEZOMIB
- I-673 TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA/VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) CAUSED BY SUSCEPTIBLE ISOLATES OF S. AUREUS (INCLUDING METHICILLIN-SUSCEPTIBLE AND RESISTANT ISOLATES) WHEN ALTERNATIVE TREATMENTS ARE NOT SUITABLE
- I-674 TREATMENT OF PATIENTS WITH DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I DISORDER (BIPOLAR DEPRESSION) AS MONOTHERAPY AND AS ADJUNCTIVE THERAPY WITH LITHIUM OR VALPROATE
- I-675 MAINTENANCE TREATMENT OF MAJOR DEPRESSIVE DISORDER
- I-676 FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC ADENOCARCINOMA OF THE PANCREAS, IN COMBINATION WITH GEMCITABINE
- I-677 TREATMENT OF PATIENTS WITH LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, DIFFERENTIATED THYROID CARCINOMA (DTC) THAT IS REFRACTORY TO RADIOACTIVE IODINE TREATMENT
- I-678 TRAMETINIB, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- I-679 RISK REDUCTION OF REBLEEDING OF GASTRIC OR DUODENAL ULCERS FOLLOWING THERAPEUTIC ENDOSCOPY IN ADULTS
- I-680 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- I-681 PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT) WHICH MAY LEAD TO PULMONARY EMBOLISM (PE), IN ADULT PATIENTS WHO HAVE UNDERGONE HIP OR KNEE REPLACEMENT
- I-682 TREATMENT OF DEEP VEIN THROMBOSIS (DVT) AND PULMONARY EMBOLISM (PE) IN PATIENTS WHO HAVE BEEN TREATED WITH A PARENTERAL ANTICOAGULANT FOR 5-10 DAYS
- I-683 TO REDUCE THE RISK OF RECURRENCE OF DVT AND PE IN PATIENTS WHO HAVE BEEN PREVIOUSLY TREATED
- I-684 PREVENTION OF ACUTE NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF EMETOGENIC CANCER CHEMOTHERAPY, INCLUDING HIGHLY EMETOGENIC CANCER CHEMOTHERAPY IN PEDIATRIC PATIENTS AGED 1 MONTH TO LESS THAN 17 YEARS
- I-685 EXPANDED INDICATION OF RASAGILINE AS AN ADD-ON THERAPY TO STABLE DOSES OF DOPAMINE AGONISTS IN THE TREATMENT OF EARLY PARKINSON'S DISEASE
- I-686 INDICATED FOR THE TREATMENT OF DIABETIC MACULAR EDEMA IN PATIENTS WHO ARE PSEUDOPHAKIC OR ARE PHAKIC AND SCHEDULED FOR CATARACT SURGERY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

| | |
|-------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| I-687 | GUIDING SENTINEL LYMPH NODE BIOPSY, USING A HAND-HELD GAMMA COUNTER IN PATIENTS WITH CLINICALLY NODE NEGATIVE SQUAMOUS CELL CARCINOMA OF THE ORAL CAVITY |
| I-688 | GADAVIST IS INDICATED WITH MRI TO DETECT THE PRESENCE AND EXTENT OF MALIGNANT BREAST DISEASE |
| I-689 | TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WITH 17P DELETION |
| I-690 | INDICATED FOR THE TREATMENT OF DEEP VEIN THROMBOSIS (DVT) |
| I-691 | INDICATED TO REDUCE THE RISK OF RECURRENT DEEP VEIN THROMBOSIS (DVT) AND PULMONARY EMBOLISM (PE) FOLLOWING INITIAL THERAPY |
| I-692 | INDICATED FOR MANAGEMENT OF OSTEOARTHRITIS PAIN. |
| I-693 | TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (CRPC) |
| I-694 | TREATMENT OF PATIENTS WITH MODERATE TO SEVERE PLAQUE PSORIASIS WHO ARE CANDIDATES FOR PHOTOTHERAPY OR SYSTEMIC THERAPY |
| I-695 | REVISED INDICATION FOR BORTEZOMIB IN THE TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA |
| I-696 | USE AS MONOTHERAPY IN THE TREATMENT OF PARTIAL-ONSET SEIZURES IN PATIENTS WITH EPILEPSY AGE 17 YEARS AND OLDER |
| I-697 | FOR USE IN COMBINATION WITH SOFOSBUVIR FOR THE TREATMENT OF PATIENTS WITH CHRONIC HEPATITIS C VIRUS GENOTYPE 1 INFECTION |
| I-698 | SCHIZOAFFECTIVE DISORDER AS MONOTHERAPY AND AS AN ADJUNCT TO MOOD STABILIZERS OR ANTIDEPRESSANTS |
| I-699 | FOR TREATMENT OF PATIENTS WITH POLYCYTHEMIA VERA WHO HAVE HAD AN INADEQUATE RESPONSE TO OR ARE INTOLERANT OF HYDROXYUREA |
| I-700 | TREATMENT OF PEDIATRIC PATIENTS WITH TOURETTE'S DISORDER (6-18 YEARS) |
| I-701 | FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE, WELL-OR MODERATELY-DIFFERENTIATED, LOCALLY ADVANCED OR METASTATIC GASTROENTEROPANCREATIC NEUROENDOCRINE TUMORS (GEP-NETS) TO IMPROVE PROGRESSION FREE SURVIVAL |
| I-702 | FOR THE TREATMENT OF PATIENTS WITH WALDENSTROM MACROGLOBULINEMIA |
| I-703 | MODERATE TO SEVERE BINGE EATING DISORDER (BED) |
| I-704 | EXPANDED INDICATION TO INCLUDE PATIENTS WHO ARE VIROLOGICALLY-SUPPRESSED (HIV-1 RNA <50 COPIES/ML) ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS WITH NO HISTORY OF TREATMENT FAILURE IN ORDER TO REPLACE THEIR CURRENT REGIMEN |
| I-705 | TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 6 YEARS AND OLDER WHO HAVE AN R117H MUTATION IN THE CFTR GENE |
| I-706 | EXPANDED INDICATION FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA |
| I-707 | POMALYST, IN COMBINATION WITH DEXAMETHASONE, IS INDICATED FOR PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST 2 PRIOR THERAPIES AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY |
| I-708 | DAILY TREATMENT OF ASTHMA IN PATIENTS AGED 18 YEARS AND OLDER |
| I-709 | TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS |
| I-710 | ADJUNCTIVE THERAPY FOR THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC (PG TC) SEIZURES IN PATIENTS WITH EPILEPSY 12 YEARS OF AGE OR OLDER. |
| I-711 | INCLUSION OF PEDIATRIC PATIENTS AGES 6 YRS AND OLDER FOR THE TREATMENT OF THROMBOCYTOPENIA IN PATIENTS WITH CHRONIC ITP WHO HAVE HAD AN INSUFFICIENT RESPONSE TO CORTICOSTEROIDS, IMMUNOGLOBULINS, OR SPLENECTOMY. |
| I-712 | EXPANDED INDICATION FOR USE IN COMBINATION WITH LENALIDOMIDE AND DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE PRIOR LINES OF THERAPY |
| I-713 | REVISIONS TO THE LABELING TO PERMIT THE USE OF ZUBSOLV AS INITIAL ("INDUCTION") TREATMENT OF OPIOID DEPENDENCE |
| I-714 | EXTENDS THE 2011 APPROVAL OF BRILINTA FOR USE BEGINNING WITH ACS TO USE BEGINNING MORE REMOTE FROM MYOCARDIAL INFARCTION |
| I-715 | FOR THE ADDITION OF THE INDICATION FOR MONOTHERAPY TREATMENT IN PARTIAL-ONSET SEIZURES IN ADULTS. |
| I-716 | REVISED INDICATION TO INCLUDE LANGUAGE ABOUT THE BENEFITS OF USING LETAIRIS IN |

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

COMBINATION WITH TADALAFIL TO REDUCE THE RISK OF DISEASE PROGRESSION AND HOSPITALIZATION FOR WORSENING PAH AND TO IMPROVE EXERCISE ABILITY, BASED ON THE AMBITION STUDY

- I-717 EXPANDED INDICATION TO INCLUDE THE TREATMENT OF CHRONIC HEPATITIS C GENOTYPE 4
- I-718 EXPANDED INDICATION TO INCLUDE SUBJECTS INFECTED WITH CHRONIC HEPATITIS C, GENOTYPE 6 VIRUS INFECTION BASED UPON THE RESULTS OF THE ELECTRON- 2 STUDY
- I-719 EXPANDED INDICATION TO INCLUDE THE TREATMENT OF SUBJECTS WITH GENOTYPE 5 CHRONIC HEPATITIS C VIRUS INFECTION BASED ON THE RESULTS FROM STUDY GS-US-337-119.
- I-720 EXPANDED INDICATION TO INCLUDE TREATMENT OF GENOTYPE 4, CHRONIC HEPATITIS C VIRUS INFECTION BASED UPON THE RESULTS FROM STUDIES ION-4 AND GS-US-337-119.
- I-721 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA WHO HAVE RECEIVED A PRIOR ANTHRACYCLINE-CONTAINING REGIMEN.
- I-722 REVISED INDICATION FOR USE IN COMBINATION WITH DEXAMETHASONE OR WITH LENALIDOMIDE PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY.
- I-723 AS A SINGLE AGENT FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE OR MORE LINES OF THERAPY
- I-724 TREATMENT OF ADULT PATIENTS WITH PROGRESSIVE, WELL DIFFERENTIATED, NON-FUNCTIONAL NEUROENDOCRINE TUMORS (NET) OF GI OR LUNG ORIGIN WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC DISEASE
- I-725 TREATMENT OF HORMONE RECEPTOR (HR)-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN COMBINATION THERAPY WITH PALBOCICLIB AND FULVESTRANT IN WOMEN WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY.
- I-726 EXPANSION OF THE PATIENT POPULATION TO INCLUDE PATIENTS WITH RECURRENCE OF HEPATITIS C VIRUS (HCV) GENOTYPE 1 OR 3 AFTER LIVER TRANSPLANTATION
- I-727 EXPANSION OF THE INDICATION TO INCLUDE TREATMENT OF SUBJECTS WITH GENOTYPE-1 CHRONIC HEPATITIS C VIRUS INFECTION, INCLUDING SUBJECTS WHO ARE CO-INFECTED WITH THE HUMAN IMMUNODEFICIENCY VIRUS (HIV-1) BASED ON THE RESULTS FROM THE ALLY-2 CLINICAL TRIAL
- I-728 EXPANDED INDICATION FOR USE IN ULTRASONOGRAPHY OF THE LIVER FOR CHARACTERIZATION OF FOCAL LIVER LESIONS IN ADULT AND PEDIATRIC PATIENTS
- I-729 PROVIDES FOR THE FRONTLINE INDICATION FOR THE TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA
- I-730 NEW INDICATION FOR THE TREATMENT OF PATIENTS WITH METASTATIC, SQUAMOUS, NON-SMALL CELL LUNG CANCER PROGRESSING AFTER PLATINUM-BASED CHEMOTHERAPY
- I-731 FOR USE IN MAGNETIC RESONANCE ANGIOGRAPHY IN ADULT AND PEDIATRIC PATIENTS (INCLUDING TERM NEONATES) TO EVALUATE KNOWN OR SUSPECTED SUPRA-AORTIC OR RENAL ARTERY DISEASE
- I-732 TREATMENT OF PEDIATRIC PATIENTS 7 TO 17 YEARS OF AGE WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA TO REDUCE LDL-C, TOTAL C, NONHDL-C AND APOB AS AN ADJUNCT TO DIET, EITHER ALONE OR WITH OTHER LIPID-LOWERING TREATMENTS
- I-733 USE OF CANAGLIFLOZIN FOR INITIAL THERAPY IN COMBINATION WITH METFORMIN
- I-734 EXPANDED INDICATION FOR THE USE OF LENVIMA IN COMBINATION WITH EVEROLIMUS FOR THE TREATMENT OF PATIENTS WITH ADVANCED RCC FOLLOWING ONE PRIOR ANTI-ANGIOGENIC THERAPY.
- I-735 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHEN TREATMENT WITH BOTH CANAGLIFLOZIN AND METFORMIN IS APPROPRIATE
- I-736 REVISED INDICATION TO INCLUDE THE TREATMENT OF PATIENTS WITH SMALL LYMPHOCYTIC LEUKEMIA (SLL)
- I-737 REVISED INDICATION TO INCLUDE THE TREATMENT OF PATIENTS WITH SMALL LYMPHOCYTIC LEUKEMIA (SLL) WITH 17P DELETION
- I-738 REVISIONS TO THE INDICATIONS AND USAGE SECTION WITH RESPECT TO COMPLICATED INTRA-ABDOMINAL INFECTIONS
- I-739 TO REDUCE THE RISK OF CARDIOVASCULAR DEATH IN ADULT PATIENTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE
- I-740 EXPANDED INDICATION FOR THE TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 6 YEARS AND OLDER TO INCLUDE THE G1244E, G1349D, G178R, G551S, S1251N, S1255P, S549N, OR S549R MUTATION IN THE CFTR GENE

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

| | |
|-------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| I-741 | TREATMENT OF PATIENTS WITH MARGINAL ZONE LYMPHOMA (MZL) WHO REQUIRE SYSTEMIC THERAPY AND HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED THERAPY |
| I-742 | TREATMENT OF NODAL MARGINAL ZONE LYMPHOMA |
| I-743 | INFORMATION ADDED TO THE LABELING FOR THE ADDITION OF THE TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 4 (GT4) INFECTED PATIENTS WITH COMPENSATED CIRRHOSIS BASED ON RESULTS FROM STUDY M11-665 |
| I-744 | TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA (HCC) WHO HAVE BEEN PREVIOUSLY TREATED WITH SORAFENIB |
| I-745 | MEKINIST, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST. |
| I-746 | NEW INDICATION OF MAINTENANCE MONOTHERAPY TREATMENT OF BIPOLAR I DISORDER IN ADULTS |
| I-747 | FOR REDUCING THE RISK OF GRAFT REJECTION WHEN USED WITH HIGH-DOSE BUSULFAN AND CYCLOPHOSPHAMIDE AS A PREPARATIVE REGIMEN FOR ALLOGENEIC HEMATOPOIETIC PROGENITOR (STEM) CELL TRANSPLANTATION FOR PEDIATRIC PATIENTS WITH CLASS 3 BETA-THALASSEMIA |
| I-748 | TO REDUCE THE ACUTE COMPLICATIONS OF SICKLE CELL DISEASE IN ADULT AND PEDIATRIC PATIENTS FIVE YEARS OF AGE AND OLDER |
| I-749 | MONOTHERAPY FOR THE TREATMENT OF HORMONE RECEPTOR (HR) POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE, ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN NOT PREVIOUSLY TREATED WITH ENDOCRINE THERAPY |
| I-750 | REDUCE THE RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE |
| I-751 | TREATMENT OF TARDIVE DYSKINESIA |
| I-752 | CORONARY COMPUTED TOMOGRAPHY ANGIOGRAPHY (CCTA) TO ASSIST DIAGNOSTIC EVALUATION OF PATIENTS WITH SUSPECTED CORONARY ARTERY DISEASE |
| I-753 | TREATMENT OF ADULT PATIENTS WITH CHRONIC GRAFT VERSUS HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY |
| I-754 | TO REDUCE THE FREQUENCY OF SHORT-ACTING SOMATOSTATIN ANALOG RESCUE THERAPY WHEN USED FOR THE TREATMENT OF ADULTS WITH CARCINOID SYNDROME |
| I-755 | ADJUVANT TREATMENT OF ADULT PATIENTS AT HIGH RISK OF RECURRENT RENAL CELL CARCINOMA (RCC) FOLLOWING NEPHRECTOMY |
| I-756 | EXPANDED THE APPROVED INDICATION BY REMOVING THE RESTRICTION FOR USE ONLY IN PATIENTS WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB |
| I-757 | TREATMENT OF PATIENTS WITH ERDHEIM-CHESTER DISEASE WITH BRAF V600 MUTATION |
| I-758 | FOR USE WITH RILPIVIRINE AS A COMPLETE REGIMEN TO REPLACE THE CURRENT ARV REGIMEN IN VIROLOGICALLY SUPPRESSED PATIENTS ON A STABLE ARV REGIMEN FOR AT LEAST 6 MONTHS WITH NO HISTORY OF TX FAILURE OR KNOWN SUBSTITUTIONS ASSOC. WITH RESISTANCE TO EITHER ARV |
| I-759 | TREATMENT OF ADULT PATIENTS WITH NEWLY-DIAGNOSED CHRONIC PHASE (CP) PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOGENOUS LEUKEMIA (PH+CML) |
| I-760 | FOR THE TREATMENT OF PATIENTS WITH ADVANCED RENAL CELL CARCINOMA |
| I-761 | TREATMENT OF ADULT PATIENTS WITH ACTIVE PSORIATIC ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO METHOTREXATE OR OTHER NON-BIOLOGIC DISEASE-MODIFYING ANTIRHEUMATIC DRUGS |
| I-762 | TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATED, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2-NEGATIVE METASTATIC BREAST CANCER WHO HAVE BEEN TREATED WITH CHEMOTHERAPY IN THE NEOADJUVANT, ADJUVANT OR METASTATIC SETTING |
| I-763 | TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WHOSE TUMORS HAVE NON-RESISTANT EPIDERMAL GROWTH FACTOR RECEPTOR MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST |
| I-764 | TREATMENT IN ADULT PATIENTS FOR IRRITABLE BOWEL SYNDROME WITH CONSTIPATION (IBS-C) |
| I-765 | ABIRATERONE ACETATE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC HIGH-RISK CASTRATION-SENSITIVE PROSTATE CANCER |
| I-766 | TREATMENT OF MINIMALLY TO MODERATELY THICK ACTINIC KERATOSIS OF THE UPPER |

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

- EXTREMITIES IN CONJUNCTION WITH A BLUE LIGHT PHOTODYNAMIC THERAPY ILLUMINATOR
- I-767 TREATMENT OF IRON DEFICIENCY ANEMIA IN ADULT PATIENTS WHO HAVE INTOLERANCE TO ORAL IRON OR HAVE HAD UNSATISFACTORY RESPONSE TO ORAL IRON
- I-768 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- I-769 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- I-770 TREATMENT OF ACUTE OTITIS EXTERNA IN PATIENTS 6 MONTHS OF AGE AND OLDER DUE TO PSEUDOMONAS AERUGINOSA AND STAPHYLOCOCCUS AUREUS
- I-771 REVISION OF THE INDICATION SECTION OF THE PACKAGE INSERT REGARDING AN INTERSCALENE BRACHIAL PLEXUS NERVE BLOCK TO PRODUCE POSTSURGICAL REGIONAL ANALGESIA
- I-772 FOR THE MAINTENANCE TREATMENT OF ADULT PATIENTS WITH RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- I-773 FOR THE ADJUNCTIVE TREATMENT OF ADULT AND PEDIATRIC PATIENTS AGE 2 YEARS AND OLDER WITH TUBEROUS SCLEROSIS COMPLEX (TSC)-ASSOCIATED PARTIAL-ONSET SEIZURES
- I-774 TO ALLOW FOR FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS, AS DETECTED BY AN FDA APPROVED TEST
- I-775 REVISED INDICATION FOR FIXED-DOSE COMBINATION OF FLUTICASONE FUROATE, UMECLIDINIUM, AND VILANTEROL TO TREAT AIRFLOW OBSTRUCTION IN CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) AND TO REDUCE COPD EXACERBATIONS IN PTS WITH HISTORY OF EXACERBATIONS
- I-776 FIRSTLINE MAINTENANCE TX IN PTS W/ DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE, SOMATIC BRCA-MUTATED ADVANCED EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CA WHO ARE IN COMPLETE OR PARTIAL RESPONSE TO FIRSTLINE PLATINUM-BASED CHEMOTHERAPY
- I-777 CO-ADMINISTRATION THERAPY OF MIRABEGRON WITH SOLIFENACIN SUCCINATE FOR TREATMENT OF OVERACTIVE BLADDER WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY
- I-778 DABRAFENIB IN COMBINATION WITH TRAMETINIB FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- I-779 USE OF TOLVAPTAN TO SLOW KIDNEY FUNCTION DECLINE IN ADULTS AT RISK OF RAPIDLY PROGRESSING AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE (ADPKD)
- I-780 TREATMENT OF ADULT PATIENTS WITH MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS (UC)
- I-781 DABRAFENIB IN COMBINATION WITH TRAMETINIB FOR THE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ANAPLASTIC THYROID CANCER (ATC) WITH BRAF V600E MUTATION AND WITH NO SATISFACTORY LOCOREGIONAL TREATMENT OPTIONS
- I-782 REVISIONS TO INDICATION FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL), WITH OR WITHOUT 17P DELETION, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- I-783 EXPANDED INDICATION TO INCLUDE RIBOCICLIB WITH AN AROMATASE INHIBITOR IN PRE/PERIMENOPAUSAL WOMEN WITH HORMONE RECEPTOR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER, AS INITIAL ENDOCRINE-BASED THERAPY
- I-784 RIBOCICLIB WITH FULVESTRANT FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER, AS INITIAL ENDOCRINE BASED THERAPY OR FOLLOWING DISEASE PROGRESSION ON ENDOCRINE THERAPY
- I-785 TREATMENT OF PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- I-786 TREATMENT OF PATIENTS WITH NON-METASTATIC CASTRATION-RESISTANT PROSTATE CANCER
- I-787 FIRST-LINE TREATMENT OF PATIENTS WITH UNRESECTABLE HEPATOCELLULAR CARCINOMA (HCC)
- I-788 NEW INDICATION FOR CANAGLIFLOZIN TO REDUCE THE RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS (CARDIOVASCULAR DEATH, NONFATAL MYOCARDIAL INFARCTION AND

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY INDICATION**

NONFATAL STROKE) IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE (CVD)

- I-789 VENETOCLAX IN COMBO WITH AZACITIDINE OR DECITABINE OR LOW-DOSE CYTARABINE FOR THE TX OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA IN ADULTS WHO ARE AGE 75 YEARS OR OLDER, OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- I-791 TREATMENT OF PEDIATRIC PATIENTS ONE YEAR OF AGE AND OLDER WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE (PH+) ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN COMBINATION WITH CHEMOTHERAPY

EXCLUSIVITY MISCELLANEOUS

- M-1 INFORMATION REGARDING SUPERIORITY CLAIM OVER RANITIDINE FOR DAY AND NIGHT HEARTBURN ADDED TO CLINICAL STUDIES SECTION
- M-2 APPROVAL FOR ADDITION TO CLINICAL PHARMACOLOGY SECTION OF THE LABEL REGARDING (1) IMPROVEMENT IN BONE MINERAL DENSITY IN CHILDHOOD-ONSET ADULT GROWTH HORMONE DEFICIENT PATIENTS AND (2) INCREASES IN SERUM ALKALINE PHOSPHATASE
- M-3 ADDITION OF EFFICACY AND SAFETY INFORMATION IN WHICH FOSAMAX WAS USED CONCOMITANTLY WITH ESTROGEN ALONE OR WITH ESTROGEN PLUS PROGESTIN
- M-4 CHANGES TO PEDIATRIC USE SECTION TO PROVIDE INFORMATION REGARDING SAFETY AND EFFICACY IN PEDIATRIC PATIENTS AS YOUNG AS 2 YEARS OLD
- M-5 INFORMATION REGARDING EFFECTS IN PATIENTS WITH ASTHMA ON CONCOMITANT INHALED CORTICOSTEROIDS IN CLINICAL PHARMACOLOGY SECTION
- M-6 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES DONE WITH GLUCOPHAGE/GLYBURIDE COMBINATION ADDED TO CLINICAL PHARMACOLOGY AND DOSING AND ADMINISTRATION
- M-7 CLINICAL PHARMACOLOGY IN PEDIATRIC PATIENTS; DOSAGE AND ADMINISTRATION INFORMATION
- M-8 ADDITIONAL INFORMATION FOR THE USE OF SONATA CAPSULES FOR UP TO 5 WEEKS (35 NIGHTS) OF TREATMENT IN A CONTROLLED TRIAL SETTING
- M-9 ADDITION TO THE CLINICAL STUDIES SECTION OF THE LABELING OF TEXT AND TWO TABLES CONTAINING INFORMATION FOR THE PRESCRIBING PHYSICIAN ON BLOOD PRESSURE, HEART RATE, AND HEART RATE VARIABILITY
- M-10 INFORMATION REGARDING MAINTENANCE OF AN ANTIDEPRESSANT EFFECT UP TO 1 YEAR OF DOSING
- M-11 USE FOR LONG-TERM TREATMENT OF POSTTRAUMATIC STRESS DISORDER
- M-12 NEW LANGUAGE FOR PEDIATRIC USE
- M-13 INFORMATION FROM PEDIATRIC STUDIES ADDED TO CLINICAL PHARMACOLOGY, PRECAUTIONS, AND DOSAGE AND ADMINISTRATION
- M-14 ADDITIONAL CLINICAL TRIAL INFORMATION ADDED TO PEDIATRIC USE SUBSECTION
- M-15 LONGER TERM EFFICACY INFORMATION FOR RISPERIDONE IN THE TREATMENT OF SCHIZOPHRENIA
- M-16 CHANGE IN WORDING OF THE PEDIATRIC SECTION OF THE PACKAGE INSERT
- M-17 INFORMATION REGARDING USE OF ULTANE IN PEDIATRIC PATIENTS WITH CONGENITAL HEART DISEASE
- M-18 INFORMATION DENOTING THE EFFICACY OF REMERON IN MAINTAINING A RESPONSE IN PATIENTS WITH MAJOR DEPRESSIVE DISORDER (MDD)
- M-19 INFORMATION REGARDING USE IN PEDIATRIC PATIENTS TWO YEARS OF AGE AND OLDER
- M-20 LABELING REVISIONS RELATED TO MCCUNE ALBRIGHT SYNDROME
- M-21 COMPARISON DATA ON THE ANTIHYPERTENSIVE EFFECTS OF ATACAND AND COZAAR
- M-22 CHANGE IN TIME TO ONSET OF ACTION
- M-23 INFORMATION REGARDING ELIMINATION ADDED TO CLINICAL PHARMACOLOGY, STUDY RESULTS IN PATIENTS WITH HEPATIC AND RENAL IMPAIRMENT
- M-24 INFORMATION ON RESULTS OF A LONG TERM LONGITUDINAL GROWTH STUDY AND PEDIATRIC SAFETY INFORMATION
- M-25 ADDITIONAL SAFETY AND PHARMACOKINETICS INFORMATION IN CHILDREN 6 MONTHS TO LESS THAN 6 YEARS OF AGE ADDED TO PACKAGE INSERT

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- M-26 INCORPORATION OF INFORMATION CONTAINED IN THE PEG-INTRON PACKAGE INSERT INTO THE REBETOL PACKAGE INSERT AND MEDGUIDE-PEG-INTRON WAS APPROVED FOR USE IN COMBINATION WITH REBETOL FOR TREATMENT OF CHRONIC HEPATITIS C VIRUS INFECTION ON 8/7/01
- M-27 INFORMATION DESCRIBING ASPIRIN ENDOSCOPY STUDY AND THE MAXIMUM RECOMMENDED DOSE FOR PATIENTS WITH MODERATE HEPATIC INSUFFICIENCY
- M-28 INFORMATION FROM A STUDY IN PEDIATRIC PATIENTS IN ASSOCIATION WITH A NEUROLOGICAL CONDITION
- M-29 LABELING CHANGES TO PROVIDE INFORMATION IN THE MANAGEMENT OF OBESITY IN ADOLESCENTS AGED 12 TO 16 YEARS
- M-30 CHANGES TO CLINICAL PHARMACOLOGY, PRECAUTIONS, AND DOSAGE AND ADMINISTRATION SECTIONS OF LABELING CONCERNING USE OF LOTENSIN IN PEDIATRIC PATIENTS WITH HYPERTENSION
- M-31 INFORMATION FOR USE IN PEDIATRIC PATIENTS WITH CHRONIC KIDNEY DISEASE STAGE 5 (END-STAGE RENAL DISEASE)
- M-32 ADDITIONAL LANGUAGE TO CLINICAL PHARMACOLOGY AND CLINICAL STUDIES
- M-33 INFORMATION FOR USE OF ADVAIR DISKUS 100/50 IN CHILDREN 4 TO 11 YEARS OF AGE WITH ASTHMA
- M-34 EXPANDED INFORMATION TO PEDIATRIC USE SUBSECTION OF LABELING IN RESPONSE TO PEDIATRIC WRITTEN REQUEST
- M-35 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES DONE WITH ACTOS IN COMBINATION WITH METFORMIN, A SULFONYLUREA, OR INSULIN ADDED TO CLINICAL PHARMACOLOGY
- M-36 ADDITION OF INFORMATION TO CLINICAL STUDIES REGARDING PREVENTION OF CARDIOVASCULAR DISEASE
- M-37 INFORMATION ADDED TO THE LABELING THAT DETAILS INFORMATION RELATIVE TO STUDIES DONE IN PEDIATRIC POPULATIONS IN THE CLINICAL PHARMACOLOGY AND PEDIATRIC USE SUBSECTIONS
- M-38 SAFETY AND IOP-LOWERING EFFECTS OF TRUSOPT HAVE BEEN DEMONSTRATED IN PEDIATRIC PATIENTS IN A 3 MONTH, MULTI-CENTER DOUBLE MASKED ACTIVE-TREATMENT-CONTROLLED TRIAL
- M-39 FOR LABELING CHANGES BASED ON RESULTS OF THE SPD422-202 CLINICAL STUDY REPORT (CSR) SUBMITTED IN RESPONSE TO THE WRITTEN REQUEST
- M-40 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES PERFORMED IN PEDIATRIC PATIENTS WITH LEUKEMIA ADDED TO PRECAUTIONS
- M-41 REVISION TO THE PEDIATRIC USE PRECAUTIONS OF THE PRESCRIBING INFORMATION TO INCORPORATE THE RESULTS FROM THE CAPPS-169 STUDY ENTITLED "THE EFFECT OF ORTHO TRICYCLEN ON BONE MINERAL DENSITY IN PEDIATRIC SUBJECTS WITH ANOREXIA NERVOSA"
- M-42 ADDITION OF A GERIATRIC USE SUBSECTION TO THE PRECAUTIONS SECTION OF THE PACKAGE INSERT AND GERIATRIC DOSING INFORMATION
- M-43 INCLUSION OF RESULTS OF STUDY "PLACEBO-CONTROLLED STUDY TO EVALUATE SAFETY AND PILOT EFFICACY OF ILOPROST AS ADD ON THERAPY WITH BOSENTAN IN SUBJECTS WITH PULMONARY ARTERIAL HYPERTENSION"
- M-44 CLINICAL INFORMATION ADDED TO THE PEDIATRIC USE SUBSECTION OF PRECAUTIONS REGARDING THE USE OF NOVOLOG IN ADOLESCENTS WITH TYPE I DIABETES AGE 6 TO 18
- M-45 INFORMATION ADDED TO CLINICAL TRIALS SECTION OF LABELING, "EFFECTS OF HUMATROPE TREATMENT IN ADULTS WITH GROWTH HORMONE DEFICIENCY"
- M-46 PROVISION OF RESULTS OF STUDY AND PROPOSED REVISIONS TO PACKAGE INSERT SEE SECTION ON CARDIAC ELECTROPHYSIOLOGY
- M-47 PROVIDES FOR USE OF ANTARA WITHOUT REGARD TO MEALS
- M-48 CHANGES TO THE LABELING DESCRIBING THE RESULTS OF A STUDY OF THE USE OF NOVOLOG MIX 70/30 WITH ORAL ANTIDIABETIC AGENTS IN PATIENTS WITH TYPE 2 DIABETES
- M-49 CLINICAL DATA ADDED TO THE CLINICAL PHARMACOLOGY SECTION REGARDING EFFECT OF SINGULAIR ON GROWTH RATES IN PREPUBERTAL CHILDREN
- M-50 NEW INFO TO THE CLINICAL STUDIES, ADULT GROWTH HORMONE DEFICIENCY (GHD) SUBSECTION OF THE NUTROPIN AQ PACKAGE INSERT DESCRIBING THE EFFECTS OF SOMATROPIN ON VISCERAL ADIPOSE TISSUE IN THE ADULT GROWTH HORMONE DEFICIENT PATIENT POPULATION
- M-51 INFORMATION ADDED TO LABELING REGARDING OSTEOGENESIS IMPERFECTA STUDY

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- M-52 INFORMATION ADDED TO THE CLINICAL PHARMACOLOGY/CLINICAL STUDIES SECTION REGARDING THE USE OF RISEDRONATE ADMINISTERED ONCE A WEEK IN THE PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- M-53 FOR LABELING CHANGES TO THE QUALITY OF LIFE (QOL) STATEMENT IN THE APPROVED PACKAGE INSERT
- M-54 INFORMATION FROM PEDIATRIC STUDIES ADDED TO LABEL
- M-55 INFORMATION ON RESULTS OF A STUDY OF THE USE OF SANDOSTATIN LAR DEPOT IN PEDIATRIC PATIENTS WITH HYPOTHALAMIC OBESITY.
- M-56 INFORMATION ADDED TO CLINICAL TRIAL SECTION WITH INFORMATION ON "GEMINI" TRIAL
- M-57 CLINICAL DATA ADDED TO THE CLINICAL PHARMACOLOGY SECTION REGARDING THE PHARMACOKINETICS OF EZETIMIBE IN ASIAN SUBJECTS
- M-58 CHANGES TO THE CLINICAL STUDIES, PRIMARY HYPERCHOLESTEROLEMIA, VYTORIN SUBSECTION OF THE PACKAGE INSERT TO ADD EFFICACY DATA FOR THE EZETIMIBE/SIMVASTATIN COMBINATION PRODUCT AND FOR AN ATORVASTATIN PRODUCT ON LDL-C AND OTHER LIPID PRMTRS
- M-59 RESULTS OF THE T20-310 STUDY WHICH EVALUATED THE PHARMACOKINETICS, SAFETY, AND ANTIVIRAL ACTIVITY OF FUZEON IN TREATMENT EXPERIENCED PEDIATRIC SUBJECTS AND ADOLESCENTS WAS ADDED TO THE PEDIATRIC SUBSECTION OF PRECAUTIONS
- M-60 CHANGES TO CLINICAL STUDIES, PRIMARY HYPERCHOLESTEROLEMIA, TO ADD EFFICACY DATA FOR THE EZETIMIBE/SIMVASTATIN COMBINATION PRODUCT AND FOR A ROSUVASTATIN PRODUCT ON LDL-C AND OTHER LIPID PARAMETERS IN PATIENTS WTH HYPERCHOLESTEROLEMIA
- M-61 REVISIONS TO LABELING BASED ON DATA SUBMITTED IN RESPONSE TO PEDIATRIC WRITTEN REQUEST
- M-62 CLINICAL INFORMATION FROM ONE CLINICAL STUDY INVESTIGATING THE USE OF AVANDAMET PLUS INSULIN IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHO HAVE NOT ACHIEVED ADEQUATE GLYCEMIC CONTROL WITH PREVIOUS ANTI-DIABETIC THERAPIES
- M-63 DETAILED INFORMATION ON AN INCONCLUSIVE PEDIATRIC STUDY
- M-64 CHANGES TO CLINICAL PHARMACOLOGY DETAILING STUDY RESULTS
- M-65 ADDITION OF INFORMATION TO LABEL TO INCLUDE INFORMATION REGARDING USE IN PATIENTS WITH HIV-ASSOCIATED ADIPOSE REDISTRIBUTION SYNDROME (HARS)
- M-66 USE IN SPECIFIC POPULATIONS - PATIENTS WITH CONCOMITANT ILLNESS SUBSECTION OF THE LABELING REGARDING USE OF STRATTERA IN PATIENTS WITH ADHD WHO HAVE COMORBID TIC DISORDER
- M-67 INDICATION EXPANDED TO INCLUDE PATIENTS ON PERITONEAL DIALYSIS
- M-68 DESCRIPTION OF RESULTS OF STUDY OF INITIAL THERAPY IN COMBINATION WITH METFORMIN WHEN DIET AND EXERCISE DO NOT PROVIDE GLYCEMIC CONTROL
- M-69 RESULTS OF STUDY OF COMBINATION THERAPY AND NON-INFERIORITY STUDY
- M-70 PROVISION OF INFORMATION OF THE RESULTS OF A PHASE 2 RANDOMIZED TRIAL OF SPRYCEL 70MG TWICE DAILY OR IMATINIB 800MG DAILY
- M-71 REVISIONS TO PROVIDE FOR RESULTS OF MAINTENANCE DATA IN ADULT PATIENTS WITH MAJOR DEPRESSIVE DISORDER
- M-72 INFORMATION ABOUT USE OF INSPIRA (EPLERENONE) FOR HYPERTENSION IN PEDIATRIC PATIENTS
- M-73 NEW INFORMATION ADDED REGARDING THE TUMOR SHRINKING POTENTIAL OF SANDOSTATIN LAR DEPOT INJECTION ON GH - SECRETING PITUITARY ADENOMAS
- M-74 REVISIONS TO CLINICAL STUDIES - CHILDREN AND ADOLESCENTS BASED ON CLINICAL TRIAL DATA TO SUPPORT A DURATION OF ACTION CLAIM UP TO 12 HOURS
- M-75 PROVISION FOR USE OF ARGATROBAN IN CERTAIN PEDIATRIC PATIENTS WITH HEPARIN-INDUCED THROMBOCYTOPENIA (HIT) OR HEPARIN-INDUCED THROMBOCYTOPENIA WITH THROMBOSIS (HITTS)
- M-76 REMOVAL OF SCREEN REQUIREMENT IN PTS WITH G6PD DEFICIENCY PRIOR TO INITIATING ACZONE TREATMENT; REMOVAL OF BLOOD COUNT & RETICULOCYTE MONITORING DURING TREATMENT IN G6PD DEFICIENT PTS AND IN PATIENTS WITH HISTORY OF ANEMIA
- M-77 USE IN COMBINATION WITH THE NEW AKTILITE CL128 LAMP FOR THE TREATMENT OF THIN AND MODERATELY THICK, NON-HYPERKERATOTIC, NON-PIGMENTED ACTINIC KERATOSES OF THE FACE AND SCALP IN IMMUNOCOMPETENT PATIENTS
- M-78 CLINICAL TRIAL INFO ON USE OF STRATTERA IN PATIENTS WITH ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) AND COMORBID ANXIETY DISORDER WITHOUT CAUSING

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

WORSENING OF ANXIETY

- M-79 LABELING REVISIONS RELATED TO SMOKING AND ERLOTINIB EXPOSURE
- M-80 ADDITIONAL TIME POINT OF 30 MINUTES (0.5 HOUR) IN CHILDREN AGED 6-12 YEARS WITH A DIAGNOSIS OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER (ADHD)
- M-81 ADDITIONAL INFO FOR PEDIATRIC USE FOR CASODEX (STUDIED IN COMBINATION WITH ARIMIDEX) IN THE PEDIATRIC POPULATION, SPECIFICALLY BOYS WITH FAMILIAL MALE-LIMITED PRECOCIOUS PUBERTY (TESTOTOXICOSIS)
- M-82 LABELING REVISIONS RELATED TO CLINICAL STUDIES
- M-83 ADDITIONAL INFORMATION ADDED TO LABELING REGARDING ESTABLISHMENT OF EFFICACY IN ADDITIONAL CLINICAL TRIALS AND ONE MAINTENANCE TRIAL
- M-84 STUDY INFORMATION ADDED TO LABEL REGARDING BONE MINERAL DENSITY
- M-85 INFORMATION ADDED TO LABELING REGARDING USE OF PREVACID IN PATIENTS LESS THAN 1 YEAR WITH SYMPTOMATIC GERD
- M-86 LABELING CHANGES SUBMITTED IN RESPONSE TO PEDIATRIC WRITTEN REQUEST FOR INFANTS AGES BIRTH TO 11 MONTH INCLUSIVE REFLECTING LACK OF EFFICACY FOR GERD INDICATION FOR THIS PATIENT POPULATION
- M-87 INCLUSION OF RESULTS FROM TWO DRUG INTERACTION STUDIES WITH LIPITOR AND CRESTOR IN CLINICAL PHARMACOLOGY SECTION
- M-88 ADDITION OF INFORMATION REGARDING ABUSE POTENTIAL OF CONCERTA VERSUS IMMEDIATE-RELEASE METHYLPHENIDATE
- M-89 PROVIDES FOR REVISIONS TO MULTIPLE SECTIONS OF THE PACKAGE INSERT TO REFLECT RESULTS OF CLINICAL TRIALS 205.235 (UPLIFT) AND 205.266 (VA STUDY) IN SUPPORT OF EXACERBATION CLAIM
- M-90 LABELING CHANGES BASED ON DATA FROM CLINICAL STUDIES NV20235 AND NV20236 STUDIES OF SEASONAL PROPHYLAXIS OF INFLUENZA IN IMMUNOCOMPROMISED PATIENTS AND CHILDREN AGES 1-12
- M-91 UPDATED LABELING BASED UPON STUDY: A SINGLE-DOSE, SINGLE-BLIND, PLACEBO-AND MOXIFLOXACIN-CONTROLLED 2-PERIOD, RANDOMIZED, CROSSOVER, 3RD PERIOD SEQUENTIAL STUDY OF SIDE EFFECTS OF TEMSIROLIMUS ON CARDIAC REPOLARIZATION IN HEALTHY SUBJECTS
- M-92 UPDATES TO THE PACKAGE INSERT BASED UPON THE TRIAL ENTITLED "A PHASE I PHARMACOKINETIC AND PHARMACODYNAMIC STUDY OF TEMSIROLIMUS IN PATIENTS WITH ADVANCED MALIGNANCIES AND NORMAL AND IMPAIRED LIVER FUNCTION"
- M-93 EXPANSION OF LABELING TO INCLUDE INFORMATION ON SAFETY AND EFFICACY OF CREON IN PATIENTS AGES 7 YEARS THROUGH 11 YEARS WITH PANCREATIC EXOCRINE INSUFFICIENCY DUE TO CYSTIC FIBROSIS
- M-94 INFO ADDED TO LABEL RELATED TO NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME POSITIVE (PH+) CHRONIC MYELOID LEUKEMIA IC CHRONIC PHASE
- M-95 INFORMATION FOR TREATMENT OF CHRONIC HEPATITIS B (CHB) IN ADULT PATIENTS WITH DECOMPENSATED LIVER DISEASE BASED ON DATA FROM CLINICAL TRIAL GS-US-174-0108
- M-96 UPDATED INFORMATION IN THE CLINICAL STUDIES SECTION RELATED TO THE LOSS AND RECOVERY OF BONE MINERAL DENSITY IN ADOLESCENT GIRLS DURING AND FOLLOWING THE USE OF DEPO-PROVERA CONTRACEPTIVE INJECTION
- M-97 LABELING CHANGES IN RESPONSE TO PEDIATRIC STUDIES - NOT INDICATED FOR USE IN PEDIATRIC POPULATION
- M-98 NEW INFORMATION FROM A STUDY WHICH EVALUATED THE SAFETY AND EFFICACY OF FAMVIR IN TREATING RECURRENT GENITAL HERPES IN IMMUNOCOMPETENT BLACK/AFRICAN AMERICAN SUBJECTS.
- M-99 ADDITION OF FINDINGS FROM A SINGLE PEDIATRIC CLINICAL TRIAL (P04292) OF NASONEX NASAL SPRAY IN THE TREATMENT OF NASAL POLYPS IN PATIENTS 6 TO <18 YEARS OF AGE TO THE PACKAGE INSERT.
- M-100 INFORMATION ADDED TO LABEL BASED UPON COMPLETED CLINICAL TRIAL REPORTS
- M-101 INCLUSION OF DATA FROM AN ADDITIONAL 19 SUBJECTS WITH HYPERCALCEMIA FROM PARATHYROID CARCINOMA TO THE INFORMATION CURRENTLY PRESENTED IN THE LABEL
- M-102 INFORMATION FROM PEDIATRIC STUDY REPORT ML16633, "INTRAVENOUS GRANISETRON (KYTRIL) IN THE PREVENTION OF POST-OPERATIVE NAUSEA AND VOMITING (PONV) IN PEDIATRIC SUBJECTS UNDERGOING TONSILLECTOMY OR ADENOTONSILLECTOMY."
- M-103 SAFETY, EFFICACY AND PHARMACOKINETIC INFO FOR FASLODEX IN THE PEDIATRIC

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- POPULATION, SPECIFICALLY FOR GIRLS WITH PROGRESSIVE PRECOCIOUS PUBERTY ASSOCIATED WITH MCCUNE-ALBRIGHT SYNDROME ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING
- M-104 INFORMATION ADDED TO DOSING AND ADMINISTRATION REGARDING A 26 WEEK STUDY
- M-105 NEW LANGUAGE ADDED TO CLINICAL STUDIES REGARDING USE IN SMOKERS WITH CARDIOVASCULAR DISEASE, CHRONIC OBSTRUCTIVE PULMONARY DISEASE, AND USE ACCORDING TO AN ALTERNATIVE SET OF DIRECTIONS FOR SETTING A QUIT DATE
- M-106 ADDITION OF THE T1-WEIGHTED GD-ENHANCED LESION EFFICACY VARIABLE IN THE CLINICAL STUDIES SECTION 14 OF THE PACKAGE INSERT
- M-107 INFORMATION TO THE CLINICAL STUDIES SECTION OF THE LUPRON DEPOT-PED 1-MONTH BASED UPON THE PHASE 3/4 COMPLETED CLINICAL STUDY REPORT FOR STUDY M90-516 ENTITLED "STUDY OF LUPRON DEPOT IN THE TREATMENT OF CENTRAL PRECOCIOUS PUBERTY".
- M-108 CHANGES ARE BASED ON RESULTS FROM STUDY CV181057
- M-109 CHANGES TO THE PACKAGE INSERT TO REFLECT THE RESULTS OF THE STUDY OF HEART AND RENAL PROTECTION (SHARP) TRIAL
- M-110 CHANGES TO THE PACKAGE INSERT TO REFLECT THE RESULTS OF THE STUDY OF HEART AND RENAL PROTECTION (SHARP) TRIAL
- M-111 LABELING CHANGES BASED ON STUDY HW80-EW-GWCI ENTITLED A PLACEBO AND POSITIVE CONTROLLED STUDY OF THE ELECTROPHYSIOLOGICAL EFFECTS OF A SINGLE 10 MCG DOSE OF EXENATIDE ON THE 12 LEAD ELECTROCARDIOGRAM QT INTERVAL IN HEALTHY SUBJECTS
- M-112 REVISIONS TO THE PEDIATRIC USE SECTION OF THE PACKAGE INSERT TO ADD INFORMATION FROM A PEDIATRIC STUDY IN PATIENTS AGED 12 YEARS TO LESS THAN 18 YEARS OF AGE WITH RECURRENT HERPES LABIALIS
- M-113 LABELING CHANGES BASED ON STUDY H80-US-GWCO ENTITLED A RANDOMIZED TRIAL COMPARING EXENATIDE WITH PLACEBO IN SUBJECTS WITH TYPE 2 DIABETES ON INSULIN GLARGINE WITH OR WITHOUT ORAL ANTIHYPERGLYCEMIC MEDICATIONS
- M-114 CHANGES IN SECTION 14 OF THE PACKAGE INSERT TO INCLUDE DATA FROM THE SWITCHMRK STUDIES (SWITCH OF SUPPRESSED SUBJECTS FROM LOPINAVIR/RITONAVIR TO RALTEGRAVIR)
- M-115 REVISIONS TO THE PI BASED ON RESULTS FROM STUDY NN2211-1842, ENTITLED THE EFFECT OF INSULIN DETEMIR IN COMBINATION WITH LIRAGLUTIDE AND METFORMIN COMPARED TO LIRAGLUTIDE AND METFORMIN IN SUBJECTS WITH TYPE 2 DIABETES
- M-116 LABELING CHANGES BASED ON RESULTS FROM CLINICAL STUDY 01-06-TL-OPIMET-008
- M-117 ADDITION OF RESULTS OF PEDIATRIC TRIAL TO LABEL
- M-118 LABELING CHANGES BASED UPON SAFETY AND EFFICACY RESULTS FROM TRIAL 1218.36
- M-119 LABELING CHANGES REGARDING MISSED DOSES
- M-120 CHANGES TO CLINICAL TRIALS DETAILING STUDY RESULTS
- M-121 LABELING CHANGES BASED UPON SAFETY AND EFFICACY RESULTS FROM TRIAL 1218.43
- M-122 LABELING CHANGES TO INCLUDE THE RESULTS OF THE PARAMOUNT TRIAL
- M-123 UPDATED RESULTS OF OVERALL SURVIVAL FROM 'CONFIRM' STUDY
- M-124 LONG TERM SAFETY AND EFFICACY DATA FROM STUDY CLDT600A2303 FOR SUBJECTS PREVIOUSLY ENROLLED IN THE ORIGINAL TWO YEAR GLOBE (NV-02B-007/CLDT600A2302) AND NV02B-015 STUDIES WHO CONTINUED TELBIVUDINE TREATMENT FOR UP TO 208 WEEKS
- M-125 LABELING CHANGES TO INCLUDE LACK OF EFFICACY IN CHILDREN 6 MONTHS TO 4 YEARS OF AGE
- M-126 UPDATES TO THE CLINICAL STUDIES SECTION 14, OF THE PACKAGE INSERT (PI), WITH THE RESULTS OF CLINICAL TRIAL P06086
- M-127 REVISIONS TO THE PEDIATRIC USE SECTION OF THE PACKAGE INSERT TO REFLECT THE RESULTS FROM CLINICAL STUDY C-10-004
- M-128 CLINICAL TRIAL STUDY RESULTS
- M-129 RESULTS OF A CLINICAL STUDY REPORT WHICH ASSESSES THE SAFETY AND EFFICACY IN CHILDREN AGES 6 TO 12 YEARS OF AGE
- M-130 ADDITION OF INFORMATION ON LONG-TERM TREATMENT WITH VPRIV IN THE CLINICAL TRIALS SECTION OF THE PACKAGE INSERT
- M-131 INFORMATION FROM STUDIES CONDUCTED IN PEDIATRIC PATIENTS WITH NEWLY DIAGNOSED NON-DISSEMINATED DIFFUSED INTRINSIC BRAINSTEM GLIOMAS
- M-132 REVISIONS TO THE CLINICAL TRIALS SECTION IN THE INOMAX LABEL TO REFLECT RESULTS

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

FROM THE PEDIATRIC STUDY REPORTS

- M-133 INFORMATION ADDED TO THE LABELING REGARDING THE ADDITION OF SILDENAFIL TO BOSENTAN THERAPY
- M-134 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES PERFORMED WITH SAXAGLIPTIN IN COMBINATION WITH METFORMIN AND A SULFONYLUREA ADDED TO THE LABELING
- M-135 ADDITION OF INFORMATION TO THE CLINICAL STUDIES - RADIOGRAPHIC RESPONSE SECTION OF THE PACKAGE INSERT
- M-136 ADDITIONAL INFORMATION ADDED TO THE USE IN SPECIFIC POPULATIONS SECTION OF THE LABELING REGARDING POST-OPERATIVE NAUSEA AND VOMITING STUDIES IN PEDIATRIC PATIENTS
- M-137 LABELING REVISIONS RESULTING FROM A MAINTENANCE TRIAL IN PEDIATRIC PATIENTS WITH IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER
- M-138 INFORMATION ADDED TO THE 8.4 PEDIATRIC USE SECTION ON THE USE OF MEMANTINE IN CHILDREN AGES 6-12 YEARS WITH AUTISM SPECTRUM DISORDER
- M-139 INFORMATION ADDED TO THE DOSING AND ADMINISTRATION SECTION OF THE PACKAGE INSERT REGARDING RETREATMENT WITH VELCADE FOR PATIENTS WITH MULTIPLE MYELOMA
- M-140 INFORMATION ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING REGARDING USE OF LATISSE IN PATIENTS WHO WERE POST-CHEMOTHERAPY OR HAD ALOPECIA AREATA, AND ADOLESCENTS WHO HAD HYPERTRICHOSIS WITH NO ASSOCIATED MEDICAL CONDITION
- M-141 REVISIONS TO THE PEDIATRIC USE SECTION OF THE LABELING TO INCORPORATE STUDY RESULTS FOR TREATMENT OF MAJOR DEPRESSIVE DISORDER IN ADOLESCENTS (AGES 12-17)
- M-142 ADDITIONS TO THE LABELING DESCRIBING RESULTS FROM STUDY H6P-MC-HDAY
- M-143 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSATION IN PATIENTS WITH CURRENT OR PAST HISTORY OF MAJOR DEPRESSIVE DISORDER
- M-144 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSATION IN PATIENTS WHO HAD BEEN PREVIOUSLY TREATED WITH VARENICLINE
- M-145 ADDITION OF INFORMATION ABOUT LONG-TERM TREATMENT OF PULMONARY ARTERIAL HYPERTENSION TO THE CLINICAL STUDIES SECTION OF THE LABELING
- M-146 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION ON INITIAL COMBINATION THERAPY WITH LINAGLIPTIN AND METFORMIN VS. LINAGLIPTIN MONOTHERAPY IN TREATMENT NAIVE PATIENTS
- M-147 OTC USE FOR TEMPORARY RELIEF OF OCULAR SYMPTOMS DUE TO HAY FEVER OR OTHER UPPER RESPIRATORY ALLERGIES
- M-148 LABELING CHANGES BASED ON STUDY H80-EW-GWDM
- M-149 INFORMATION ADDED TO THE LABELING REGARDING MAINTENANCE MONOTHERAPY FOR ADHD
- M-150 ADDITION OF THE RESULTS OF A CONTROLLED CLINICAL STUDY TREATING ADULT PATIENTS WITH SCHIZOPHRENIA EXPERIENCING AN ACUTE RELAPSE
- M-151 REVISIONS TO THE LABELING BASED ON THE OUTCOMES OF PEDIATRIC STUDIES CONDUCTED TO ASSESS THE SAFETY AND EFFICACY OF XOPENEX IN SUBJECTS LESS THAN 6 YEARS OF AGE
- M-152 INFORMATION ADDED TO THE CLINICAL PHARMACOLOGY SECTION OF THE LABELING REGARDING A SAFETY STUDY IN PEDIATRIC SUBJECTS AGES 6 MONTHS TO 4 YEARS OF AGE WITH AN ACTIVE HEAD LICE INFESTATION
- M-153 ADDITION OF INFORMATION REGARDING THE INTRANASAL ABUSE POTENTIAL OF OXYCONTIN
- M-154 UPDATE TO THE LABELING TO REFLECT THE RESULTS OF A LONG-TERM MAINTENANCE TREATMENT STUDY OF ADHD IN CHILDREN AND ADOLESCENTS AGES 6-17.
- M-155 ADDITION OF CLINICAL FINDINGS FROM AN OBSERVATIONAL STUDY IN A PEDIATRIC AGE GROUP GREATER THAN 2 MONTHS TO 18 YEARS IN SECTION 8.4 PEDIATRIC USE OF THE PACKAGE INSERT
- M-156 UPDATE TO THE LABELING WITH INFORMATION REGARDING A CLINICAL TRIAL IN CHILDREN LESS THAN 4 YEARS OF AGE.
- M-157 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF DAPAGLIFLOZIN 10MG ONCE DAILY IN PATIENTS WITH TYPE 2 DIABETES WHO HAVE INADEQUATE GLYCEMIC CONTROL ON A BACKGROUND COMBINATION OF METFORMIN AND SULFONYLUREA

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- M-158 UPDATES TO THE LABELING TO REFLECT SAFETY RESULTS FROM CLINICAL TRIALS IN SCHIZOPHRENIA ADOLESCENT PATIENTS AGED 12 TO 17 YEARS
- M-159 ADDITION OF PED SAFETY INFORMATION DERIVED FROM A MAINTENANCE TREATMENT STUDY OF BIPOLAR 1 DISORDER TO DELAY THE TIME TO OCCURRENCE OF MOOD EPISODES IN PATIENTS (> THAN OR = TO 13 YRS OF AGE) TREATED FOR ACUTE MOOD EPISODES WITH STANDARD THERAPY
- M-160 UPDATED LABELING WITH DATA FROM A RANDOMIZED, DOUBLE-BLIND ACTIVE-CONTROLLED STUDY COMPARING EMPAGLIFLOZIN TO GLIMEPIRIDE IN PATIENTS WITH TYPE 2 DIABETES AND INSUFFICIENT GLYCEMIC CONTROL DESPITE METFORMIN TREATMENT
- M-161 UPDATED LABELING WITH DATA FROM A RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDY OF EMPAGLIFLOZIN IN PATIENTS WITH TYPE 2 DIABETES MELLITUS AND INSUFFICIENT GLYCEMIC CONTROL ON A MULTIPLE DAILY INJECTION INSULIN REGIMEN ALONE OR WITH METFORMIN
- M-162 INCLUSION OF EFFICACY AND SAFETY DATA TO THE PRESCRIBING INFORMATION OF BYDUREON BASED ON STUDY GWDE
- M-163 INFORMATION ADDED TO THE LABELING REGARDING PREVIOUSLY UNTREATED ALK-POSITIVE METASTATIC NON SMALL CELL LUNG CANCER (NSCLC)
- M-164 REVISES THE CLINICAL TRIALS SECTION OF THE PRESCRIBING INFORMATION TO INCORPORATE THE RESULTS FROM STUDY E7273-G000-401 ENTITLED "PHASE IV RANDOMIZED STUDY OF TWO DOSE LEVELS OF TARGRETIN CAPSULES IN SUBJECTS WITH REFRACTORY CUTANEOUS T-CELL LYMPHOMA"
- M-165 PROVIDES FOR UPDATES TO THE PEDIATRIC USE SECTION BASED ON THE PEDIATRIC STUDY REPORT ENTITLED, "A PHASE II PILOT TRIAL OF BORTEZOMIB IN COMBINATION WITH INTENSIVE RE-INDUCTION THERAPY IN CHILDREN WITH RELAPSED ACUTE LYMPHOBLASTIC LYMPHOMA (LL) "
- M-166 UPDATE TO LABELING WITH WEEK 48 RESULTS FROM VIKING-4 IN ANTIRETROVIRAL THERAPY (ART) - EXPERIENCED INTEGRASE STRAND TRANSFER INHIBITOR (INSTI) - RESISTANT SUBJECTS
- M-167 APPROVED FOR REVISIONS TO THE LABELING BASED ON THE CLINICAL STUDY ENTITLED "BRONCHOPULMONARY DYSPLASIA (BPD) IN PRETERM INFANTS REQUIRING MECHANICAL VENTILATION OR POSITIVE PRESSURE SUPPORT ON DAYS 5 TO 14 AFTER BIRTH".
- M-168 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING REGARDING THE RE-NOVATE AND RE-NOVATE LL STUDIES (PROPHYLAXIS OF DEEP VEIN THROMBOSIS AND PULMONARY EMBOLISM FOLLOWING HIP REPLACEMENT SURGERY)
- M-169 UPDATES TO LABELING DESCRIBING RESPONSE TO A REPEAT COURSE OF PICATO GEL 0.015% ON THE FACE OR SCALP IF AN INCOMPLETE RESPONSE IS OBSERVED AT A FOLLOW-UP EXAMINATION.
- M-170 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION REGARDING USE FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- M-171 UPDATES TO LABELING WITH RESULTS TO THE TIGER CLINICAL TRIAL
- M-172 UPDATES TO THE CLINICAL TRIALS SECTION OF THE LABELING TO INCLUDE RESULTS OF STUDIES PERFORMED TO EVALUATE THE BENEFIT OF ADDING INCRUSE ELLIPTA TO PATIENTS WHO ARE ON BACKGROUND THERAPY WITH BREO ELLIPTA AND ADVAIR DISKUS
- M-173 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING DESCRIBING THE EFFECTS OF STIOLTO RESPIMAT ON COPD PATIENTS
- M-174 INFORMATION ADDED TO CLINICAL STUDIES SECTION OF THE LABELING REGARDING INITIAL COMBINATION THERAPY OF EMPAGLIFLOZIN WITH METFORMIN
- M-175 INFORMATION ADDED TO THE LABELING DESCRIBING SAVOR, A PHASE IV TRIAL EVALUATING THE EFFECT OF SAXAGLIPTIN ON THE INCIDENCE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION OR ISCHAEMIC STROKE IN PATIENTS WITH TYPE 2 DIABETES
- M-176 INFORMATION ADDED TO THE LABELING DESCRIBING TRIAL NN2211-3916, A TRIAL EVALUATING THE SAFETY AND EFFICACY OF LIRAGLUTIDE IN SUBJECTS WITH TYPE 2 DIABETES AND MODERATE RENAL IMPAIRMENT
- M-177 INFORMATION ADDED TO THE LABELING DESCRIBING EXAMINE, A TRIAL EVALUATING CARDIOVASCULAR ISCHEMIC RISKS ASSOCIATED WITH ALOGLIPTIN USE IN PATIENTS WITH TYPE 2 DIABETES AT HIGH RISK OF ISCHEMIC CARDIOVASCULAR DISEASE
- M-178 INFORMATION ADDED TO THE LABELING REGARDING MAINTENANCE OF REMISSION IN CROHN'S DISEASE IN PEDIATRIC PATIENTS
- M-179 UPDATES TO THE PRODUCT LABELING WITH STUDY REPORTS FROM THE OPTIMIST-1 AND

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

OPTIMIST-2 CLINICAL TRIALS

- M-180 INFORMATION ADDED TO THE LABELING REGARDING THE ADDITION OF MAINTENANCE TREATMENT IN PATIENTS WITH SCHIZOPHRENIA
- M-181 UPDATE TO THE DOSAGE AND ADMINISTRATION, PATIENT SELECTION (2.1), SECTION OF THE PACKAGE INSERT TO INCLUDE THE USE OF AN FDA-APPROVED PLASMA TEST FOR THE IDENTIFICATION OF EGFR EXON 19 DELETION OR EXON 21 (L858R) SUBSTITUTION MUTATIONS
- M-182 UPDATES TO THE PRODUCT LABELING BASED ON THE RESULTS OF STUDY H7T-MC-TADO TITLED, "A PHASE 3 DOUBLE-BLIND, RANDOMIZED, MULTICENTER, EFFICACY AND SAFETY STUDY OF PRASUGREL COMPARED TO PLACEBO IN PEDIATRIC PATIENTS WITH SICKLE CELL DISEASE"
- M-183 CHANGES TO THE DOSAGE AND ADMINISTRATION AND CLINICAL STUDIES SECTIONS OF THE LABELING TO SUPPORT THE REDUCE-TO-QUIT PARADIGM
- M-184 UPDATES MADE TO THE LABELING TO INCLUDE INFORMATION FROM STUDY MO25743 ON THE ANTI-TUMOR ACTIVITY OF VEMURAFENIB IN THE TREATMENT OF PATIENTS WITH BRAF V600E MUTATION-POSITIVE MELANOMA WITH BRAIN METASTASES
- M-185 UPDATES TO THE LABELING TO INCLUDE RESULTS OF A TRIAL TO EVALUATE THE SAFETY OF MOXIFLOXACIN IN PEDIATRIC PATIENTS WITH COMPLICATED INTRA-ABDOMINAL INFECTIONS
- M-186 UPDATES TO THE PRODUCT INFORMATION REGARDING MAINTENANCE TREATMENT OF SCHIZOPHRENIA IN ADULTS BASED UPON THE RESULTS FROM STUDY 331-10-232
- M-187 ADDITION OF CLINICAL INFORMATION OBTAINED FROM A PEDIATRIC TRIAL TO SECTION 8.4 OF THE LABELING
- M-188 PROVIDES FOR DATA SUPPORTING THE SAFETY AND EFFECTIVENESS FOR THE MAINTENANCE TREATMENT OF MODERATE TO SEVERE BINGE EATING DISORDER (BED)
- M-189 LABELING DESCRIBING THE EXPECTED REDUCTION OF ABUSE OF SINGLE-ENTITY MORPHINE BY THE INTRANASAL ROUTE OF ADMINISTRATION DUE TO PHYSICOCHEMICAL PROPERTIES
- M-190 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING REGARDING THE LACK OF EFFICACY OF TARCEVA IN MAINTENANCE TREATMENT OF PATIENTS WITHOUT EGFR MUTATIONS
- M-191 ADDITION OF DATA BASED ON PEDIATRIC STUDIES TO FULFILL THE POSTMARKETING REQUIREMENT 1857-2
- M-192 PROVIDES FOR DATA EVALUATING THE NEUROPSYCHIATRIC SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSATION IN SUBJECTS WITH AND WITHOUT A HISTORY OF PSYCHIATRIC DISORDERS
- M-193 INFORMATION ADDED TO THE LABELING REGARDING A 15-WEEK, RANDOMIZED, DOUBLE-BLIND, PARALLEL-GROUP, PLACEBO-CONTROLLED FLEXIBLE-DOSE SAFETY AND EFFICACY STUDY OF PREGABALIN IN ADOLESCENTS (12 THROUGH 17 YEARS OLD) WITH FIBROMYALGIA
- M-194 INFORMATION ADDED TO THE LABELING REGARDING USE OF REGADENOSON ADMINISTRATION FOLLOWING AN INADEQUATE EXERCISE STRESS TEST AS COMPARED TO REGADENOSON ALONE
- M-195 REVISIONS TO THE PEDIATRIC USE SECTION OF THE LABELING REFLECTING LACK OF EFFICACY FOR IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER IN PEDIATRIC PATIENTS AGES 6-17
- M-196 REVISIONS TO THE PACKAGE INSERT BASED ON DATA FROM A RANDOMIZED, PLACEBO CONTROLLED, MULTICENTER STUDY OF INTRAVENOUS ACETAMINOPHEN FOR THE TREATMENT OF ACUTE PAIN IN PEDIATRIC PATIENTS TO FULFILL THE POST-MARKETING REQUIREMENT 1704-1
- M-197 NEW CLINICAL DATA ADDED TO THE PRESCRIBING INFORMATION REGARDING CANAGLIFLOZIN ADD-ON COMBINATION THERAPY WITH METFORMIN AND A DIPEPTIDYL-PEPTIDASE-4 INHIBITOR
- M-198 PACKAGE INSERT UPDATED WITH RESULTS FROM STUDY CV181168, A MULTICENTER, RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED, PARALLEL GROUP, PHASE 3 TRIAL TO EVALUATE THE SAFETY AND EFFICACY OF SAXAGLIPTIN ADDED TO DAPAGLIFLOZIN AND METFORMIN
- M-199 INFORMATION ADDED TO LABELING REGARDING THE TREATMENT OF PATIENTS WITH ALK-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAD NOT RECEIVED PRIOR SYSTEMIC THERAPY FOR METASTATIC DISEASE.
- M-200 CLINICAL INFORMATION ADDED TO THE USE IN SPECIFIC POPULATIONS SECTION OF THE LABELING.
- M-201 REVISIONS TO THE PACKAGE INSERT BASED ON DATA FROM AN OPEN LABEL, MULTI-CENTER STUDY OF CABAZITAXEL IN PEDIATRIC PATIENTS WITH REFRACTORY SOLID TUMORS INCLUDING TUMORS OF THE CENTRAL NERVOUS SYSTEM.

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

- M-202 INCLUSION OF DATA FROM THE SUMMIT STUDY FOR BREO ELLIPTA (FLUTICASONE FUROATE/VILANTEROL TRIFENATATE) INHALATION POWDER IN THE PACKAGE INSERT.
- M-203 PROVIDES FOR REVISIONS TO THE PACKAGE INSERT TO REFLECT RESULTS OF TWO POSTMARKETING REQUIREMENT STUDIES ROP111662 AND ROP111569
- M-204 CLINICAL INFORMATION ADDED TO THE PACKAGE INSERT REGARDING USE OF ATORVASTATIN IN CHILDREN AND ADOLESCENTS AGES 10-17 WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH)
- M-205 INFORMATION ADDED TO THE LABELING REGARDING RANDOMIZED, MULTICENTER, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDIES ON PATIENTS WITH SEVERE RENAL IMPAIRMENT
- M-206 INFORMATION ADDED TO LABELING REGARDING 48 WEEK EFFICACY, RESISTANCE AND SAFETY DATA ON VIROLOGICALLY SUPPRESSED HIV-1 INFECTED ADULTS SWITCHING FROM COMPLERA TO ODEFSEY
- M-207 INFORMATION ADDED TO LABELING REGARDING 48 WEEK EFFICACY, RESISTANCE AND SAFETY DATA ON VIROLOGICALLY SUPPRESSED HIV-1 INFECTED ADULTS SWITCHING FROM ATRIPLA TO ODEFSEY
- M-208 INFORMATION ADDED TO THE LABELING TO INCLUDE RESULTS OF A RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDY IN PATIENTS WITH SEVERE COPD ASSOCIATED WITH CHRONIC BRONCHITIS AND A HISTORY OF EXACERBATIONS
- M-209 INFORMATION ADDED TO THE LABELING REGARDING CABAZITAXEL AT 20 MG/M2 BASED ON THE RESULTS OF THE PROSELICA STUDY
- M-210 INFORMATION ADDED TO LABELING TO SUPPORT THE USE OF SYMBICORT TO REDUCE EXACERBATIONS IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- M-211 PROVIDES FOR LABELING CHANGES REGARDING THE USE OF DAPTOMYCIN IN THE PEDIATRIC POPULATION FOR STAPHYLOCOCCUS AUREUS BACTEREMIA (SAB) BASED ON RESULTS OF A TRIAL IN PEDIATRIC PATIENTS 1 TO 17 YEARS OF AGE
- M-212 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF DAPAGLIFLOZIN IN PATIENTS WITH TYPE 2 DIABETES WHO HAVE INADEQUATE GLYCEMIC CONTROL ON A BACKGROUND COMBINATION OF METFORMIN AND EXENATIDE EXTENDED RELEASE
- M-213 INFORMATION ADDED TO THE LABELING TO INCLUDE THE EFFICACY AND SAFETY OF CARIPRAZINE RELATIVE TO PLACEBO IN THE PREVENTION OF RELAPSE OF SYMPTOMS IN PATIENTS WITH SCHIZOPHRENIA
- M-214 INFORMATION ADDED TO THE CLINICAL TRIALS SECTION OF THE LABELING REGARDING A POSTMARKETING SAFETY AND EFFICACY STUDY EVALUATING THE RISK OF SERIOUS ASTHMA-RELATED EVENTS
- M-215 INFORMATION ADDED TO THE LABELING REGARDING THE COMPARISON OF PALIPERIDONE PALMITATE COMPARED WITH ORAL ANTIPSYCHOTIC TREATMENT IN DELAYING TIME TO TREATMENT FAILURE IN ADULTS WITH SCHIZOPHRENIA WHO HAVE BEEN INCARCERATED
- M-216 UPDATE THE PRESCRIBING INFORMATION AND PATIENT LABELING WITH FINDINGS FROM STUDY RP103-08 CONDUCTED IN TREATMENT-NAIVE NEPHROPATHIC CYSTINOSIS PATIENTS TO EXPAND THE INDICATED POPULATION TO PATIENTS 1 YEAR AND OLDER
- M-217 INCORPORATION OF THE LABELING REVISIONS PROVIDED FOR IN NDA 022253/S-039 AND NDA 022255/S-022 INTO THE LACOSAMIDE INJECTION LABELING
- M-218 ADDITIONAL INFORMATION ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING REGARDING A NEW CLINICAL TRIAL IN PATIENTS AGED 6 THROUGH 11 YEARS (TRIAL 4)
- M-219 INFORMATION ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING REGARDING A NEW CLINICAL TRIAL IN PATIENTS 7 TO 14 YEARS OF AGE WITH DUCHENNE MUSCULAR DYSTROPHY
- M-220 ADDITIONAL INFORMATION ADDED TO THE LABELING FROM STUDY PC B308/13 REGARDING THE USE OF BLUE LIGHT CYSTOSCOPY WITH CYSVIEW AS AN ADJUNCT TO WHITE LIGHT CYSTOSCOPY
- M-221 DRUG FACTS LABELING CHANGES UNDER THE DIRECTIONS HEADING TO REVISE THE STATED PREPARATION TIME OF A DRY SITE FROM 120 SECONDS SCRUBBING AND 90 SECONDS DRYING TO 30 SECONDS SCRUBBING AND 30 SECONDS DRYING
- M-222 ADDITION OF DATA BASED ON THE ASSESSMENT OF SAFETY AND EFFICACY IN PEDIATRIC PATIENTS WITH MAJOR DEPRESSIVE DISORDER TO FULFILL POSTMARKETING STUDY REQUIREMENT 1229-1
- M-223 INFORMATION ADDED TO SECTION 8.1 OF THE LABELING REGARDING PREGNANT PATIENTS WHO ARE ALREADY ON A STABLE RILPIVIRINE REGIMEN PRIOR TO PREGNANCY AND WHO ARE VIROLOGICALLY SUPPRESSED (HIV-1 RNA LESS THAN 50 COPIES/ML)
- M-224 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF EXENATIDE EXTENDED RELEASE AS ADD-ON IN PATIENTS WITH TYPE 2 DIABETES WHO HAVE INADEQUATE

PATENT AND EXCLUSIVITY TERMS**EXCLUSIVITY MISCELLANEOUS**

| | |
|-------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| | GLYCEMIC CONTROL ON BASAL INSULIN GLARGINE WITH OR WITHOUT METFORMIN |
| M-225 | REVISIONS TO SECTION 8.4 OF THE PRESCRIBING INFORMATION TO INCLUDE A SAFETY AND EFFICACY STUDY IN PEDIATRIC PATIENTS AGES >=6 YEARS TO <18 YEARS WITH CHRONIC IDIOPATHIC CONSTIPATION |
| M-226 | CHANGES TO THE LABELING BASED ON RESULTS FROM A CONTROLLED CLINICAL TRIAL IN PATIENTS WITH LATER-ONSET SPINAL MUSCULAR ATROPHY |
| M-227 | ADDITION TO THE CLINICAL STUDIES SECTION OF THE LABELING WITH THE SUBSECTION ENTITLED DIGIT SYMBOL SUBSTITUTION TEST IN MAJOR DEPRESSIVE DISORDER |
| M-228 | INFORMATION ADDED TO THE PACKAGE INSERT REGARDING THE REVISION OF THE MONOTHERAPY INDICATION OF VENETOCLAX |
| M-229 | REVISED LABELING TO INCORPORATE THE PEDIATRIC USE OF LOTEPREDNOL ETABONATE GEL IN PATIENTS FOR THE TREATMENT OF POSTOPERATIVE INFLAMMATION FOLLOWING OCULAR SURGERY |
| M-230 | REVISIONS TO THE GLECAPREVIR/PIBRENTASVIR COMBINATION PRODUCT PRESCRIBING INFORMATION TO INCLUDE SAFETY AND EFFICACY DATA FROM THE HCV/HIV-1 COINFECTION STUDY M14-730 AND FROM THE LIVER AND RENAL TRANSPLANT STUDY M13-596 |
| M-231 | REVISIONS TO THE USE IN SPECIFIC POPULATIONS SECTION (SECTION 8.3) OF THE PACKAGE INSERT WITH THE RESULTS OF CLINICAL TRIAL WV25651, CONDUCTED TO EVALUATE THE EFFECT OF VALGANCYCLOVIR ON SPERMATOGENESIS AND TO FULFILL PMR 1670-3 |
| M-232 | INFORMATION ADDED TO SECTION 8.4 OF THE LABELING TO DESCRIBE THE RESULTS FROM PEDIATRIC STUDIES |
| M-233 | INFORMATION ADDED TO THE LABELING TO DESCRIBE FIXED-DOSE COMBINATION OF TIOTROPIUM BROMIDE AND OLODATEROL TO INCLUDE REDUCTION OF COPD EXACERBATIONS |
| M-234 | UPDATE TO THE PRESCRIBING INFORMATION FOR VORTIOXETINE ON TREATMENT-EMERGENT SEXUAL DYSFUNCTION COMPARING VORTIOXETINE AND SSRIS |
| M-235 | INFORMATION ADDED TO SECTION 14 OF THE LABELING TO DESCRIBE STUDY LAP016A2307 TO FULFILL POSTMARKETING STUDY REQUIREMENT 1586-1 |

ORPHAN DRUG EXCLUSIVITY

| | |
|--------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| ODE-1 | TO REDUCE CHRONIC DROOLING IN PATIENTS AGED 3 - 16 WITH NEUROLOGIC CONDITIONS ASSOCIATED WITH PROBLEM DROOLING (E.G. CEREBRAL PALSY) |
| ODE-2 | FOR TREATMENT OF NON-INFECTIOUS UVEITIS AFFECTING THE POSTERIOR SEGMENT OF THE EYE |
| ODE-3 | TO TREAT INFANTILE SPASMS |
| ODE-4 | TREATMENT OF PATIENTS WITH SUBEPENDYMAL GIANT CELL ASTROCYTOMA (SEGA) ASSOCIATED WITH TUBEROUS SCLEROSIS WHO REQUIRE THERAPEUTIC INTERVENTION BUT ARE NOT CANDIDATES FOR CURATIVE SURGICAL RESECTION |
| ODE-5 | FOR SEQUENTIAL USE FOR THE TREATMENT OF CYANIDE POISONING THAT IS JUDGED TO BE LIFE-THREATENING |
| ODE-6 | FOR THE MANAGEMENT OF POSTHERPETIC NEURALGIA |
| ODE-7 | TO REDUCE THE RISK OF PRETERM BIRTH IN WOMEN WITH SINGLETON PREGNANCY WHO HAVE A HISTORY OF SINGLETON SPONTANEOUS PRETERM BIRTH |
| ODE-8 | TREATMENT OF SEVERE HYPERCALCEMIA IN PATIENTS WITH PRIMARY HYPERPARATHYROIDISM WHO ARE UNABLE TO UNDERGO PARATHYROIDECTOMY |
| ODE-9 | TREATMENT OF ASYMPTOMATIC OR PROGRESSIVE MEDULLARY THYROID CANCER IN PATIENTS WITH UNRESECTABLE LOCALLY ADVANCED OR METASTATIC DISEASE |
| ODE-10 | FOR USE IN COMBINATION CHEMOTHERAPY WITH 5-FLUOROURACIL IN THE PALLIATIVE TREATMENT OF PATIENTS WITH ADVANCED METASTATIC COLORECTAL CANCER |
| ODE-11 | TREATMENT OF PROGRESSIVE NEUROENDOCRINE TUMORS OF PANCREATIC ORIGIN (PNET) IN PATIENTS WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC DISEASE |
| ODE-12 | TREATMENT OF PERIPHERAL T-CELL LYMPHOMA (PTCL) IN PATIENTS WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY |
| ODE-13 | TREATMENT OF UNRESECTABLE OR METASTATIC MELANOMA WITH THE BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST |
| ODE-14 | TREATMENT OF ACUTE ATTACKS OF HEREDITARY ANGIOEDEMA IN ADULTS 18 YEARS OF AGE AND OLDER |

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-15 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) THAT IS ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE AS DETECTED BY AN FDA APPROVED TEST
- ODE-16 TREATMENT OF PATIENTS WITH TRANSFUSIONAL IRON OVERLOAD DUE TO THALASSEMIA SYNDROMES WHEN CURRENT CHELATION THERAPY IS INADEQUATE
- ODE-17 ADJUNCTIVE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME IN PATIENTS 2 YEARS OF AGE OR OLDER
- ODE-18 ADJUNCTIVE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME INPATIENTS 2 YEARS OF AGE OR OLDER
- ODE-19 TREATMENT OF PATIENTS WITH INTERMEDIATE OR HIGH-RISK MYELOFIBROSIS, INCLUDING PRIMARY MYELOFIBROSIS, POST-POLYCYTHEMIA VERA MYELOFIBROSIS AND POST-ESSENTIAL THROMBOCYTHEMIA MYELOFIBROSIS
- ODE-20 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE A G551D MUTATION IN THE CFTR GENE.
- ODE-21 AN ADJUNCT TO AB EXTERNO GLAUCOMA SURGERY
- ODE-22 FOR THE CONTROL OF HYPERGLYCEMIA SECONDARY TO HYPERCORTISOLISM IN ADULT PATIENTS WITH ENDOGENOUS CUSHING'S SYNDROME WHO HAVE TYPE 2 DIABETES MELLITUS OR GLUCOSE INTOLERANCE AND HAVE FAILED SURGERY OR ARE NOT CANDIDATES FOR SURGERY
- ODE-23 ADVANCED SOFT TISSUE SARCOMA (STS) WHO HAVE RECEIVED PRIOR CHEMOTHERAPY
- ODE-24 TREATMENT OF ADULTS WITH RENAL ANGIOMYOLIPOMA AND TUBEROUS SCLEROSIS COMPLEX (TSC) NOT REQUIRING IMMEDIATE SURGERY
- ODE-25 MANAGEMENT OF POSTHERPETIC NEURALGIA IN ADULTS.
- ODE-26 TREATMENT OF ENDOGENOUS ANTERIOR UVEITIS
- ODE-27 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES INCLUDING BORTEZOMIB AND AN IMMUNOMODULATORY AGENT AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY
- ODE-28 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME-NEGATIVE (PH-) ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN SECOND OR GREATER RELAPSE OR WHOSE DISEASE HAS PROGRESSED FOLLOWING TWO OR MORE ANTI-LEUKEMIA THERAPIES
- ODE-29 LOCALIZATION OF LYMPH NODES DRAINING A PRIMARY TUMOR IN PATIENTS WITH MELANOMA WHEN USED WITH A HAND-HELD GAMMA COUNTER
- ODE-30 TREATMENT OF ADULT PATIENTS WITH CHRONIC, ACCELERATED OR BLAST PHASE PHILADELPHIA CHROMOSOME-POSITIVE (PH+) CHRONIC MYELOGENOUS LEUKEMIA (CML) WITH RESISTANCE, OR INTOLERANCE TO PRIOR THERAPY
- ODE-31 TREATMENT OF CORNEAL CYSTINE CRYSTAL ACCUMULATION IN PATIENTS WITH CYSTINOSIS
- ODE-32 TREATMENT OF ADULT PATIENTS WITH CHRONIC OR ACCELERATED PHASE CHRONIC MYELOID LEUKEMIA (CML) WITH RESISTANCE AND/OR INTOLERANCE TO TWO OR MORE TYROSINE KINASE INHIBITORS (TKI)
- ODE-33 TREATMENT OF PROGRESSIVE, METASTATIC MEDULLARY THYROID CANCER (MTC)
- ODE-34 TREATMENT OF ADULT PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- ODE-35 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (PH+ALL) THAT IS RESISTANT OR INTOLERANT TO PRIOR TYROSINE KINASE INHIBITOR THERAPY.
- ODE-36 ADJUNCT TO A LOW-FAT DIET AND OTHER LIPID-LOWERING TREATMENTS, INCLUDING LDL APHERESIS WHERE AVAILABLE, TO REDUCE LDL-C, TC, APOLIPOPROTEIN B, & NON-HDL-C IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- ODE-37 TREATMENT OF ADULT PATIENTS WITH SHORT BOWEL SYNDROME (SBS) WHO ARE DEPENDENT ON PARENTERAL SUPPORT
- ODE-38 PART OF COMBINATION THERAPY IN ADULTS (GREATER THAN OR EQUAL TO 18 YEARS) WITH PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS (MDR-TB)
- ODE-39 TREATMENT OF CHRONIC IRON OVERLOAD IN PATIENTS 10 YRS. & OLDER WITH NON-TRANSFUSION DEPENDENT THALASSEMIA (NTDT) SYNDROMES AND WITH A LIVER IRON CONCENTRATION OF AT LEAST 5 MG OF IRON PER GRAM OF LIVER DRY WEIGHT & SERUM FERRITIN GREATER THAN 300 MCG/L.
- ODE-40 TREATMENT OF PEDIATRIC PATIENTS WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (PH+ ALL) IN COMBINATION WITH

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

CHEMOTHERAPY, APPROVED UNDER NDA #21588/S-037

- ODE-41 ADJUNCT TO LIPID-LOWERING MEDICATIONS AND DIET TO REDUCE LDL-C, APOLIPOPROTEIN B (APO B), TOTAL CHOLESTEROL (TC), AND NON-HIGH DENSITY LIPOPROTEIN-CHOLESTEROL (NON-HDL-C) IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFH)
- ODE-42 USE AS A NITROGEN-BINDING ADJUNCTIVE THERAPY FOR CHRONIC MGMT OF ADULT AND PEDIATRIC PATIENTS AT LEAST 2 YRS WITH UREA CYCLE DISORDERS THAT CANNOT BE MANAGED BY DIETARY PROTEIN RESTRICTION AND/OR AMINO ACID SUPPLEMENTATION ALONE
- ODE-43 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES INCLUDING LENALIDOMIDE AND BORTEZOMIB AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY.
- ODE-44 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED, UNRESECTABLE OR METASTATIC GASTROINTESTINAL STROMAL TUMOR (GIST) WHO HAVE BEEN PREVIOUSLY TREATED WITH IMATINIB MESYLATE AND SUNITINIB MALATE.
- ODE-45 MANAGEMENT OF NEPHROPATHIC CYSTINOSIS IN ADULTS AND CHILDREN AGES 6 YEARS AND OLDER.
- ODE-46 IMPROVEMENT OF NEUROLOGICAL OUTCOME BY REDUCING THE INCIDENCE AND SEVERITY OF ISCHEMIC DEFICITS IN ADULT PATIENTS WITH SUBARACHNOID HEMORRHAGE FROM RUPTURED INTRACRANIAL BERRY ANEURYSMS REGARDLESS OF THEIR POST-ICTUS NEUROLOGICAL CONDITION
- ODE-47 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATION AS DETECTED BY AN FDA APPROVED TEST.
- ODE-48 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA APPROVED TEST
- ODE-49 TREATMENT OF MANTLE CELL LYMPHOMA WHOSE DISEASE HAS RELAPSED OR PROGRESSED AFTER TWO PRIOR THERAPIES, ONE OF WHICH INCLUDED BORTEZOMIB
- ODE-50 FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST.
- ODE-51 TOPICAL TREATMENT OF STAGE 1A AND 1B MYCOSIS FUNGOIDES-TYPE CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO HAVE RECEIVED PRIOR SKIN-DIRECTED THERAPY
- ODE-52 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS AS FIRST-LINE TREATMENT, IN COMBINATION WITH GEMCITABINE.
- ODE-53 TREATMENT OF ADULTS WITH PULMONARY ARTERIAL HYPERTENSION (PAH) WHO GROUP 1, TO IMPROVE EXERCISE CAPACITY, WHO FUNCTIONAL CLASS AND TO DELAY CLINICAL WORSENING.
- ODE-54 TX OF PAH TO DELAY DISEASE PROGRESSION. DISEASE PROGRESSION INCLUDED: DEATH, INITIATION OF IV OR SC PROSTANOIDS, OR CLINICAL WORSENING OF PAH (DECREASED 6-MINUTE WALK DISTANCE, WORSENERD PAH SYMPTOMS AND NEED FOR ADDITIONAL PAH TREATMENT).
- ODE-55 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA (MCL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-56 TREATMENT OF PATIENTS WITH LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, DIFFERENTIATED THYROID CARCINOMA (DCT) THAT IS REFRACTORY TO RADIOACTIVE IODINE TREATMENT.
- ODE-57 TRAMETINIB IN COMBO WITH DABRAFENIB FOR TX. OF PTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST. THIS INDICATION IS BASED ON THE DEMONSTRATION OF DURABLE RESPONSE RATE
- ODE-58 DABRAFENIB IN COMBO WITH TRAMETINIB FOR TX. OF PTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST. THIS INDICATION IS BASED ON THE DEMONSTRATION OF DURABLE RESPONSE RATE
- ODE-59 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER
- ODE-60 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-61 TREATMENT OF NEUROGENIC SYMPTOMATIC ORTHOSTATIC HYPOTENSION IN PATIENTS WITH PRIMARY AUTONOMIC FAILURE, DOPAMINE-BETA-HYDROXYLASE DEFICIENCY, AND NONDIABETIC AUTONOMIC NEUROPATHY
- ODE-62 TREATMENT OF PROLIFERATING INFANTILE HEMANGIOMA REQUIRING SYSTEMIC THERAPY.

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

- ODE-63 TREATMENT OF VISCERAL LEISHMANIASIS DUE TO LEISHMANIA DONOVANI; CUTANEOUS LEISHMANIASIS DUE TO LEISHMANIA BRAZILIENSIS, LEISHMANIA GUYANENSIS, AND LEISHMANIA PANAMENSIS; AND MUCOSAL LEISHMANIASIS DUE TO LEISHMANIA BRAZILIENSIS.
- ODE-64 SELECTIVE HEPATIC INTRA-ARTERIAL USE FOR IMAGING TUMORS IN ADULTS WITH KNOWN HEPATOCELLULAR CARCINOMA (HCC)
- ODE-65 TREATMENT OF PATIENTS WITH ACUTE LYMPHOBLASTIC LEUKEMIA AS PART OF A COMBINATION REGIMEN.
- ODE-66 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB.
- ODE-67 GUIDING SENTINEL LYMPH NODE BIOPSY, USING A HAND-HELD GAMMA COUNTER IN PATIENTS WITH CLINICALLY NODE NEGATIVE SQUAMOUS CELL CARCINOMA OF THE ORAL CAVITY
- ODE-68 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA
- ODE-69 TREATMENT OF MALIGNANT HYPERTHERMIA IN CONJUNCTION WITH APPROPRIATE SUPPORTIVE MEASURES AND FOR THE PREVENTION OF MALIGNANT HYPERTHERMIA IN PATIENTS AT HIGH RISK
- ODE-70 RELAPSED CLL, IN COMBO. WITH RITUXIMAB, IN PATIENTS FOR WHOM RITUXIMAB ALONE WOULD BE CONSIDERED APPROPRIATE THERAPY DUE TO OTHER CO-MORBIDITIES; AND RELAPSED SLL IN PATIENTS WHO HAVE RECEIVED AT LEAST 2 PRIOR SYSTEMIC THERAPIES
- ODE-71 RELAPSED FOLLICULAR B-CELL NON-HODGKIN LYMPHOMA (FL) IN PATIENTS WHO HAVE RECEIVED AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- ODE-72 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA WITH 17P DELETION WHO HAVE NOT RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-73 LONG-TERM TREATMENT OF ADULT PATIENTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 EXTENSIVE METABOLIZERS (EMS), INTERMEDIATE METABOLIZERS (IMS), OR POOR METABOLIZERS (PMS) AS DETECTED BY AN FDA-CLEARED TEST.
- ODE-74 TREATMENT OF PATIENTS WITH SEVERE APLASTIC ANEMIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO IMMUNOSUPPRESSIVE THERAPY
- ODE-75 TREATMENT OF PATIENTS WITH SEVERE APLASTIC ANEMIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO IMMUNOSUPPRESSIVE THERAPY.
- ODE-76 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE NOT RECEIVED AT LEAST 1 PRIOR THERAPY
- ODE-77 TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS
- ODE-78 TREATMENT OF HYPERCALCEMIA IN ADULT PATIENTS WITH PRIMARY HYPERPARATHYROIDISM FOR WHOM PARATHYROIDECTOMY WOULD BE INDICATED ON THE BASIS OF SERUM CALCIUM LEVELS, BUT WHO ARE UNABLE TO UNDERGO PARATHYROIDECTOMY.
- ODE-79 TREATMENT OF PATIENTS WITH POLYCYTHEMIA VERA WHO HAVE HAD AN INADEQUATE RESPONSE TO OR ARE INTOLERANT OF HYDROXYUREA
- ODE-80 TREATMENT OF PEDIATRIC PATIENTS WITH TOURETTE'S
- ODE-81 TREATMENT OF PATIENTS WITH ACROMEGALY WHO HAVE HAD AN INADEQUATE RESPONSE TO SURGERY AND/OR FOR WHOM SURGERY IS NOT AN OPTION
- ODE-82 TREATMENT OF PATIENTS WITH UNRESECTABLE, WELL- OR MODERATELY-DIFFERENTIATED LOCALLY ADVANCED OR METASTATIC GASTROENTEROPANCREATIC NEUROENDOCRINE TUMORS TO IMPROVE PROGRESSION-FREE SURVIVAL
- ODE-83 USE OF AS MONOTHERAPY FOR PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA MUTATED (AS DETECTED BY AN FDA-APPROVED TEST) ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH THREE OR MORE PRIOR LINES OF CHEMOTHERAPY
- ODE-84 TREATMENT OF MOTOR FLUCTUATIONS IN PATIENTS WITH ADVANCED PARKINSON'S DISEASE
- ODE-85 AS A REPLACEMENT SOLUTION IN CONTINUOUS RENAL REPLACEMENT THERAPY (CRRT) AND IN CASE OF DRUG POISONING WHEN CRRT IS USED TO REMOVE DIALYZABLE SUBSTANCES
- ODE-86 TREATMENT OF PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA
- ODE-87 TREATMENT OF PATIENTS WITH LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, RADIOACTIVE IODINE REFRACTORY DIFFERENTIATED THYROID CANCER
- ODE-88 FOR USE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE NOT RECEIVED AT LEAST ONE PRIOR THERAPY (FIRST LINE TREATMENT)
- ODE-89 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST 2 PRIOR REGIMENS, INCLUDING BORTEZOMIB AND AN IMMUNOMODULATORY AGENT

PATENT AND EXCLUSIVITY TERMS

ADB 46 of 133

ORPHAN DRUG EXCLUSIVITY

- ODE-90 TREATMENT OF INVASIVE MUCORMYCOSIS IN PATIENTS 18 YEARS OF AGE AND OLDER
- ODE-91 TREATMENT OF BILE ACID SYNTHESIS DISORDERS DUE TO SINGLE ENZYME DEFECTS
- ODE-92 TREATMENT OF LYMPHANGIOLEIOMYOMATOSIS (LAM)
- ODE-93 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 12 YEARS AND OLDER WHO ARE HOMOZYGOUS FOR F508DEL MUTATION IN THE CFTR GENE
- ODE-94 PROPHYLAXIS OF ORGAN REJECTION IN KIDNEY TRANSPLANT PATIENTS CONVERTED FROM TACROLIMUS IMMEDIATE-RELEASE FORMULATIONS IN COMBINATION WITH OTHER IMMUNOSUPPRESSANTS
- ODE-95 FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- ODE-96 TREATMENT OF PRIMARY HYPERKALEMIC PERIODIC PARALYSIS, PRIMARY HYPOKALEMIC PERIOD PARALYSIS, AND RELATED VARIANTS
- ODE-97 TO EXPAND THE INDICATION TO PEDIATRIC PATIENTS 2-6 YEARS OF AGE WITH NEPHROPATHIC CYSTINOSIS
- ODE-98 TREATMENT OF HEREDITARY OROTIC ACIDURIA
- ODE-99 FOR USE IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN, FOR THE TREATMENT OF PATIENTS WITH METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED FOLLOWING GEMCITABINE-BASED THERAPY
- ODE-100 FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA OR LEIOMYOSARCOMA WHO RECEIVED A PRIOR ANTHRACYCLINE-CONTAINING REGIMEN
- ODE-101 FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATION, IN COMBINATION WITH VEMURAFENIB. COTELLIC IS NOT INDICATED FOR TREATMENT OF PATIENTS WITH WILD-TYPE BRAF MELANOMA
- ODE-102 FOR TREATMENT OF PATIENTS WITH METASTATIC EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) T790M MUTATION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC), AS DETECTED BY AN FDA-APPROVED TEST, WHO HAVE PROGRESSED ON OR AFTER EGFR TKI THERAPY
- ODE-103 USE IN COMBINATION WITH LENALIDOMIDE AND DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-104 EMERGENCY TX OF PTS FOLLOWING A FU OR CAPECITABINE OD, OR WHO EXHIBIT EARLY-ONSET, SEVERE OR LIFE-THREATENING TOXICITY AFFECTING THE CARDIAC SYSTEM OR CNS, AND/OR EARLY-ONSET, UNUSUALLY SEVERE AR W/IN 96 HRS FOLLOWING THE END OF FU OR CAPECITABINE ADMIN.
- ODE-105 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- ODE-106 FOR USE OF UPTRAVI (SELEXIPAG) TABLETS, 200, 400, 600, 800, 1000, 1200, 1400, AND 1600 MCG FOR TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH, WHO GROUP I) TO REDUCE THE RISKS OF DISEASE PROGRESSION AND HOSPITALIZATION FOR PAH
- ODE-107 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA WHO HAVE RECEIVED A PRIOR ANTHRACYCLINE-CONTAINING REGIMEN
- ODE-108 TREATMENT OF ADULT PATIENTS WITH PROGRESSIVE, WELL-DIFFERENTIATED, NON-FUNCTIONAL, NEUROENDOCRINE TUMORS (NET) OF GASTROINTESTINAL (GI) OR LUNG ORIGIN, (EXCLUDING PANCREATIC) WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC DISEASE
- ODE-109 INDICATED FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA WITHOUT 17P DELETION WHO HAVE NOT RECEIVED AT LEAST ONE PRIOR THERAPY (FIRST LINE THERAPY)
- ODE-110 FOR HIGH-DOSE CONDITIONING TREATMENT PRIOR TO HEMATOPOIETIC PROGENITOR (STEM) CELL TRANSPLANTATION IN PATIENTS WITH MULTIPLE MYELOMA
- ODE-111 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WHOSE TUMORS ARE ROS-1 POSITIVE.
- ODE-112 FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH HEPATIC VENO-OCCLUSIVE DISEASE (VOD), ALSO KNOWN AS SINUSOIDAL OBSTRUCTION SYNDROME (SOS), WITH RENAL OR PULMONARY DYSFUNCTION FOLLOWING HEMATOPOIETIC STEM CELL TRANSPLANTATION (HSCT).
- ODE-113 FOR TREATMENT OF PEDIATRIC AND ADULT PATIENTS WITH ACQUIRED METHEMOGLOBINEMIA.
- ODE-114 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WITH 17P DELETION, AS DETECTED BY AN FDA APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

THERAPY

- ODE-115 TREATMENT OF PATIENTS WITH METASTATIC, SQUAMOUS, NON-SMALL CELL LUNG CANCER PROGRESSING AFTER PLATINUM-BASED CHEMOTHERAPY
- ODE-116 TREATMENT OF PROGRESSIVE KERATOCONUS
- ODE-117 FOR TREATMENT OF PATIENTS WITH SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- ODE-118 AN ADJUNCT TO DIET TO REDUCE LDL-C, TOTAL-C, NONHDL-C AND APOB IN CHILDREN AND ADOLESCENTS 7 TO 17 YEARS OF AGE WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA, EITHER ALONE OR WITH OTHER LIPID-LOWERING TREATMENTS (E.G., LDL APHERESIS)
- ODE-119 TREATMENT OF PRIMARY BILIARY CHOLANGITIS (PBC) IN COMBINATION WITH URSODEOXYCHOLIC ACID (UDCA) IN ADULTS WITH AN INADEQUATE RESPONSE TO UDCA, OR AS MONOTHERAPY IN ADULTS UNABLE TO TOLERATE UDCA
- ODE-120 FOR USE AFTER RADIOLABELING WITH GA 68, WITH POSITRON EMISSION TOMOGRAPHY (PET) FOR LOCALIZATION OF SOMATOSTATIN RECEPTOR POSITIVE NEUROENDOCRINE TUMORS (NETS) IN ADULT AND PEDIATRIC PATIENTS.
- ODE-121 TREATMENT OF CORNEAL ECTASIA FOLLOWING REFRACTIVE SURGERY
- ODE-122 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- ODE-123 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6-11 YEAR OLD WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE
- ODE-124 REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE FORMULATIONS, WHEN ORAL ADMINISTRATION IS TEMPORARILY NOT FEASIBLE, IN ADULTS WITH THE FOLLOWING SEIZURE TYPES: PARTIAL WITH COMPLEX SYMPTOMOLOGY, GENERALIZED CLONIC-TONIC, AND MIXED
- ODE-125 INDICATED IN PEDIATRIC PATIENTS 10 YEARS AND OLDER FOR THE PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGES 3 AND 4 AND CKD STAGE 5 IN PATIENTS ON HEMODIALYSIS OR PERITONEAL DIALYSIS
- ODE-126 AS MONOTHERAPY FOR THE TREATMENT OF PATIENTS WITH DELETERIOUS BRCA MUTATION (GERMLINE AND/OR SOMATIC) ASSOCIATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH TWO OR MORE CHEMOTHERAPIES
- ODE-127 TREATMENT OF SPINAL MUSCULAR ATROPHY IN PEDIATRIC AND ADULT PATIENTS
- ODE-128 TREATMENT OF PATIENTS WITH MARGINAL ZONE LYMPHOMA (MZL) WHO REQUIRE SYSTEMIC THERAPY AND HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED THERAPY
- ODE-129 INDICATED FOR REDUCING THE RISK OF GRAFT REJECTION WHEN USED IN CONJUNCTION WITH HIGH-DOSE BUSULFAN & CYCLOPHOSPHAMIDE AS A PREPARATIVE REGIMEN FOR ALLOGENIC HEMATOPOIETIC PROGENITOR CELL TRANSPLANTATION FOR PEDIATRIC PATIENTS WITH CLASS 3 BETA-THALASSEMIA
- ODE-130 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY IN PATIENTS 5 YEARS OF AGE AND OLDER
- ODE-131 TREATMENT OF MULTIPLE MYELOMA (MM), AS MAINTENANCE FOLLOWING AUTOLOGOUS HEMATOPOIETIC STEM CELL TRANSPLANTATION (AUTO-HSCT)
- ODE-132 TREATMENT OF CARCINOID SYNDROME DIARRHEA IN COMBINATION WITH SOMATOSTATIN ANALOG (SSA) THERAPY IN ADULTS INADEQUATELY CONTROLLED BY SSA THERAPY
- ODE-133 INDICATED FOR MAINTENANCE TREATMENT OF ADULT PATIENTS WITH RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- ODE-134 TREATMENT OF CHOREA ASSOCIATED WITH HUNTINGTON'S DISEASE
- ODE-135 TREATMENT OF CHRONIC HCV GENOTYPE 2 OR 3 INFECTION IN PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER OR WEIGHING AT LEAST 35 KG WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-136 TREATMENT OF PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER OR WEIGHING AT LEAST 35 KG WITH CHRONIC HEPATITIS C VIRUS GENOTYPE 1, 4, 5, OR 6 INFECTION WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS
- ODE-137 TREATMENT OF OLIGOARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PERSISTENT OLIGOARTHRITIS, PSORIATIC JUVENILE IDIOPATHIC ARTHRITIS, ENTHESITIS-RELATED ARTHRITIS, OR UNDIFFERENTIATED ARTHRITIS) & POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS IN CHILDREN 0-16 YRS
- ODE-138 TREATMENT OF PEDIATRIC PATIENTS WITH ACUTE LYMPHOBLASTIC LEUKEMIA AS A COMPONENT OF A COMBINATION CHEMOTHERAPY MAINTENANCE REGIMEN
- ODE-139 TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA (HCC OR LIVER CANCER) WHO

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

HAVE BEEN PREVIOUSLY TREATED WITH THE DRUG SORAFENIB.

- ODE-140 TREATMENT OF ADULT PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM), SYSTEMIC MASTOCYTOSIS WITH ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN), OR MAST CELL LEUKEMIA (MCL)
- ODE-141 TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) THAT IS FLT3 MUTATION-POSITIVE AS DETECTED BY AN FDA APPROVED TEST, IN COMBINATION WITH STANDARD CYTARABINE AND DAUNORUBICIN INDUCTION AND CYTARABINE CONSOLIDATION
- ODE-142 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- ODE-143 TO DECREASE THE RECURRENCE OF PNEUMOTHORAX IN ADULTS
- ODE-144 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS)
- ODE-145 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS ARE ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE AS DETECTED BY AN FDA-APPROVED TEST
- ODE-146 OPTICAL IMAGING AGENT INDICATED IN PATIENTS WITH GLIOMA (SUSPECTED WORLD HEALTH ORGANIZATION GRADES III OR IV ON PREOPERATIVE IMAGING) AS AN ADJUNCT FOR THE VISUALIZATION OF MALIGNANT TISSUE DURING SURGERY
- ODE-147 DABRAFENIB IN COMBINATION WITH TRAMETINIB, FOR THE TX. OF PTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-148 TRAMETINIB IN COMBINATION WITH DABRAFENIB, FOR THE TX. OF PTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-149 TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH CENTRAL PRECOCIOUS PUBERTY
- ODE-150 TO REDUCE THE ACUTE COMPLICATIONS OF SICKLE CELL DISEASE IN ADULT AND PEDIATRIC PATIENTS 5 YEARS OF AGE AND OLDER.
- ODE-151 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA WITH AN ISOCITRATE DEHYDROGENASE-2 (IDH2) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-152 TREATMENT OF ADULT PATIENTS WITH CHRONIC GRAFT VERSUS HOST DISEASE (CGVHD)
- ODE-153 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- ODE-154 FOR USE IN CHILDREN AGES 2 TO 12 YEARS OLD WITH CHAGAS DISEASE
- ODE-155 TREATMENT OF ADULT PATIENTS WITH RELAPSED FOLLICULAR LYMPHOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- ODE-156 TREATMENT OF ADULTS WITH CARCINOID SYNDROME; WHEN USED, IT REDUCES THE FREQUENCY OF SHORT-ACTING SOMATOSTATIN ANALOG RESCUE THERAPY
- ODE-157 FOR USE AS A NITROGEN-BINDING AGENT FOR CHRONIC MANAGEMENT OF PEDIATRIC PATIENTS >=2 MONTHS AND < 2 YEARS OF AGE WITH UREA CYCLE DISORDERS (UCDS) WHO CANNOT BE MANAGED BY DIETARY PROTEIN RESTRICTION AND/OR AMINO ACID SUPPLEMENTATION ALONE
- ODE-158 TREATMENT OF PATIENTS WITH ERDHEIM-CHESTER DISEASE WITH BRAF V600 MUTATION
- ODE-159 FOR TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK) POSITIVE, METASTATIC NON-SMALL-CELL LUNG CANCER (NSCLC) AS DETECTED BY AN FDA APPROVED TEST, EXCLUDING PATIENTS WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- ODE-160 FOR TREATMENT OF SCURVY IN ADULT AND PEDIATRIC PATIENTS AGE 5 MONTHS AND OLDER FOR WHOM ORAL ADMINISTRATION IS NOT POSSIBLE, INSUFFICIENT OR CONTRAINDICATED
- ODE-161 TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1) IN PEDIATRIC PATIENTS AGED 3 YRS AND OLDER WITH IDIOPATHIC OR CONGENITAL PAH TO IMPROVE PULMONARY VASCULAR RESISTANCE (PVR), WHICH IS EXPECTED TO RESULT IN AN IMPROVEMENT IN EXERCISE ABILITY
- ODE-162 TREATMENT OF NEPHROPATHIC CYSTINOSIS IN PEDIATRIC PATIENTS 1 YEAR OF AGE TO LESS THAN 2 YEARS OF AGE
- ODE-163 TREATMENT OF ADULT PATIENTS WITH NEWLY-DIAGNOSED CHRONIC PHASE (CP) PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOGENOUS LEUKEMIA (PH+ CML)
- ODE-164 TREATMENT OF PEDIATRIC PATIENTS WITH PHILADELPHIA CHROMOSOME-POSITIVE (PH+)

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

CHRONIC MYELOID LEUKEMIA (CML) IN CHRONIC PHASE

- ODE-165 PROPHYLAXIS OF CYTOMEGALOVIRUS (CMV) INFECTION AND DISEASE IN ADULT CMV-SEROPOSITIVE RECIPIENTS [R+] OF AN ALLOGENEIC HEMATOPOIETIC STEM CELL TRANSPLANT (HSCT)
- ODE-166 TREATMENT OF SOMATOSTATIN RECEPTOR-POSITIVE GASTROENTEROPANCREATIC NEUROENDOCRINE TUMORS (GEP-NETS) INCLUDING FOREGUT, MIDGUT, AND HINDGUT NEUROENDOCRINE TUMORS IN ADULTS
- ODE-167 ARSENIC TRIOXIDE FOR USE IN COMBINATION WITH TRETINOIN FOR TREATMENT OF ADULTS WITH NEWLY-DIAGNOSED LOW-RISK ACUTE PROMYELOCYTIC LEUKEMIA (APL) WHOSE APL IS CHARACTERIZED BY THE PRESENCE OF THE T(15;17) TRANSLOCATION OR PML/RAR-ALPHA GENE EXPRESSION
- ODE-168 FOR THE MAINTENANCE TREATMENT OF ADULT PATIENTS WITH RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- ODE-169 FOR THE ADJUNCTIVE TREATMENT OF ADULT AND PEDIATRIC PATIENTS AGED 2 YEARS AND OLDER WITH TUBEROUS SCLEROSIS COMPLEX (TSC)-ASSOCIATED PARTIAL-ONSET SEIZURES
- ODE-170 FOR THE DIAGNOSIS OF ADULT GROWTH HORMONE DEFICIENCY (AGHD)
- ODE-171 TREATMENT OF PEDIATRIC PATIENTS GREATER THAN OR EQUAL TO 1 YEAR OF AGE WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH+CML) IN CHRONIC PHASE
- ODE-172 TREATMENT OF PEDIATRIC PATIENTS GREATER THAN OR EQUAL TO 1 YEAR OF AGE WITH CHRONIC PHASE PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA WITH RESISTANCE OR INTOLERANCE TO PRIOR TYROSINE-KINASE INHIBITOR THERAPY
- ODE-173 TREATMENT OF PATIENTS WITH CYSTIC FIBROSIS AGED 12 YEARS AND OLDER WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR WHO HAVE AT LEAST ONE MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR GENE RESPONSIVE TO TEZACAFTOR/IVACAFTOR
- ODE-174 FOR THE TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC IMMUNE THROMBOCYTOPENIA (ITP) WHO HAVE HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT
- ODE-175 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA (MCL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-176 INDICATED FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 L858R MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST
- ODE-177 TO REDUCE THE FREQUENCY OF PAINFUL CRISES AND TO REDUCE THE NEED FOR BLOOD TRANSFUSIONS IN PEDIATRIC PATIENTS, 2 YEARS OF AGE AND OLDER, WITH SICKLE CELL ANEMIA WITH RECURRENT MODERATE TO SEVERE PAINFUL CRISIS
- ODE-178 INDICATED TO SLOW KIDNEY FUNCTION DECLINE IN ADULTS AT RISK OF RAPIDLY PROGRESSING AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE (ADPKD)
- ODE-179 TREATMENT OF PATIENTS WITH CLL AND TREATMENT OF PATIENTS WITH INDOLENT B-CELL NHL THAT HAS PROGRESSED DURING OR WITHIN SIX MONTHS OF TREATMENT WITH RITUXIMAB OR A RITUXIMAB CONTAINING REGIMEN
- ODE-180 MAINTENANCE TREATMENT OF ADULT PATIENTS WITH RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER, WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- ODE-181 TREATMENT OF ADULT PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH THREE OR MORE PRIOR LINES OF CHEMOTHERAPY
- ODE-182 TRAMETINIB IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- ODE-183 TRAMETINIB AND DABRAFENIB IN COMBINATION, FOR THE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ANAPLASTIC THYROID CANCER (ATC) WITH BRAF V600E MUTATION AND WITH NO SATISFACTORY LOCOREGIONAL TREATMENT OPTIONS
- ODE-184 INDICATED IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF PEDIATRIC PATIENTS WITH HIV-1 INFECTION
- ODE-185 INDICATED FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)

PATENT AND EXCLUSIVITY TERMS**ORPHAN DRUG EXCLUSIVITY**

OR SMALL LYMPHOCYTIC LYMPHOMA (SLL), WITH OR WITHOUT 17P DELETION, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY

- ODE-186 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE ONE OF THE FOLLOWING MUTATIONS IN THE CFTR GENE: G1244E, G1349D, G178R, G551S, S1251N, S1255P, S549N, OR S549R
- ODE-187 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE AN R117H MUTATION IN THE CFTR GENE
- ODE-188 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGES 2 TO LESS THAN 6 YEARS WHO HAVE ONE OF THE FOLLOWING MUTATIONS IN THE CFTR GENE: G551D, G1244E, G1349D, G178R, G551S, S1251N, S1255P, S549N, S549R, AND R117H
- ODE-189 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 2 YEARS AND OLDER WHO HAVE ONE OF THE FOLLOWING MUTATIONS IN THE CFTR GENE: 711+3A-G, E831X, 2789+5G-A, 3272-26A-G, AND 3849+10KBC-T
- ODE-190 TX OF CF IN PTS 2 YRS AND OLDER WHO HAVE ONE OF THE FOLLOWING MUTATIONS IN THE CFTR GENE: E56K, P67L, R74W, D110E, D110H, R117C, E193K, L206W, R347H, R352Q, A455E, D579G, S945L, S977F, F1052V, K1060T, A1067T, G1069R, R1070Q, R1070W, F1074L, D1152H, D1270N
- ODE-191 TO DECREASE THE RECURRENCE OF MALIGNANT PLEURAL EFFUSIONS IN SYMPTOMATIC PATIENTS FOLLOWING MAXIMAL DRAINAGE OF THE PLEURAL EFFUSION
- ODE-192 INDICATED TO INDUCE CONTROLLED CARDIAC SEPTAL INFRACTION TO IMPROVE EXERCISE CAPACITY IN ADULTS WITH SYMPTOMATIC HYPERTROPHIC OBSTRUCTIVE CARDIOMYOPATHY WHO ARE NOT CANDIDATES FOR SURGICAL MYECTOMY
- ODE-193 INDICATED FOR THE TREATMENT OF ONCHOCERCIASIS DUE TO ONCHOCERCA VOLVULUS IN PATIENTS AGED 12 YEARS AND OLDER
- ODE-194 ENCORAFENIB IS INDICATED IN COMBINATION WITH BINIMETINIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH A BRAF V600E OR V600K MUTATION, AS DETECTED BY AN FDA-APPROVED TEST
- ODE-195 FOR THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 2 THROUGH 5 YEARS OLD WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE
- ODE-196 INDICATED FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH UNRESECTABLE HEPATOCELLULAR CARCINOMA (HCC)
- ODE-197 INDICATED FOR THE TREATMENT OF THE POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS IN ADULTS
- ODE-198 INDICATED FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH DRAVET SYNDROME (DS) IN PATIENTS 2 YEARS OF AGE AND OLDER TAKING CLOBAZAM
- ODE-199 THE TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 12 MONTHS AND OLDER WHO HAVE ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR POTENTIATION BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA
- ODE-200 INDICATED FOR THE TREATMENT OF HUMAN SMALLPOX DISEASE CAUSED BY VARIOLA VIRUS IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 13 KG
- ODE-201 INDICATED FOR THE RADICAL CURE (PREVENTION OF RELAPSE) OF PLASMODIUM VIVAX MALARIA IN PATIENTS AGED 16 YEARS AND OLDER WHO ARE RECEIVING APPROPRIATE ANTIMALARIAL THERAPY FOR ACUTE P. VIVAX INFECTION
- ODE-202 INDICATED AS A SOURCE OF CALORIES AND FATTY ACIDS IN PEDIATRIC PATIENTS WITH PARENTERAL NUTRITION-ASSOCIATED CHOLESTASIS (PNAC)
- ODE-203 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) WITH A SUSCEPTIBLE ISOCITRATE DEHYDROGENASE-1 (IDH1) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-204 TREATMENT OF ADULT AND PEDIATRIC PATIENTS 12 YEARS AND OLDER WITH IOBENGUANE SCAN POSITIVE, UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC PHEOCHROMOCYTOMA OR PARAGANGLIOMA WHO REQUIRE SYSTEMIC ANTICANCER THERAPY
- ODE-205 INDICATED FOR THE TREATMENT OF ADULTS WITH A CONFIRMED DIAGNOSIS OF FABRY DISEASE AND AN AMENABLE GALACTOSIDASE ALPHA GENE (GLA) VARIANT BASED ON IN VITRO ASSAY DATA
- ODE-206 FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETION OR EXON 21 L858R SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- ODE-207 TREATMENT OF STATUS EPILEPTICUS IN ADULTS
- ODE-208 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY CHRONIC LYMPHOCYTIC

PATENT AND EXCLUSIVITY TERMS

ADB 51 of 133

ORPHAN DRUG EXCLUSIVITY

- LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) AFTER AT LEAST TWO PRIOR THERAPIES
- ODE-209 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY FOLLICULAR LYMPHOMA (FL) AFTER AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- ODE-210 INDICATED IN COMBINATION WITH STANDARD IMMUNOSUPPRESSIVE THERAPY FOR THE FIRST-LINE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 2 YEARS AND OLDER WITH SEVERE APLASTIC ANEMIA
- ODE-211 INDICATED IN COMBO WITH AZACITIDINE, OR DECITABINE, OR LOW-DOSE CYTARABINE FOR THE TX OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA IN ADULTS WHO ARE AGE 75 YEARS OR OLDER, OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY
- ODE-212 INDICATED FOR THE TREATMENT OF THE POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS IN ADULTS
- ODE-213 INDICATED FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH EPIDERMAL GROWTH FACTOR (EGFR) EXON 19 DELETION OR EXON 21 L858R SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- ODE-214 TX OF MAC LUNG DISEASE IN ADULTS WITH LIMITED OR NO ALTERNATIVE TX OPTIONS AS PART OF A COMBO ANTIBACTERIAL DRUG REGIMEN WHO DO NOT ACHIEVE NEGATIVE SPUTUM CULTURES AFTER A MINIMUM OF 6 CONSECUTIVE MONTHS OF A MULTIDRUG BACKGROUND REGIMEN THERAPY
- ODE-215 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH SOLID TUMORS THAT HAVE A NEUROTROPHIC RECEPTOR TYROSINE KINASE (NTRK) GENE FUSION WITHOUT A KNOWN ACQUIRED RESISTANCE MUTATION
- ODE-216 INDICATED FOR THE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME (LGS) OR DRAVET SYNDROME (DS) IN PATIENTS 2 YEARS OF AGE AND OLDER
- ODE-217 INDICATED FOR THE TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)- POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE DISEASE HAS PROGRESSED ON CRIZOTINIB AND AT LEAST ONE OTHER ALK INHIBITOR FOR METASTATIC DIESASE
- ODE-218 INDICATED FOR THE TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)- POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE DISEASE HAS PROGRESSED ON ALECTINIB AS THE FIRST ALK INHIBITOR THERAPY FOR METASTATIC DISEASE
- ODE-219 INDICATED FOR THE TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)- POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE DISEASE HAS PROGRESSED ON CERITINIB AS THE FIRST ALK INHIBITOR THERAPY FOR METASTATIC DISEASE
- ODE-220 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH SOLID TUMORS THAT ARE METASTATIC OR WHERE SURGICAL RESECTION IS LIKELY TO RESULT IN SEVERE MORBIDITY
- ODE-221 INDICATED FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH SOLID TUMORS THAT HAVE NO SATISFACTORY ALTERNATIVE TREATMENTS OR THAT HAVE PROGRESSED FOLLOWING TREATMENT
- ODE-222 INDICATED FOR THE TREATMENT OF ADULT PATIENTS WHO HAVE RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) WITH A FMS-LIKE TYROSINE KINASE 3 (FLT3) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-223 INDICATED FOR THE TREATMENT OF LAMBERT-EATON MYASTHENIC SYNDROME (LEMS) IN ADULTS
- ODE-224 INDICATED, IN COMBINATION WITH LOW-DOSE CYTARABINE, FOR THE TREATMENT OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) IN ADULT PATIENTS WHO ARE ≥ 75 YEARS OLD OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY

PATENT USE

- U-1 PREVENTION OF PREGNANCY
- U-2 TREATMENT OR PROPHYLAXIS OF ANGINA PECTORIS AND ARRHYTHMIA
- U-3 TREATMENT OF HYPERTENSION
- U-4 PROVIDING PREVENTION AND TREATMENT OF EMESIS AND NAUSEA IN MAMMALS
- U-5 METHOD OF PRODUCING BRONCHODILATION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-6 METHOD OF PRODUCING SYMPATHOMIMETIC EFFECTS

U-7 INCREASING CARDIAC CONTRACTILITY

U-8 ACUTE MYOCARDIAL INFARCTION

U-9 CONTROL OF EMESIS ASSOCIATED WITH ANY CANCER CHEMOTHERAPY AGENT

U-10 DIAGNOSTIC METHOD FOR DISTINGUISHING BETWEEN HYPOTHALAMIC MALFUNCTIONS OR LESIONS IN HUMANS

U-11 TREATMENT OR PROPHYLAXIS OF CARDIAC DISORDERS

U-12 METHOD OF TREATING [A] HUMAN SUFFERING FROM DEPRESSION

U-13 A METHOD FOR TREATING ANXIETY IN A HUMAN SUBJECT IN NEED OF SUCH TREATMENT

U-14 ADJUNCTIVE THERAPY FOR THE PREVENTION AND TREATMENT OF HYPERAMMONEMIA IN THE CHRONIC MANAGEMENT OF PATIENTS WITH UREA CYCLE ENZYMOPATHIES

U-15 METHOD OF LOWERING INTRAOCULAR PRESSURE

U-16 USE IN LUNG SCANNING PROCEDURES

U-17 TREATMENT OF VENTRICULAR AND SUPRAVENTRICULAR ARRHYTHMIAS

U-18 METHOD FOR INHIBITING GASTRIC SECRETION IN MAMMALS

U-19 TREATMENT OF INFLAMMATION

U-20 A PROCESS FOR TREATING A PATIENT SUFFERING FROM PARKINSON'S SYNDROME AND IN NEED OF TREATMENT

U-21 TREATMENT OF HUMANS SUFFERING UNDESIRE UROTOXIC SIDE EFFECTS CAUSED BY CYTOSTATICALLY ACTIVE ALKYLATING AGENTS

U-22 METHOD OF COMBATTING PATHOLOGICALLY REDUCED CEREBRAL FUNCTIONS AND PERFORMANCE WEAKNESSES, CEREBRAL INSUFFICIENCY AND DISORDERS IN CEREBRAL CIRCULATION AND METABOLISM IN WARM-BLOODED ANIMALS

U-23 METHOD FOR TREATING PROSTATIC CARCINOMA COMPRISING ADMINISTERING FLUTAMIDE

U-24 METHOD FOR TREATING PROSTATE ADENOCARCINOMA COMPRISING ADMINISTERING AN ANTIANDROGEN INCLUDING FLUTAMIDE AND AN LHRH AGONIST

U-25 REDUCING CHOLESTEROL IN CHOLELITHIASIS PATIENTS

U-26 REDUCING CHOLESTEROL GALLSTONES AND/OR FRAGMENTS THEREOF

U-27 DISSOLVING CHOLESTEROL GALLSTONES AND/OR FRAGMENTS THEREOF

U-28 CEREBRAL, CORONARY, PERIPHERAL, VISCERAL AND RENAL ARTERIOGRAPHY, AORTOGRAPHY AND LEFT VENTRICULOGRAPHY

U-29 CT IMAGING OF THE HEAD AND BODY, AND INTRAVENOUS EXCRETORY UROGRAPHY

U-30 CEREBRAL ANGIOGRAPHY, AND VENOGRAPHY

U-31 INTRA-ARTERIAL DIGITAL SUBTRACTION ANGIOGRAPHY

U-32 PALLIATIVE TREATMENT OF PATIENTS WITH OVARIAN CARCINOMA RECURRENT AFTER PRIOR CHEMOTHERAPY, INCLUDING PATIENTS WHO HAVE BEEN PREVIOUSLY TREATED WITH CISPLATIN

U-33 TREATING VIRAL INFECTIONS IN A MAMMAL

U-34 TREATING VIRAL INFECTIONS IN A WARM-BLOODED ANIMAL

U-35 TREATING CYTOMEGALOVIRUS IN A HUMAN WITH AN INJECTABLE COMPOSITION

U-36 METHODS OF TREATING BACTERIAL ILLNESSES

U-37 METHOD OF TREATING GASTROINTESTINAL DISEASE

U-38 TREATMENT OF PAROXYSMAL SUPRAVENTRICULAR TACHYCARDIA

U-39 ANGINA PECTORIS

U-40 METHOD OF TREATMENT OF BURNS

U-41 METHOD OF TREATING CARDIAC ARRHYTHMIAS

U-42 ADJUVANT TREATMENT IN COMBINATION WITH FLUOROURACIL AFTER SURGICAL RESECTION IN PATIENTS WITH DUKES' STAGE C COLON CANCER

U-43 MANAGEMENT OF CHRONIC PAIN IN PATIENTS REQUIRING OPIOID ANALGESIA

U-44 RELIEF OF NAUSEA AND VOMITING

U-45 TREATMENT OF INFLAMMATION AND ANALGESIA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-46 TREATMENT OF PANIC DISORDER

U-47 STIMULATION OF THE RELEASE OF GROWTH HORMONE

U-48 ANALGESIA

U-49 SYMPTOMATIC CANCER-RELATED HYPERCALCEMIA

U-50 USE IN TREATING INFLAMMATORY DERMATOSES

U-51 BLOOD POOL IMAGING, INCLUDING CARDIAC FIRST PASS AND GATED EQUILIBRIUM IMAGING AND FOR DETECTION OF SITES OF GASTROINTESTINAL BLEEDING

U-52 TREATMENT OF ADULT AND PEDIATRIC PATIENTS (OVER SIX MONTHS OF AGE) WITH ADVANCED HIV INFECTION

U-53 HYPERCALCEMIA OF MALIGNANCY

U-54 REVERSAL AGENT OR ANTAGONIST OF NONDEPOLARIZING NEUROMUSCULAR BLOCKING AGENTS

U-55 TREATMENT OF PAIN

U-56 AID TO SMOKING CESSATION

U-57 OPHTHALMIC USE OF NORFLOXACIN

U-58 METHOD OF TREATING INFLAMMATORY INTESTINAL DISEASES

U-59 METHOD OF TREATING HYPERCHOLESTEROLEMIA

U-60 NASAL ADMINISTRATION OF BUTORPHANOL

U-61 CEREBRAL AND PERIPHERAL ARTERIOGRAPHY AND CT IMAGING OF THE HEAD

U-62 CORONARY ARTERIOGRAPHY, LEFT VENTRICULOGRAPHY, CT IMAGING OF THE BODY, INTRAVENOUS EXCRETORY UROGRAPHY, INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY AND VENOGRAPHY

U-63 ISOPRENALINE ANTAGONISM ON THE HEART RATE OR BLOOD PRESSURE

U-64 TREATMENT OF VIRAL INFECTIONS

U-65 METHOD OF TREATMENT OF A PATIENT INFECTED WITH HIV

U-66 TRIPHASIC REGIMEN

U-67 METHOD OF INDUCING ANESTHESIA IN A WARM BLOODED ANIMAL

U-68 TREATMENT OF ACTINIC KERATOSIS

U-69 TREATMENT OF PNEUMOCYSTIS CARINII INFECTIONS

U-70 TREATMENT OF TRANSIENT INSOMNIA

U-71 METHOD OF TREATMENT OF HEART FAILURE

U-72 TREATMENT OF MIGRAINE

U-73 METHOD OF TREATING DISEASES OR INFECTIONS CAUSED BY MYCETES

U-74 METHOD OF PROVIDING HYPNOTIC EFFECT

U-75 RELIEF OF OCULAR ITCHING DUE TO SEASONAL ALLERGIC CONJUNCTIVITIS

U-76 USE TO IMAGE A SUBJECT WITH A MAGNETIC RESONANCE IMAGING SYSTEM

U-77 TREATMENT OF SYMPTOMS OF SEASONAL ALLERGIC RHINITIS

U-78 ULCERATIVE COLITIS

U-79 SYMPTOMATIC TREATMENT OF PATIENTS WITH NOCTURNAL HEARTBURN DUE TO GERD

U-80 METHOD OF TREATING OCULAR BACTERIAL INFECTIONS

U-81 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS

U-82 TREATMENT FOR DEMENTIA IN PATIENTS WITH ALZHEIMER'S DISEASE

U-83 TREATMENT OF SEIZURES

U-84 A METHOD OF BLOCKING THE UPTAKE OF MONOAMINES BY BRAIN NEURONS IN ANIMALS

U-85 NASAL TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS

U-86 METHOD OF TREATING CERTAIN FORMS OF EPILEPSY

U-87 METHOD FOR NONINVASIVE ADMINISTRATION OF SEDATIVES, ANALGESICS, AND ANESTHETICS

U-88 TREATMENT OF MODERATE PLAQUE PSORIASIS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-89 TREATMENT OR PROPHYLAXIS OF EMESIS

U-90 TREATMENT OF PSYCHOTIC DISORDERS

U-91 ALTERNATIVE THERAPY TO TRIMETHOPRIM-SULFAMETHOXAZOLE FOR TREATMENT OF MODERATE-TO-SEVERE PNEUMOCYSTIS CARINII PNEUMONIA IN IMMUNOCOMPROMISED AND AIDS PATIENTS

U-92 TREATMENT OF DIABETIC NEPHROPATHY IN PATIENTS WITH TYPE I INSULIN DEPENDENT DIABETES MELLITUS AND RETINOPATY

U-93 USE AS AN ANTIHISTAMINE/DECONGESTANT

U-94 TREATMENT-ADULTS W/ ADVANCED HIV, INTOLERANT OF APPROVED THERAPIES, INTOLERANT OF APPROVED THERAPIES W/PROVEN BENEFIT OR HAVE EXPERIENCED CLINICAL/IMMUNOLOGICAL DETERIORATION WHILE RECEIVING..OR FOR WHOM SUCH THERAPIES-CONTRAINDICATED

U-95 SHORT TERM MANAGEMENT OF MODERATE PRURITIS IN ADULTS WITH ATOPIC DERMATITIS AND LICHEN SIMPLEX CHRONICUS

U-96 METHOD OF TREATING VARICELLA ZOSTER (SHINGLES) INFECTIONS

U-97 A METHOD OF TREATING A PATIENT IN NEED OF MEMORY ENHANCEMENT

U-98 A METHOD OF INDUCING REGRESSION OF LEUKEMIA CELL GROWTH IN A MAMMAL

U-99 METHOD OF PROVIDING POTASSIUM TO A SUBJECT IN NEED OF POTASSIUM

U-100 METHOD OF TREATING OCULAR INFLAMMATION

U-101 ADJUNCT TO CONVENTIONAL CT OR MRI IMAGING IN THE LOCALIZATION OF STROKE IN PATIENTS IN WHOM STROKE HAS ALREADY BEEN DIAGNOSED

U-102 METHOD OF HORMONALLY TREATING MENOPAUSAL OR POST-MENOPAUSAL DISORDERS IN WOMEN

U-103 TREATMENT OF OCULAR HYPERTENSION

U-104 TREATMENT OF AQUEOUS HUMOR FORMATION AND INTRAOCULAR PRESSURE

U-105 EMESIS

U-106 TREATMENT OF EPILEPSY

U-107 TREATMENT OF HYPERTENSION AND ANGINA PECTORIS

U-108 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER, GASTROESOPHAGEAL REFLUX DISEASE (GERD), SEVERE EROSIIVE ESOPHAGITIS, POORLY RESPONSIVE SYMPTOMATIC GERD AND PATHOLOGIAL HYPERSECRETORY CONDITIONS AND MAINTENANCE HEALING OF EROSIIVE ESOPHAGITIS

U-109 ADJUNCT DIET IN THE TX OF ELEVATED TOTAL CHOLESTEROL AND LDL-C LEVELS IN PTS W/PRIMARY HYPERCHOLESTEROLEMIA WHOSE RESPONSE TO DIETARY RESTRICTION OF SAT FAT AND CHOLESTEROL AND OTHER NONPHARMACOLOGICAL MEASURES HAS NOT BEEN ADEQUATE

U-110 USE AS A RETRIEVABLE PESSARY

U-111 DIABETES

U-112 CONTRACEPTION

U-113 METHOD OF CONDUCTING RADIOLOGICAL EXAMINATION OF A PATIENT BY ADMINISTERING TO SAID PATIENT A RADIOPAQUE AMOUNT OF IOPROMIDE

U-114 USE FOR INHIBITING BONE RESORPTION

U-115 USE OF VASODILATORS TO EFFECT AND ENHANCE AN ERECTION (AND THUS TREAT ERECTILE DYSFUNCTION), BY INJECTION INTO THE PENIS

U-116 METHOD OF MYOCARDIAL IMAGING

U-117 TREATMENT OF OCULAR ALLERGIC RESPONSE IN HUMAN EYES

U-118 METHOD OF LOWERING BLOOD SUGAR LEVEL

U-119 TREATMENT OF NASAL HYPERSECRETION

U-120 CONTROLLING OR PREVENTING POST-OPERATIVE INTRAOCULAR PRESSURE RISES ASSOCIATED WITH OPHTHALMIC LASER SURGICAL PROCEDURES

U-121 METHOD OF TREATING CONDITIONS MEDIATED THROUGH HISTAMINE H2-RECEPTORS

U-122 A THERAPEUTIC METHOD FOR CONTROLLING THROMBOSIS

U-123 METHOD FOR CONTROLLING THROMBOSIS AND DECREASING BLOOD HYPERCOAGULATION AND HEMORRHAGING RISKS

U-124 TREATMENT OF ACNE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-125 TREATMENT NEUROGENERATIVE DISEASES

U-126 TREATMENT OF GASTRITIS

U-127 METHOD OF PRODUCING NEUROMUSCULAR BLOCKADE

U-128 METHOD FOR TREATMENT OF TUMORS

U-129 METHOD TO DESTROY OR IMPAIR TARGET CELLS

U-130 MANAGEMENT OF PATIENTS WITH MASTOCYTOSIS

U-131 PHOTODAMAGED SKIN

U-132 INHIBITING HIV PROTEASE

U-133 MANAGEMENT OF OBESITY INCLUDING WEIGHT LOSS AND MAINTENANCE IN PATIENTS ON A REDUCED-CALORIE DIET

U-134 TREATMENT OF ACNE VULGARIS

U-135 ANTITUMOR AGENT

U-136 PROCESS FOR WASTE NITROGEN REMOVAL

U-137 METHOD OF TREATING BACTERIAL VAGINOSIS

U-138 TREATMENT OF ALLERGIC RHINITIS

U-139 TREATMENT OF ALLERGIC REACTIONS

U-140 USE OF NORVIR TO INHIBIT HIV PROTEASE OR TO INHIBIT AN HIV INFECTION

U-141 TREATMENT OF ULCERATIVE COLITIS

U-142 METHOD OF TREATING ALLERGIC REACTIONS IN A MAMMAL BY USING THIS ACTIVE METABOLITE

U-143 BIODEGRADABLE SUPERPARAMAGNETIC METAL OXIDES AS CONTRAST AGENTS FOR MR IMAGING

U-144 BIOLOGICALLY DEGRADABLE SUPERPARAMAGNETIC MATERIALS FOR USE IN CLINICAL APPLICATIONS

U-145 BIOLOGICALLY DEGRADABLE SUPERPARAMAGNETIC PARTICLES FOR USE AS NUCLEAR MAGNETIC RESONANCE IMAGING AGENTS

U-146 METHOD OF TREATING SUSCEPTIBLE NEOPLASMS IN MAMMALS

U-147 DETECTION OF GASTROINTESTINAL DISORDERS AND THE SUBSEQUENT BREATH COLLECTION AND MEASUREMENT OF $^{13}\text{CO}_2$

U-148 DEVICE FOR COLLECTING A BREATH SAMPLE

U-149 METHOD OF TREATING AN ANIMAL, INCLUDING A HUMAN SUFFERING FROM OR SUSCEPTIBLE TO PSYCHOSIS, ACUTE MANIA OR MILD ANXIETY STATES

U-150 METHOD OF USE FOR CONTROLLING HYPERGLYCEMIA BY ADMINISTRATION OF THIS SUSTAINED RELEASE DOSAGE FORM OF GLIPIZIDE

U-151 RELIEF OF SYMPTOMS OF THE COMMON COLD

U-152 METHOD OF TREATING ANXIETY RELATED DISORDERS INCLUDING OBSESSIVE COMPULSIVE DISORDER

U-153 TREATMENT OF INITIAL EPISODE GENITAL HERPES

U-154 METHOD OF TREATING ANIMALS SUFFERING FROM AN APPETITE DISORDER

U-155 TREATMENT OF ERECTILE DYSFUNCTION

U-156 METHOD OF PROVIDING ANESTHESIA

U-157 TREATMENT OF A HUMAN SUFFERING FROM VITAMIN B12 DEFICIENCY

U-158 ANGINA

U-159 TREATMENT OF INTERSTITIAL CYSTITIS

U-160 TREATMENT OF BACTERIAL INFECTIOUS DISEASE

U-161 METHOD OF INHIBITING CHOLESTEROL BIOSYNTHESIS IN A PATIENT

U-162 METHOD OF USE TO INHIBIT CHOLESTEROL SYNTHESIS IN A HUMAN SUFFERING FROM HYPERCHOLESTEROLEMIA

U-163 METHOD OF USING TROGLITAZONE TO TREAT IMPAIRED GLUCOSE TOLERANCE TO PREVENT OR DELAY THE ONSET OF NONINSULIN-DEPENDENT DIABETES MELLITUS

U-164 METHOD OF USING TROGLITAZONE TO PREVENT OR DELAY THE ONSET OF NONINSULIN-

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

DEPENDENT DIABETES MELLITUS IN A DEFINED POPULATION OF PATIENTS

U-165 TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA

U-166 TREATMENT OF H.PYLORI-ASSOCIATED DUODENAL ULCER

U-167 METHOD FOR TREATING HIV-1 INFECTION

U-168 METHOD OF INHIBITING LIPOXYGENASE ACTIVITY IN A MAMMAL WHICH IS THE MODE OF ACTION IN THE TREATMENT OF ASTHMA

U-169 METHODS OF USING THE COMPOUND/DRUG PRODUCT AS A CONTRAST AGENT IN MAGNETIC RESONANCE IMAGING

U-170 METHOD OF OBTAINING AN MR IMAGE USING THE COMPOSITION/DRUG PRODUCT AS A CONTRAST AGENT

U-171 METHODS OF USING THE COMPOUND/DRUG PRODUCT AS AN ORAL CONTRAST AGENT IN MAGNETIC RESONANCE IMAGING OF THE GASTROINTESTINAL TRACT

U-172 TREATMENT OF GENITAL WARTS

U-173 ADMINISTRATION TO A HOST SUFFERING FROM GESTATIONAL DIABETES

U-174 USE AS AN ANTIHISTAMINE AGENT

U-175 METHOD OF TREATING MALIGNANT TUMORS

U-176 METHOD OF TREATING A PATIENT SUFFERING FROM LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSES

U-177 FUNGICIDE

U-178 FACILITATED ADHERENCE OF AGENTS TO SKIN

U-179 ENHANCED CUTANEOUS PENETRATION OF A DERMALLY-APPLIED PHARMACOLOGICALLY ACTIVE AGENT

U-180 TREATMENT OF ADULT AND PEDIATRIC PATIENTS (OVER 6 MONTHS OF AGE) WITH ADVANCED HIV INFECTION

U-181 PRODUCING ALPHA ADRENERGIC ANTAGONISTIC ACTION IN A HOST

U-182 USE OF SALMETEROL IN PATIENTS WITH REVERSIBLE AIRWAY OBSTRUCTION

U-183 TREATMENT OF CONDITIONS CAUSED BY DISTURBANCE OF NEURONAL 5HT FUNCTION

U-184 TREATING ALLERGIC EYE DISEASES IN HUMANS

U-185 METHOD OF TREATING HYPERTENSION

U-186 METHOD FOR TREATING GI DISORDERS CAUSED BY H. PYLORI WHICH COMPRISES ADMINISTRATION OF RANITIDINE BISMUTH CITRATE AND CLARITHROMYCIN FOR A GREATER THAN ADDITIVE EFFECT

U-187 THERAPEUTIC TREATMENT OF CALCIFIC TUMORS

U-188 TREATMENT OF H.PYLORI ASSOCIATED DUODENAL ULCER

U-189 ENHANCEMENT OF THE BIOAVAILABILITY OF THE DRUG SUBSTANCE

U-190 USE OF RITONAVIR IN COMBINATION WITH ANY REVERSE TRANSCRIPTASE INHIBITOR

U-191 METHOD OF TREATMENT FOR CONTROLLING AND LOWERING INTRAOCULAR PRESSURE IN A HUMAN

U-192 USE IN TREATING ALLERGIC REACTIONS

U-193 PSORIASIS

U-194 TREATING ANGINA PECTORIS AND HIGH BLOOD PRESSURE

U-195 METHOD FOR THE DIAGNOSIS OF GASTROINTESTINAL DISORDERS BY UREA ISOTOPE OR NITROGEN LABELED CARBON

U-196 TREATMENT OF METASTATIC BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH ESTROGEN RECEPTOR POSITIVE TUMORS

U-197 USE IN COMBINATION WITH CERTAIN LHRH ANALOGUES FOR THE TREATMENT OF ADVANCED PROSTATE CANCER

U-198 TREATMENT METASTATIC CARCINOMA OF OVARY AFTER 1ST LINE FAILURE OR SUBSEQUENT CHEMOTHERAPY, TREATMENT OF BREAST CANCER AFTER FAILURE OF COMBINATION CHEMOTHERAPY FOR METASTATIC DISEASE AND 2ND LINE TREATMENT OF AIDS RELATED KAPOSI'S SARCOMA

U-199 METHOD OF TREATING INFECTIOUS UPPER GI TRACT DISORDERS CAUSED BY CAMPYLOBACTER PYLORIDIS INFECTION COMPRISING ADMINISTRATION OF A BISMUTH AGENT AND AN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

ANTIMICROBIAL AGENT

- U-200 METHOD OF TREATING GI DISORDERS COMPRISING ADMINISTRATION OF A BISMUTH-CONTAINING AGENT AND H2 RECEPTOR BLOCKING ANTI-SECRETORY AGENT
- U-201 METHOD OF TREATING GI DISORDERS COMPRISING ADMINISTRATION OF CAMPYLOBACTER-INHIBITING ANTIMICROBIAL AGENT AND H2 RECEPTOR BLOCKING ANTI-SECRETORY AGENT
- U-202 METHOD OF TREATING PEPTIC ULCER DISEASE CAUSED BY CAMPYLOBACTER PYLORIDIS COMPRISING ORAL ADMINISTRATION OF 50 TO 5,000MG BISMUTH DAILY FOR 3-56 DAYS
- U-203 TREATMENT OF ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH DISEASE PROGRESSION FOLLOWING ANTIESTROGEN THERAPY
- U-204 USE OF TAXOL IN COMBINATION WITH G-CSF FOR TREATMENT OF PATIENTS WITH AIDS-RELATED KAPOSI'S SARCOMA
- U-205 METHOD FOR TREATING HEARTBURN
- U-206 METHOD OF USING FSH ALONE, WITHOUT THE PRESENCE OF EXOGENEOUS LH, IN IN VITRO FERTILIZATION
- U-207 USE AS NASAL SPRAY
- U-208 VAGINAL ADMINISTRATION USING SPECIFIED FORMULATION
- U-209 VAGINAL ADMINISTRATION OF PROGESTERONE USING SPECIFIED FORMULATION
- U-210 METHOD OF TREATING CONGESTIVE HEART FAILURE
- U-211 USE IN PATIENTS WITH REVERSIBLE AIRWAY OBSTRUCTION
- U-212 METHOD OF TREATMENT OF PARKINSON'S DISEASE
- U-213 METHOD OF INHIBITING CHOLESTEROL BIOSYNTHESIS AND TREATING HYPERCHOLESTEROLEMIA AND METHOD FOR TREATING HYPERLIPIDEMIA
- U-214 USE AS A BLOOD GLUCOSE-LOWERING AGENT
- U-215 TREATMENT OF EPILEPSY TWICE DAILY. TREATING A PATIENT BY ADMINISTERING CARBAMAZEPINE IN A DOSAGE FORM CAPABLE OF MAINTAINING BLOOD CONCENTRATION FROM 4-12MCG/ML OVER 12 HOURS
- U-216 TREATMENT OF ADENOCARCINOMA, INCLUDING STAGE B2-C BY ADMINISTERING AN AGONIST OF LH-RH AND FLUTAMIDE
- U-217 METHOD OF PRODUCING ANESTHESIA
- U-218 METHOD FOR LIMITING THE POTENTIAL FOR MICROBIAL GROWTH IN THE DRUG PRODUCT
- U-219 TREATMENT OF PARKINSON'S DISEASE
- U-220 METHOD OF DIAGNOSIS
- U-221 SELECTIVE VASODILATION BY CONTINUOUS ADENOSINE INFUSION
- U-222 METHOD OF TREATING PAGET'S DISEASE USING ACTONEL
- U-223 TREATMENT OF BACTERIAL CONJUNCTIVITIS CAUSED BY SUSCEPTIBLE STRAINS OF MICROORGANISMS
- U-224 CONTROLLING INTRAOCULAR PRESSURE
- U-225 METHOD FOR DELIVERY
- U-226 METHOD OF ENHANCING THE DISSOLUTION PROFILE OF A PHARMACEUTICAL FROM A SOLID DOSAGE FORM CONTAINING THE PHARMACEUTICAL AND SIMETHICONE
- U-227 NASAL ADMINISTRATION
- U-228 ASTHMA
- U-229 CARDIAC INSUFFICIENCY (CONGESTIVE HEART FAILURE)
- U-230 PREVENTION OF ACUTE CARDIAC ISCHEMIC EVENTS
- U-231 USE IN PARKINSON'S DISEASE
- U-232 METHOD OF TREATING MIGRAINE
- U-233 DECREASING MORTALITY CAUSED BY CONGESTIVE HEART FAILURE
- U-234 METHOD OF USING RIBAVIRIN TO TREAT VIRAL INFECTIONS IN MAMMALS
- U-235 METHOD OF MODULATING TH1 AND TH2 RESPONSE IN ACTIVATED T CELLS OF A HUMAN COMPRISING ADMINISTERING RIBAVIRIN TO THE T CELLS IN A DOSAGE WHICH PROMOTES THE TH1 RESPONSE AND SUPPRESSES THE TH2 RESPONSE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-236 TREATING MALE PATTERN BALDNESS WITH 0.05 TO 3.0MG/DAY
- U-237 METHOD OF PERFORMING NMR IMAGING WITH A PATIENT COMPRISING ADMINISTERING TO THE PATIENT AN EFFECTIVE AMOUNT OF CONTRAST AGENT DISCLOSED IN THE CLAIMS
- U-238 IMAGING A BODY TISSUE AND SUBJECTING TO NMR TOMOGRAPHY, ADMINISTERING AN AMOUNT OF PHARMACEUTICAL AGENT FOR AFFECTING THE RELAXATION TIMES OF ATOMS IN BODY TISSUES UNDERGOING NMR DIAGNOSIS, WHEREBY THE IMAGE CONTRAST IS ENHANCED....
- U-239 TREATING OR CONTROLLING OCULAR INFLAMMATION WHICH COMPRISES TOPICALLY ADMINISTERING TO AFFECTED EYE A COMPOSITION COMPRISING AN NSAID, A POLYMERIC QUATERNARY AMMONIUM COMPOUND AND BORIC ACID
- U-240 TREATMENT OF ACUTE MIGRAINE ATTACKS
- U-241 FOR SHORT-TERM TREATMENT ACTIVE DUODENAL ULCER, MAINTENANCE THERAPY FOR DUODENAL ULCER PATIENTS AT REDUCED DOSAGE AFTER HEALING OF ACTIVE ULCER, SHORT-TERM TREATMENT ACTIVE BENIGN GASTRIC ULCER & GERD, PATHOLOGICAL HYPERSECRETORY CONDITIONS
- U-242 USE OF FOLLITROPIN ALPHA ALONE IN IN-VITRO FERTILIZATION
- U-243 TOPICAL ADMINISTRATION
- U-244 PLATELET AGGREGATION INHIBITORS
- U-245 TREATMENT OF SEBORRHEA DERMATITIS IN HUMANS
- U-246 PHOSPHATE BINDING
- U-247 TREATMENT OF RHEUMATOID ARTHRITIS
- U-248 TREATMENT OF HIV
- U-249 METHOD OF TREATING ALLERGIC OR NON-ALLERGIC RHINITIS IN PATIENTS BY ADMINISTERING AEROSOLIZED PARTICLES OF MOMETASONE FUROATE
- U-250 TREATMENT OF HEPATITIS B INFECTION
- U-251 USE OF TROGLITAZONE IN COMBINATION WITH SULFONYLUREAS IN THE TREATMENT OF TYPE II DIABETES
- U-252 METHOD OF TREATING A HUMAN SUBJECT HAVING GAUCHER'S DISEASE
- U-253 ORAL TRANSMUCOSAL USE
- U-254 USE OF AGGRASTAT IN COMBINATION WITH HEPARIN
- U-255 IMPROVED WAKEFULNESS IN PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY
- U-256 TREATMENT OF HIV INFECTION IN COMBINATION WITH ONE OR MORE ADDITIONAL HIV ANTIVIRAL AGENTS
- U-257 TREATMENT OF HIV INFECTION
- U-258 TREATMENT OF NEURODEGENERATIVE DISEASES
- U-259 TREATMENT OF ANDROGENIC ALOPECIA BY ORAL ADMINISTRATION DRUG SUBSTANCE
- U-260 REDUCTION OF INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA AND OCULAR HYPERTENSION WHO ARE INTOLERANT OF OTHER IOP LOWERING MEDICATIONS OR INSUFFICIENTLY RESPONSIVE TO ANOTHER IOP LOWERING MEDICATION
- U-261 TREATING BENIGN PROSTATIC HYPERPLASIA WITH A GENUS OF COMPOUNDS, INCLUDING FINASTERIDE
- U-262 TREATING BENIGN PROSTATIC HYPERTROPHY WITH FINASTERIDE
- U-263 METHOD OF TREATING A MALIGNANT CONDITION THROUGH INTRAVASCULAR ADMINISTRATION OF BUSULFAN. METHOD FOR TREATING LEUKEMIA OR LYMPHOMA IN A PATIENT UNDERGOING A BONE MARROW TRANSPLANT THROUGH INTRAVENOUS ADMINISTRATION OF BUSULFAN
- U-264 METHOD OF TREATING A MALIGNANT DISEASE THROUGH PARENTERAL ADMINISTRATION OF BUSULFAN. METHOD FOR TREATING A PATIENT UNDERGOING A BONE MARROW TRANSPLANT THROUGH INTRAVASCULAR ADMINISTRATION OF BUSULFAN
- U-265 USE AS LAXATIVE
- U-266 RELIEF OF THE SIGNS AND SYMPTOMS OF OSTEOARTHRITIS; RELIEF OF THE SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS IN ADULTS; MANAGEMENT OF ACUTE PAIN IN ADULTS; TREATMENT OF PRIMARY DYSMENORRHEA; ACUTE TREATMENT OF MIGRAINE ATTACKS IN ADULTS
- U-267 PREVENTING HEARTBURN EPISODES FOLLOWING INGESTION OF HEARTBURN-INDUCING FOOD/BEVERAGE, COMPRISING ADMIN TO PT, 30 MIN PRIOR TO CONSUMPTION BY THE PT THE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

FOOD/BEVERAGE, A COMPOSITION COMPRISING 10MG FAMOTIDINE

U-268 ACROMEGALY

U-269 EXCESS GH-SECRETION OR GASTRO-INTESTINAL DISORDERS

U-270 METHOD OF IMPROVING THE TIME FOR ADMINISTRATION OR THE TIME BETWEEN CHANGES OF GIVING SETS FOR THE DRUG PRODUCT

U-271 METHOD OF TREATING TUMORS

U-272 METHOD OF TREATING CARCINOMA

U-273 CUTANEOUS T-CELL LYMPHOMA

U-274 ZANAMIVIR FOR INHALATION

U-275 METHOD OF USE OF THE DRUG SUBSTANCE

U-276 METHOD OF USE OF LEVOBUPIVACAINE

U-277 NEUROLOGICAL AND OTHER DISORDERS (TREATMENT OF EPILEPSY, BID ORAL DOSING)

U-278 METHOD OF USE OF THE INDICATION OF THE DRUG PRODUCT

U-279 METHOD OF USE OF THE APPROVED PRODUCT

U-280 TREATING PRECIPITATED ACUTE URINARY RETENTION WITH FINASTERIDE

U-281 ANTIMYCOTIC USES, SPECIFICALLY TREATMENT OF ONYCHOMYCOSIS

U-282 METHOD OF TREATING BACTERIAL INFECTIONS

U-283 METHOD FOR TREATING MENOPAUSAL SYMPTOMS IN A POSTMENOPAUSAL FEMALE

U-284 MENOPAUSAL AND POSTMENOPAUSAL DISORDERS (INCLUDING VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE, AND VULVAR AND VAGINAL ATROPHY) AND OSTEOPOROSIS

U-285 DEPRESSION AND SOCIAL ANXIETY DISORDER/SOCIAL PHOBIA

U-286 DEPRESSION

U-287 TREATMENT OR PREVENTION OF OSTEOPOROSIS

U-288 THERAPY OF INFLUENZA

U-289 TREATMENT OF NON-HYPERKERATOTIC ACTINIC KERATOSES OF FACE AND SCALP

U-290 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS)

U-291 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH CYCLOSPORIN

U-292 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH AZATHIOPRINE

U-293 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH A CORTICOSTEROID

U-294 TREATMENT OF HYPERPIGMENTARY DISORDERS

U-295 TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS

U-296 TREATING MIGRAINE PAIN AND ONE OR MORE OF A CLUSTER OF SYMPTOMS CHARACTERISTIC OF A MIGRAINE ATTACK SYMPTOMS BEING SELECTED FROM PHOTOPHOBIA, PHONOPHOBIA NAUSEA AND FUNCTIONAL DISABILITY

U-297 PREVENTION OR TREATMENT OF REVERSIBLE VASOCONSTRICTION BY THE INHALATION OF NITRIC OXIDE WITH AN OXYGEN CONTAINING GAS

U-298 METHOD OF COMBATING BACTERIA IN A PATIENT

U-299 TREATMENT OF ADENOMATOUS POLYPS

U-300 INDICATED FOR THE REDUCTION OF ELEVATED TOTAL AND LDL CHOLESTEROL LEVELS IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA

U-301 USE OF TROGLITAZONE IN COMBINATION WITH SULFONYLUREAS AND BIGUANIDES IN THE TREATMENT OF TYPE II DIABETES

U-302 TO REDUCE THE RISK OF STROKE IN PATIENTS WHO HAVE HAD TRANSIENT ISCHEMIA OF THE BRAIN OR COMPLETED ISCHEMIC STROKE DUE TO THROMBOSIS

U-303 METHOD OF USE PATENT-PRODUCT APPROVED FOR TREATMENT OF OSTEOPOROSIS, PAGET'S DISEASE, PREVENTION AND TREATMENT OF GLUCOCORTICOID INDUCED OSTEOPOROSIS

U-304 A METHOD OF TREATMENT OF A CONDITION INVOLVING AN ANTIBODY ANTIGEN REACTION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-305 METHODS FOR USING THE DRUG PRODUCT

U-306 TREATMENT OF POST-MENOPAUSAL UROGENITAL SYMPTOMS ASSOCIATED WITH ESTROGEN DEFICIENCY

U-307 CLAIMS AN OLANZAPINE POLYMORPH USEFUL FOR TREATING ANY NUMBER OF LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSES, EMPLOYING OLANZAPINE AS PER THE INDICATION OF THIS NDA

U-308 CLAIMS A SOLID ORAL FORMULATION INCLUDING TABLETS AND GRANULES OF OLANZAPINE USEFUL FOR TREATING ANY NUMBER OF LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSES, EMPLOYING OLANZAPINE AS PER THE INDICATIONS OF THIS NDA

U-309 TREATING SJOEGREN SYNDROME

U-310 TREATMENT OF XEROSTOMIA

U-311 HORMONE REPLACEMENT

U-312 PANIC DISORDER, OBSESSIVE-COMPULSIVE DISORDER, POSTTRAUMATIC STRESS DISORDER

U-313 TREATMENT OF CONGESTIVE HEART FAILURE

U-314 METHOD FOR TREATING HYPERPARATHYROIDISM WHICH COMPRISES SUPPRESSING PARATHYROID ACTIVITY

U-315 METHOD FOR ADMINISTERING DRUG TO GASTROINTESTINAL TRACT

U-316 METHOD OF TREATING A SUBJECT SUFFERING FROM PROSTATE CANCER

U-317 METHOD OF USING TROGLITAZONE TO TREAT PATIENTS HAVING INSULIN RESISTANCE

U-318 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER WITH SYMPTOMS OF URINARY FREQUENCY, URGENCY, OR URGE INCONTINENCE

U-319 TREATMENT OF MICROBIAL INFECTIONS

U-320 INHIBITING OR ELIMINATING ACUTE MYELOID LEUKEMIA

U-321 REDUCTION OF ELEVATED IPTH LEVELS IN THE MGT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS UNDERGONG CHRONIC RENAL DIALYSIS

U-322 TREATMENT OF ALZHEIMER'S DEMENTIA

U-323 USE AS A BILE ACID SEQUESTRANT

U-324 METHOD OF TREATING AN ANIMAL, INCLUDING A HUMAN, SUFFERING FROM OR SUSCEPTIBLE TO PSYCHOSIS OR ACUTE MANIA EMPLOYING OLANZAPINE

U-325 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED CONDITIONS, INCLUDING "BIPOLAR DISORDER NOS" EMPLOYING OLANZAPINE

U-326 METHOD OF TREATING SCHIZOPHRENIA AND BIPOLAR DISORDER

U-327 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED PSYCHOTIC CONDITONS EMPLOYING OLANZAPINE

U-328 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED CONDITIONS INCLUDING "A PSYCHOTIC CONDITION" EMPLOYING AN OLANZAPINE POLYMORPH

U-329 USE OF AVANDIA AS MONOTHERAPY, IN COMBINATION WITH METFORMIN, AND IN COMBINATION WITH SULFONYLUREAS TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS

U-330 TREATMENT OF NAUSEA AND VOMITING

U-331 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-332 TREATMENT OR PREVENTION OF BRONCHOSPASM

U-333 METHOD OF TREATING OCULAR HYPERTENSION

U-334 TREATMENT OF EXCESSIVE FEMALE FACIAL HAIR

U-335 USE OF PRAVASTATIN SODIUM FOR SECONDARY PREVENTION OF CORONARY EVENTS IN MEN AND WOMEN WHO HAVE HAD A MYOCARDIAL INFARCTION AND HAVE NORMAL CHOLESTEROL LEVELS

U-336 DIAGNOSTIC RADIOIMAGING

U-337 USE OF CARDIOLITE/MIRALUMA KIT FOR THE PREPARATION OF TC99M SESTAMIBI

U-338 METHODS FOR TREATING DISTURBANCES OF MOOD, DISTURBANCES OF APPETITE, DEPRESSED MOOD, OR CARBOHYDRATE CRAVING ALL ASSOCIATED WITH PREMENSTRUAL SYNDROME

U-339 PREVENTION OF CARDIO-TOXICITY CAUSED BY THE ADMINISTRATION OF DOXORUBICIN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-340 THE LONG TERM TREATMENT OF GROWTH FAILURE DUE TO LACK OF ADEQUATE ENDOGENOUS GROWTH HORMONE SECRETION IN CHILDREN

U-341 METHOD FOR ENHANCING THE TREATMENT OF ... LATE LUTEAL PHASE DYSPHORIC DISORDER

U-342 METHOD FOR TREATMENT OF LATE LUTEAL PHASE DYSPHORIC DISORDER

U-343 REDUCTION OF INTESTINAL GAS, CRAMPING AND ANORECTAL IRRITATION

U-344 METHOD FOR INHIBITING HIV INFECTION BY ADMINISTERING RITONAVIR IN COMBINATION WITH ANOTHER HIV PROTEASE INHIBITOR

U-345 RITONAVIR AND ANOTHER HIV PROTEASE INHIBITOR FOR CONCOMITANT ADMINISTRATION FOR THE TREATMENT OF AN HIV INFECTION

U-346 METHOD FOR INHIBITING CYTOCHROME P450 MONOOXYGENASE WITH RITONAVIR AND A METHOD FOR IMPROVING THE PHARMACOKINETICS OF A DRUG THAT IS METABOLIZED BY CYTOCHROME P450 MONOOXYGENASE BY ADMIN THE DRUG AND RITONAVIR

U-347 METHOD OF USE IN COMBINATION WITH REVERSE TRANSCRIPTASE INHIBITORS

U-348 METHOD OF USE FOR INHIBITING HIV INFECTION

U-349 METHOD OF USE WHICH IS SUBJECT OF THE APPLICATION

U-350 PREPARATION OF A PHARMACEUTICAL COMPOSITION FOR CONCOMITANT ADMIN WITH A REVERSE TRANSCRIPTASE INHIBITOR

U-351 INHIBITING PROTEASE WITH LOPINAVIR AND INHIBITING AN HIV INFECTION WITH LOPINAVIR

U-352 INHIBITING HIV INFECTION BY ADMINISTERING RITONAVIR IN COMBINATION WITH A REVERSE TRANSCRIPTASE INHIBITOR

U-353 PREVENTION AND TREATMENT OF OSTEOPOROSIS

U-354 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID WITHOUT CAUSING TREATMENT-LIMITING ELEVATIONS IN URIC ACID OR GLUCOSE LEVELS OR CAUSING LIVER DAMAGE, BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-355 METHOD OF ASSISTING PERSON TO QUIT SMOKING...TRANSDERMALLY ADMIN NICOTINE VIA..PATCH ADHERED TO SKIN AT DOSING RATE APPROX SAME AS ABSORBED FROM SMOKING

U-356 DELIVERING A MEDICINAL AEROSOL FORMULATION USING CFC-FREE PROPELLANT 134A.

U-357 USE OF THE DRUG PRODUCT IN PHOTODYNAMIC THERAPEUTIC PROTOCOLS FOR THE TREATMENT OF AGE-RELATED MACULAR DEGENERATION AND RELATED CONDITIONS INVOLVING UNWANTED NEOVASCULATURE IN THE EYE

U-358 DEPRESSION, OBSESSIVE COMPULSIVE DISORDER, PANIC DISORDER AND SOCIAL ANXIETY DISORDER

U-359 METHOD OF USE OF VISICOL

U-360 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF PATHOLOGICAL PSYCHOLOGICAL CONDITIONS INCLUDING MENTAL DISORDERS EMPLOYING OLANZAPINE AS PER THE INDICATION WHICH IS THE SUBJECT MATTER OF THIS SNDA-011

U-361 MANAGEMENT OF ANXIETY DISORDERS AND THE SHORT-TERM RELIEF OF THE SYMPTOMS OF ANXIETY

U-362 USE OF APPROVED FORMULATIONS TO TREAT ALL APPROVED DISEASE INDICATIONS

U-363 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF PATHOLOGICAL PSYCHOLOGICAL CONDITIONS THAT RELATE TO THE USE OF A PSYCHOACTIVE SUBSTANCE EMPLOYING OLANZAPINE AS PER THE INDICATION THE SUBJECT MATTER OF SUPPLEMENT 011

U-364 TREATING A PATIENT SUFFERING FROM OR SUSCEPTIBLE TO ANY NUMBER OF LISTED CONDITIONS INCLUDING PSYCHOSIS, EMPLOYING OLANZAPINE AS PER THE INDICATION WHICH IS THE SUBJECT MATTER OF THIS SNDA-011

U-365 METHOD FOR THE TREATMENT OF CARDIOVASCULAR DISEASE THROUGH THE ADMINISTRATION OF A CALCIUM BLOCKING VASODILATOR IN OUR EXTENDED, CONTROLLED RELEASE FORMULATION

U-366 METHOD FOR THE TREATMENT OF CARDIOVASCULAR DISEASE THROUGH THE ADMINISTRATION OF A CALCIUM BLOCKING VASODILATOR IN A DELAYED RELEASE FORMULATION

U-367 TREATMENT OF CARDIOVASCULAR DISORDERS

U-368 HEARTBURN

U-369 METHOD OF CONTROLLING AND LOWERING INTRAOCULAR PRESSURE

U-370 INTRAVAGINAL TREATMENT OF VAGINAL INFECTIONS WITH BUFFERED METRONIDAZOLE COMPOSITIONS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-371 APPROVAL FOR MARKETING ONLY UNDER A SPECIAL RESTRICTION PROGRAM APPROVED BY FDA CALLED "SYSTEM FOR THALIDOMIDE EDUCATION AND PRESCRIBING SAFETY" (S.T.E.P.S.)

U-372 METHOD FOR ADMINISTERING A BENEFICIAL DRUG TO THE GI TRACT OF AN ANIMAL, WHICH METHOD COMPRISES ADMITTING AN OSMOTIC DEVICE ORALLY INTO THE ANIMAL...

U-373 GENERAL USE CLAIM SUBMITTED FOR 12 NEXIUM PATIENTS STATING "PERTINENT TO THE CAPSULE FORMULATION FOR NEXIUM AND ITS INDICATIONS FOR THE TREATMENT OF GERD AND ERADICATION OF H.PYLORI TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE

U-374 KIT ADAPTED AND DESIGNED TO PROVIDE BOTH DATA ON THE CURRENT REPRODUCTIVE STATUS OF A PATIENT AND CONTRACEPTION FOR THOSE WHO ARE NOT PREGNANT, BUT RECENTLY ENGAGED IN UNPROTECTED SEX

U-375 METHOD OF USING RIBAVIRIN FOR TREATING A DISEASE RESPONSIVE TO RIBAVIRIN, E.G. HEPATITIS C

U-376 TREATMENT OF INFLUENZA

U-377 METHOD OF TREATING PT WITH CHRONIC HEPATITIS C HAVING HCV GENOTYPE 1 AND VIRAL LOAD GREATER THAN 2 MILLION COPIES/ML TO ERADICATE DETECTABLE HCV-RNA BY ADMIN COMBINATION OF RIBAVIRIN AND INTERFERON ALFA-2B FOR A LEAST 24 WEEKS

U-378 METHOD FOR TREATING INCONTINENCE

U-379 METHOD OF TREATING ONYCHOMYCOSIS

U-380 COMBINATIONS OF TAXOL (PACLITAXEL) AND CISPLATIN WHICH ARE SUITABLE FOR THE TREATMENT OF OVARIAN AND NON-SMALL CELL LUNG CARCINOMAS

U-381 TREATMENT OF HYPERPHOSPHATEMIA

U-382 METHOD OF STABILIZING PROSTAGLANDIN

U-383 METHOD FOR TREATING GLAUCOMA AND OCULAR HYPERTENSION

U-384 TREATMENT OF CMV RETINITIS

U-385 TREATMENT OF PEPTIC ULCERS

U-386 TREATMENT OF PATIENTS SUFFERING FROM A LATE ASTHMATIC REACTION OR LATE PHASE ASTHMA

U-387 TREATMENT OF PATIENTS WITH RESPIRATORY DISORDERS

U-388 SMOKING CESSATION AID APPLIED TO THE SKIN

U-389 SMOKING CESSATION AID APPLIED TO THE SKIN ON WAKING AND REMOVED PRIOR TO SLEEP AFTER ABOUT 16 HOURS

U-390 METHOD OF USING THE DRUG TO TREAT NEUROIMMUNOLOGIC DISEASES (INCLUDING MULTIPLE SCLEROSIS)

U-391 USE OF CASODEX IN COMBINATION WITH LHRH AGONISTS FOR THE TREATMENT OF PROSTATE CANCER

U-392 TREATMENT OF PATIENTS FOR INFLAMMATION

U-393 MANAGEMENT OF INCONTINENCE, MGT OF HORMONE REPLACEMENT THERAPY, TREATMENT OF INVOLUNTARY INCONTINENCE, MGT OVERACTIVE BLADDER AND INCREASING COMPLIANCE IN SUCH PT

U-394 METHOD OF USE OF ALPHAGAN

U-395 METHOD OF USE OF ALPHAGAN P

U-396 METHOD OF TREATING PEOPLE SUFFERING FROM DEPRESSION

U-397 METHOD OF TREATING PEOPLE SUFFERING FROM DEPRESSION WITHOUT AN INCREASE IN NAUSEA

U-398 TREATMENT OF GENERALIZED ANXIETY DISORDER

U-399 IN-THE-EYE USE OF CHLORINE DIOXIDE CONTAINING COMPOSITIONS

U-400 USE OF RIBAVIRIN TO INCREASE TYPE 1 CYTOKINE RESPONSE AND SUPPRESS TYPE 2 CYTOKINE RESPONSE TO LYMPHOCYTES, INCLUDING METHODS THAT TAKE ADVANTAGE OF SUCH MODULATION TO TREAT INFECTIONS AND INFESTATIONS

U-401 USE OF LOPINAVIR IN COMBINATION WITH REVERSE TRANSCRIPTASE INHIBITORS FOR TREATING HIV INFECTION AND IN COMBO WITH OTHER HIV PROTEASE INHIBITORS

U-402 TREATMENT OF ACTINIC KERATOSES

U-403 ANTI-ALLERGIC FOR VARIOUS ALLERGIC DISEASES

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-404 TREATMENT OF ALLERGIC CONJUNCTIVITIS

U-405 FOR WOMEN WITH SEVERE DIARRHEA-PREDOMINANT IRRITABLE BOWEL SYNDROME (IBS)

U-406 METHOD OF USE OF ATOVAQUONE AND PROGUANIL

U-407 METHOD OF TREATING OTOPATHY

U-408 FOR INDUCING OVULATION IN CONJUNCTION WITH A GONADOTROPIN RELEASING FACTOR ANTAGONIST AND RECRUITING OOCYTES FOR IN-VITRO FERTILIZATION

U-409 METHOD OF TREATING INFLAMMATION USING DRUG SUBSTANCE

U-410 METHOD OF REDUCING AMOUNT OF RESPECTIVE ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (INCLUDING PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

U-411 METHOD OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (WHICH INCLUDES PIOGLITAZONE) IN COMBINATION WITH AN INSULIN PREPARATION

U-412 TREATMENT OF TYPE 2 DIABETES

U-413 USE OF THE ACTIVE INGREDIENT FOR INHIBITING THE BIOSYNTHESIS OF CHOLESTEROL AND TREATMENT OF ATHEROSCLEROSIS

U-414 A METHOD OF TREATING GLYCOMETABOLISM DISORDERS BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE

U-415 A METHOD FOR REDUCING THE AMOUNT OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE AS SAID ACTIVE COMPONENTS

U-416 A METHOD FOR REDUCING SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE AS SAID ACTIVE COMPONENTS

U-417 COMBINATION USE OF AD-4833 WITH A BIGUANIDE

U-418 A METHOD OF TREATING LIPID METABOLISM DISORDERS BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (WHICH INCLUDES PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

U-419 A METHOD OF TREATING LIPID METABOLISM DISORDERS BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE

U-420 METHOD OF TREATMENT OF TYPE II DIABETES

U-421 USE FOR SEDATION

U-422 METHOD OF TREATING AT LEAST ONE OF ATTENTION DEFICIT DISORDER AND ATTENTION DEFICIT HYPERACTIVITY DISORDER

U-423 METHOD OF TREATING AT LEAST ONE OF ATTENTION DEFICIT DISORDER, ATTENTION DEFICIT HYPERACTIVITY DISORDER, OR AIDS RELATED DEMENTIA

U-424 FOR ONCE DAILY, BOLUS ADMINISTRATION TO A PATIENT IN ORDER TO ENGENDER TREATMENT FOR A NERVOUS DISORDER FOR SUBSTANTIALLY AN ENTIRE DAY ON A CHRONIC BASIS

U-425 METHOD OF REDUCING SIDE EFFECTS OF ACTIVE COMPONENTS ADMIN TO A DIABETIC BY ADMIN A CHEMICAL COMPOUND HAVING FORMULA (INCL PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER

U-426 PREVENTION OF PREMATURE LH SURGES IN WOMEN UNDERGOING CONTROLLED OVARIAN STIMULATION

U-427 METHOD OF TREATING ALLERGIC REACTIONS IN MAMMALS

U-428 METHOD OF TREATING ALLERGY IN A MAMMAL USING THIS ACTIVE METABOLITE

U-429 METHOD OF USING DESLORATADINE TO TREAT ALLERGIC RHINITIS

U-430 METHOD OF TREATING A DIABETIC BY ADMINISTERING AN INSULIN SENSITIZER IN COMBINATION WITH AN INSULIN SECRETION ENHANCER, AND A DRUG PRODUCT COMPRISING AN INSULIN SENSITIZER AND AN INSULIN SECRETION ENHANCER

U-431 POSTTRAUMATIC STRESS DISORDER

U-432 REDUCTION OF ATHEROSCLEROTIC EVENTS (MYOCARDIAL INFARCTION, STROKE, AND VASCULAR DEATH) IN PATIENTS WITH ATHEROSCLEROSIS DOCUMENTED BY RECENT STROKE, RECENT MYOCARDIAL INFARCTION OR ESTABLISHED PERIPHERAL ARTERIAL DISEASE

U-433 USE OF LEVOCARNITINE IN PREVENTION AND TREATMENT OF CARNITINE DEFICIENCY IN PATIENTS WITH END STAGE RENAL DISEASE WHO ARE UNDERGOING DIALYSIS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-434 CONTROLLED SYMPTOMS OF DIARRHEA, BLOATING PRESSURE AND CRAMPS, COMMONLY REFERRED TO AS GAS

U-435 A TITRATION DOSING REGIMEN FOR THE TREATMENT OF PAIN USING AN INITIAL DOSE OF ABOUT 25MG

U-436 ACUTE TREATMENT OF MIGRAINE ATTACKS WITH OR WITHOUT AURA IN ADULTS

U-437 METHOD OF USE EQUAL TO PROCESS OF PREPARATION

U-438 TREATMENT/PREVENTION OF NEURODEGENERATIVE DISEASE

U-439 TREATMENT OF OBESITY

U-440 METHOD FOR TRANSDERMAL ADMINISTRATION OF A DRUG THROUGH NON-SCROTAL SKIN USING A TRANSDERMAL DRUG DELIVERY DEVICE CONTAINING THE DRUG AND HAVING AN ADHESIVE SURFACE

U-441 METHOD OF TREATING MS BY ADMINISTERING COPAXONE

U-442 METHOD FOR DELIVERING A DRUG TO A PATIENT IN NEED OF THE DRUG, WHILE AVOIDING THE OCCURRENCE OF AN ADVERSE SIDE EFFECT KNOWN OR SUSPECTED OF BEING CAUSED BY SAID DRUG

U-443 MANAGEMENT OF MODERATE TO SEVERE PAIN WHEN A CONTINUOUS, AROUND-THE-CLOCK ANALGESIC IS NEEDED FOR AN EXTENDED PERIOD OF TIME

U-444 TREATMENT OF MIGRAINE

U-445 USE AS AN ANTIMYCOTIC AGENT

U-446 TOPICAL TREATMENT OF OCULAR HYPERTENSION AND GLAUCOMA

U-447 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-448 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID WITHOUT CAUSING TREATMENT-LIMITING ELEVATIONS IN URIC ACID OR GLUCOSE LEVELS OR CAUSING LIVER DAMAGE, BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-449 USE IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR THE TREATMENT OF METASTATIC COLORECTAL CANCER WHERE THE DOSE OF LEUCOVORIN IS AT LEAST 200MG PER SQUARE METER

U-450 INTERMEDIATE REL NICOTINIC ACID FORMULATIONS HAVING UNIQUE URINARY METAB PROFILES RESULTING FROM ABSORPTION PROFILES OF NICOTINIC ACID FROM THE INTERMEDIATE NICOTINIC ACID FORMULATIONS, SUITABLE FOR TX HYPERLIPIDEMIA FOLLOWING QD DOSING

U-451 TREATMENT OF DEPRESSION AND GENERALIZED ANXIETY DISORDER

U-452 USE OF LANSOPRAZOLE FOR COMBATTING DISEASES CAUSED BY THE GENUS CAMPYLOBACTER (C.PYLORI=H.PYLORI)

U-453 TREATMENT OF PLATELET ASSOCIATED ISCHEMIC DISORDERS

U-454 METHOD OF TX A PT SUSPECTED OF HAVING HEPATITIS C BY ADMIN, IN COMBINATION, A CONJUGATE COMPRISING PEG 12000 & INTERFERON ALFA-2B IN AN AMT OF FROM 0.5MCG/KG TO 2MCG/KG, ONCE WEEKLY, AND RIBAVIRIN

U-455 TREATMENT OF PULMONARY HYPERTENSION WITH UT-15

U-456 METHOD OF DECREASING THE PRODUCTION OF A-BETA USING A COMPOSITION WHICH DECREASES BLOOD CHOLESTEROL IN PATIENTS AT RISK OF OR EXHIBITING SYMPTOMS OF ALZHEIMER'S DISEASE

U-457 METHOD OF TREATING A VAGINAL FUNGAL INFECTION IN A FEMALE HUMAN

U-458 METHOD OF USE OF IMAGENT

U-459 TREATMENT OF DEPRESSION AND GENERALIZED ANXIETY DISORDER

U-460 METHOD OF TREATING PSYCHIATRIC SYMPTOMS ASSOCIATED WITH PREMENSTRUAL DISORDERS USING SERTRALINE

U-461 METHOD OF TREATMENT OF LATE LUTEAL PHASE DYSPHORIC DISORDER (PMDD) USING SERTRALINE

U-462 SIGNS AND SYMPTOMS OF OSTEOARTHRITIS AND ADULT RHEUMATOID ARTHRITIS AND TREATMENT OF PRIMARY DYSMENORRHEA

U-463 VENOGRAPHY

U-464 PERIPHERAL ARTERIOGRAPHY

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-465 CT IMAGING OF THE HEAD

U-466 TREATMENT OF IRRITABLE BOWEL SYNDROME

U-467 USE OF EPLERENONE IN COMBINATION WITH AN ANGIOTENSIN CONVERTING ENZYME (ACE) INHIBITOR FOR TREATING HYPERTENSION

U-468 METHOD OF USING FEXOFENADINE HCL IN TREATING ALLERGIC RHINITIS

U-469 TREATMENT OF GASTROESOPHAGEAL REFLEX DISEASE (GERD) AND ERADICATION OF H.PYLORI TO REDUCE RISK OF DUODENAL ULCER RECURRENCE

U-470 THERAPY IN CHRONIC HEPATITIS B VIRUS INFECTION

U-471 METHOD OF TREATING A PATIENT SUFFERING FROM DIABETES MELLITUS

U-472 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER USING METHYLPHENIDATE BI-MODAL RELEASE PROFILE EXTENDED-RELEASE CAPSULES

U-473 TO REDUCE PLASMA CHOLESTEROL LEVELS IN A MAMMAL

U-474 TO REDUCE PLASMA CHOLESTEROL LEVELS BY ADMIN EZETIMIBE IN COMBO WITH CHOLESTEROL BIOSYNTHESIS INHIB SELECTED FROM GROUP CONSISTING OF HMG COA REDUCTASE INHIBITORS INCL SIMVASTATIN

U-475 TREATMENT OF CUTANEOUS MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO ARE REFRACTORY TO AT LEAST ONE PRIOR SYSTEMIC THERAPY

U-476 METHOD OF TREATING ANDROGEN RESPONSIVE/MEDIATED CONDITION IN MAMMAL BY ADMIN A SAFE, EFFECTIVE AMOUNT OF DUTASTERIDE OR PHARMACEUTICALLY ACCEPTABLE DERIVATIVE THEREOF..CONDITIONS INCLUDE BENIGN PROSTATIC HYPERTROPHY

U-477 METHOD OF INHIBITING 5 ALPHA TESTOSTERONE REDUCTASE ENZYME WITH DUTASTERIDE OR ITS DERIVATIVE AND TREATING ANDROGEN RESPONSIVE/MEDIATED DISEASE INCLUDING BENIGN PROSTATIC HYPERPLASIA

U-478 METHOD OF TREATING HEPATITIS C VIRAL INFECTION BY CONTINUOUS PARENTERAL ADMIN INTERFERON ALPHA 2-10 MILLION IU WEEKLY, SUBCUTANEOUSLY, INJECTION OF POLYMER-INTERFERON ALPHA CONJUGATE-POLYMER IS PEG-INTERFERON IS ALPHA 2B

U-479 METHOD OF USING PEG-INTRON/REBETOL COMBINATION THERAPY AND INTRON/REBETOL COMBINATION THERAPY

U-480 CONTRAST AGENT FOR MRI

U-481 DISUBSTITUTED ACETYLENES BEARING HETEROAROMATIC AND HETEROBICYCLIC GROUPS HAVING RETINOID-LIKE ACTIVITY

U-482 METHOD OF IN VITRO FERTILIZATION THERAPY INCLUDING MEANS FOR INDUCING OVULATION....

U-483 METHOD FOR THE ADMINISTRATION OF DRUGS USING THAT COMPOUND

U-484 METHOD OF TREATING A SKIN DISEASE WITH A CORTICOSTEROID-CONTAINING PHARMACEUTICAL COMPOSITION

U-485 METHOD AND COMPOSITION FOR REDUCING NERVE INJURY PAIN ASSOCIATED WITH SHINGLES (HERPES ZOSTER AND POST-HERPETIC NEURALGIA)

U-486 EXTERNAL PREPARATION FOR APPLICATION TO THE SKIN CONTAINING LIDOCAINE-DRUG RETAINING LAYER PLACED ON SUPPORT AND COMPRISES ADHESIVE GEL BASE 1-10% BY WEIGHT OF LIDOCAINE

U-487 METHOD AND COMPOSITION FOR REDUCING NERVE INJURY PAIN ASSOCIATED WITH SHINGLES (HERPES ZOSTER AND POST-HERPETIC NEURALGIA)

U-488 METHOD FOR REDUCING THE PAIN ASSOCIATED WITH HERPES-ZOSTER AND POST-HERPETIC NEURALGIA

U-489 EXPECTORANT

U-490 TESTOSTERONE REPLACEMENT THERAPY IN MALES FOR CONDITIONS ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE

U-491 METHOD OF DELIVERING A DRUG TO THE LUNG

U-492 METHOD FOR THE TREATMENT OF SKIN, SUFFERING FROM A CONDITION SELECTED FROM A GROUP CONSISTING OF NONACNE INFLAMMATORY DERMATOSES... COMPRISING APPLYING TO AFFECTED AREA. A THERAPEUTICALLY EFFECTIVE AMT AZELAIC ACID

U-493 TREATMENT OF TYPE 2 DIABETES MELLITUS

U-494 TREATMENT OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER

U-495 PERITONEAL DIALYSIS SOLUTION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-496 METHOD FOR TREATING CHRONIC RENAL FAILURE

U-497 RELIEF OF THE SIGNS AND SYMPTOMS OF OSTEOARTHRITIS AND RHEUMATOID ARTHRITIS

U-498 INTRA-ARTERIAL AND INTRAVENOUS USES OF ULTRAVIST

U-499 METHOD OF USING REBETOL CAPSULES IN COMBINATION WITH A CONJUGATE COMPRISING POLYETHYLENE GLYCOL(PEG) AND AN ALPHA INTERFERON, INCLUDING, FOR EXAMPLE, PEG-INTRON POWDER FOR INJECTION

U-500 USE AS AN ANTIHYPERTENSIVE AGENT

U-501 TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) IN ADULTS

U-502 PITYRIASIS VERSICOLOR

U-503 GENERATOR MUST BE USED WITH INFUSION SYSTEM SPECIFICALLY LABELED FOR USE WITH GENERATOR

U-504 TINEA PEDIS, TINEA CRURIS, TINEA CORPORIS

U-505 ULTRASOUND CONTRAST AGENT

U-506 PHARM PRODUCT CONTAINER 1ST CHAMBER IS DISPOSED AQUEOUS DILUENT SOL 2ND CHAMBER PHARM ACTIVE AGENT COMPRISING ACETYLCHOLINE,BUFFER IN 1ST CHAM IS SUFFICIENT TO BUFFER PH OF MIXED SOL RESULTING MIXTURE OF AQUEOUS DILUENT SOL & PHARM ACTIVE..

U-507 ACROMEGALY IN PATIENTS W/INADEQUATE RESPONSE TO SURGERY AND/OR RADIATION THERAPY AND/OR MEDICAL THERAPIES, OR FOR WHOM THESE THERAPIES ARE NOT APPROPRIATE

U-508 METHOD OF RELEASING 17-BETA OESTRADIOL PRECURSOR IN A SUBSTANTIALLY ZERO ORDER PATTERN FOR AT LEAST THREE WEEKS

U-509 TREATMENT OF CUTANEOUS MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO ARE REFRACTORY TO AT LEAST ONE PRIOR SYSTEMIC THERAPY

U-510 TOPICAL TREATMENT OF CUTANEOUS LESIONS IN PATIENTS WITH CUTANEOUS T-CELL LYMPHOMA (STAGE IA AND IB) WHO HAVE REFRACTORY OR PERSISTENT DISEASE AFTER OTHER THERAPIES OR WHO HAVE NOT TOLERATED OTHER THERAPIES

U-511 USE OF QUINOLONE COMPOUNDS AGAINST ANAEROBIC PATHOGENIC BACTERIA

U-512 USE OF QUINOLONE COMPOUNDS AGAINST ATYPICAL UPPER RESPIRATORY PATHOGENIC BACTERIA

U-513 METHODS OF USE OF ANTIMICROBIAL COMPOUNDS AGAINST PATHOGENIC MYCOPLASMA BACTERIA

U-514 PREVENTION OF OVULATION IN A WOMAN

U-515 TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES AND HAVE DEMONSTRATED DISEASE PROGRESSION ON THE LAST THERAPY

U-516 METHOD OF TREATING A PSYCHOTIC DISEASE

U-517 STABLE GEL FORMULATION FOR TOPICAL TREATMENT OF SKIN CONDITIONS

U-518 OBSESSIVE COMPULSIVE DISORDER

U-519 POST OPERATIVE NAUSEA AND VOMITING

U-520 PREMENOPAUSAL OSTEOPOROSIS

U-521 METHOD OF USING RIBAVIRIN IN COMBINATION WITH INTRON A (INTERFERON ALPHA-2 B RECOMBINANT) INJECTION TO TREAT PATIENTS WITH CHRONIC HEPATITIS C

U-522 TREATMENT OF CMV RETINITIS BY INTRAVITREAL ADMIN OF A PHOSPHOROTHIOATE OLIGONUCLEOTIDE CAPABLE OF HYBRIDIZING WITH CMV MRNA

U-523 METHOD OF TREATING INFECTION BY CRYPTOSPORIDIUM PARVUM IN AN IMMUNOCOMPROMISED MAMMAL

U-524 METHOD OF TREATING DIARRHEA

U-525 METHOD OF TREATING PARASITIC INFECTIONS

U-526 METHOD OF PROVIDING CONTROLLED RELEASE OF A TREATING AGENT USING A CONTROLLED RELEASE COMPOSITION

U-527 METHOD OF DELIVERING AN ACTIVE INGREDIENT USING A PROGRESSIVE HYDRATION BIOADHESIVE

U-528 PREVENTION OF CHEMOTHERAPY-INDUCED NAUSEA AND VOMITING

U-529 ONCE DAILY TREATMENT OF ASTHMA WITH NEBULIZED BUDESONIDE

U-530 TREATMENT OF HERPES ZOSTER, TREATMENT OF GENITAL HERPES, TREATMENT OF COLD

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

SORES, SUPPRESSION OF GENITAL HERPES IN IMMUNOCOPETENT AND HIV-INFECTED INDIVIDUALS, REDUCTION OF RISK OF HETEROSEXUAL TRANSMISSION OF GENITAL HERPES

U-531 TREATMENT OF PATIENTS WITH ESSENTIAL HYPERTENSION. MAY BE USED ALONE OR GIVEN WITH OTHER CLASSES OF ANTIHYPERTENSIVES, ESPECIALLY THIAZIDE DERIVATIVES

U-532 TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD IN PATIENTS REQUIRING MORE THAN ONE BRONCHO DILATOR

U-533 ERECTILE DYSFUNCTION

U-534 HUMALOG IS AN INSULIN ANALOG THAT IS INDICATED IN THE TREATMENT OF PATIENTS WITH DIABETES MELLITUS FOR THE CONTROL OF HYPERGLYCEMIA

U-535 TREATMENT OF SOCIAL ANXIETY DISORDER

U-536 CONTRAST AGENT FOR MAGNETIC RESONACE IMAGING

U-537 TREATMENT OF CONDITIONS RELATED TO HYPERALDOSTERONISM SUCH AS HYPERTENSION AND CARDIAC INSUFFICIENCY, WITH EPLERENONE

U-538 FIRST LINE TREATMENT OF SEVERE HYPERTENSION, IN PATIENTS WITH HYPERTENSION SEVERE ENOUGH THAT THE VALUE OF ACHIEVING PROMPT BLOOD PRESSURE CONTROL EXCEEDS THE RISK OF INITIATING COMBINATION THERAPY IN THESE PATIENTS

U-539 TREATMENT OF MODERATE TO SEVERE DEMENTIA OF THE ALZHEIMER'S TYPE

U-540 TREATMENT OF FUNGAL INFECTIONS

U-541 METHOD OF TREATMENT OF ADULTS INFECTED WITH HIV-1

U-542 METHOD OF TREATING PATIENT WITH TYPE 2 DIABETES BY ONCE DAILY ADMINISTRATION

U-543 TREATMENT OF SCHIZOPHRENIA

U-544 TREATMENT OF OVERACTIVE BLADDER. TREATMENT OF URINARY INCONTINENCE.

U-545 METHOD FOR THE PREVENTION AND/OR TREATMENT OF THROMBOTIC EPISODES, SUCH AS MYOCARDIAL INFARCTION, IN A HUMAN PATIENT AND METHOD FOR THE PREVENTION OF VENOUS THROMBOSIS IN A POSTOPERATIVE HUMAN PATIENT

U-546 USE OF REPAGLINIDE IN COMBINATION WITH METFORMIN TO LOWER BLOOD GLUCOSE

U-547 MAINTENANCE MONOTHERAPY FOR BIPOLAR DISORDER

U-548 A METHOD OF REDUCING FLUSH IN AN INDIVIDUAL BEING TREATED FOR A LIPIDEMIC DISORDER AND EFFECTIVELY TREATING THE LIPIDEMIC DISORDER

U-549 USE IN THE TREATMENT OF MEN WITH ADVANCED SYMPTOMATIC PROSTATE CANCER

U-550 TREATMENT OF BIPOLAR DISORDER AND SCHIZOPHRENIA

U-551 METHOD FOR REDUCING TOXICITY OF ALIMTA TREATED PATIENTS BY ADMINISTERING FOLIC ACID

U-552 TREATMENT OF HYPERTENSION AND HYPERLIPIDEMIA WITH A SINGLE COMPOSITION

U-553 MANAGEMENT OF PAIN AND DISCOMFORT ASSOCIATED WITH PERIDONTAL SCALING AND ROOT PLANNING PROCEDURES BY APPLICATION OF AN EUTECTIC MIXTURE OF LOCAL ANESTHETICS TO PERIDONTAL POCKETS

U-554 TREATING HIV INFECTION WITH INDINAVIR SULFATE IN COMBINATION WITH ANTIRETROVIRAL AGENTS

U-555 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS AND ACUTE UNCOMPLICATED PYELONEPHRITIS

U-556 USE AS ADJUNCT DIAGNOSTIC FOR SERUM THYROGLOBULIN (TG) TESTING

U-557 NASAL TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS

U-558 INDICATED FOR THE RELIEF OF BRONCHOSPASM IN PATIENTS 2-12 YEARS OF AGE WITH ASTHMA (REVERSIBLE OBSTRUCTIVE AIRWAY DISEASE)

U-559 METHOD OF DECREASING OR REDUCING PARATHYROID HORMONE LEVEL; METHOD OF MODULATING PARATHYROID HORMONE SECRETION;METHOD OF TREATING HYPERPARATHYROIDISM; METHOD OF REDUCING SERUM IONIZED CALCIUM LEVEL

U-560 METHOD OF DECREASING PARATHYROID HORMONE LEVEL;METHOD OF TREATING HYPERPARATHYROIDISM

U-561 COSOPT IS INDICATED FOR THE REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION WHO ARE INSUFFICIENTLY RESPONSIVE TO BETA BLOCKERS

U-562 TOPICAL TREATMENT OF CUTANEOUS LESIONS IN PATIENTS WITH AIDS-RELATED KAPOSI'S

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

SARCOMA

- U-563 MARINOL IS INDICATED FOR, INTER ALIA, ANOREXIA ASSOCIATED WITH WEIGHT LOSS IN PATIENTS WITH AIDS
- U-564 TREATMENT OF HIV IN CONCOMITANT THERAPY
- U-565 TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS, AND CHRONIC URTICARIA
- U-566 FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- U-567 METHOD OF TREATING INFERTILITY
- U-568 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION
- U-569 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION AND WHEREIN THEREAFTER AN OVULATORY INDUCING AMOUNT OF HCG IS ADMINISTERED
- U-570 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION AND WHEREIN THE DAILY AMOUNT OF FSH IS ABOUT 5-10 IU/KG
- U-571 TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA AND BIPOLAR I MANIA
- U-572 INTENSIVE CARE UNIT SEDATION
- U-573 TREATMENT OF ACUTE PROMYELOGENOUS LEUKEMIA (APL)
- U-574 PROPHYLAXIS AND TREATMENT OF THE NASAL SYMPTOMS OF SEASONAL ALLERGIC RHINITIS AND TREATMENT OF THE NASAL SYMPTOMS OF PERENNIAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-575 LOTEMAX OPHTHALMIC SUSPENSION IS INDICATED FOR THE TREATMENT OF STEROID RESPONSIVE CONDITIONS OF THE PALPEBRAL BULBAR CONJUNCTIVA, CORNEA AND ANTERIOR SEGMENT OF THE GLOBE.
- U-576 ALREX OPHTHALMIC SUSPENSION IS INDICATED FOR THE TEMPORARY RELIEF OF THE SIGNS AND SYMPTOMS OF SEASONAL ALLERGIC CONJUNCTIVITIS.
- U-577 TREATMENT OF BENIGN PROSTATIC HYPERPLASIA WITH FINASTERIDE IN COMBINATION WITH DOXAZOSIN
- U-578 TREATMENT OF COMMUNITY ACQUIRED PNEUMONIA, ACUTE EXACERBATION OF CHRONIC BRONCHITIS, AND ACUTE BACTERIAL SINUSITIS CAUSED BY SUSCEPTIBLE STRAINS OF DESIGNATED MICROORGANISMS IN PATIENTS 18 YEARS AND OLDER.
- U-579 TREATMENT OF EPILEPSY AND/OR MIGRAINE.
- U-580 TREATMENT OF DISORDERS OF THE SEROTONERGIC SYSTEM SUCH AS DEPRESSION AND ANXIETY-RELATED DISORDERS
- U-581 METHOD OF TREATING A CONDITION CAPABLE OF TREATMENT BY INHALATION, E.G. ASTHMA, COMPRISING ADMINISTRATION OF A FORMULATION CLAIMED IN US PATENT NO. 6743413
- U-582 METHOD FOR THE TREATMENT OF A RESPIRATORY DISORDER, E.G. ASTHMA, COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AN AEROSOL COMPOSITION TO A PATIENT FROM A METERED DOSE INHALER SYSTEM AS CLAIMED IN US PATENT NO. 6253762
- U-583 METHOD FOR THE TREATMENT OF A RESPIRATORY DISORDER, E.G. ASTHMA, COMPRISING ADMINISTERING TO A PATIENT BY INHALATION, A METERED AEROSOL DOSE OF A DRUG FORMULATION FROM THE METERED DOSE INHALER SYSTEM CLAIMED IN US 6546928
- U-584 SINGLE-DOSE ADMINISTRATION BY THE EPIDURAL ROUTE, AT THE LUMBAR LEVEL, FOR THE TREATMENT OF PAIN FOLLOWING MAJOR SURGERY
- U-585 TO PROMOTE WEIGHT GAIN AFTER WEIGHT LOSS IN CERTAIN TYPES OF PATIENTS
- U-586 AN INTERMEDIATE RELEASE NICOTINIC ACID FORMULATION SUITABLE FOR ORAL ADMINISTRATION ONCE-A-DAY AS A SINGLE DOSE FOR TREATING HYPERLIPIDEMIA WITHOUT CAUSING DRUG-INDUCED HEPATOTOXICITY OR ELEVATIONS IN URIC ACID OR GLUCOSE OR BOTH
- U-587 USE OF EPLERENONE IN COMBINATION WITH AN ANGIOTENSIN CONVERTING ENZYME (ACE) INHIBITOR (AND OPTIONALLY A DIURETIC) FOR TREATING CONGESTIVE HEART FAILURE AND HYPERTENSION
- U-588 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER; TREATMENT OF HEARTBURN AND OTHER SYMPTOMS ASSOCIATED WITH GERD; SHORT-TERM TREATMENT OF EROSIIVE ESOPHAGITIS; MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS
- U-589 METHOD FOR TREATMENT OF A RESPIRATORY DISORDER, E.G., BRONCHOSPASM, COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AN AEROSOL COMPOSITION TO A PATIENT FROM A

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- METERED DOSE INHALER SYSTEM AS CLAIMED IN U.S. PATENT NO. 6131966
- U-590 METHOD FOR TREATMENT OF A RESPIRATORY DISORDER, E.G., BRONCHOSPASM, COMPRISING ADMINISTERING TO A PATIENT BY ORAL OR NASAL INHALATION A DRUG FORMULATION BY USING THE METERED DOSE INHALER SYSTEM AS CLAIMED IN US PATENT NO. 6532955
- U-591 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER USING A DOSAGE FORM WHICH PROVIDES ONCE-DAILY ORAL ADMINISTRATION OF A PHENIDATE DRUG
- U-592 TREATMENT OF PRIMARY HYPERCHOLESTEROLEMIA, MIXED HYPERLIPIDEMIA AND/OR HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFH)
- U-593 TREATMENT OF PRIMARY HYPERCHOLESTEROLEMIA, MIXED HYPERLIPIDEMIA AND/OR HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFH)
- U-594 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- U-595 35 MG ORALLY ONCE A WEEK FOR PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN; 35 MG ORALLY ONCE A WEEK FOR TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- U-596 TREATMENT OF HORMONE RECEPTOR POSITIVE METASTATIC BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH DISEASE PROGRESSION FOLLOWING ANTIESTROGEN THERAPY
- U-597 FORTEO IS INDICATED FOR THE TREATMENT OF POST MENOPAUSAL WOMEN WITH OSTEOPOROSIS WHO ARE AT HIGH RISK FOR FRACTURE
- U-598 PROPHYLACTIC TREATMENT OF MIGRAINE
- U-599 METHOD FOR TREATING ALLERGIC CONJUNCTIVITIS
- U-600 A METHOD OF TREATING A PATIENT IN NEED OF OPHTHALMIC ANTIMICROBIAL THERAPY WITH LEVOFLOXACIN
- U-601 TREATMENT OF BIPOLAR DISORDER
- U-602 SIGNS AND SYMPTOMS OF OSTEOARTHRITIS, RHEUMATOID ARTHRITIS IN ADULTS, AND/OR PAUCIARTICULAR OR POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS, ACUTE PAIN IN ADULTS; PRIMARY DYSMENORRHEA; AND/OR ACUTE MIGRAINE ATTACKS IN ADULTS
- U-603 METHOD OF TREATING INFECTIONS COMPRISING ORALLY ADMINISTERING AN EFFECTIVE AMOUNT OF THE FDA APPROVED ORAL SUSPENSION
- U-604 METHOD OF LOWERING BLOOD GLUCOSE BY ONCE DAILY ADMINISTRATION
- U-605 TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD); ALTHOUGH THE MECHANISM OF THE ANTIDEPRESSANT ACTION OF DULOXETINE IN HUMANS IS UNKNOWN, IT IS BELIEVED TO BE RELATED TO ITS POTENTIATION OF SERATONERGIC AND NORADRENERGIC ACTIVITY IN THE CNS
- U-606 USE OF IRINOTECAN IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR THE TREATMENT OF METASTATIC COLORECTAL CANCER
- U-607 CANCIDAS IS INDICATED FOR EMPIRICAL THERAPY FOR PRESUMED FUNGAL INFECTIONS IN FEBRILE, NEUTROPENIC PATIENTS.
- U-608 USE OF QUINOLONE COMPOUNDS AGAINST PNEUMOCOCCAL PATHOGENIC BACTERIA
- U-609 USE OF QUINOLONE COMPOUNDS AGAINST QUINOLONE-RESISTANT PNEUMOCOCCAL PATHOGENIC BACTERIA
- U-610 ATROVENT HFA (IPRATROPIUM BROMIDE HFA) INHALATION AEROSOL IS INDICATED AS A BRONCHODILATOR FOR MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE, INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA.
- U-611 METHOD OF USING DESLORATADINE TO TREAT SEASONAL AND PERENNIAL ALLERGIC RHINITIS, PRURITIS, AND CHRONIC IDIOPATHIC URTICARIA IN PATIENTS 2 YEARS OF AGE AND OLDER
- U-612 TREATMENT OF SEASONAL ALLERGY SYMPTOMS WITH NASAL CONGESTION IN ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER
- U-613 REDUCTION OF SERUM PHOSPHATE
- U-614 TREATMENT OF SEXUAL DYSFUNCTION
- U-615 ADJUNCTIVE THERAPY TO DIET IN ADULTS TO REDUCE LDL-C, TOTAL-C, TRIGLYCERIDES AND APO B, AND INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA OR MIXED DYSLIPIDEMIA (TYPES IIA, IIB) AND TO TREAT HYPERTRIGLYCERIDEMIA (TYPES IV, V)
- U-616 MANAGEMENT OF PERSISTENT, MODERATE TO SEVERE PAIN IN PATIENTS REQUIRING CONTINUOUS, AROUND-THE-CLOCK ANALGESIA WITH A HIGH POTENCY OPIOID FOR AN EXTENDED PERIOD OF TIME GENERALLY WEEKS TO MONTHS OR LONGER
- U-617 TREATMENT OF ACUTE PROMYELOGENOUS LEUKEMIA (APL)
- U-618 USE OF ROSUVASTATIN CALCIUM TO REDUCE ELEVATED TOTAL-C, LDL-C, APOB, NONHDL-C OR

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

TG LEVELS; TO INCREASE HDL-C IN ADULT PATIENTS WITH PRIMARY HYPERLIPIDEMIA OR MIXED DYSLIPIDEMIA; AND TO SLOW THE PROGRESSION OF ATHEROSCLEROSIS.

- U-619 TREATMENT OF MALIGNANT NEOPLASM
- U-620 TREATMENT OF INSOMNIA
- U-621 METHOD OF TREATING CANCER
- U-622 TREATMENT OF VEGF MEDIATED OCULAR DISEASE.
- U-623 SHORT TERM TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
- U-624 REDUCTION OF RISK OF UPPER GASTROINTESTINAL BLEEDING IN CRITICALLY ILL PATIENTS
- U-625 ALLERGIC RHINITIS OR NASAL POLYPS
- U-626 CLOLAR IS INDICATED FOR THE TREATMENT OF PEDIATRIC PATIENTS 1 TO 21 YEARS OLD WITH RELAPSED OR REFRACTORY ACUTE LYMPHOBLASTIC LEUKEMIA AFTER AT LEAST TWO PRIOR REGIMENS
- U-627 TREATMENT OF PATIENTS USING EXTENDED-RELEASE CARBAMAZEPINE
- U-628 USE OF AVANDIA IN COMBINATION WITH A SULFONYLUREA, AND IN COMBINATION WITH METFORMIN AND A SULFONYLUREA TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-629 METHOD OF INDUCING A HYPNOTIC OR SEDATIVE EFFECT IN A HUMAN BY ADMINISTERING ESZOPICLONE
- U-630 TREATING URINARY INCONTINENCE BY ADMINISTERING AN EXTENDED-RELEASE FORM OF DARIFENACIN
- U-631 TREATING A DISEASE OF ALTERED MOTILITY OR TONE OF SMOOTH MUSCLE BY ADMINISTERING A MUSCARINIC RECEPTOR ANTAGONIZING AMOUNT OF DARIFENACIN
- U-632 METHOD OF TREATMENT OF CANCER BY ADMINISTERING PARTICLES OF PACLITAXEL THAT HAVE A PROTEIN COATING
- U-633 METHOD FOR TREATMENT OF TUMORS BY ADMINISTERING PACLITAXEL AT A DOSE IN THE RANGE OF ABOUT 30MG/METER SQUARE TO ABOUT 100MG/METER SQUARE IN A PHARMACEUTICALLY ACCEPTABLE FORMULATION THAT DOES NOT CONTAIN CREMOPHOR
- U-634 METHOD FOR DELIVERY OF A BIOLOGIC (INCLUDING ANTINEOPLASTIC AGENTS) BY ADMINISTERING TO A PATIENT AN EFFECTIVE AMOUNT OF A BIOLOGIC AS A SOLID OR LIQUID WITH A POLYMERIC BIOCOMPATIBLE MATERIAL
- U-635 TREATMENT OF GERD, MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS AND RISK REDUCTION OF NSAID ASSOCIATED GASTRIC ULCERS
- U-636 TREATMENT OR PREVENTION OF BRONCHOSPASM OR ASTHMATIC SYMPTOMS
- U-637 TREATMENT OF DIABETES WITH AN AMYLIN AGONIST
- U-638 TREATMENT OF DIABETES WITH AN AMYLIN AGONIST, INCLUDING WITH INSULIN
- U-639 TREATMENT OF A MAMMAL HAVING A NEED OF OR REDUCED ABILITY TO PRODUCE INSULIN WITH AN INSULIN AND AN AMYLIN SUCH AS PRAMLINTIDE
- U-640 USE OF AN AMYLIN AGONIST TO REDUCE GASTRIC MOTILITY AND TREAT POST PRANDIAL HYPERGLYCEMIA
- U-641 USE OF AN AMYLIN AGONIST HAVING SPECIFIED BINDING ACTIVITY TO REDUCE GASTRIC MOTILITY, INCLUDING USE THROUGH PARENTERAL ADMINISTRATION
- U-642 TREATMENT AND PREVENTION OF OSTEOPOROSIS
- U-643 THE SHORT TERM TREATMENT (UP TO 10 DAYS) IN PTS HAVING GASTROESOPHAGEAL REFLUX DISEASE (GERD) AS AN ALTERNATIVE TO ORAL THERAPY IN PTS WHEN THERAPY WITH NEXIUM CAPSULES IS NOT POSSIBLE OR APPROPRIATE
- U-644 TREATMENT OF SEASONAL ALLERGIC RHINITIS
- U-645 TREATMENT OF ASTHMA
- U-646 METHOD OF TREATING OTITIS
- U-647 TREATMENT OF OSTEOPOROSIS IN POST MENOPAUSAL WOMEN AND/OR THE TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
- U-648 THE TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN AND/OR THE TREATMENT TO INCREASE BONE MASS IN MEN
- U-649 A METHOD FOR TREATING A TUMOR DISEASE
- U-650 TREATMENT OF ESOPHAGEAL CANDIDIASIS AND PROPHYLAXIS OF CANDIDA INFECTIONS IN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

HSCT PATIENTS

- U-651 TREATMENT OF ACUTE PROMYELOCYTIC LEUKEMIA (APL)
- U-652 TREATMENT OF CARDIAC ARRHYTHMIA
- U-653 STIMULATING INSULIN RELEASE BY ADMINISTERING EXENATIDE
- U-654 LOWERING PLASMA GLUCAGON IN A SUBJECT IN NEED THEREOF, INCLUDING ONE WITH TYPE 2 DIABETES, BY ADMINISTERING AN EXENDIN OR ANALOG, SUCH AS EXENDIN-4
- U-655 TREATMENT OF MILD TO MODERATE ACTIVE CHROHN'S DISEASE INVOLVING THE ILEUM AND/OR THE ASCENDING COLON AND THE MAINTENANCE OF CLINICAL REMISSION OF MILD TO MODERATE CROHN'S DISEASE INVOLVING THE ILEUM AND/OR ASCENDING COLON FOR UP TO 3 MONTHS
- U-656 REDUCING GASTRIC MOTILITY OR DELAYING GASTRIC EMPTYING BY ADMINISTERING AN EXENDIN, SUCH AS EXENDIN-4
- U-657 PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- U-658 TREATMENT OF ADVANCED HORMONE-DEPENDENT BREAST CANCER
- U-659 TREATMENT OF LOCALLY ADVANCED OR METASTATIC NON SMALL-CELL LUNG CANCER (NSCLC) AFTER FAILURE OF AT LEAST ONE PRIOR CHEMOTHERAPY REGIMEN
- U-660 TREATMENT OF HYPERTENSION AND TREATMENT OF HEART FAILURE
- U-661 TREATMENT OF SEIZURE DISORDER
- U-662 TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- U-663 THE TREATMENT OF UNCOMPLICATED URINARY TRACT INFECTIONS
- U-664 TREATMENT OF CONDITIONS FOR WHICH AN ALDOSTERONE RECEPTOR BLOCKER IS INDICATED, SUCH AS HYPERTENSION, HEART FAILURE, AND POST-MYOCARDIAL INFARCTION
- U-665 METHOD OF USING THE DRUG SUBSTANCE/DRUG PRODUCT FOR ULTRASOUND IMAGING
- U-666 METHOD OF TREATING ADHD
- U-667 MANAGEMENT OF INCONTINENCE; METHOD FOR TREATING INCONTINENCE
- U-668 LEVEMIR IS A LONG-ACTING BASAL INSULIN ANALOG THAT IS INDICATED IN THE TREATMENT OF PATIENTS WITH DIABETES MELLITUS
- U-669 INDICATION OF TYPE II DIABETES
- U-670 TREATMENT OF HIV-1 INFECTION BY THE CO-ADMINISTRATION OF TIPRANAVIR AND RITONAVIR.
- U-671 PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGE 3 AND 4
- U-672 TREATMENT OF INFLAMMATION OR AN INFLAMMATION-ASSOCIATED DISORDER
- U-673 METHOD OF TREATMENT WITH ONCE-DAILY DOSES OF 625MG/5ML
- U-674 METHOD OF TREATING INSOMNIA CHARACTERIZED BY DIFFICULTY WITH SLEEP ONSET
- U-675 PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA; RELIEF OF SYMPTOMS OF ALLERGIC RHINITIS
- U-676 METHOD OF TREATING ATTENTION DEFICIT DISORDER USING ORAL ADMINISTRATION OF A BI-MODAL OR PULSATILE RELEASE COMPOSITION
- U-677 A METHOD OF TREATING DISEASE AMENABLE TO TREATMENT WITH A PHENIDATE DRUG BY ONCE DAILY ORAL ADMINISTRATION OF AN EXTENDED RELEASE DOSAGE FORM
- U-678 METHOD OF TREATING ATTENTION DEFICIT DISORDER AND/OR ATTENTION DEFICIT HYPERACTIVITY DISORDER
- U-679 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES WHO ARE ALREADY TREATED WITH A PIOGLITAZONE AND METFORMIN
- U-680 A METHOD OF TREATING DYSLIPIDEMIA AND DYSLIPOPTEINEMIA USING A DOSAGE FORM THAT CAN PROVIDE AN EFFECTIVE AMOUNT OF FENOFIBRATE TO A PATIENT IN A FASTED STATE WHICH IS AT LEAST 90% OF THE AUC AMOUNT PROVIDED BY THE DOSAGE FORM
- U-681 TREATMENT OF PRIMARY IGF-1 DEFICIENCY
- U-682 NON-BENZODIAZEPINE HYPNOTIC AGENT INDICATED FOR TREATMENT OF INSOMNIA, CHARACTERIZED BY DIFFICULTIES WITH SLEEP ONSET AND/OR SLEEP MAINTENANCE
- U-683 PREVENTION OR TREATMENT OF ISCHEMIC HEART DISEASE
- U-684 TREATMENT OF UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER

- U-685 EXPECTORANT AND COUGH SUPPRESSANT
- U-686 EXPECTORANT AND NASAL DECONGESTANT
- U-687 REDUCING FOOD INTAKE IN A SUBJECT WITH TYPE 2 DIABETES BY ADMINISTERING AN EXENDIN, SUCH AS EXENDIN-4
- U-688 TREATMENT OF HIV-INFECTION IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS
- U-689 TREATMENT OF PATIENTS WITH T-CELL ACUTE LYMPHOBLASTIC LEUKEMIA WHOSE DISEASE HAS NOT RESPONDED TO OR HAS RELAPSED FOLLOWING TREATMENT WITH AT LEAST TWO CHEMOTHERAPY REGIMENS
- U-690 TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-691 USE AS A MONOTHERAPY, IN COMBINATION WITH A SULFONYLUREA, METFORMIN OR INSULIN OR IN COMBINATION WITH A SULFONYLUREA PLUS METFORMIN TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-692 USE OF VALSARTAN TO REDUCE CARDIOVASCULAR MORTALITY IN CLINICALLY STABLE PATIENTS WITH LEFT VENTRICULAR FAILURE OR LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION
- U-693 THE RECOMMENDED INITIAL DOSE OF EQUETRO IS 400MG/DAY GIVEN IN DIVIDED DOSES, TWICE DAILY. THE DOSE SHOULD BE ADJUSTED IN 200MG DAILY INCREMENTS TO ACHIEVE OPTIMAL CLINICAL RESPONSE.
- U-694 LENALIDOMIDE IS AN ANALOGUE OF THALIDOMIDE. THALIDOMIDE IS A KNOWN HUMAN TERATOGEN THAT CAUSES SEVERE LIFE-THREATENING HUMAN BIRTH DEFECTS. IF LENALIDOMIDE IS TAKEN DURING PREGNANCY, IT MAY CAUSE BIRTH DEFECTS OR DEATH TO AN UNBORN BABY.
- U-695 TREATMENT OF PATIENTS WITH T-CELL ACUTE LYMPHOBLASTIC LEUKEMIA AND T-CELL LYMPHOBLASTIC LYMPHOMA WHOSE DISEASE HAS NOT RESPONDED TO OR HAS RELAPSED FOLLOWING TREATMENT WITH AT LEAST TWO CHEMOTHERAPY REGIMENS
- U-696 TREATMENT OF PATIENTS WITH T-CELL LYMPHOBLASTIC LYMPHOMA WHOSE DISEASE HAS NOT RESPONDED TO OR HAS RELAPSED FOLLOWING TREATMENT WITH AT LEAST TWO CHEMOTHERAPY REGIMENS
- U-697 A METHOD OF USING RINFABATE RECOMBINANT (RHIGFBP-3) WITH MECASERMIN RECOMBINANT (RHIGF-1) TO PROMOTE LINEAR GROWTH IN THE TREATMENT OF PRIMARY IGF-1 DEFICIENCY
- U-698 METHOD OF USING ANTAGONIST OF ARGININE VASOPRESSIN (AVA) V1A AND V2 RECEPTORS FOR INTRAVENOUS TREATMENT OF PATIENTS WITH EUVOLEMIC HYONATREMIA
- U-699 NASAL TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS
- U-700 TREATMENT AND PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- U-701 TREATMENT OF HYPERCHOLESTEROLEMIA AND/OR HYPERTRIGLYCERIDEMIA
- U-702 TOPICAL AEROSOL HAIR REGROWTH TREATMENT
- U-703 TREATMENT OF PROTEIN KINASE RELATED DISORDERS, SUCH AS GASTROINTESTINAL STROMAL TUMOR AND RENAL CELL CARCINOMA WITH SUNITINIB
- U-704 METHOD OF ADMINISTERING INSULIN VIA INHALATION
- U-705 TREATING CHRONIC ANGINA BY ADMINISTERING AN EXTENDED RELEASE FORM OF RANOLAZINE
- U-706 TREATMENT OF BENIGN PROSTATIC HYPERPLASIA
- U-707 ALLERGIC RHINITIS
- U-708 TREATMENT OF CHRONIC NON-INFECTIOUS UVEITIS AFFECTING THE POSTERIOR SEGMENT OF THE EYE
- U-709 METHOD OF COMBATING BACTERIA IN A PATIENT
- U-710 A METHOD OF TREATING RESPIRATORY DISORDERS, E.G., ASTHMA, WHICH COMPRISES ADMINISTRATION BY INHALATION OF AN EFFECTIVE AMOUNT OF A PHARMACEUTICAL FORMULATION AS CLAIMED IN US PATENT NO. 5658549
- U-711 ACUTE AND LONGER-TERM TREATMENT OF MAJOR DEPRESSIVE DISORDER
- U-712 A METHOD OF USING A NICOTINIC ACID FORMULATION TO REDUCE ELEVATED TC, LDL-C AND TG LEVELS, AND RAISE HDL-C LEVELS IN PATIENTS WITH HYPERLIPIDEMIA
- U-713 TREATMENT OF MILD TO MODERATE DEMENTIA OF THE ALZHEIMER'S TYPE
- U-714 TOPICAL TREATMENT OF INTERDIGITAL TINEA PEDIS AND TINEA CORPORIS DUE TO TRICHOPHYTON RUBRUM, TRICHOPHYTON MENTAGROPHYTES OR EPIDERMOPHYTON FLOCCOSUM

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-715 FOR CLEANSING THE BOWEL IN PREPARATION FOR COLONOSCOPY, IN ADULTS 18 YEARS OF AGE OR OLDER

U-716 THE TREATMENT OR PREVENTION OF BRONCHOSPASM IN ADULTS AND CHILDREN 4 YEARS OF AGE AND OLDER WITH REVERSIBLE OBSTRUCTIVE AIRWAYS DISEASE AND THE PREVENTION OF EXERCISED-INDUCED BRONCHOSPASM IN PATIENTS 4 YEARS OF AGE AND OLDER

U-717 METHOD OF RELIEVING OR PREVENTING CONSTIPATION IN A HUMAN CONSTIPATED PATIENT

U-718 TREATMENT OF FUNGAL INFECTIONS

U-719 TREATMENT OF PSYCHOSIS

U-720 TREATMENT OF NEUROLEPTIC DISEASES

U-721 TREATMENT OF INFLUENZA

U-722 PROPHYLAXIS OF INFLUENZA

U-723 PROPHYLACTIC TREATMENT OF MIGRAINE

U-724 METHOD OF TREATING SEIZURES

U-725 ALLERGIC RHINITIS AND URTICARIA

U-726 ALLERGIC RHINITIS

U-727 FOR THE TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)

U-728 METHOD FOR TREATING BACTERIAL INFECTION

U-729 TREATMENT OF GASTROESOPHAGEAL REFLUX DISEASE (GERD), RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER, H. PYLORI ERADICATION TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE

U-730 USE AS A NASAL SPRAY FOR TREATMENT OF THE SYMPTOMS OF SEASONAL ALLERGIC RHINITIS AND VASOMOTOR RHINITIS

U-731 USE IN COMBINATION WITH DEXAMETHASONE IS INDICATED FOR THE TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED MULTIPLE MYELOMA

U-732 ACUTE TREATMENT OF THE CUTANEOUS MANIFESTATIONS OF MODERATE TO SEVERE ERYTHEMA NODOSUM LEPROSUM (ENL)

U-733 MAINTENANCE THERAPY FOR PREVENTION AND SUPPRESSION OF THE CUTANEOUS MANIFESTATIONS OF ENL RECURRENCE

U-734 FIRST LINE THERAPY FOR TYPE 2 DIABETES MELLITUS

U-735 METHOD OF TREATING CHRONIC IRON OVERLOAD

U-736 METHOD FOR IONTOPHORETIC TRANSDERMAL DELIVERY OF FENTANYL HYDROCHLORIDE

U-737 DISINFECTION OF PATIENT SKIN PRIOR TO AN INVASIVE PROCEDURE

U-738 INDICATED FOR THE LONG-TERM, TWICE-DAILY MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS 12 YEARS OF AGE OR OLDER

U-739 METHOD FOR TREATING CONSTIPATION BY OPENING CIC CHANNELS IN A MAMMALIAN SUBJECT

U-740 FOR THE TREATMENT OF PATIENTS WITH PRIMARY BILIARY CIRRHOSIS

U-741 COMBINATION THERAPY WITH CISPLATIN FOR THE TREATMENT OF LATE STAGE CERVICAL CANCER

U-742 TWICE DAILY TOPICAL TREATMENT OF MODERATE TO SEVERE PLAQUE PSORIASIS.

U-743 ONCE A DAY TOPICAL TREATMENT OF THE INFLAMMATORY LESIONS OF ROSACEA

U-744 TREATMENT OF HIV INFECTION IN ANTIRETROVIRAL TREATMENT-EXPERIENCED ADULT PATIENTS

U-745 TREATMENT OR PREVENTION OF EMESIS

U-746 PREVENTION OR TREATMENT OF NAUSEA OR EMESIS INDUCED BY A CANCER CHEMOTHERAPEUTIC AGENT

U-747 PREVENTION OR TREATMENT OF POST-OPERATIVE NAUSEA AND VOMITING

U-748 A METHOD FOR THE TREATMENT OF A PROTEIN TYROSINE KINASE-ASSOCIATED DISORDER

U-749 METHOD OF CONTRACEPTION

U-750 TREATMENT OF HIV-1 INFECTION IN ADULTS

U-751 ONCE DAILY DOSING OF BUDESONIDE VIA NEBULIZER FOR THE TREATMENT OF ASTHMA

U-752 SUNSCREEN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-753 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES

U-754 USE FOR THE LONG-TERM MAINTENANCE TREATMENT OF ASTHMA

U-755 TREATMENT OF ANOREXIA, CACHEXIA, OR AN UNEXPLAINED, SIGNIFICANT WEIGHT LOSS IN PATIENTS WITH A DIAGNOSIS OF ACQUIRED IMMUNODEFICIENCY SYNDROME (AIDS)

U-756 ADDITION OF ONCE-WEEKLY DOSING FOR THE TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS

U-757 USE AS A BILE ACID SEQUESTANT FOR LOWERING CHOLESTEROL

U-758 TREATMENT OF SYMPTOMS OF PREMENSTRUAL DYSPHORIC DISORDER

U-759 METHOD OF USE OF ADMINISTERING LEVOTHYROXINE

U-760 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS AND TREATMENT OF OROPHARYNGEAL CANDIDIASIS

U-761 TREATMENT OF SCHIZOPHRENIA INCLUDING MAINTAINING STABILITY IN PATIENTS WITH SCHIZOPHRENIA

U-762 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE

U-763 ADMINISTRATION OF ARIPIPIRAZOLE BY INJECTION

U-764 TREATMENT OF SCHIZOPHRENIA

U-765 METHOD OF TREATING ALLERGIC CONJUNCTIVITIS

U-766 TREATMENT OF SEIZURES

U-767 MANAGEMENT OF BREAKTHROUGH PAIN IN PATIENTS WITH CANCER

U-768 A METHOD OF REDUCING THE CAPACITY OF EXTENDED RELEASE NICOTINIC ACID TO PROVOKE A FLUSHING REACTION BY PRETREATING AN INDIVIDUAL WITH A FLUSH INHIBITING AGENT PRIOR TO THE ADMINISTRATION OF THE EXTENDED RELEASE NICOTINIC ACID

U-769 REVLIMID (LENALIDOMIDE) IN COMBINATION WITH DEXAMETHASONE IS INDICATED FOR THE TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY

U-770 LONG-TERM TREATMENT OF PATHOLOGICAL HYPERSECRETORY CONDITIONS

U-771 METHOD FOR THE TREATMENT OF DIABETES MELLITUS, SUCH AS TYPE 1 DIABETES MELLITUS OR TYPE 2 DIABETES MELLITUS, IN A HUMAN PATIENT

U-772 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN CHILDREN 2 TO 11 YEARS AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA IN CHILDREN 6 MONTHS TO 11 YEARS

U-773 PATHOLOGICAL HYPERSECRETORY CONDITIONS

U-774 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR

U-775 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH METFORMIN AND/OR A SULFONYLUREA

U-776 TREATMENT OF CUTANEOUS MANIFESTATION IN PATIENTS WITH CUTANEOUS T-CELL LYMPHOMA (CTCL) WHO HAVE PROGRESSIVE, PERSISTENT OR RECURRENT DISEASE ON OR FOLLOWING TWO SYSTEMIC THERAPIES.

U-777 DECREASING MORTALITY CAUSED BY CONGESTIVE HEART FAILURE

U-778 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION

U-779 A METHOD FOR TREATMENT OF A CANCER, WHEREIN THE CANCER IS CHRONIC MYELOGENOUS LEUKEMIA

U-780 A METHOD FOR THE TREATMENT OF CANCER

U-781 FOR TREATMENT OF ADULT PATIENTS WITH TYPE 2 DIABETES MELLITUS WHO ARE NAIVE TO PHARMACOLOGIC THERAPY

U-782 TREATMENT OF CHRONIC HEPATITIS B IN ADULT PATIENTS WITH EVIDENCE OF VIRAL REPLICATION AND EITHER EVIDENCE OF PERSISTANT ELEVATIONS IN SERUM AMINOTRANSFERASES (ALT OR AST) OR HISTOLOGICALLY ACTIVE DISEASE

U-783 DESONATE GEL IS INDICATED FOR THE TREATMENT OF MILD TO MODERATE ATOPIC DERMATITIS IN PATIENTS 3 MONTHS OF AGE AND OLDER

U-784 TREATMENT OF MODERATE TO SEVERE PRIMARY RESTLESS LEGS SYNDROME (RLS)

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-785 USE AS REPLACEMENT SOLUTION, HEMOFILTRATION SOLUTION OR HEMODIAFILTRATION SOLUTION IN CONTINUOUS RENAL REPLACEMENT THERAPY
- U-786 PRODUCT IS APPROVED FOR THE TOPICAL TREATMENT OF TINEA PEDIS
- U-787 MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN ADULT AND PEDIATRIC PATIENTS SIX YEARS OF AGE OR OLDER, INCLUDING PATIENTS REQUIRING ORAL CORTICOSTEROID THERAPY FOR ASTHMA
- U-788 METHOD OF TREATING PSYCHIATRIC SYMPTOMS ASSOCIATED WITH PREMENSTRUAL DISORDERS USING PAROXETINE
- U-789 TREATMENT OF KNOWN OR SUSPECTED CYANIDE POISONING
- U-790 FORTEO IS INDICATED FOR THE TREATMENT OF POST MENOPAUSAL WOMEN WITH OSTEOPOROSIS WHO ARE AT RISK FOR FRACTURE. FORTEO CAN BE USED BY PEOPLE WHO HAVE HAD A FRACTURE RELATED TO OSTEOPOROSIS
- U-791 GLEEVEC IS ALSO INDICATED FOR THE TREATMENT OF PATIENTS WITH KIT (CD117) POSITIVE UNRESECTABLE AND/OR METASTATIC MALIGNANT GASTROINTESTINAL STROMAL TUMORS (GIST)
- U-792 TREATMENT OF SEBORRHEA DERMATITIS IN HUMANS
- U-793 FOR THE LONG TERM TREATMENT, TWICE DAILY (MORNING AND EVENING) MAINTENANCE TREATMENT OF BRONCHOCONSTRICTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- U-794 CLOSURE OF A CLINICALLY SIGNIFICANT PATENT DUCTUS ARTERIOSUS IN PREMATURE INFANTS WEIGHING BETWEEN 500 AND 1500G, WHO ARE NO MORE THAN 32 WEEKS GESTATIONAL AGE WHEN USUAL MEDICAL MANAGEMENT IS INEFFECTIVE
- U-795 METHOD FOR INHIBITING NOREPINEPHRINE UPTAKE
- U-796 METHOD OF TREATING DEPRESSION
- U-797 METHOD OF TREATING ANXIETY
- U-798 TREATMENT AND PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN BY ONCE-MONTHLY ORAL ADMINISTRATION OF IBANDRONATE SODIUM MONOHYDRATE EQUIVALENT TO 150MG OF IBANDRONIC ACID
- U-799 METHOD FOR INHIBITING SEROTONIN UPTAKE
- U-800 TREATMENT OF PATIENTS WITH ADVANCED OR METASTATIC BREAST CANCER WHOSE TUMORS OVEREXPRESS HER2 AND WHO HAVE RECEIVED PRIOR THERAPY INCLUDING ANTHRACYCLINE, A TAXANE AND TRASTUZUMAB
- U-801 METHOD OF TREATING CANCER
- U-802 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR
- U-803 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH METFORMIN
- U-804 TREATMENT OF ACTINIC KERATOSES BY PHOTODYNAMIC THERAPY
- U-805 TREATMENT OF IMPETIGO DUE TO STAPHYLOCOCCUS AUREUS OR STREPTOCOCCUS PYOGENES
- U-806 INTRATHECAL TREATMENT OF LYMPHOMATOUS MENINGITIS
- U-807 PREVENTION OF EXERCISE-INDUCED BRONCHOCONSTRICTION
- U-808 THE TREATMENT OF THE SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS IN PATIENTS 2 YEARS OF AGE AND OLDER
- U-809 TREATMENT OF CHRONIC IDIOPATHIC URTICARIA
- U-810 METHOD OF TREATMENT TO ALLEVIATE INFLAMMATION OF THE EYE
- U-811 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS AND TREATMENT OF THE UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA
- U-812 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS
- U-813 MAINTENANCE TREATMENT OF BRONCHOCONSTRICTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-814 TREATMENT OF SCHIZOPHRENIA
- U-815 TREATS COLD SORES/FEVER BLISTERS ON THE FACE OR LIPS. SHORTENS HEALING TIME AND DURATION OF SYMPTOMS: TINGLING, PAIN, BURNING AND/OR ITCHING

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-816 DEPRESSION, PANIC DISORDER, PREMENSTRUAL DISORDERS AND SOCIAL ANXIETY DISORDER

U-817 NASAL ADMINISTRATION OF CYANOCOBALAMIN

U-818 TOPICAL TREATMENT OF ACNE VULGARIS

U-819 MANAGEMENT OF FIBROMYALGIA

U-820 IMPROVED WAKEFULNESS IN PATIENTS WITH EXCESSIVE SLEEPINESS ASSOCIATED WITH NARCOLEPSY, OBSTRUCTIVE SLEEP APNEA/HYPOPNEA SYNDROME, AND SHIFT WORK SLEEP DISORDER

U-821 METHOD OF INHIBITING ENTHOTHELIN RECEPTORS BY ADMINISTERING AMBRISENTAN TO A PATIENT TO TREAT PULMONARY ARTERIAL HYPERTENSION.

U-822 USE IN LIPID MANAGEMENT

U-823 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS AND FOR THE TREATMENT OF UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA IN CHILDREN 6 TO 11 YEARS OF AGE

U-824 METHOD OF TREATING PATIENTS INFECTED WITH CCR5-TROPIC HIV-1

U-825 USE FOR PREVENTION OF BREAST CANCER

U-826 RELIEF OF MODERATE TO SEVERE PAIN

U-827 USE FOR TREATMENT OF DIABETES, PARTICULARLY TYPE 2 DIABETES

U-828 PREVENTION OF PREGNANCY IN WOMEN WHO ELECT TO USE ORAL CONTRACEPTIVES AS A METHOD OF CONTRACEPTION

U-829 TREATMENT OF EXTRAVASATION RESULTING FROM IV ANTHRACYCLINE CHEMOTHERAPY

U-830 TREATMENT OF RELAPSED SMALL CELL LUNG CANCER

U-831 METHOD OF ADMINISTERING LANREOTIDE ACETATE

U-832 ZINGO IS INDICATED FOR THE USE ON INTACT SKIN TO PROVIDE LOCAL ANALGESIA PRIOR TO VENIPUNCTURE OR INTRAVENOUS CANNULATION.

U-833 METHOD OF TREATING PAIN USING A PHARMACEUTICALLY ACCEPTABLE SALT OF ROPIVACAINE AND ADMINISTERING A COMPOSITION CONTAINING LESS THAN 0.25% BY WEIGHT OF ROPIVACAINE

U-834 INVIRASE IN COMBINATION WITH RITONAVIR AND OTHER ANTIRETROVIRAL AGENTS IS INDICATED FOR THE TREATMENT OF HIV INFECTION

U-835 RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF ATOPIC DERMATITIS IN PATIENTS ONE YEAR OF AGE OR OLDER

U-836 A METHOD FOR THE TREATMENT OF LEUKEMIAS

U-837 GASTROINTESTINAL LAVAGE INDICATED FOR CLEANSING OF THE COLON AS A PREPARATION FOR COLONOSCOPY IN ADULTS

U-838 METHOD OF TREATING PAIN USING A PHARMACEUTICALLY ACCEPTABLE SALT OF ROPIVACAINE AND ADMINISTERING A COMPOSITION CONTAINING LESS THAN 0.5% BY WEIGHT OF ROPIVACAINE

U-839 TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)

U-840 TREATMENT FOR TYPE 2 DIABETES MELLITUS

U-841 INDICATED FOR THE LONG-TERM, MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS 12 YEARS OF AGE AND OLDER

U-842 INDICATED FOR THE TREATMENT OF ATTENTION-DEFICIT/HYPERACTIVITY DISORDER (ADHD)

U-843 METHOD FOR ADMINISTRATION OF TESTOSTERONE

U-844 PREFEST IS INDICATED IN WOMEN WHO HAVE A UTERUS FOR THE TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE; TREATMENT OF VULVAR AND VAGINAL ATROPHY; PREVENTION OF OSTEOPOROSIS

U-845 TREATMENT OF PATIENTS WITH CANDIDEMIA, ACUTE DISSEMINATED CANDIDIASIS, CANDIDA PERITONITIS AND ABSCESES

U-846 USE FOR DELINEATION (VISUALIZATION) DURING A VITRECTOMY SURGICAL PROCEDURE

U-847 ADJUNCTIVE THERAPY TO DIET IN ADULTS TO REDUCE LDL-C, TRIGLYCERIDES AND APO B, AND INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA OR MIXED DYSLIPIDEMIA (TYPES IIA, IIB) AND TO TREAT HYPERTRIGLYCERIDEMIA (TYPES IV, V)

U-848 ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-849 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION WHO REQUIRE ADJUNCTIVE OR REPLACEMENT THERAPY DUE TO INADEQUATELY CONTROLLED IOP. DOSE IS ONE DROP OF COMBIGAN IN THE AFFECTED EYE TWICE DAILY
- U-850 PREVENTION OR TREATMENT OF NAUSEA OR EMESIS INDUCED BY A CANCER CHEMOTHERAPEUTIC AGENT
- U-851 TREATMENT OF TYPE 2 DIABETES MELLITUS
- U-852 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS
- U-853 TREATMENT OR PREVENTION OF EMESIS
- U-854 PREVENTION OF CMV DISEASE IN KIDNEY, HEART, AND KIDNEY-PANCREAS TRANSPLANT PATIENTS AT HIGH RISK (DONOR CMV SEROPOSITIVE/RECIPIENT CMV SERONEGATIVE)
- U-855 METHOD TO INDUCE NATRIURESIS, DIURESIS AND/OR VASODILATION
- U-856 SUPPORT EMBRYO IMPLANTATION AND EARLY PREGNANCY BY SUPPLEMENTATION OF CORPUS LUTEAL FUNCTION AS PART OF AN ASSISTED REPRODUCTIVE TECHNOLOGY (ART) TREATMENT PROGRAM FOR INFERTILE WOMEN
- U-857 INHIBITION OF TRANSPLANT REJECTION
- U-858 PEDIATRIC USE AGED 1-11 YEARS, GERD AND EROSIIVE ESOPHAGITIS
- U-859 EROSIIVE ESOPHAGITIS, HYPERSECRETORY CONDITIONS INCLUDING ZOLLINGER-ELLISON SYNDROME, MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS AND REDUCTION OF SYMPTOMS IN PATIENTS WITH GERD
- U-860 FOR THE APPROVED USES AND CONDITIONS OF USE, INCLUDING DEPRESSION
- U-861 RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF CORTICOSTEROID RESPONSIVE DERMATOSES IN PATIENTS 12 YEARS OF AGE OR OLDER
- U-862 ADJUNCT TO DIET TO REDUCE ELEVATED TOTAL-C, LDL-C, NON-HDL-C, APO B, TG, AND LP(A) LEVELS AND TO INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA, MIXED DYSLIPIDEMIA, AND HYPERTRIGLYCERIDEMIA
- U-863 TAKING ASPIRIN OR NON-STEROIDAL ANTI-INFLAMMATORY MEDICATIONS APPROXIMATELY 30 MINUTES BEFORE DOSING CAN MINIMIZE FLUSHING, A COMMON SIDE EFFECT OF NIACIN THERAPY
- U-864 PEDIATRIC USE AGES 1-2 YEARS, GERD AND EROSIIVE ESOPHAGITIS
- U-865 TREATMENT OF A WOMAN WITH OSTEOPOROSIS AND A HIGH RISK FOR BONE FRACTURE BY REDUCING THE RISK OF VERTEBRAL AND NONVERTEBRAL BONE FRACTURE
- U-866 THE LABEL REFERENCES THE EFFECTS OF THE ACTIVE INGREDIENT OF REVLIMID UPON CYTOKINES
- U-867 TREATMENT OF MIGRAINE
- U-868 METHOD OF USING ANTAGONIST OF ARGININE VASOPRESSIN (AVA) V1A AND V2 RECEPTORS FOR INTRAVENOUS TREATMENT OF PATIENTS WITH HYPERVOLEMIC HYPONATREMIA
- U-869 METHOD FOR STIMULATING CORONARY VASODILATION FOR PURPOSES OF IMAGING THE HEART
- U-870 METHOD OF PRODUCING CORONARY VASODILATION WITHOUT PERIPHERAL VASODILATION
- U-871 METHOD OF REDUCING RISK OF MYOCARDIAL INFARCTION, STROKE AND DEATH
- U-872 TWICE DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA. TO REDUCE EXACERBATIONS OF COPD IN PATIENTS WITH A HISTORY OF EXACERBATIONS
- U-873 METHOD OF TREATING CONSTIPATION IN A PATIENT WITH IRRITABLE BOWEL SYNDROME BY OPENING CHLORIDE CHANNELS (CIC)
- U-874 METHOD OF TREATING CONSTIPATION IN A PATIENT WITH IRRITABLE BOWEL SYNDROME
- U-875 FIRST-LINE TREATMENT OF LOCALLY ADVANCED UNRESECTABLE OR METASTATIC PANCREATIC CANCER, IN COMBINATION WITH GEMCITABINE
- U-876 TREATMENT OF MIGRAINE WITH OR WITHOUT AURA
- U-877 FOR USE AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PEPTIC ULCER
- U-878 A METHOD FOR BINDING A PERIPHERAL OPIOID RECEPTOR
- U-879 A METHOD OF TREATING OR PREVENTING ILEUS
- U-880 ENDOMETRIN IS A PROGESTERONE INDICATED TO SUPPORT EMBRYO IMPLANTATION AND EARLY PREGNANCY BY SUPPLEMENTATION OF CORPUS LUTEAL FUNCTION AS PART OF AN ASSISTED

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

REPRODUCTIVE TECHNOLOGY (ART) TREATMENT PROGRAM FOR INFERTILE WOMEN

U-881 TREATMENT OF NON-SMALL CELL LUNG CANCER

U-882 MANAGEMENT OF FIBROMYALGIA (FM)

U-883 TREATMENT OF GASTROINTESTINAL STROMAL TUMOR WITH SUNITINIB

U-884 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA

U-885 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST 1 PRIOR THERAPY

U-886 ADMINISTERING DESLORATADINE TO TREAT THE SYMPTOMS OF PERENNIAL ALLERGIC RHINITIS, SEASONAL ALLERGIC RHINITIS, OR CHRONIC IDIOPATHIC URTICARIA

U-887 TREATMENT AND PREVENTION OF OSTEOPOROSIS

U-888 FEMALE HORMONE REPLACEMENT THERAPY FOR POSTMENOPAUSAL WOMEN

U-889 MENOPAUSAL AND POSTMENOPAUSAL DISORDERS (INCLUDING VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE)

U-890 REDUCTION OF SERUM PHOSPHATE IN PATIENTS WITH END STAGE RENAL DISEASE

U-891 USE AS AN INTRAOCULAR IRRIGATING SOLUTION DURING SURGICAL PROCEDURES INVOLVING PERFUSION OF THE EYE

U-892 TREATMENT OF CUTANEOUS MANIFESTATIONS IN PATIENTS WITH CUTANEOUS T-CELL LYMPHOMA (CTCL)

U-893 CLEVIPREX IS A DIHYDROPYRIDINE CALCIUM CHANNEL BLOCKER INDICATED FOR THE REDUCTION OF BLOOD PRESSURE WHEN ORAL THERAPY IS NOT FEASIBLE OR NOT DESIRABLE

U-894 TREATMENT OF COLD SORES IN PEDIATRIC PATIENTS TWELVE YEARS OF AGE AND OLDER

U-895 TREATMENT OF HIV INFECTION IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS

U-896 TREATMENT OF NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS IN ADULTS AND CHILDREN TWO YEARS OF AGE AND OLDER

U-897 METHOD OF TREATING TONSILLITIS AND/OR PHARYNGITIS SECONDARY TO STREPTOCOCCUS PYOGENES IN A ONCE-A-DAY AMOXICILLIN PRODUCT

U-898 USE OF GLUTAMINE TOGETHER WITH GROWTH HORMONE FOR THE TREATMENT OF PATIENTS WITH SHORT BOWEL SYNDROME

U-899 USE OF THALIDOMIDE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED MULTIPLE MYELOMA

U-900 INTEGRASE INHIBITION FOR THE TREATMENT OF HIV INFECTION

U-901 PREVENTION OF POSTOPERATIVE NAUSEA AND VOMITING

U-902 USE IN THE TREATMENT OF THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA (BPH)

U-903 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) IN ADULT PATIENTS

U-904 TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE

U-905 TREATMENT OF MODERATE TO SEVERE VAGINAL DRYNESS AND PAIN WITH INTERCOURSE, SYMPTOMS OF VULVAR AND VAGINAL ATROPHY, ASSOCIATED WITH MENOPAUSE

U-906 PROPHYLAXIS OF ORGAN REJECTION IN KIDNEY, LIVER AND HEART ALLOGENIC TRANSPLANTS; TREATMENT OF PATIENTS WITH SEVERE ACTIVE, RHEUMATOID ARTHRITIS; TREATMENT OF ADULT, NONIMMUNOCOMPROMISED PATIENTS WITH SEVERE, RECALCITRANT, PLAQUE PSORIASIS

U-907 FOR THE MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS IN SUBJECTS 18 YEARS OF AGE AND OLDER

U-908 PROPHYLAXIS OF ORGAN REJECTION IN PATIENTS RECEIVING ALLOGENEIC RENAL TRANSPLANTS

U-909 TREATMENT OF CYSTIC FIBROSIS PATIENTS WITH PSEUDOMONAS AERUGINOSA

U-910 TREATMENT OF METASTATIC CARCINOMA OF THE OVARY AFTER FAILURE OF INITIAL OR SUBSEQUENT CHEMOTHERAPY

U-911 METHOD OF TREATING, AS ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER WHEN ORAL TREATMENT IS TEMPORARILY NOT FEASIBLE

U-912 SEDATION OF NON-INTUBATED PATIENTS PRIOR TO AND/OR DURING SURGICAL AND OTHER PROCEDURES

U-913 TREATMENT OF OVERACTIVE BLADDER WITH SYMPTOMS OF URGE URINARY INCONTINENCE,

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

URGENCY, AND FREQUENCY

- U-914 METHOD OF TREATING, AS ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER
- U-915 TREATMENT OF MUSCULOSKELETAL CONDITIONS
- U-916 TOPICAL TREATMENT OF ACNE VULGARIS IN PATIENTS 12 YEARS OR OLDER
- U-917 TREATMENT OF INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS
- U-918 TO TREAT OR PREVENT INFECTIONS CAUSED BY SUSCEPTIBLE BACTERIA USING DELAYED-RELEASE TABLETS CONSISTING OF DOXYCYCLINE HYCLATE COATED PELLETS IN A TABLET
- U-919 FOR THE TREATMENT OF DERMATITIS
- U-920 STEROID-RESPONSIVE INFLAMMATORY OCULAR CONDITIONS FOR WHICH A CORTICOSTEROID IS INDICATED AND WHERE SUPERFICIAL BACTERIAL OCULAR INFECTION OR A RISK OF BACTERIAL OCULAR INFECTION EXISTS
- U-921 TREATMENT OF ACNE VULGARIS
- U-922 FOR THE TREATMENT OF FUNGAL INFECTIONS
- U-923 METHOD OF TREATING OPHTHALMIC INFLAMMATION AND INFECTION
- U-924 TREATMENT OF MILD TO MODERATE INFECTION CAUSED BY SUSCEPTIBLE STRAINS
- U-925 TREATMENT OF ONLY INFLAMMATORY LESIONS (PAPULES AND PUSTULES) OF ROSACEA
- U-926 MGT SPECIFIC BACTERIAL INFECTIONS. TREATMENT PTS W/ COMMUNITY ACQUIRED PNEUMONIA OR BACTERIAL SINUSITIS DUE TO CONFIRMED, OR SUSPECTED B-LACTAMASE PRODUCING PATHOGENS & S. PNEUMONIAE WITH REDUCED SUSCEPTIBILITY TO PENICILLIN (MIC=2MC/ML)
- U-927 METHOD FOR INCREASING TEAR PRODUCTION
- U-928 TREATMENT OF BACTERIAL INFECTIOUS DISEASE
- U-929 TREATMENT OF OBSESSIVE COMPULSIVE DISORDER TREATABLE WITH AN SSRI
- U-930 TREATMENT OF IDIOPATHIC THROMBOCYTOPENIC PURPURA (ITP)
- U-931 RELIEF OF MODERATE TO SEVERE ACUTE PAIN
- U-932 PYLERA CAPSULES, IN COMBINATION WITH OMEPRAZOLE ARE INDICATED FOR THE TREATMENT OF PATIENTS WITH HELICOBACTER PYLORI INFECTION AND DUODENAL ULCER DISEASE TO ERADICATE H. PYLORI
- U-933 FOR THE TREATMENT OF PATIENTS WITH HELICOBACTER PYLORI INFECTION AND DUODENAL ULCER DISEASE TO ERADICATE H. PYLORI. THE ERADICATION OF HELICOBACTER PYLORI HAS BEEN SHOWN TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE
- U-934 IN COMBINATION WITH GRANULOCYTE-COLONY STIMULATING FACTOR (G-CSF) TO MOBILIZE HEMATOPOIETIC STEM CELL TO THE PERIPHERAL BLOOD FOR COLLECTION AND SUBSEQUENT AUTOLOGOUS TRANSPLANTATION WITH NON-HODGKINS LYMPHOMA AND MULTIPLE MYELOMA
- U-935 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION IN ADULT PATIENTS, AND TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) IN PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER
- U-936 USE IN COMBINATION WITH GRANULOCYTE-COLONY STIMULATING FACTOR (G-CSF) TO MOBILIZE HEMATOPOIETIC STEM CELLS TO PERIPHERAL BLOOD FOR COLLECTION & SUBSEQUENT AUTOLOGOUS TRANSPLANTATION IN PATIENTS WITH NON-HODGKIN'S LYMPHOMA & MULTIPLE MYELOMA
- U-937 TREATMENT OF PROSTATE CANCER
- U-938 TREATMENT OF HAIR LOSS AND HYPOTRICHOSIS OF THE EYELASHES BY INCREASING THEIR GROWTH INCLUDING LENGTH, THICKNESS AND DARKNESS
- U-939 TREATMENT OF HYPOTRICHOSIS OF THE EYELASHES BY INCREASING AND STIMULATING THEIR GROWTH INCLUDING LENGTH, THICKNESS AND DARKNESS
- U-940 METHOD TO TREAT AIDS-RELATED KAPOSI'S SARCOMA
- U-941 METHOD TO TREAT OVARIAN CANCER
- U-942 METHOD TO TREAT MULTIPLE MYELOMA
- U-943 GNRH ANTAGONIST INDICATED FOR TREATMENT OF PATIENTS WITH ADVANCED PROSTATE CANCER
- U-944 TREATMENT OF PATIENTS WITH B-CELL CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)
- U-945 SEDATIVE-HYPNOTIC AGENT INDICATED FOR MONITORED ANESTHESIA CARE (MAC) SEDATION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-946 TREATMENT OF BREAST CANCER

U-947 WHEN PATIENTS ARE UNABLE TO TAKE THE ORAL FORMULATIONS, PREVACID IV, FOR INJECTION IS INDICATED AS AN ALTERNATIVE FOR THE SHORT-TERM TREATMENT (UP TO 7 DAYS) OF ALL GRADES OF EROSIIVE ESOPHAGITIS

U-948 TREATMENT OF DIABETES MELLITUS

U-949 HEALING OF ALL GRADES OF EROSIIVE ESOPHAGITIS (EE) FOR UP TO 8 WEEKS

U-950 MAINTAIN HEALING OF EROSIIVE ESOPHAGITIS (EE) FOR UP TO 6 MONTHS

U-951 TREATMENT OF HEARTBURN ASSOCIATED WITH NON-EROSIVE GASTROESOPHAGEAL REFLUX DISEASE (GERD) FOR 4 WEEKS

U-952 USE AS AN ANALGESIC

U-953 METHOD OF TREATING OPHTHALMIC INFLAMMATION AND INFECTION

U-954 CHRONIC MANAGEMENT OF HYPERURICEMIA IN PATIENTS WITH GOUT. NOT RECOMMENDED FOR THE TREATMENT OF ASYMPTOMATIC HYPERURICEMIA

U-955 PROPHYLACTIC TREATMENT OF MIGRAINE

U-956 TREATMENT OF PATIENTS WITH H. PYLORI INFECTION AND DUODENAL ULCER DISEASE

U-957 A METHOD OF TREATING CANCER IN A PATIENT COMPRISING ADMINISTERING IXABEPILONE OR PHARMACEUTICAL COMPOSITIONS COMPRISING IXABEPILONE

U-958 METHOD OF TREATING PATIENT COMPRISING MIXING FIRST AND SECOND VIALS OF PRODUCT COMPRISING LYOPHILIZED IXABEPILONE TO PROVIDE AN EPOTHILONE ANALOG SOLUTION, DILUTING SOLUTION WITH A SUITABLE DILUENT TO PREPARE INTRAVENOUS FORMULATION FOR PT

U-959 METHOD OF TREATING CANCER, IV ADMIN, LYOPHILIZED IXABEPILONE DILUTED, EVERY WEEK OR 3 WEEKS; LYOPHILIZED IXABEPILONE WITH SOLVENT (DEHYDRATED ETHANOL) DILUTED TO CONCENTRATION OF 0.1MG/ML TO 0.9MG/ML

U-960 METHOD OF TREATING CANCER IN A PATIENT COMPRISING INTRAVENOUSLY ADMINISTERING TO THE PATIENT IXABEPILONE DILUTED IN A PARENTERAL DILUENT

U-961 METHOD OF TREATING BREAST CANCER BY ADMINISTERING IXABEPILONE; A METHOD OF TREATING A CANCER RESPONSIBLE TO MICROTUBULE STABILIZATION BY ADMINISTERING IXABEPILONE

U-962 SYMBYAX IS INDICATED FOR THE ACUTE TREATMENT OF TREATMENT RESISTANT DEPRESSION IN ADULTS

U-963 PROZAC AND OLANZAPINE IN COMBINATION FOR THE ACUTE TREATMENT OF TREATMENT RESISTANT DEPRESSION IN ADULTS

U-964 ZYPREXA ZYDIS AND FLUOXETINE IN COMBINATION FOR THE ACUTE TREATMENT OF TREATMENT RESISTANT DEPRESSION IN ADULTS

U-965 USE OF IXABEPILONE IN COMBINATION WITH CAPECITABINE IN TREATMENT OF METASTASIS BREAST CANCER

U-966 TREATMENT OF ASTHMA (MAINTENANCE AND PROPHYLACTIC THERAPY)

U-967 A METHOD OF REVERSING SOFT-TISSUE ANESTHESIA I.E. ANESTHESIA OF THE LIP AND TONGUE, AND THE ASSOCIATED FUNCTIONAL DEFICITS RESULTING FROM AN INTRAORAL SUBMUCOSAL INJECTION OF A LOCAL ANESTHETIC

U-968 A METHOD FOR IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

U-969 TREATMENT OF MIGRAINE

U-970 TOPICAL TREATMENT OF LICE INFESTATIONS

U-971 INDICATED FOR THE ACUTE TREATMENT OF ADULTS WITH SCHIZOPHRENIA

U-972 MONOTHERAPY OR AS ADJUNCTIVE THERAPY TO LITHIUM OR VALPROATE FOR THE MAINTENANCE TREATMENT OF BIPOLAR I DISORDER

U-973 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO ARE ALREADY TREATED WITH PIOGLITAZONE AND METFORMIN OR WHO HAVE INADEQUATE GLYCEMIC CONTROL ON PIOGLITAZONE OR METFORMIN ALONE

U-974 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES WHO ARE ALREADY TREATED WITH A PIOGLITAZONE AND METFORMIN

U-975 TREATMENT OF PULMONARY HYPERTENSION

U-976 IMPROVEMENT OF GLYCEMIC CONTROL IN INDIVIDUALS WITH TYPE 2 DIABETES

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-977 TREATMENT OF ACUTE, UNCOMPLICATED MALARIA INFECTION DUE TO PLASMODIUM FALCIPARUM IN PATIENTS OF 5KG BODYWEIGHT AND ABOVE

U-978 METHOD OF TREATING HYPONATREMIA

U-979 RELIEF OF MUSCLE SPASM

U-980 NONSTEROIDAL ANTI-INFLAMMATORY DRUG INDICATED FOR RELIEF OF MILD TO MODERATE ACUTE PAIN

U-981 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS, REDUCTION IN FEVER THROUGH ANTI-INFLAMMATORY, ANALGESIC, AND ANTIPYRETIC ACTIVITY

U-982 A METHOD OF TREATING OSTEOPOROSIS

U-983 METHOD OF TREATING OSTEOPOROSIS IN A POST-MENOPAUSAL WOMAN AT RISK FOR FRACTURE

U-984 METHOD FOR THE TREATMENT OF A WOMAN WITH OSTEOPOROSIS AND AT RISK FOR BONE FRACTURE

U-985 TREATMENT OF MACULAR EDEMA FOLLOWING BRANCH RETINAL VEIN OCCLUSION (BRVO) OR CENTRAL RETINAL VEIN OCCLUSION (CRVO)

U-986 TREATMENT OF PATIENTS INFECTED WITH PEDICULUS HUMANUS CAPITIS (HEAD LICE AND THEIR OVA) OF THE SCALP HAIR

U-987 TREATMENT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS WITH CHRONIC KIDNEY DISEASE ON DIALYSIS

U-988 TREATMENT OF RHINITIS COMPRISING THE NASAL APPLICATION OF A PHARMACEUTICAL FORMULATION AS CLAIMED IN US PATENT 7541350

U-989 FOR REDUCING BLOOD PHENYLALANINE LEVELS IN A HUMAN SUFFERING FROM HYPERPHENYLALANINEMIA

U-990 TREATMENT OF PROTOZOAL INFECTION

U-991 TREATMENT OR PROPHYLAXIS OF THROMBOSIS OR EMBOLISMS

U-992 REDUCTION OF THE RISK OF CARDIOVASCULAR HOSPITALIZATION

U-993 METHOD OF TREATING INFERTILITY

U-994 METHOD OF TREATMENT OF OSTEOPOROSIS WHEREIN THE OSTEOPOROSIS IS STEROID-INDUCED

U-995 METHOD FOR TREATING TYPE II DIABETES BY ADMINISTERING SAXAGLIPTIN

U-996 AN ADJUNCTIVE THERAPY TO DIET TO REDUCE ELEVATED TOTAL CHOLESTEROL (TC), LOW-DENSITY LIPOPROTEIN CHOLESTEROL, APOLIPOPROTEIN B, TRIGLYCERIDES, AND TO INCREASE HDL-C IN ADULT PATIENTS WITH PRIMARY HYPERLIPIDEMIA OR MIX DYSLIPIDEMIA

U-997 TREATMENT OF MAJOR DEPRESSIVE DISORDER BY DOSING AT INTERVALS OF 24 HOURS

U-998 ADJUNCTIVE THERAPY TO DIET TO REDUCE ELEVATED TOTAL CHOLESTEROL, LOW-DENSITY LIPOPROTEIN CHOLESTEROL, APOLIPOPROTEIN B, TRIGLYCERIDES AND TO INCREASE HDL-C IN ADULT PATIENTS WITH PRIMARY HYPERLIPIDEMIA OR MIXED DYSLIPIDEMIA

U-999 TREATMENT OF CHRONIC HEPATITIS B IN ADULT PATIENTS

U-1000 ADJUNCTIVE THERAPY TO DIET IN PATIENTS WITH HYPERLIPIDEMIAS

U-1001 METHOD FOR DELIVERING DRUG TO LUNG OF MAMMAL, COMPRISING ADMINISTERING DRUG PRODUCT BY INHALATION. TREATING A MAMMAL HAVING A CONDITION CAPABLE OF TREATMENT BY INHALATION, COMPRISING ADMINISTERING TO THE LUNG THE DRUG PRODUCT BY INHALATION

U-1002 METHOD OF TREATING INFLAMMATORY CONDITIONS

U-1003 A METHOD OF MYOCARDIAL PERFUSION IMAGING AND INCREASING CORONARY BLOOD FLOW

U-1004 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA

U-1005 METHOD OF TREATING A STAPHYLOCOCCAL INFECTION

U-1006 NEW COMBINATION PRODUCT FOR THE EARLY TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) TO REDUCE THE LIKELIHOOD OF ULCERATIVE COLD SORES AND TO SHORTEN THE LESION HEALING TIME IN ADULTS AND ADOLESCENTS (12 YEARS OF AGE AND OLDER)

U-1007 METHOD OF TREATING GOUT FLARES

U-1008 APPLICATION OF ANTISEPTIC WITH MOISTURIZERS FOR SURGICAL AND HEALTHCARE PERSONNEL SKIN DISINFECTION

U-1009 METHOD FOR ADMINISTRATION OF TESTOSTERONE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1010 TO REDUCE BLOOD PHENYLALANINE LEVELS IN PATIENTS WITH HYPERPHENYLALANINEMIA DUE TO TETRA HYDROBIOPTERIN RESPONSIVE PHENYLKETONURIA. KUVAN SHOULD BE TAKEN ORALLY WITH FOOD TO INCREASE ABSORPTION

U-1011 USE OF GRANISETRON TRANSDERMAL SYSTEM TO TREAT/PREVENT CHEMOTHERAPY INDUCED NAUSEA AND VOMITING

U-1012 METHOD FOR TREATING INSOMNIA WHILE REDUCING THE RISK OF AN ADVERSE DRUG INTERACTION

U-1013 METHOD OF USING RIBAVIRIN IN COMBINATION WITH PEGYLATED INTERFERON ALPHA-2B TO TREAT PATIENTS WITH CHRONIC HEPATITIS C

U-1014 METHOD OF USING RIBAVIRIN IN COMBINATION WITH INTERFERON ALPHA-2B (PEGYLATED AND NONPEGYLATED) TO TREAT PATIENTS WITH CHRONIC HEPATITIS C

U-1015 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA

U-1016 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT-EXPERIENCED ADULT PATIENTS, WHO HAVE EVIDENCE OF VIRAL REPLICATION AND HIV-1 STRAINS RESISTANT TO AN NNRTI AND OTHER ANTIRETROVIRAL AGENTS

U-1017 A METHOD OF TREATING NASAL AND NON-NASAL SYMPTOMS OF SEASONAL ALLERGIC RHINITIS

U-1018 TREATMENT OF PULMONARY HYPERTENSION BY INHALATION

U-1019 TREATMENT OF PULMONARY HYPERTENSION

U-1020 METHOD OF USING COLCHICINE FOR THE PROPHYLAXIS OF GOUT FLARES

U-1021 SHORT-TERM TREATMENT (4-8 WEEKS) OF ACTIVE BENIGN GASTRIC ULCER

U-1022 FOR THE PREPARATION OF SKIN PRIOR TO SURGERY; HELPS REDUCE BACTERIA THAT CAN POTENTIALLY CAUSE SKIN INFECTION

U-1023 TREATMENT OF ATROPHIC VAGINITIS DUE TO MENOPAUSE

U-1024 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION WHO REQUIRE ADJUNCTIVE OR REPLACEMENT THERAPY DUE TO INADEQUATELY CONTROLLED IOP

U-1025 TREATING FREQUENT HEARTBURN

U-1026 A METHOD OF TREATING HUMAN SUFFERING FROM OR SUSCEPTIBLE TO PSYCHOSIS.

U-1027 REDUCTION OF ELEVATED PLASMA STEROL AND/OR STANOL LEVELS IN A MAMMAL

U-1028 A METHOD OF DISTRIBUTING SODIUM OXYBATE UNDER CONTROL OF A CENTRAL PHARMACY

U-1029 METHOD FOR TREATING ACUTE ELEVATIONS OF BLOOD PRESSURE IN HUMAN SUBJECT IN NEED THEREOF

U-1030 IMPROVEMENT OF WALKING IN PATIENTS WITH MULTIPLE SCLEROSIS (MS)

U-1031 IMPROVE RESPIRATORY SYMPTOMS IN CYSTIC FIBROSIS IN PATIENTS WITH PSEUDOMONAS AERUGINOSA

U-1032 USE OF ROSUVASTATIN CALCIUM FOR THE PRIMARY PREVENTION OF CARDIOVASCULAR DISEASE IN INDIVIDUALS WITHOUT CLINICALLY EVIDENT CORONARY HEART DISEASE BUT WITH INCREASED RISK FACTORS

U-1033 TOPICAL TREATMENT OF ACNE VULGARIS

U-1034 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) IN ADULTS

U-1035 NONSTEROIDAL ANTI-INFLAMMATORY DRUG INDICATED FOR RELIEF OF MILD TO MODERATE ACUTE PAIN

U-1036 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH INSULIN

U-1037 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH A PPAR-GAMMA AGONIST

U-1038 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH METFORMIN AND A PPAR-GAMMA AGONIST

U-1039 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH METFORMIN

U-1040 INHIBITION OF THROMBIN IN A PATIENT

U-1041 TREATMENT OF DISORDERS RESPONSIVE TO GROWTH HORMONE

U-1042 METHOD FOR STIMULATING CORONARY VASODILATION FOR PURPOSES OF IMAGING THE HEART

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1043 MANAGEMENT OF MODERATE TO SEVERE PAIN

U-1044 TOPICAL TREATMENT OF SCALP PSORIASIS

U-1045 MAINTENANCE TREATMENT IN PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NSCLC WHO HAVE NOT PROGRESSED ON 1ST-LINE TREATMENT WITH PLATINUM-BASED CHEMOTHERAPY

U-1046 MAINTENANCE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NSCLC WHOSE DISEASE HAS NOT PROGRESSED AFTER FOUR CYCLES PLATINUM-BASED CHEMOTHERAPY

U-1047 TREATMENT OF BIOPSY-CONFIRMED, PRIMARY SUPERFICIAL BASAL CELL CARCINOMA (SBCC)

U-1048 WORKS THROUGH THE INDUCTION OF INTERFERON AND OTHER CYTOKINES

U-1049 PROPHYLAXIS OF ORGAN REJECTION IN ADULT PATIENTS AT LOW-MODERATE IMMUNOLOGIC RISK RECEIVING A RENAL TRANSPLANT

U-1050 USE OF METAXALONE FOR TREATMENT OF MUSCULOSKELETAL CONDITIONS

U-1051 TREATMENT OF OROPHARYNGEAL CANDIDIASIS

U-1052 RELIEF OF SIGNS AND SYMPTOMS OF ARTHRITIS AND RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER

U-1053 RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER

U-1054 ONYCHOMYCOSIS OF THE TOENAIL CAUSED BY TRICOPHYTON RUBRUM OR TRICHOPHYTON MENTAGROPHYTES, ONCE DAILY USE FOR 12 CONSECUTIVE WEEKS

U-1055 AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO ARE ALREADY TREATED WITH A THIAZOLIDINEDIONE (TZD) AND METFORMIN OR WHO HAVE INADEQUATE GLYCEMIC CONTROL ON A TZD OR METFORMIN ALONE

U-1056 TREATMENT OF PAIN USING A NASAL SPRAY OF KETOROLAC TROMETHAMINE

U-1057 TREATMENT OF INFLAMMATION AND PAIN USING A NASAL SPRAY OF KETOROLAC TROMETHAMINE

U-1058 USE OF THALIDOMIDE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED MULTIPLE MYELOMA

U-1059 ADJUNCTIVE THERAPY TO DIET TO PATIENTS WITH HYPERTRIGLYCERIDEMIA

U-1060 ADJUNCTIVE THERAPY TO DIET IN PATIENTS WITH ELEVATED CHOLESTEROL AND/OR LIPID LEVELS

U-1061 ADJUNCTIVE THERAPY TO DIET IN PATIENTS WITH MIXED DYSLIPIDEMIA

U-1062 ADMINISTRATION OF APPROVED PRODUCT FOR TREATMENT OF ALZHEIMER'S DISEASE

U-1063 TREATMENT OF ONLY INFLAMMATORY LESIONS (PAPULES AND PUSTULES) OF ROSACEA

U-1064 TREATMENT OF BIPOLAR DISORDER AND SCHIZOPHRENIA

U-1065 METHOD OF TREATING ANDROGEN RESPONSIVE OR MEDICATED CONDITION IN A MAMMAL BY ADMINISTERING A SAFE & EFFECTIVE AMOUNT OF DUTASTERIDE OR A PHARMACEUTICALLY ACCEPTABLE SOLVATE THEREOF.. CONDITIONS INCLUDE BENIGN PROSTATIC HYPERTROPHY

U-1066 METHOD OF TREATING AN ANDROGEN RESPONSE OR MEDIATED DISEASE IN A MAMMAL BY ADMINISTERING AN EFFECTIVE ANDROGEN RESPONSIVE OR MEDICATED DISEASE AMOUNT OF DUTASTERIDE..CONDITIONS INCLUDE BENIGN PROSTATIC HYPERPLASIA

U-1067 TREATMENT OF CANCER

U-1068 TREATMENT OF ASTHMA

U-1069 A METHOD OF TREATING A PATIENT WITH A PRESCRIPTION DRUG USING AN EXCLUSIVE COMPUTER DATABASE IN A COMPUTER SYSTEM FOR DISTRIBUTION

U-1070 A METHOD TO CONTROL ABUSE OF A SENSITIVE DRUG BY CONTROLLING WITH A COMPUTER PROCESSOR THE DISTRIBUTION OF THE SENSITIVE DRUG VIA AN EXCLUSIVITY CENTRAL PHARMACY THAT MAINTAINS A CENTRAL DATABASE

U-1071 METHOD OF TREATING BLADDER DYSFUNCTION WITH ONCE A DAY TROSPIDIUM SALT FORMULATION

U-1072 THE MANAGEMENT OF MODERATE TO SEVERE CHRONIC PAIN IN PATIENTS REQUIRING A CONTINUOUS, AROUND-THE-CLOCK OPIOID ANALGESIC FOR AN EXTENDED PERIOD OF TIME

U-1073 USE FOR THE TREATMENT OF ASTHMA AND COPD

U-1074 USE OF EXENATIDE MAY RESULT IN REDUCTION IN BODY WEIGHT

U-1075 USE FOR THE TREATMENT OF ASTHMA

U-1076 REDUCE CHRONIC SEVERE DROOLING (I.E., SIALORRHEA) IN PATIENTS WITH NEUROLOGIC CONDITIONS ASSOCIATED WITH PROBLEM DROOLING

U-1077 PRETREATMENT OF PATIENTS WITH VITAMIN B12 AND FOLIC ACID PRIOR TO PEMETREXED

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

DISODIUM ADMINISTRATION

- U-1078 TREATMENT OF ACNE
- U-1079 REVLIMID (LENALIDOMIDE) IS INDICATED FOR THE TREATMENT OF PATIENTS WITH TRANSFUSION-DEPENDENT ANEMIA IN MYELODYSPLASTIC SYNDROMES (MDS)
- U-1080 METHOD TO TREAT PULMONARY HYPERTENSION BY ADMINISTERING AMBRISENTAN TO A PATIENT
- U-1081 LUMIGAN IS A PROSTAGLANDIN ANALOG INDICATED FOR THE REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- U-1082 USE OF A COMBINATION OF TOBRAMYCIN AND DEXAMETHASONE TO TREAT OCULAR INFLAMMATION WHERE AN INFECTION OR RISK OF INFECTION EXISTS
- U-1083 ACUTE TREATMENT OF MIGRAINE ATTACKS, WITH OR WITHOUT AURA, AND THE TREATMENT OF CLUSTER HEADACHE EPISODES
- U-1084 RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF CORTICOSTEROID RESPONSIVE DERMATOSES IN PATIENTS 12 YEARS OF AGE OR OLDER
- U-1085 METHOD FOR TREATING IRRITABLE BOWEL SYNDROME AND METHOD FOR TREATING ABDOMINAL DISCOMFORT ASSOCIATED WITH IRRITABLE BOWEL SYNDROME
- U-1086 TREATMENT OF AUTOIMMUNE DISEASE
- U-1087 DETECTION OF NON-MUSCLE INVASIVE PAPILLARY CANCER OF THE BLADDER BY PHOTODYNAMIC CYSTOSCOPY
- U-1088 RELIEF OF MUSCLE SPASM
- U-1089 INHIBITION OF THROMBIN
- U-1090 LO LOESTRIN FE IS INDICATED FOR THE PREVENTION OF PREGNANCY IN WOMEN WHO ELECT TO USE ORAL CONTRACEPTIVES AS A METHOD OF CONTRACEPTION
- U-1091 ASSESSMENT OF BRONCHIAL HYPERRESPONSIVENESS IN PATIENTS 6 YEARS OF AGE OR OLDER WHO DO NOT HAVE CLINICALLY APPARENT ASTHMA
- U-1092 TREATMENT OF BREAST CANCER
- U-1093 TREATMENT OF PSEUDBULBAR AFFECT
- U-1094 MANAGEMENT OF CHRONIC MUSCULOSKELETAL PAIN
- U-1095 METHOD OF TREATING OCULAR INFLAMMATION
- U-1096 TREATMENT OF PATIENTS WITH METASTATIC BREAST CANCER
- U-1097 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHEN TREATMENT WITH BOTH SAXAGLIPTIN AND METFORMIN IS APPROPRIATE
- U-1098 METHOD OF TREATING HYPERPARATHYROIDISM; METHOD OF TREATING HYPERCALCEMIA
- U-1099 TREATMENT OF PAIN, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY, POSTHERPETIC NEURALGIA, AND FIBROMYALGIA
- U-1100 REDUCTION OF EXCESS ABDOMINAL FAT IN HIV-INFECTED PATIENTS WITH LIPODYSTROPHY
- U-1101 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS IN PATIENTS WITH NARCOLEPSY
- U-1102 METHOD OF TREATING CATAPLEXY IN PATIENTS WITH NARCOLEPSY
- U-1103 TESTOSTERONE REPLACEMENT THERAPY IN MALES FOR CONDITIONS ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE
- U-1104 USE OF TRAMADOL FOR THE MANAGEMENT OF MODERATE TO MODERATELY SEVERE CHRONIC PAIN
- U-1105 TOPICAL TREATMENT OF HEAD LICE INFESTATION IN PATIENTS FOUR (4) YEARS OF AGE AND OLDER
- U-1106 TREATING HYPERTRIGLYCERIDEMIAS WITH REDUCTION OF FOOD EFFECT
- U-1107 TREATING HYPERCHOLESTEROLEMIAS WITH REDUCTION OF FOOD EFFECT
- U-1108 TREATING TYPE 2 DIABETES MELLITUS WITH EXENATIDE BY STIMULATING INSULIN RELEASE
- U-1109 TREATMENT OF CUTANEOUS MANIFESTATIONS OF ERYTHEMA NODOSUM LEPROSUM (ENL) IN CONNECTION WITH A SPECIAL PROGRAM APPROVED BY FDA CALLED "SYSTEM FOR THALIDOMIDE EDUCATION AND PRESCRIBING SAFETY" (S.T.E.P.S.)
- U-1110 METHOD OF TREATING A PATIENT WITH A PRESCRIPTION DRUG USING A COMPUTER DATABASE IN A COMPUTER SYSTEM FOR DISTRIBUTION
- U-1111 NONSTEROIDAL ANTI-INFLAMMATORY DRUG INDICATED FOR RELIEF OF MILD TO MODERATE ACUTE PAIN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1112 METHOD OF MR IMAGING OF A MAMMAL

U-1113 TREATMENT AND PROPHYLAXIS OF INFLUENZA

U-1114 TREATMENT WITH GABAPENTIN, INCLUDING TREATMENT OF NEUROPATHIC PAIN, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH POSTHERPETIC NEURALGIA

U-1115 TREATMENT TO REDUCE THE RISK OF COPD EXACERBATIONS IN PATIENTS WITH SEVERE COPD ASSOCIATED WITH CHRONIC BRONCHITIS AND A HISTORY OF EXACERBATIONS

U-1116 METHOD OF ADMINISTERING COLCHICINE TO FAMILIAL MEDITERRANEAN FEVER PATIENTS

U-1117 TREATMENT OF BREAST CANCER

U-1118 USE FOR THE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA

U-1119 CONTRAST AGENT FOR MAGNETIC RESONANCE IMAGING

U-1120 TO REDUCE GASTROINTESTINAL SIDE EFFECTS ADMINISTER WITH A MEAL; AS STARTING DOSE ADMINISTER ONCE DAILY WITH EVENING MEAL

U-1121 METHOD OF TREATING TRAVELERS' DIARRHEA

U-1122 TREATMENT OF SECONDARILY INFECTED TRAUMATIC SKIN LESIONS DUE TO S. AUREUS AND S. PYOGENES

U-1123 TREATMENT OF ALCOHOL DEPENDENCE

U-1124 PREVENTION OF RELAPSE TO OPIOID DEPENDENCE, FOLLOWING OPIOID DETOXIFICATION

U-1125 METHOD FOR THE DETECTION OF NEUROENDOCRINE TUMORS

U-1126 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAVE RECEIVED PRIOR CHEMOTHERAPY CONTAINING DOCETAXEL

U-1127 TREATMENT OF PATENT DUCTUS ARTERIOSUS

U-1128 TREATMENT OF CHRONIC HEPATITIS C (CHC) GENOTYPE 1 INFECTION IN COMBINATION WITH PEGINTERFERON ALFA AND RIBAVIRIN IN ADULT PATIENTS (≥ 18 YEARS OF AGE) WITH COMPENSATED LIVER DISEASE

U-1129 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN

U-1130 SECONDARY PREVENTION OF CARDIOVASCULAR EVENTS BY DOSING ONCE PER DAY IN THE EVENING OR A NIGHT WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN

U-1131 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN

U-1132 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-1133 SECONDARY PREVENTION OF CARDIOVASCULAR EVENTS BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-1134 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-1135 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A) AND INCREASE OF HDL-C

U-1136 SECONDARY PREVENTION OF CARDIOVASCULAR EVENTS BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C

U-1137 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C

U-1138 TREATMENT OF PRIMARY AND MIXED DYSLIPIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-1139 REDUCTION IN RISK OF RECURRENT NONFATAL MYOCARDIAL INFARCTION BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-1140 REDUCTION IN ELEVATED TC AND LDL-C BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-1141 REDUCTION IN TG BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

U-1142 TREATMENT OF PRIMARY AND MIXED DYSLIPIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN

PATENT AND EXCLUSIVITY TERMS

ADB 86 of 133

PATENT USE

- U-1143 REDUCTION IN RISK OF RECURRENT NONFATAL MYOCARDIAL INFARCTION BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1144 REDUCTION IN ELEVATED TC AND LDL-C BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1145 REDUCTION IN TG BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1146 REDUCTION IN TG WITH REDUCED FLUSHING BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1147 TREATMENT OF PRIMARY AND MIXED DYSLIPIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C
- U-1148 REDUCTION IN RISK OF RECURRENT NONFATAL MYOCARDIAL INFARCTION BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C
- U-1149 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1150 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION IN TOTAL-C, LDL-C, TG, LP(A), AND INCREASE OF HDL-C
- U-1151 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION IN TOTAL-C, LDL-C, LP(A), AND INCREASE OF HDL-C
- U-1152 CYANOCOBALAMIN ADMINISTRATION THROUGH NASAL INFUSION
- U-1153 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS, IS INDICATED FOR THE TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS TYPE 1 (HIV-1) INFECTION IN ANTIRETROVIRAL TREATMENT-NAIVE ADULT PATIENTS, AS SET FORTH IN THE LABELING, INCLUDING I&U SECTION
- U-1154 TREATMENT OF PROTEIN KINASE RELATED DISORDERS, SUCH AS GASTROINTESTINAL STROMAL TUMORS, RENAL CELL CARCINOMA AND ADVANCED PANCREATIC NEUROENDOCRINE TUMORS, WITH SUNITINIB
- U-1155 USE OF THALIDOMIDE IN TREATMENT OF CUTANEOUS MANIFESTATIONS OF ERYTHEMA NODOSUM LEPROSUM (ENL)
- U-1156 TO REDUCE BLOOD PHENYLALANINE (PHE) LEVELS IN PATIENTS WITH HYPERPHENYLALANINEMIA (HPA)
- U-1157 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES IN ADULTS AND CHILDREN 2 YEARS OF AGE AND OLDER AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH HIVES (URTICARIA) IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER
- U-1158 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH HIVES (URTICARIA) IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER
- U-1159 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES, SWELLING OF THE NASAL PASSAGES AND SINUS CONGESTION AND PRESSURE IN ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER
- U-1160 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH HIVES (URTICARIA) IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER AND 12 YEARS OF AGE AND OLDER
- U-1161 FOR THE TREATMENT AND PROPHYLAXIS OF GOUT FLARES & THE TREATMENT OF FAMILIAL MEDITERRANEAN FEVER
- U-1162 TREATMENT OF SEBORRHEIC DERMATITIS OF THE SCALP
- U-1163 METHOD OF TREATING THROMBOSIS
- U-1164 METHOD OF TREATING AN ARGATROBAN TREATABLE CONDITION
- U-1165 USE FOR THE TREATMENT OF MULTIPLE MYELOMA
- U-1166 A METHOD FOR TREATMENT OF GOUT FLARES DURING PROPHYLAXIS
- U-1167 PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT)
- U-1168 THE LONG TERM, ONCE-DAILY MAINTENANCE BRONCHODILATOR TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA
- U-1169 MANAGEMENT OF BREAKTHROUGH PAIN IN CANCER PATIENTS 18 YEARS OF AGE AND OLDER WHO ARE RECEIVING AND TOLERANT TO OPIOID THERAPY FOR THEIR UNDERLYING PERSISTENT

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

CANCER PAIN

- U-1170 TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-1171 REDUCTION OF THE RATE OF THROMBOTIC EVENTS IN PATIENTS WITH ACUTE CORONARY SYNDROME
- U-1172 TO REDUCE ELEVATED TOTAL-C, APO B, AND NON-HDL-C IN PATIENTS WITH PRIMARY HYPERLIPIDEMIA BY ADMINISTRATION OF EZETIMIBE IN COMBINATION WITH A STATIN
- U-1173 TO REDUCE ELEVATED TOTAL-C, LDL-C, APO B AND NON-HDL-C IN PATIENTS WITH PRIMARY HYPERLIPIDEMIA BY ADMINISTRATION OF EZETIMIBE ALONE OR IN COMBINATION WITH A STATIN OR WITH FENOFIBRATE
- U-1174 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH STERILE WATER FOR INJECTION, 0.9% SODIUM CHLORIDE INJECTION, OR FLOLAN STERILE DILUENT FOR INJECTION PRIOR TO ADMINISTRATION
- U-1175 REDUCTION OF CARDIAC TISSUE DAMAGE ASSOCIATED WITH MYOCARDIAL INFARCTION
- U-1176 TREATMENT OR PREVENTION OF STROKE
- U-1177 REDUCTION OF CARDIAC TISSUE DAMAGE ASSOCIATED WITH MYOCARDIAL INFARCTION
- U-1178 RELIEF OF MODERATE TO SEVERE CHRONIC PAIN
- U-1179 TREATMENT OF A CANCER MEDIATED BY AN ANAPLASTIC LYMPHOMA KINASE (ALK)
- U-1180 TREATMENT OF THE FOLLOWING INFECTIONS: COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS AND STAPHYLOCOCCUS AUREUS BLOODSTREAM INFECTIONS (BACTEREMIA) INCLUDING THOSE WITH RIGHT-SIDED INFECTIVE ENDOCARDITIS
- U-1181 A METHOD OF TREATING OR PREVENTING OCULAR PAIN IN A PATIENT
- U-1182 TREATMENT OF CYCLIC HEAVY MENSTRUAL BLEEDING
- U-1183 A METHOD FOR ADMINISTERING FOLLICLE STIMULATING HORMONE (FSH) FOR OVARIAN FOLLICLE OR TESTICULAR STIMULATION IN THE HUMAN
- U-1184 TREATMENT OF ERECTILE DYSFUNCTION AND THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA
- U-1185 TREATMENT OF OPIOID-INDUCED CONSTIPATION
- U-1186 ADMINISTRATION OF AN INHALABLE POWDER COMPRISING TIOTROPIUM VIA DEVICE
- U-1187 TREATMENT OF PATHOLOGICAL STATE BY ANTAGONIZING BRADYKININ RECEPTOR INCLUDING TREATMENT OF ACUTE ATTACKS OF HEREDITARY ANGIOEDEMA (HAE)
- U-1188 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE
- U-1189 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH METFORMIN
- U-1190 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH INSULIN
- U-1191 METHOD OF TX TYPE 2 DM IN PTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBO WITH AN AGENT ACTING ON AN ATP-DEPENDENT CHANNEL IN BETA CELLS SUCH AS A SULFYONYLUREA(INCL GLIPIZIDE, GLIMEPIRIIDE & GLYBURIDE)
- U-1192 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH A SULFONYLUREA (SUCH AS GLIPIZIDE, GLIMEPIRIDE AND GLYBURIDE)
- U-1193 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH A PPAR-GAMMA AGONIST (SUCH AS PIOGLITAZONE AND ROSIGLITAZONE)
- U-1194 METHOD FOR TREATING INSOMNIA
- U-1195 PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGE 5, WHICH MAY RESULT IN RENAL OSTEODYSSTROPHY, WHILE AVOIDING HYPERPHOSPHATEMIA
- U-1196 RELIEF OF SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS AND OSTEOARTHRITIS AND TO DECREASE RISK OF DEVELOPING UPPER GASTROINTESTINAL ULCERS IN PATIENTS WHO ARE TAKING IBUPROFEN FOR THOSE INDICATIONS
- U-1197 METHOD OF TREATMENT OF CHILDREN WITH CENTRAL PRECOCIOUS PUBERTY
- U-1198 RECTIV IS A NITRATE VASODILATOR INDICATED FOR THE TREATMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH CHRONIC ANAL FISSURE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1199 TREATMENT AND PREVENTION OF POSTMENOPAUSAL OR GLUCOCORTICOID-INDUCED OSTEOPOROSIS AND TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS

U-1200 REDUCING THE RISK OF STROKE AND SYSTEMIC EMBOLISM

U-1201 FOR THE TREATMENT OF INTERMEDIATE OR HIGH-RISK MYELOFIBROSIS

U-1202 METHOD FOR RELIEVING OR TREATING CONSTIPATION IN A PATIENT WITH IRRITABLE BOWEL SYNDROME

U-1203 METHOD FOR RELIEVING OR TREATING CONSTIPATION IN A HUMAN CONSTIPATED PATIENT

U-1204 TREATMENT OF UVEITIS

U-1205 TREATMENT OF MACULAR EDEMA

U-1206 DELIVERING AN OCULAR IMPLANT AS DESCRIBED IN THE DOSAGE AND ADMINISTRATION SECTION OF THE APPROVED LABELING OF OZURDEX

U-1207 INFANT USE AGED 1 MONTH TO LESS THAN ONE YEAR, GERD AND EROSIIVE ESOPHAGITIS

U-1208 TREATMENT OF HYPOTRICHOSIS OF THE EYELASHES BY INCREASING THEIR GROWTH INCLUDING LENGTH, THICKNESS AND DARKNESS

U-1209 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION IN ADULT PATIENTS, AND TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION IN PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER

U-1210 USE OF REVLIMID (LENALIDOMIDE) WHILE PREVENTING THE EXPOSURE OF A FETUS OR OTHER CONTRAINDICATED INDIVIDUAL TO REVLIMID (LENALIDOMIDE)

U-1211 USE OF REVLIMID (LENALIDOMIDE) TO INHIBIT THE SECRETION OF PRO-INFLAMMATORY CYTOKINES, INCLUDING TUMOR NECROSIS FACTOR ALPHA

U-1212 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF MULTIPLE MYELOMA AND TRANSFUSION-DEPENDENT ANEMIA IN MYELOYDYSPLASTIC SYNDROMES (MDS)

U-1213 TOPICAL TREATMENT OF SEBORRHEIC DERMATITIS IN IMMUNOCOMPETENT PATIENTS 12 YEARS OF AGE AND OLDER

U-1214 METHOD FOR RELIEVING CONSTIPATION IN A HUMAN PATIENT THAT COMPRISES ADMINISTERING TO THE PATIENT A DOSAGE UNIT COMPRISING (I) 24MCG+/- 10% OF A DRUG SUBSTANCE AND (II) A PHARMACEUTICALLY SUITABLE EXCIPIENT

U-1215 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF TRANSFUSION-DEPENDENT ANEMIA IN MYELOYDYSPLASTIC SYNDROMES (MDS)

U-1216 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF MULTIPLE MYELOMA

U-1217 METHOD OF INCREASING HAIR GROWTH

U-1218 METHOD OF STIMULATING HAIR GROWTH

U-1219 METHOD OF INCREASING THE NUMBER OF HAIRS

U-1220 TREATMENT OF RENAL CELL CARCINOMA

U-1221 TO STIMULATE THE IMMUNE SYSTEM TO INDUCE T CELL PROLIFERATION

U-1222 TO INHIBIT THE PROLIFERATIVE ACTIVITY OF NEOPLASTIC CELLS

U-1223 METHOD FOR TREATING TYPE 2 DIABETES USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENATIDE

U-1224 REDUCTIONS IN BODY WEIGHT ARE OBSERVED WITH EXENATIDE

U-1225 ACCELERATING THE TIME TO UPPER AND LOWER GASTROINTESTINAL RECOVERY FOLLOWING PARTIAL LARGE OR SMALL BOWEL RESECTION SURGERY WITH PRIMARY ANASTOMOSIS

U-1226 A METHOD OF PROVIDING A PREDETERMINED CONCENTRATION OF NITRIC OXIDE TO A PATIENT

U-1227 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND METFORMIN HCL EXTENDED RELEASE IS APPROPRIATE

U-1228 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND METFORMIN HCL EXTENDED RELEASE IS APPROPRIATE ALONE OR IN COMBINATION WITH INSULIN

U-1229 TREATMENT OF MILDLY TO MODERATELY ACTIVE ULCERATIVE COLITIS IN MALE PATIENTS

U-1230 A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT

U-1231 TREATMENT OF MODERATE-TO-SEVERE PRIMARY RESTLESS LEG SYNDROME IN ADULTS

U-1232 USE AS ANTICOAGULANT IN PTS W/ UNSTABLE ANGINA UNDERGOING PTCA; W/ PROVISIONAL USE OF GLYCOPROTEIN IIB/IIIA INHIBITOR, AS ANTICOAGULANT IN PTS UNDERGOING PCI AND FOR PTS W/, OR AT RISK OF, HIT/HITTS UNDERGOING PCI.INTENDED FOR USE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

W/ASPIRIN

- U-1233 TREATMENT OF CHRONIC HEPATITIS C (CHC) GENOTYPE 1 INFECTION, ADMINISTERED WITH FOOD
- U-1234 FOR REDUCING TOTAL CHOLESTEROL (TOTAL-C), LDL-C, APO-LIPOPROTEIN B, OR TOTAL TRIGLYCERIDES, AND TREATING HYPERTRIGLYCERIDEMIA
- U-1235 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION
- U-1236 USE OF THALOMID (THALIDOMIDE) FOR THE TREATMENT OF MULTIPLE MYELOMA
- U-1237 COMBO W/ OTHER ANTIRETROVIRALS FOR TX OF HIV-1 IN ANTIRETROVIRAL TX-EXPERIENCED PT 6 YEARS UP, WHO HAVE EVIDENCE OF VIRAL REPLICATION AND HIV-1 STRAINS RESISTANT TO NON-NUCLEOSIDE REVERSE TRANSCRIPTASE INHIBITOR AND OTHER ANTIRETROVIRALS
- U-1238 TREATMENT OF ANEMIA DUE TO CHRONIC KIDNEY DISEASE
- U-1239 MAGNETIC RESONANCE IMAGING OF THE LIVER
- U-1240 TREATMENT OF HEAVY MENSTRUAL BLEEDING IN WOMEN WITHOUT ORGANIC PATHOLOGY WHO CHOOSE TO USE AN ORAL CONTRACEPTIVE AS THEIR METHOD OF CONTRACEPTION
- U-1241 MANAGEMENT OF MODERATE TO SEVERE PAIN BY ORALLY ADMINISTERING AN INTACT COMPOSITION AS CLAIMED
- U-1242 PREVENTION OF RESPIRATORY DISTRESS (RDS) IN PREMATURE INFANTS
- U-1243 WITH DRY HANDS, GENTLY REMOVE THE SUPRENZA (PHENTERMINE HYDROCHLORIDE ODT) TABLET FROM THE BOTTLE. IMMEDIATELY PLACE THE SUPRENZA TABLET ON TOP OF THE TONGUE WHERE IT WILL DISSOLVE, THEN SWALLOW WITH OR WITHOUT WATER
- U-1244 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH SULFONYLUREA
- U-1245 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH PIOGLITAZONE
- U-1246 SINGLE DOSE ADMINISTRATION INTO THE SURGICAL SITE TO PRODUCE POSTSURGICAL ANALGESIA
- U-1247 MANAGEMENT OF POSTHERPETIC NEURALGIA (PHN) IN ADULTS
- U-1248 USE OF TOPICAL DICLOFENAC ON THE KNEE AND A SECOND TOPICAL MEDICATION ON THE SAME KNEE
- U-1249 TREATMENT OF MALE PATIENT HAVING A DISEASE OR CONDITION RESPONSIVE TO A TERATOGENIC DRUG
- U-1250 TREATMENT OF PAIN, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY OR SPINAL CORD INJURY, POSTHERPETIC NEURALGIA, AND FIBROMYALGIA
- U-1251 A METHOD OF CONTROLLING POSTOPERATIVE OCULAR PAIN AND BURNING/STINGING IN A PATIENT
- U-1252 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE
- U-1253 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY
- U-1254 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY CONTROLLING WEIGHT GAIN
- U-1255 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY
- U-1256 TREATMENT OF SEBORRHEIC DERMATITIS
- U-1257 TREATMENT OF OPHTHALMIC DISORDERS
- U-1258 VISUALIZATION DURING VITRECTOMY PROCEDURES
- U-1259 PROPHYLAXIS OF HIV-1 INFECTION
- U-1260 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES INCLUDING BORTEZOMIB AND AN IMMUNOMODULATORY AGENT AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY
- U-1261 REDUCTION OF THE RISK OF HOSPITALIZATION FOR ATRIAL FIBRILLATION
- U-1262 USE OF QSYMIA (PHENTERMINE AND TOPIRAMATE) FOR WEIGHT MANAGEMENT, INCLUDING, BUT NOT LIMITED TO EFFECTING WEIGHT LOSS, TREATING OBESITY, AND/OR TREATING OVERWEIGHT

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1263 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) OR CHRONIC BRONCHITIS

U-1264 TREATMENT OF A RESPIRATORY DISEASE

U-1265 PATENTED METHOD OF USING REPAGLINIDE IN COMBINATION WITH METFORMIN AS INDICATED FOR IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

U-1266 METHOD OF TREATING MIDDLE-OF-THE-NIGHT INSOMNIA

U-1267 TREATMENT OF RHEUMATOID ARTHRITIS BY DELAYED RELEASE FORMULATION OF 1MG OR 2MG OF PREDNISONE

U-1268 TREATMENT OF PULMONARY, GASTROINTESTINAL AND/OR RHEUMATOLOGICAL DISEASES OR CONDITIONS BY USE OF DELAYED RELEASE FORMULATIONS OF 1MG OR 2MG PREDNISONE

U-1269 TREATMENT OF RHEUMATOLOGIC, ALLERGIC, PULMONARY, GASTROINTESTINAL, DERMATOLOGIC DISEASES OR CONDITIONS BY THE USE OF A DELAYED RELEASE 5MG PREDNISONE TABLET

U-1270 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH INSULIN (WITH OR WITHOUT METFORMIN AND/OR PIOGLITAZONE)

U-1271 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME-NEGATIVE (PH-) ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN SECOND OR GREATER RELAPSE OR WHOSE DISEASE HAS PROGRESSED FOLLOWING TWO OR MORE ANTI-LEUKEMIA THERAPIES

U-1272 TREATMENT OF SIGNS AND SYMPTOMS OF PARKINSON'S DISEASE BY APPLICATION OF CLAIMED TRANSDERMAL SYSTEM

U-1273 TREATMENT OF RESTLESS LEGS SYNDROME BY APPLICATION OF CLAIMED TRANSDERMAL DELIVERY SYSTEM

U-1274 TREATMENT OF EXOCRINE PANCREATIC INSUFFICIENCY DUE TO CYSTIC FIBROSIS OR OTHER CONDITIONS

U-1275 TREATMENT OF CHRONIC HEPATITIS B IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER

U-1276 MANAGEMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY

U-1277 METHOD OF INCREASING EYELASH GROWTH INCLUDING LENGTH, THICKNESS, DARKNESS AND/OR NUMBER OF EYELASHES BY ADMINISTERING BIMATOPROST TO AN EYELID MARGIN

U-1278 METHOD OF TREATING IRRITABLE BOWEL SYNDROME WITH CONSTIPATION IN ADULTS

U-1279 TREATMENT OF HIV INFECTION USING A COMPOSITION CONTAINING A PHARMACOKINETIC ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENASE

U-1280 USE OF A CALCIPOTRIENE CONTAINING FOAM FOR THE TREATMENT OF PSORIASIS

U-1281 THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAVE PREVIOUSLY RECEIVED DOCETAXEL

U-1282 PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING

U-1283 A METHOD OF TREATING CHRONIC MYELOGENOUS LEUKEMIA

U-1284 A METHOD OF TREATING A NEOPLASM

U-1285 TREATMENT OF PATIENTS WITH RELAPSING FORMS OF MULTIPLE SCLEROSIS

U-1286 A METHOD OF REDUCING THE RISK OF PULMONARY EDEMA IN PATIENTS IN NEED OF TREATMENT WITH INHALED NITRIC OXIDE

U-1287 METHOD OF REDUCING TG LEVELS IN PATIENT SUFFERING FROM SEVERE HYPERTRIGLYCERIDEMIA

U-1288 TREATMENT OF ERECTILE DYSFUNCTION BY ADMINISTERING A FILM-COATED TABLET

U-1289 MANAGEMENT OF MODERATE TO SEVERE ACUTE PAIN

U-1290 TREATMENT OF LUNG CANCER

U-1291 TREATMENT OF ACUTE PROMYELOCYTIC LEUKEMIA (APL) IN PATIENTS WHOSE APL IS CHARACTERIZED BY THE PRESENCE OF THE (15;17) TRANSLOCATION OR PML/RAR-ALPHA GENE EXPRESSION

U-1292 TREATMENT OF DISEASES OR CONDITIONS BY THE USE OF A DELAYED RELEASE 1, 2, OR 5 MG PREDNISONE TABLET

U-1293 A METHOD OF LOWERING INTRAOCULAR PRESSURE IN A PATIENT WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION

U-1294 METHOD OF TREATING GLAUCOMA IN A PATIENT

U-1295 A METHOD OF TREATING A PATIENT WITH GLAUCOMA OR OCULAR HYPERTENSION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1296 USE OF PEMETREXED WITH PRIOR AND/OR REPEATED VITAMIN B12 AND FOLIC ACID ADMINISTRATION

U-1297 TREATMENT OF PULMONARY ARTERIAL HYPERTENSION BY INHIBITING ENDOTHELIN RECEPTORS

U-1298 ADJUNCTIVE THERAPY IN THE TREATMENT OF PARTIAL SEIZURES

U-1299 TREATMENT OF PATIENTS WITH LEUKEMIA INCLUDING CHRONIC MYELOID/MYELOGENOUS LEUKEMIA (CML)

U-1300 TREATMENT OF PATIENTS WITH TYROSINE KINASE INHIBITOR (TKI) RESISTANT OR INTOLERANT CHRONIC MYELOID/MYELOGENOUS LEUKEMIA (CML)

U-1301 TREATMENT OF DEEP VEIN THROMBOSIS (DVT)

U-1302 TREATMENT OF PULMONARY EMBOLISM (PE)

U-1303 REDUCTION IN THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS (DVT) AND PULMONARY EMBOLISM

U-1304 USE OF ONCE-A-DAY AMOXICILLIN PRODUCT TO TREAT TONSILLITIS AND/OR PHARYNGITIS SECONDARY TO STREPTOCOCCUS PYOGENES

U-1305 TREATMENT OF HIV-1 INFECTION IN ADULT PATIENTS, AND TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER, CO-ADMINISTERED WITH RITONAVIR (PREZISTA/RITONAVIR) AND WITH OTHER ANTIRETROVIRAL AGENTS

U-1306 TREATMENT OF THROMBOCYTOPENIA IN PATIENTS WITH CHRONIC HEPATITIS C TO ALLOW THE INITIATION AND MAINTENANCE OF INTERFERON-BASED THERAPY

U-1307 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT-NAIVE ADULT PATIENTS WITH HIV-1 RNA LESS THAN OR EQUAL TO 100,000 AT THE START OF THERAPY

U-1308 MULTIPLE MYELOMA

U-1309 BONE METASTASES

U-1310 FOR THE MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS

U-1311 METHOD OF TREATING CYSTIC FIBROSIS

U-1312 USE FOR THE TREATMENT OF HYPERGLYCEMIA

U-1313 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

U-1314 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER

U-1315 THE LONG TERM TREATMENT OF PROPHYLACTIC MANAGEMENT OF OCULAR HYPERTENSION AND GLAUCOMA

U-1316 A DOSING REGIMEN FOR THE TREATMENT OF HYPERCHOLESTEROLEMIA AND HYPERLIPIDEMIA IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA USING AT LEAST THREE STEP-WISE INCREASING DOSES

U-1317 TREATMENT OF HYPERCHOLESTEROLEMIA, HYPERLIPIDEMIA AND HYPERLIPOPROTEINEMIA IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA

U-1318 TREATMENT OF HYPERCHOLESTEROLEMIA BY DECREASING THE AMOUNT OR ACTIVITY OF MICROSOMAL TRIGLYCERIDE TRANSFER PROTEIN IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA

U-1319 SYMPTOMATIC RELIEF OF NON-INFECTIOUS DIARRHEA

U-1320 TREATMENT OF ADULT PATIENTS WITH SHORT BOWEL SYNDROME WHO ARE DEPENDENT ON PARENTERAL SUPPORT

U-1321 TREATMENT OF PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS

U-1322 METHOD OF REDUCING OCULAR HYPERTENSION

U-1323 REDUCING THE RISK OF STROKE

U-1324 MANAGEMENT OF CYSTIC FIBROSIS PATIENTS

U-1325 INDUCTION OF REMISSION IN PATIENTS WITH ACTIVE, MILD TO MODERATE ULCERATIVE COLITIS

U-1326 METHOD OF INDUCING CONTRACEPTION IN A FEMALE OF REPRODUCTIVE AGE WHO HAS NOT YET REACHED PREMENOPAUSE

U-1327 METHOD FOR TREATING ACUTE MIGRAINE IN ADULTS, WITH OR WITHOUT AURA, COMPRISING IONTOPHORETIC TRANSDERMAL DELIVERY OF SUMATRIPTAN OR A SALT THEREOF, USING A FLOWABLE HYDROGEL FORMULATION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-1328 METHOD FOR TREATING ACUTE MIGRAINE IN ADULTS, WITH OR WITHOUT AURA, COMPRISING IONTOPHORETIC TRANSDERMAL DELIVERY OF SUMATRIPTAN OR A SALT THEREOF
- U-1329 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER
- U-1330 METHODS OF TREATING LIPID METABOLISM AND GLYCOMETABOLISM DISORDERS COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-1331 METHODS OF REDUCING THE AMOUNT OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-1332 METHODS OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-1333 METHODS OF LOWERING ELEVATED POST PRANDIAL BLOOD GLUCOSE LEVELS COMPRISING ADMINISTERING A DIPEPTIDYL PEPTIDASE INHIBITOR
- U-1334 METHODS OF TREATING DIABETES COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-1335 METHODS OF MODIFYING GLUCOSE METABOLISM AND TREATING DIABETES COMPRISING ADMINISTERING A DIPEPTIDYL PEPTIDASE INHIBITOR AND ONE OR MORE OTHER THERAPEUTIC AGENTS SUCH AS METFORMIN
- U-1336 METHODS OF TREATING DIABETES COMPRISING ADMINISTERING A DIPEPTIDYL PEPTIDASE INHIBITOR AND METFORMIN
- U-1337 METHOD OF TREATING DIABETES COMPRISING ADMINISTERING ALOGLIPTIN
- U-1338 METHOD OF TREATING DIABETES COMPRISING ADMINISTERING A COMPOUND SUCH AS ALOGLIPTIN
- U-1339 METHODS OF TREATING DIABETES COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1340 METHODS OF TREATING LIPID METABOLISM DISORDERS COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1341 METHODS OF TREATING GLYCOMETABOLISM DISORDERS COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1342 METHODS OF REDUCING THE AMOUNT OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1343 METHODS OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1344 METHODS OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN PREPARATION
- U-1345 USE IN RELIEVING OR PREVENTING CONSTIPATION IN A HUMAN PATIENT WITH A DOSAGE UNIT COMPRISING 24MICROG+/- 10% OF A DRUG SUBSTANCE AND A PHARMACEUTICALLY SUITABLE EXCIPIENT
- U-1346 USE OF FEBUXOSTAT FOR THE MANAGEMENT OF HYPERURICEMIA IN PATIENTS SUFFERING FROM GOUT AND, WHEN USED WITH THEOPHYLLINE WITHOUT THE NEED FOR DOSE ADJUSTMENT OF THEOPHYLLINE
- U-1347 TREATMENT OF A SKIN DISORDER
- U-1348 TREATMENT OF OSTEOARTHRITIS
- U-1349 TREATMENT OF JUVENILE RHEUMATOID ARTHRITIS
- U-1350 TREATMENT OF ANKYLOSING SPONDYLITIS
- U-1351 TREATMENT OF ACUTE PAIN
- U-1352 TREATMENT OF PRIMARY DYSMENORRHEA
- U-1353 ADJUNCTIVE THERAPY TO LIPID-LOWERING MEDICATIONS AND DIET TO REDUCE LOW DENSITY LIPOPROTEIN-CHOLESTEROL, APOLIPOPROTEIN B, TOTAL CHOLESTEROL, AND NON-HIGH DENSITY LIPOPROTEIN CHOLESTEROL IN PTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-1354 INHIBITION OF PREMATURE LH SURGES IN WOMEN UNDERGOING CONTROLLED OVARIAN HYPERSTIMULATION WITH FSH
- U-1355 MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN ADULT AND ADOLESCENT PATIENTS 12 YEARS OF AGE AND OLDER. PATENT CLAIMS METHOD FOR TREATING A RESPIRATORY DISEASE IN A CHILD
- U-1356 TREATMENT OF NASAL SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER. TREATMENT OF NASAL SYMPTOMS ASSOCIATED W PERENNIAL ALLERGIC RHINITIS IN ADULTS AND ADOLESCENTS 12 YEARS OF AGE AND OLDER
- U-1357 TREATMENT OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS IN ADULTS AND ADOLESCENTS 12 YEARS OF AGE AND OLDER. PATENT CLAIMS METHODS FOR TREATING A RESPIRATORY DISEASE IN A CHILD
- U-1358 TREATMENT OF BACTERIAL INFECTIONS IN THE NASAL PASSAGE OF ADULT PATIENTS AND HEALTH CARE WORKERS WITH METHICILLIN RESISTANT S. AUREUS
- U-1359 USE OF POMALIDOMIDE TO INHIBIT THE SECRETION OF PRO-INFLAMMATION CYTOKINES, INCLUDING TUMOR NECROSIS FACTOR ALPHA
- U-1360 USE OF POMALIDOMIDE FOR THE TREATMENT OF MULTIPLE MYELOMA
- U-1361 USE OF POMALIDOMIDE WHILE PREVENTING THE EXPOSURE OF A FETUS OR OTHER CONTRAINDICATED INDIVIDUAL TO POMALIDOMIDE
- U-1362 TREATMENT OF DISEASES OR CONDITIONS BY THE USE OF A DELAYED-RELEASE 1,2, OR 5MG PREDNISONE TABLET
- U-1363 A METHOD OF TREATING OR PREVENTING OCULAR PAIN AND BURNING/STINGING FOLLOWING CORNEAL SURGERY
- U-1364 MAINTENANCE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)
- U-1365 PROPHYLAXIS OF ALLOGRAFT REJECTION IN ADULT PATIENTS RECEIVING A LIVER TRANSPLANT
- U-1366 TREATMENT OF INFERTILITY THROUGH INDUCTION OF OVULATION AND PREGNANCY TO ANOVULATORY INFERTILE WOMEN
- U-1367 METHOD OF ADMINISTERING FSH FOR THE TREATMENT OF INFERTILITY THROUGH INDUCTION OF OVULATION AND PREGNANCY IN ANOVULATORY INFERTILE WOMEN
- U-1368 TREATMENT OF SOLID EXCRETORY SYSTEM TUMORS; ADVANCED RENAL CELL CARCINOMA (RCC), AFTER FAILURE OF TREATMENT WITH SUNITINIB OR SORAFENIB
- U-1369 TREATMENT OF VAGINAL SYMPTOMS OF UROGENITAL ATROPHY BY ORALLY ADMINISTERING OSPEMIFENE WITH FOOD TO ENHANCE BIOAVAILABILITY OF OSPEMIFENE
- U-1370 TREATMENT OF DYSpareunia ASSOCIATED WITH MENOPAUSE
- U-1371 REDUCTION OF INTRAOCULAR PRESSURE IN PATIENTS WITH ELEVATED INTRAOCULAR PRESSURE OR GLAUCOMA
- U-1372 ADMINISTRATION WITHOUT FOOD FOR TREATMENT OF HIV-1 INFECTION
- U-1373 METHOD OF TREATING ACETAMINOPHEN OVERDOSE WITH ACETYLCYSTEINE SOLUTIONS
- U-1374 TREATMENT OF PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH+CML)
- U-1375 ADASUVE IS A TYPICAL ANTIPSYCHOTIC INDICATED FOR THE ACUTE TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA OR BIPOLAR I DISORDER IN ADULTS
- U-1376 TREATMENT OF INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS
- U-1377 IMPROVE RESPIRATORY SYMPTOMS IN CYSTIC FIBROSIS IN PATIENTS WITH PSEUDOMONAS AERUGINOSA
- U-1378 TREATMENT OF A NITROGEN METABOLISM DISORDER
- U-1379 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO HAVE ONE OR MORE SPECIFIED CARDIOVASCULAR RISK FACTORS
- U-1380 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO HAVE ONE OR MORE SPECIFIED CARDIOVASCULAR RISK FACTORS WHEREIN THE PATIENT HAS CARDIOVASCULAR DISEASE
- U-1381 USE OF PRASUGREL AND ASPIRIN IN PATIENTS REQUIRING THE REDUCTION OF THROMBOTIC CARDIOVASCULAR EVENTS
- U-1382 TREATMENT OF NAUSEA AND VOMITING OF PREGNANCY IN WOMEN WHO DO NOT RESPOND TO CONSERVATIVE MANAGEMENT
- U-1383 DOSAGE ADJUSTMENT OF A NITROGEN SCAVENGING DRUG IN THE TREATMENT OF A UREA CYCLE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

DISORDER

- U-1384 METHOD OF TREATING MULTIPLE SCLEROSIS
- U-1385 METHOD OF TREATING AN AUTOIMMUNE DISEASE SELECTED FROM AUTOIMMUNE POLYARTHRITIS AND MULTIPLE SCLEROSIS BUT NOT TREATING PSORIATIC ARTHRITIS
- U-1386 A METHOD OF INCREASING THE TESTOSTERONE BLOOD LEVEL OF A PERSON IN NEED THEREOF
- U-1387 REDUCTION IN RISK OF HOSPITALIZATION IN PATIENTS WITH A HISTORY OF PAROXYSMAL OR PERSISTENT AF WITHOUT SEVERE HEART FAILURE AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS
- U-1388 TREATMENT OF PATIENTS WITH A HISTORY OF PAROXYSMAL OR PERSISTENT AF WITHOUT SEVERE HEART FAILURE AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS
- U-1389 ELLA IS A PROGESTERONE AGONIST/ANTAGONIST EMERGENCY CONTRACEPTION INDICATED FOR THE PREVENTION OF PREGNANCY FOLLOWING UNPROTECTED INTERCOURSE OR A KNOWN OR SUSPECTED CONTRACEPTIVE FAILURE. ELLA CAN BE TAKEN WITH OR WITHOUT FOOD
- U-1390 A METHOD OF INCREASING THE TESTOSTERONE BLOOD LEVEL OF AN ADULT MALE SUBJECT IN NEED THEREOF
- U-1391 METHOD FOR TREATING OPIOID-INDUCED CONSTIPATION
- U-1392 METHOD OF RELIEVING OR PREVENTING CONSTIPATION IN A HUMAN PATIENT WITH OPIOID-INDUCED CONSTIPATION
- U-1393 METHOD FOR RELIEVING OR TREATING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION
- U-1394 METHOD FOR RELIEVING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION THAT COMPRISES ADMINISTERING TO THE PATIENT A DOSAGE UNIT COMPRISING (I) 24MICROG +/- 10% OF A DRUG SUBSTANCE AND (II) A PHARMACEUTICALLY SUITABLE EXCIPIENT
- U-1395 USE IN RELIEVING OR PREVENTING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION WITH A DOSAGE UNIT COMPRISING 24MICROG +/- 10% OF A DRUG SUBSTANCE AND A PHARMACEUTICALLY SUITABLE EXCIPIENT
- U-1396 TREATMENT OF ADVANCED HORMONE RECEPTOR POSITIVE, HER2-NEGATIVE BREAST CANCER IN COMBINATION WITH EXEMESTANE AFTER FAILURE OF TREATMENT WITH LETROZOLE OR ANASTROZOLE
- U-1397 USE AS AN ANTISEPTIC FOR THE PREPARATION OF A PATIENT'S SKIN PRIOR TO SURGERY
- U-1398 METHOD OF TREATING CHRONIC HEPATITIS C
- U-1399 MANAGEMENT OF NEPHROPATHIC CYSTINOSIS BY ADMINISTERING A TOTAL DAILY DOSE IN TWO DIVIDED DOSES
- U-1400 FOR THE TREATMENT OF PRIMARY HYPERLIPIDEMIA, MIXED HYPERLIPIDEMIA OR HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- U-1401 INDICATED FOR LONG-TERM, ONCE-DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PTS WITH COPD, INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA, ALSO TO REDUCE EXACERBATIONS OF COPD IN PTS WITH A HISTORY OF EXACERBATIONS
- U-1402 FOR USE IN THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) AND/OR INDOLENT B-CELL NON-HODGKIN LYMPHOMA (NHL)
- U-1403 FIRST-LINE TREATMENT OF METASTATIC NON SMALL-CELL LUNG CANCER (NSCLC) WITH EGFR EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- U-1404 METHOD FOR TREATING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION BY OPENING CIC CHANNELS
- U-1405 THERAPEUTIC TREATMENT OF BONE METASTASES
- U-1406 TREATMENT OF MELANOMA
- U-1407 TREATMENT OF NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH + CML)
- U-1408 TREATMENT OF PLAQUE PSORIASIS IN PATIENTS 18 YEARS OF AGE OR OLDER
- U-1409 TREATMENT OF HIV-1 BY ONCE DAILY ADMINISTRATION
- U-1410 TREATMENT OF CORTICOSTEROID-RESPONSIVE DERMATOSES
- U-1411 THIS DRUG IS ADMINISTERED BY SUBLINGUAL ROUTE TO HUMANS FOR MAINTENANCE TREATMENT OF OPIOID DEPENDENCE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1412 TREATMENT OF ATOPIC DERMATITIS

U-1413 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH FLOLAN STERILE DILUENT FOR INJECTION PRIOR TO INFUSION

U-1414 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF MANTLE CELL LYMPHOMA (MCL)

U-1415 TREATING A PATIENT HAVING A CONDITION SUSCEPTIBLE TO TREATMENT WITH METHYLPHENIDATE, SUCH AS ADHD, BY ADMINISTERING THE FORMULATION RECITED IN CLAIMS 1 OR 2

U-1416 USE OF FENOFIBRATE FOR REDUCING ELEVATED TOTAL CHOLESTEROL (TOTAL-C), LDL-C, APO-LIPOPROTEIN B, OR TOTAL TRIGLYCERIDES

U-1417 USE FOR TREATMENT OF HELICOBACTER INFECTIONS

U-1418 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAFV600E MUTATION AS DETECTED BY AN FDA APPROVED TEST

U-1419 TREATMENT OF ACUTE CYANIDE POISONING THAT IS JUDGED TO BE LIFE THREATENING

U-1420 METHOD OF ONCE A DAY ADMINISTRATION

U-1421 SUBLINGUAL ADMINISTRATION OF A PHARMACEUTICAL COMPOSITION COMPRISING BUPRENORPHINE

U-1422 METHOD OF TREATING PATIENTS NEEDING AN IRON SUPPLEMENT

U-1423 AMYVID IS A RADIOACTIVE DIAGNOSTIC AGENT FOR POSITRON EMISSION TOMOGRAPHY (PET) IMAGING OF THE BRAIN TO ESTIMATE BETA-AMYLOID NEURITIC PLAQUE DENSITY IN ADULT PATIENTS WITH COGNITIVE IMPAIRMENT

U-1424 LONG-TERM, ONCE DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PTS WITH COPD, INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA, ALSO TO REDUCE EXACERBATIONS OF COPD IN PATIENTS WITH A HISTORY OF EXACERBATIONS

U-1425 SUBLINGUAL ADMINISTRATION OF A PHARMACEUTICAL COMPOSITION COMPRISING BUPRENORPHINE AND NALOXONE

U-1426 USE FOR TREATMENT OF DIAPER DERMATITIS COMPLICATED BY CANDIDIASIS

U-1427 ALKYLATING DRUG INDICATED FOR THE TOPICAL TREATMENT OF STAGE IA AND IB MYCOSIS FUNGOIDES-TYPE CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO HAVE RECEIVED PRIOR SKIN DIRECTED THERAPY

U-1428 TOPICAL TREATMENT OF FACIAL ERYTHEMA OF ROSACEA

U-1429 TREATMENT OF PATIENTS WITH BREAST CANCER WHOSE TUMORS OVEREXPRESS THE HER2 RECEPTOR

U-1430 TREATMENT OF ALLERGIC RHINITIS, INCLUDING SEASONAL AND PERENNIAL ALLERGIC RHINITIS

U-1431 METHOD OF TREATING HYPERGLYCEMIA TO IMPROVE GLYCEMIC CONTROL IN A PATIENT BY ORAL ADMIN OF ONCE A DAY OSMOTIC DOSAGE FORM OF GLIPIZIDE WITH POLYETHYLENE OXIDE, HYDROXYPROPYLMETHYLCELLULOSE, CELLULOSE ACETATE, AND SODIUM CHLORIDE

U-1432 METHOD OF TREATMENT OF IRON-RELATED CONDITIONS WITH AT LEAST 0.6 GRAMS OF ELEMENTAL IRON VIA AN IRON CARBOHYDRATE COMPLEX

U-1433 IMPROVEMENTS OF GLYCEMIC CONTROL IN INDIVIDUALS WITH TYPE 2 DIABETES WHO HAVE ONE OR MORE SPECIFIED CARDIOVASCULAR RISK FACTORS

U-1434 TREATMENT OF PANCREATIC CANCER

U-1435 COMBINATION USE OF TOPICAL DICLOFENAC ON THE KNEE AND ADMINISTRATION OF AN ORAL NSAID.

U-1436 USE OF TOPICAL DICLOFENAC ON THE KNEE AND A SECOND TOPICAL AGENT SELECTED FROM SUNSCREEN AND INSECT REPELLANT

U-1437 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH STERILE DILUENT FOR FLOLAN OR STERILE DILUENT FOR EPOPROSTENOL SODIUM PRIOR TO ADMINISTRATION

U-1438 ZINGO INTRADERMAL INJECTION SYSTEM IS A DRUG DELIVERY SYSTEM THAT IS CAPABLE OF DELIVERING FINE DRY POWDERED LIDOCAINE HYDROCHLORIDE MONOHYDRATE FOR LOCAL ANESTHETIC ACTION

U-1439 METHOD OF TREATING AN AFFECTIVE DISORDER SUCH AS DEPRESSION

U-1440 USE OF INGENOL MEBUTATE TO TREAT ACTINIC KERATOSIS

U-1441 A METHOD OF TREATING OR REDUCING OCULAR PAIN AND BURNING/STINGING

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1442 SUBCUTANEOUS INJECTION OF METHOTREXATE

U-1443 ACCELERATING THE TIME TO UPPER AND LOWER GASTROINTESTINAL RECOVERY FOLLOWING SURGERIES THAT INCLUDE PARTIAL BOWEL RESECTION WITH PRIMARY ANASTOMOSIS

U-1444 A DOSING REGIMEN OF AGGRASTAT (TIROFIBAN HYDROCHLORIDE) (25MCG/KG FOLLOWED BY 0.15MCG/KG/MIN INFUSION) TO REDUCE THE RATE OF THROMBOTIC CORONARY EVENTS ASSOCIATED WITH ACUTE CORONARY SYNDROME (ACS) IN PATIENTS WITH NON-ST ELEVATION ACS

U-1445 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION BY ADMINISTERING A PHARMACEUTICAL COMPOSITION COMPRISING MACITENTAN AND A POLYSORBATE, WHEREIN THE POLYSORBATE REPRESENTS 0.1 TO 1% OF THE WEIGHT OF SAID PHARMACEUTICAL COMPOSITION

U-1446 METHOD OF TREATING PULMONARY HYPERTENSION COMPRISING ADMINISTERING MACITENTAN IN COMBINATION WITH A COMPOUND HAVING PHOSPHODIESTERASE-5 INHIBITORY PROPERTIES

U-1447 TREATING PRIMARY HYPERCHOLESTEROLEMIA AND MIXED DYSLIPIDEMIA

U-1448 TREATING SEVERE HYPERTRIGLYCERIDEMIA

U-1449 METHOD OF ALLEVIATING A SKIN CONDITION

U-1450 TREATMENT OF ALLERGIC RHINITIS SYMPTOMS

U-1451 APPROVED INDICATIONS: APTIOM (ESLICARBAZEPINE ACETATE) IS INDICATED AS ADJUNCTIVE TREATMENT OF PARTIAL-ONSET SEIZURES AND APPROVED IN PATIENTS WITH EPILEPSY. PATENT CLAIMS: IN A METHOD OF TREATING A SUBJECT AFFLICTED WITH EPILEPSY

U-1452 METHOD FOR CHRONIC WEIGHT MANAGEMENT

U-1453 A METHOD OF TREATING HYPOXIC RESPIRATORY FAILURE BY VERIFYING GAS INFORMATION OF NITRIC OXIDE PRIOR TO DELIVERY TO PATIENT

U-1454 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS

U-1455 TREATMENT OF PERIANAL WARTS

U-1456 TREATMENT OF MANTLE CELL LYMPHOMA

U-1457 A METHOD OF PURGING A NITRIC OXIDE DELIVERY SYSTEM

U-1458 A METHOD OF REDUCING INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION

U-1459 TREATMENT OF CARCINOMA OF THE THYROID

U-1460 TREATMENT OF HERPES LABIALIS

U-1461 A METHOD OF GENERATING AN INJECTABLE FOAM OF CONTROLLED DENSITY AND BUBBLE SIZE

U-1462 A METHOD OF USING A SCLEROSING AGENT FOR THE TREATMENT OF INCOMPETENT GREAT SAPHENOUS VEINS, ACCESSORY SAPHENOUS VEINS AND VISIBLE VARICOSITIES OF THE GREAT SAPHENOUS (GSV) SYSTEM ABOVE AND BELOW THE KNEE

U-1463 A METHOD OF INTRAVENOUS INJECTION USING ULTRASOUND GUIDANCE, ADMINISTERED VIA A SINGLE CANNULA INTO THE LUMEN OF THE TARGET INCOMPETENT TRUNK VEINS OR BY DIRECT INJECTION INTO VARICOSITIES

U-1464 TREATMENT OF OPIOID DEPENDENCE/SUBLINGUAL OR BUCCAL APPLICATION

U-1465 USE OF THALIDOMIDE WHILE PREVENTING THE EXPOSURE OF A FETUS OR OTHER CONTRAINDICATED INDIVIDUAL TO THALIDOMIDE

U-1466 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER

U-1467 METHOD OF TREATING HEPATITIS C

U-1468 CONTROL OF PHOSPHOROUS LEVELS IN PATIENTS

U-1469 USE OF PHOSLYRA FOR REDUCTION OF SERUM PHOSPHOROUS IN PATIENTS

U-1470 FOR THE TREATMENT OF HEPATITIS C

U-1471 A METHOD FOR TREATING CARDIOVASCULAR DISEASE COMPRISING ADMINISTERING A RECONSTITUTED LYOPHILIZED PHARMACEUTICAL COMPOSITION COMPRISING EPOPROSTENOL, ARGININE AND SODIUM HYDROXIDE.

U-1472 INTENSIVE CARE UNIT SEDATION, INCLUDING SEDATION OF NON-INTUBATED PATIENTS PRIOR TO AND/OR DURING SURGICAL AND OTHER PROCEDURES

U-1473 MANAGEMENT OF RISK OF DRONEDARONE/BETA-BLOCKER INTERACTION IN PATIENTS IN SINUS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

RHYTHM WITH A HISTORY OF PAROXYSMAL OR PERSISTENT AF

U-1474 A METHOD FOR THE TREATMENT OF A PATIENT SUFFERING FROM A DISEASE TREATABLE WITH ROTIGOTINE, COMPRISING APPLYING THE CLAIMED TRANSDERMAL DELIVERY SYSTEM (TDS) TO THE SKIN OF THE PATIENT

U-1475 USE OF ORENITRAM FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1).

U-1476 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA.

U-1477 USE OF TOPICAL DICLOFENAC ON THE KNEE AND A SECOND TOPICAL PRESCRIPTION MEDICATION ON THE SAME KNEE

U-1478 METHOD OF REDUCING TG LEVELS IN PATIENT ON STATIN THERAPY SUFFERING FROM SEVERE HYPERTRIGLYCERIDEMIA

U-1479 INCREASE TEAR PRODUCTION TO TREAT PATIENTS WITH KERATOCONJUNCTIVITIS SICCA (DRY EYE).

U-1480 TREATMENT OF ADVANCED RENAL CELL CARCINOMA

U-1481 REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) RECURRENCE

U-1482 DICLOFENAC POTASSIUM FOR RELIEF OF MILD TO MODERATE ACUTE PAIN

U-1483 INCREASE TEAR PRODUCTION IN PATIENTS WITH KERATOCONJUNCTIVITIS SICCA (DRY EYE).

U-1484 COMBINATION PRODUCT FOR THE EARLY TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) TO REDUCE THE LIKELIHOOD OF ULCERATIVE COLD SORES AND TO SHORTEN THE LESION HEALING TIME IN ADULTS AND CHILDREN (6 YEARS OF AGE AND OLDER)

U-1485 TREATING A SUBJECT UNDERGOING ABDOMINAL SURGERY BY ADMINISTERING ALVIMOPAN TO ACCELERATE THE TIME TO UPPER AND LOWER GASTROINTESTINAL RECOVERY FOLLOWING SURGERIES THAT INCLUDE PARTIAL BOWEL RESECTION WITH PRIMARY ANASTOMOSIS

U-1486 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER

U-1487 METHOD OF INCREASING EYELASH GROWTH

U-1488 USE OF TOPICAL DICLOFENAC FOR TREATING PAIN

U-1489 USE OF TOPICAL DICLOFENAC ON A JOINT FOR TREATING OSTEOARTHRITIS

U-1490 FOR USE IN PATIENTS HAVING SYMPTOMATIC OR PROGRESSIVE MEDULLARY THYROID CANCER, WITH UNRESECTABLE LOCALLY ADVANCED OR METASTATIC DISEASE

U-1491 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA

U-1492 TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER

U-1493 METHOD FOR PREVENTING ITCHING ASSOCIATED WITH ALLERGIC CONJUNCTIVITIS

U-1494 SUBLINGUAL OR BUCCAL ADMINISTRATION OF A PHARMACEUTICAL COMPOSITION COMPRISING BUPRENORPHINE AND NALOXONE

U-1495 RISK REDUCTION OF REBLEEDING IN PTS FOLLOWING THERAPEUTIC ENDOSCOPY FOR ACUTE BLEEDING GASTRIC OR DUODENAL ULCERS IN ADULTS.

U-1496 METHOD TO TREAT HEMANGIOMA.

U-1497 NEURACEQ IS A RADIOACTIVE DIAGNOSTIC AGENT FOR POSITRON EMISSION TOMOGRAPHY (PET) IMAGING OF THE BRAIN TO ESTIMATE P-AMYLOID NEURITIC PLAQUE DENSITY IN ADULT PATIENTS WITH COGNITIVE IMPAIRMENT

U-1498 METHOD OF TREATING PATIENTS WITH GASTRIC RETENTIVE DOSAGE FORM

U-1499 MANAGEMENT OF ACUTE PAIN IN PATIENTS REQUIRING OPIOID ANALGESIA

U-1500 TESTOSTERONE REPLACEMENT THERAPY IN ADULT MALES FOR CONDITIONS ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE; PRIMARY HYPOGONADISM (CONGENITAL OR ACQUIRED); HYPOGONADOTROPIC HYPOGONADISM (CONGENITAL OR ACQUIRED).

U-1501 PROPHYLAXIS OF DEEP VEIN THROMBOSIS AND PULMONARY EMBOLISM

U-1502 PROPHYLAXIS OF PULMONARY EMBOLISM

U-1503 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH METFORMIN

U-1504 USE OF OTEZLA (APREMILAST) FOR INHIBITING PDE4

U-1505 USE OF OTEZLA (APREMILAST) FOR THE TREATMENT OF PSORIATIC ARTHRITIS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1506 TREATMENT OF PATIENTS WITH GASTROINTESTINAL STROMAL TUMOR (GIST), INCLUDING BUT NOT LIMITED TO PATIENTS PREVIOUSLY TREATED WITH IMATINIB AND PATIENTS WITH GIST HAVING RESISTANCE TO A KIT TYROSINE KINASE INHIBITOR

U-1507 TO MAINTAIN HEALING OF EE AND RELIEF OF HEARTBURN

U-1508 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT BY ORALLY ADMINISTERING A PLURALITY OF COMPOSITE SUBUNITS AS CLAIMED

U-1509 TREATMENT OF FREQUENT HEARTBURN BY ADMINISTERING A GASTRIC ACID REDUCER

U-1510 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT BY ORALLY ADMINISTERING AN INTACT COMPOSITION AS CLAIMED.

U-1511 TREATMENT OF HYPERTRIGLYCERIDEMIA

U-1512 REDUCTION OF THROMBOTIC CARDIOVASCULAR EVENTS

U-1513 TREATMENT OF HIV-1 INFECTION IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS

U-1514 MANAGEMENT OF BREAKTHROUGH PAIN IN PATIENTS WITH CANCER BY BUCCAL OR SUBLINGUAL ADMINISTRATION OF FENTANYL

U-1515 METHOD OF TREATING IRRITABLE BOWEL SYNDROME WITH CONSTIPATION IN ADULT PATIENTS.

U-1516 METHOD OF TREATING CHRONIC IDIOPATHIC CONSTIPATION IN ADULT PATIENTS.

U-1517 TREATMENT OF BACTERIAL INFECTIONS USING A TWO-DOSE REGIMEN OF DALBAVANCIN.

U-1518 MAINTAINING PUPIL SIZE BY PREVENTING INTRAOPERATIVE MIOSIS AND REDUCING POSTOPERATIVE OCULAR PAIN

U-1519 METHOD FOR THE LONG TERM TREATMENT OF CHRONIC CONSTIPATION IN A HUMAN SUBJECT WITH IRRITABLE BOWEL SYNDROME

U-1520 METHOD FOR THE LONG TERM TREATMENT OF CHRONIC CONSTIPATION IN A HUMAN SUBJECT

U-1521 MAINTENANCE TREATMENT OF OPIOID DEPENDENCE

U-1522 TREATMENT OF TYPE 2 DIABETES MELLITUS IN A PATIENT, WHEREIN GLYCEMIC CONTROL (HBA1C < 7.0%) IS NOT ACHIEVABLE USING ONE OR MORE OF INSULIN, METFORMIN, PIOGLITAZONE, OR ROSIGLITAZONE

U-1523 METHOD OF INDUCING TOPICAL ANESTHESIA IN THE EYE

U-1524 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE

U-1525 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS IN PATIENTS WITH NARCOLEPSY

U-1526 THE TREATMENT OF PATIENTS WITH TRAVELERS' DIARRHEA (TD) OR THE REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) RECURRENCE

U-1527 FOR THE TREATMENT OF OVERACTIVE BLADDER (OAB) WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY

U-1528 A METHOD OF LOWERING INTRAOCULAR PRESSURE

U-1529 ADJUNCTIVE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)

U-1530 USE OF ARIPIRAZOLE IN EXTENDED RELEASE INJECTABLE SUSPENSION

U-1531 METHOD FOR TRANSDERMAL DELIVERY OF TESTOSTERONE

U-1532 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS AND/OR CATAPLEXY IN NARCOLEPSY PATIENTS WITH SODIUM OXYBATE WHEN DIVALPROEX SODIUM IS CONCOMITANTLY ADMINISTERED.

U-1533 PULMONARY ADMINISTRATION OF PARTICLES COMPRISING A DIKETOPIPERAZINE AND INSULIN.

U-1534 ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN COMPLEXED WITH A DIKETOPIPERAZINE.

U-1535 ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN COMPLEXED WITH MICROPARTICLES OF A DIKETOPIPERAZINE.

U-1536 ADMINISTRATION OF A COMPOSITION COMPRISING A DIKETOPIPERAZINE AND INSULIN.

U-1537 TREATMENT OF A PATIENT HAVING DIABETES MELLITUS WITH A PRANDIAL RAPID ACTING INSULIN.

U-1538 ADMINISTRATION OF FDKP MICROPARTICLES COMPRISING INSULIN.

U-1539 PULMONARY ADMINISTRATION OF AN INSULIN COMPOSITION COMPRISING FDKP AT THE BEGINNING OF A MEAL TO A PATIENT ALSO BEING TREATED WITH A LONG-ACTING INSULIN.

U-1540 BUTRANS IS A PARTIAL OPIOID AGONIST PRODUCT INDICATED FOR THE MANAGEMENT OF PAIN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE.
- U-1541 TREATMENT OF PATIENTS WITH TUBEROUS SCLEROSIS COMPLEX (TSC) WHO HAVE SUBEPENDYMAL GIANT CELL ASTROCYTOMA (SEGA) THAT REQUIRES THERAPEUTIC INTERVENTION BUT CANNOT BE CURATIVELY RESECTED.
- U-1542 FOR USE IN THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA AND/OR NON-HODGKINS LYMPHOMA
- U-1543 TREATMENT OF A PATIENT BY ADMINISTERING THE FORMULATION RECITED IN CLAIM 1 OR CLAIM 23
- U-1544 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA (PTCL).
- U-1545 A METHOD OF TRANSDERMALLY DELIVERING TESTOSTERONE
- U-1546 FOR USE IN THE TREATMENT OF MALIGNANT HYPERTHERMIA IN CONJUNCTION WITH APPROPRIATE SUPPORTIVE MEASURES AND FOR THE PREVENTION OF MALIGNANT HYPERTHERMIA IN PATIENTS AT HIGH RISK.
- U-1547 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), CHRONIC BRONCHITIS OR EMPHYSEMA
- U-1548 FOR THE LONG-TERM, ONCE-DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH COPD, INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA, ALSO TO REDUCE EXACERBATIONS OF COPD IN PATIENTS WITH A HISTORY OF EXACERBATIONS
- U-1549 FOR THE TREATMENT OF PATIENTS WITH RELAPSED CHRONIC LYMPHOCYTIC LEUKEMIA
- U-1550 METHOD OF TREATING METASTATIC PAPILLARY RENAL CELL CARCINOMA WITH TEMSIROLIMUS.
- U-1551 METHOD OF TREATING PAPILLARY RENAL CELL CARCINOMA WITH TEMSIROLIMUS, IN THE ABSENCE OF INTERFERON ALPHA.
- U-1552 FOR HEALING OF ALL GRADES OF EROSIVE ESOPHAGITIS (EE)
- U-1553 TO MAINTAIN HEALING OF EE AND RELIEF OF HEARTBURN
- U-1554 FOR THE TREATMENT OF HEARTBURN ASSOCIATED WITH SYMPTOMATIC NON-EROSIVE GASTROESOPHAGEAL DISEASE (GERD)
- U-1555 MANAGEMENT OF MODERATE TO SEVERE PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE.
- U-1556 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE
- U-1557 A METHOD OF TESTOSTERONE REPLACEMENT THERAPY COMPRISING THE STEP OF NASALLY ADMINISTERING TO A PATIENT IN NEED OF SUCH TREATMENT AN EFFECTIVE AMOUNT OF TESTOSTERONE GEL FORMULATION.
- U-1558 FOR THE TREATMENT OF PATIENTS WITH RELAPSED FOLLICULAR B-CELL NON-HODGKIN LYMPHOMA OR [RELAPSED] SMALL LYMPHOCYTIC LYMPHOMA
- U-1559 INDICATED FOR THE ONCE-DAILY MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN PATIENTS AGED 12 YEARS OF AGE AND OLDER
- U-1560 A METHOD OF DISRUPTING LEUKOCYTE FUNCTION, INCLUDING AS AN INHIBITOR OF PI3KDELTA KINASE
- U-1561 USE OF OTEZLA (APREMILAST) FOR THE TREATMENT OF PSORIATIC ARTHRITIS
- U-1562 TREATMENT OF PATIENTS WITH HEPATIC ENCEPHALOPATHY (HE)
- U-1563 A METHOD OF TRANSDERMAL ADMINISTRATION OF A PHYSIOLOGICALLY ACTIVE AGENT TO A SUBJECT.
- U-1564 A METHOD OF TREATING GAUCHER'S DISEASE
- U-1565 METHOD OF TREATING, AS INITIAL LOADING DOSE FOR MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL ONSET-SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS OR OLDER WHEN ORAL ADMINISTRATION IS TEMPORARILY NOT FEASIBLE
- U-1566 METHOD OF TREATING, AS MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER
- U-1567 METHOD OF TREATING, AS INITIAL LOADING DOSE FOR MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL ONSET-SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS OR OLDER
- U-1568 METHOD OF TREATING, AS MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER WHEN ORAL ADMINISTRATION IS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

TEMPORARILY NOT FEASIBLE

U-1569 TREATMENT OF BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS

U-1570 TREATMENT OF BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS USING A SINGLE DOSE

U-1571 TREATMENT OF GAUCHER DISEASE TYPE 1

U-1572 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION.

U-1573 USE OF RUXOLITINIB (JAKAFI) FOR INHIBITING JANUS ASSOCIATED KINASES (JAKS) JAK1 AND/OR JAK2.

U-1574 A METHOD OF CATALYZING THE HYDROLYSIS OF GLUCOCEREBROSIDE TO GLUCOSE AND CERAMIDE.

U-1575 PATIENTS WITH SEVERE APLASTIC ANEMIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO IMMUNOSUPPRESSIVE THERAPY

U-1576 TREATMENT OF LEUKEMIA

U-1577 CONTROL OF SERUM PHOSPHOROUS LEVELS

U-1578 TREATMENT OF ACUTE OTITIS MEDIA

U-1579 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER

U-1580 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAD RECEIVED PRIOR DOCETAXEL CHEMOTHERAPY

U-1581 IN COMBINATION WITH DABRAFENIB FOR THE TREATMENT OF UNRESECTABLE OR METASTATIC MELANOMA.

U-1582 TREATMENT OF UNRESECTABLE OR METASTATIC MELANOMA

U-1583 FOR CHRONIC WEIGHT MANAGEMENT FOR TREATING OVERWEIGHT OR OBESITY

U-1584 USE OF NALTREXONE AND BUPROPION IN A LAYERED FORMULATION FOR CHRONIC WEIGHT MANAGEMENT FOR AFFECTING WEIGHT LOSS

U-1585 USE OF NALTREXONE AND BUPROPION BASED ON AN ESCALATING DOSE SCHEDULE

U-1586 FOR EFFECT ON BLOOD GLUCOSE PARAMETERS IN PATIENTS WITH INSULIN RESISTANCE

U-1587 SINGLE-DOSE INFILTRATION INTO THE SURGICAL SITE TO PRODUCE POSTSURGICAL ANALGESIA.

U-1588 THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (CRPC).

U-1589 METHOD OF USE FOR REDUCING BLOOD PHENYLALANINE LEVELS IN A HUMAN SUFFERING FROM HYPERPHENYLALANINEMIA

U-1590 KUVAN IS INDICATED TO REDUCE BLOOD PHENYLALANINE LEVELS IN PATIENTS WITH HYPERPHENYLALANINEMIA

U-1591 TREATMENT OF ASTHMA IN PATIENTS AGED 12 YEARS AND OLDER

U-1592 TO REDUCE SERUM PHOSPHATE IN PATIENTS WITH END STAGE RENAL DISEASE

U-1593 MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD, INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA, AND REDUCTION OF EXACERBATIONS IN COPD PATIENTS.

U-1594 DILATION OF THE PUPIL

U-1595 USE OF OTEZLA (APREMILAST) FOR THE TREATMENT OF PSORIASIS

U-1596 LAMICTAL IS AN ANTIEPILEPTIC DRUG (AED) INDICATED FOR: EPILEPSY-ADJUNCTIVE THERAPY IN PATIENTS GREATER THAN OR EQUAL TO 2 YEARS OF AGE: (1.1) PARTIAL SEIZURES PRIMARY GENERALIZED TONIC-CLONIC SEIZURES

U-1597 TREATMENT OF DIABETIC MACULAR EDEMA

U-1598 METHOD OF ADMINISTRATION OF CONTROLLED RELEASE OXYMORPHONE

U-1599 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS, REDUCTION IN FEVER THROUGH ANTI INFLAMMATORY, ANALGESIC, AND ANTIPYRETIC ACTIVITY

U-1600 DOSAGE MODIFICATION FOLLOWING ELEVATED LIVER ENZYMES IN TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS

U-1601 DOSE ESCALATION OVER 14 DAYS FOR TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS

U-1602 METHOD OF ADMINISTERING PIRFENIDONE CAPSULES TO TREAT A FIBROTIC CONDITION

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1603 METHOD FOR ADMINISTERING PIRFENIDONE TO REDUCE DRUG INTERACTIONS WITH FLUVOXAMINE

U-1604 METHOD FOR ADMINISTERING PIRFENIDONE TO REDUCE DRUG INTERACTIONS WITH A STRONG INHIBITOR OF CYP1A2

U-1605 METHOD FOR ADMINISTERING PIRFENIDONE TO AVOID REDUCED EFFICACY BY DISCONTINUING SMOKING OR BY DISCONTINUING OR AVOIDING ANOTHER STRONG CYP1A2 INDUCER

U-1606 METHOD FOR ADMINISTERING PIRFENIDONE WHILE AVOIDING OR DISCONTINUING CONCOMITANT USE OF A MODERATE TO STRONG INHIBITOR OF BOTH CYP1A2 AND ANOTHER CYP ENZYME INVOLVED IN PIRFENIDONE METABOLISM

U-1607 METHOD OF ADMINISTERING A DOSAGE FORM THAT INCLUDES A GRANULATE FORMULATION OF PIRFENIDONE TO TREAT A FIBROTIC CONDITION

U-1608 DOSE ESCALATION OVER 14 DAYS FOR TREATMENT OF A FIBROSIS CONDITION

U-1609 CONTINUED DOSING OR DOSAGE MODIFICATION FOLLOWING ELEVATED LIVER ENZYMES IN TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS

U-1610 CONTINUED DOSING OR DOSAGE MODIFICATION FOLLOWING ELVATED LIVER ENZYMES IN USE OF PIRFENIDONE

U-1611 METHOD FOR ADMINISTERING PIRFENIDONE TO AVOID REDUCED EFFICACY BY DISCONTINUING SMOKING OR BY DISCONTINUING A STRONG CYP1A2 INDUCER

U-1612 METHOD FOR ADMINISTERING PIRFENIDONE TO AVOID REDUCED EFFICACY BY AVOIDING SMOKING OR BY AVOIDING ANOTHER STRONG CYP1A2 INDUCER

U-1613 DOSAGE MODIFICATION IN TREATMENT WITH PIRFENIDONE TO REDUCE DRUG INTERACTIONS WITH CIPROFLOXACIN

U-1614 USE OF TOPICAL DICLOFENAC SODIUM FOR TREATING PAIN

U-1615 FOR THE TREATMENT OF PATIENTS WITH CLL, FL, OR SLL

U-1616 NASAL ADMINISTRATION OF A TESTOSTERONE GEL TO A PATIENT TO TREAT THE PATIENT FOR A CONDITION ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE

U-1617 METHOD OF TREATING MEDULLARY THYROID CANCER

U-1618 A METHOD OF TREATING A PATIENT SUFFERING FROM A PAIN ASSOCIATED SLEEP DISTURBANCE COMPRISING ADMINISTERING A LIQUID COMPOSITION FORMULATED INSIDE A SOFT GEL CAPSULE, AS CLAIMED, TO THE PATIENT

U-1619 TREATMENT OF IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP)

U-1620 METHOD OF TREATMENT OF IRON-RELATED CONDITIONS WITH AT LEAST 0.6 GRAMS OF ELEMENTAL IRON VIA AN IRON CARBOHYDRATE COMPLEX, WITH A SUBSTANTIALLY NON-IMMUNOGENIC CARBOHYDRATE COMPONENT, IN ABOUT 15 MINUTES OR LESS.

U-1621 PULMONARY ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN BOUND TO A COMPLEXING AGENT.

U-1622 FOR THE TREATMENT OF POLYCYTHEMIA VERA

U-1623 USE OF EXENATIDE MAY RESULT IN REDUCTION IN APPETITE.

U-1624 TREATMENT OF UNRESECTABLE HEPATOCELLULAR CARCINOMA, ADVANCED RENAL CELL CARCINOMA, OR DIFFERENTIATED THYROID CARCINOMA.

U-1625 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING ILOPERIDONE TO A PATIENT BY REDUCING THE DOSE IN PATIENTS WHO ARE POOR METABOLIZERS OF CYP2D6

U-1626 A METHOD OF TREATING OR PREVENTING OCULAR PAIN AND BURNING

U-1627 TREATMENT OF ACUTE UNCOMPLICATED INFLUENZA IN ADULTS

U-1628 METHOD OF TREATING DISORDERS WITH AN ETIOLOGY COMPRISING OR ASSOCIATED WITH EXCESS GH-SECRETION

U-1629 METHOD OF TREATING ACROMEGALY

U-1630 TREATMENT IN COMBINATION WITH A CORTICOID SUCH AS PREDNISONE OF PROSTATE CANCER PREVIOUSLY TREATED WITH DOCETAXEL

U-1631 TREATMENT OF INFLAMMATORY LESIONS OF ROSACEA.

U-1632 TREATMENT OF SCHIZOPHRENIA, WITH EFFICACY IN TREATING ACUTE EPISODES OF SCHIZOPHRENIA

U-1633 USE OF ARIPIPRAZOLE IN EXTENDED RELEASE INJECTABLE SUSPENSION IN TREATING ACUTE EPISODES OF SCHIZOPHRENIA

U-1634 TREATMENT OF BRCA MUTATED OVARIAN CANCER USING PARP INHIBITOR

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-1635 USE OF RITONAVIR AS A POTENT CYP3A INHIBITOR TO INCREASE PLASMA DRUG CONCENTRATION OF PARITAPREVIR AND OVERALL DRUG EXPOSURE FOR TREATMENT OF HCV INFECTION
- U-1636 USE OF DASABUVIR TO INHIBIT VIRAL REPLICATION FOR THE TREATMENT OF HCV INFECTION.
- U-1637 TREATMENT OF HCV INFECTION USING PARITAPREVIR, OMBITASVIR, RITONAVIR, AND DASABUVIR WITH RIBAVIRIN.
- U-1638 TREATMENT OF HCV INFECTION USING PARITAPREVIR
- U-1639 USE OF NALTREXONE AND BUPROPION IN EXTENDED-RELEASE FORM FOR CHRONIC WEIGHT MANAGEMENT FOR TREATING OVERWEIGHT OR OBESITY
- U-1640 TREATMENT OF MODERATE TO SEVERE CHRONIC PAIN BY ADMINISTERING AN INTACT COMPOSITION AS CLAIMED
- U-1641 MEMANTINE HCL/DONEPEZIL HCL COMBINATION FOR THE TREATMENT OF MODERATE TO SEVERE DEMENTIA OF THE ALZHEIMER'S TYPE
- U-1642 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS WITH SEVERE CHRONIC RENAL IMPAIRMENT AND FOR WHOM METFORMIN THERAPY IS INAPPROPRIATE BY ADMINISTERING LINAGLIPTIN
- U-1643 TREATING CUSHING'S SYNDROME
- U-1644 TREATMENT OF OVERACTIVE BLADDER BY APPLICATION OF OXYBUTYNIN CHLORIDE GEL TO SKIN
- U-1645 TREATMENT OF PARKINSON'S DISEASE, POST-ENCEPHALITIC PARKINSONISM, AND PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION
- U-1646 TREATMENT OF POST-ENCEPHALITIC PARKINSONISM, AND PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION
- U-1647 TREATMENT OF PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION
- U-1648 TREATMENT OF PATIENTS WITH PARKINSON'S DISEASE, POST-ENCEPHALITIC PARKINSONISM, AND PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION
- U-1649 TREATMENT OF POST-ENCEPHALITIC PARKINSONISM
- U-1650 TREATMENT OF WALDENSTROM'S MACROGLOBULINEMIA
- U-1651 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN
- U-1652 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN AND METFORMIN
- U-1653 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN (WITH OR WITHOUT METFORMIN)
- U-1654 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN (WITH OR WITHOUT INSULIN OR A SULFONYLUREA)
- U-1655 A METHOD TO ACCELERATE THE TIME TO GASTROINTESTINAL RECOVERY BY ADMINISTERING ABOUT 12 MG OF ALVIMOPAN TO THE PATIENT FROM ABOUT 30 TO 60 MINUTES PRIOR TO SURGERY
- U-1656 METHOD OF IRON ADMINISTRATION TO TREAT PATIENTS IN NEED OF IRON REPLACEMENT
- U-1657 METHOD FOR PROVIDING POST COITAL CONTRACEPTION TO A WOMAN BY ADMINISTERING ABOUT 30 MG OF ULIPRISTAL ACETATE WITHIN ABOUT 120 HOURS AFTER INTERCOURSE, WHEREIN THE WOMAN IS OVERWEIGHT HAVING A BMI OF 25 TO 29.99
- U-1658 TREATMENT OF ER-POSITIVE, HER2-NEGATIVE ADVANCED BREAST CANCER IN COMBINATION WITH LETROZOLE AS INITIAL ENDOCRINE-BASED THERAPY FOR METASTATIC DISEASE IN POSTMENOPAUSAL WOMEN
- U-1659 MANAGEMENT OF PAIN
- U-1660 TREATMENT OF HIV-1 INFECTION IN ADULTS WITH NO DARUNAVIR RESISTANCE-ASSOCIATED SUBSTITUTIONS
- U-1661 RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCERS IN PATIENTS ALSO TAKING LOW DOSE ASPIRIN
- U-1662 A METHOD OF TREATING OCULAR PAIN

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1663 TREATMENT OF HIV-1 INFECTION

U-1664 TREATMENT OF BACTERIAL VAGINOSIS WITH METRONIDAZOLE GEL

U-1665 METHOD OF TREATING ATTENTION DEFICIT HYPERACTIVITY DISORDER BY ADMINISTERING THE COMPOSITION OF CLAIM 1

U-1666 PALLIATIVE TREATMENT OF PROSTATE CANCER

U-1667 TREATMENT OF ALLERGIC RHINITIS, INCLUDING SEASONAL ALLERGIC RHINITIS

U-1668 METHOD OF TREATING DEPRESSION OR MAJOR DEPRESSIVE DISORDER

U-1669 TREATMENT OF MULTIPLE MYELOMA, IN COMBINATION WITH BORTEZOMIB AND DEXAMETHASONE

U-1670 NATROBA TOPICAL SUSPENSION IS A PEDICULICIDE INDICATED FOR THE TOPICAL TREATMENT OF HEAD LICE INFESTATION IN PATIENTS SIX (6) MONTHS OF AGE AND OLDER.

U-1671 TREATMENT OF OCULAR ITCHING ASSOCIATED WITH CONJUNCTIVITIS

U-1672 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTION

U-1673 TREATMENT OF COMPLICATED URINARY TRACT INFECTION, INCLUDING PYELONEPHRITIS

U-1674 DOSAGE MODIFICATION TO REDUCE RISKS ASSOCIATED WITH QT PROLONGATION NOT INDUCED BY OTHER DRUGS DURING TREATMENT WITH ILOPERIDONE

U-1675 USE OF TROKENDI XR FOR THE TREATMENT OF EPILEPSY

U-1676 METHODS FOR TREATING BACTERIAL INFECTIONS

U-1677 TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS (IPF)

U-1678 FOR THE TREATMENT OF PATIENTS WITH CLL, FL, OR SLL

U-1679 TREATMENT OF ACUTE OTITIS EXTERNA

U-1680 TREATMENT OF OCULAR ITCHING ASSOCIATED WITH ALLERGIC CONJUNCTIVITIS

U-1681 TREATMENT OF PATIENTS WITH PROGRESSIVE NEUROENDOCRINE TUMORS OF PANCREATIC ORIGIN (PNET) THAT ARE UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC

U-1682 TREATMENT OF BACTERIAL VAGINOSIS

U-1683 TREATMENT FOR CHRONIC LYMPHOCYTIC LEUKEMIA WITH 17P DELETION

U-1684 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA

U-1685 DOSAGE MODIFICATION TO REDUCE THE RISK ASSOCIATED WITH QT PROLONGATION NOT INDUCED BY OTHER DRUGS DURING TREATMENT WITH ILOPERIDONE

U-1686 A METHOD TO REDUCE WITHDRAWAL SYMPTOMS, INCLUDING NICOTINE CRAVING, ASSOCIATED WITH SMOKING CESSATION

U-1687 TREATMENT OF HCV INFECTION USING OMBITASVIR

U-1688 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY IN AN INDIVIDUAL WHO DOES NOT HAVE SEVERE RENAL IMPAIRMENT OR ESRD

U-1689 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY IN AN INDIVIDUAL WHO DOES NOT HAVE SEVERE RENAL IMPAIRMENT OR ESRD

U-1690 METHOD FOR REDUCTION OF SUBMENTAL FAT

U-1691 INDICATED FOR THE ONCE-DAILY INHALED TREATMENT FOR ASTHMA IN ADULTS AGED 18 YEARS AND OLDER

U-1692 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE IN AN INDIVIDUAL WHO DOES NOT HAVE SEVERE RENAL IMPAIRMENT OR ESRD

U-1693 METHOD OF TREATING ADHD IN CHILDREN 6 YEARS OF AGE AND OLDER AND ADOLESCENTS

U-1694 A METHOD FOR TREATING HEART FAILURE IN A HUMAN USING A CRYSTALLINE FORM OF IVABRADINE HYDROCHLORIDE

U-1695 METHOD FOR TREATING THYROID CARCINOMA INCLUDING DIFFERENTIATED THYROID CANCER

U-1696 TREATMENT OF UNRESECTABLE HEPATOCELLULAR CARCINOMA

U-1697 PULMONARY ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN BOUND TO A DIKETOPIPERAZINE.

U-1698 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS

U-1699 A METHOD FOR TREATING ACUTE LYMPHOBLASTIC LEUKEMIA

U-1700 A METHOD FOR TREATING PHILADELPHIA CHROMOSOME POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1701 A METHOD FOR TREATING LEUKEMIA RESULTING FROM A MUTATION IN THE BCR-ABL KINASE DOMAIN

U-1702 TREATMENT OF COPD

U-1703 TREATMENT OF RESPIRATORY COMPLAINTS

U-1704 USE FOR TREATMENT IN PATIENTS WITH DIABETES

U-1705 USE FOR TREATMENT IN PATIENTS WITH HYPERGLYCEMIA

U-1706 TREATMENT OF TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE WHEREIN THE COMBINED THERAPEUTIC EFFECT IS GREATER THAN THE ADDITIVE EFFECT OF ADMINISTERING EACH AGENT ALONE

U-1707 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS AND SYMPTOMS THEREOF.

U-1708 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS.

U-1709 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) WITH VIBERZI (ELUXADOLINE).

U-1710 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH FLUVOXAMINE

U-1711 FOR THE TREATMENT OF PATIENTS WITH CLL, FL OR SLL

U-1712 MEKINIST IN COMBINATION WITH DABRAFENIB FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA

U-1713 TAFINLAR IN COMBINATION WITH TRAMETINIB FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA

U-1714 TREATMENT OF THROMBOCYTOPENIA IN ADULT AND PEDIATRIC PATIENTS 6 YEARS AND OLDER WITH CHRONIC IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP)

U-1715 P2Y12 PLATELET INHIBITOR FOR USE AS ADJUNCT TO PERCUTANEOUS CORONARY INTERVENTION TO REDUCE RISK OF VARIOUS DISEASES/CONDITIONS IN PATIENTS NOT TREATED WITH A P2Y12 PLATELET INHIBITOR AND NOT GIVEN A GLYCOPROTEIN IIB/IIIA INHIBITOR

U-1716 TREATMENT OF COUGH AND SYMPTOMS ASSOCIATED WITH UPPER RESPIRATORY ALLERGIES OR A COMMON COLD WITH CODEINE PHOSPHATE AND CHLORPHENIRAMINE MALEATE ORALLY ADMINISTERED EXTENDED RELEASE TABLETS

U-1717 METHOD OF TREATING CYSTIC FIBROSIS IN PATIENTS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE

U-1718 METHOD OF TREATING CYSTIC FIBROSIS IN PATIENTS WHO HAVE THE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE.

U-1719 ACUTE TREATMENT OF MIGRAINE

U-1720 METHOD OF PROVIDING A THERAPEUTICALLY EFFECTIVE AND STABLE MEDIAN BLOOD PLASMA LEVEL OF LEVODOPA

U-1721 USE OF RUXOLITINIB (JAKAFI) FOR BLOCKING SIGNAL TRANSDUCTION OF JANUS ASSOCIATED KINASES (JAKS) JAK1 AND/OR JAK2

U-1722 TREATMENT OF BASAL CELL CARCINOMA

U-1723 TREATMENT OF HEART FAILURE

U-1724 METHOD OF INHIBITING HEPATITIS C VIRUS

U-1725 METHOD OF INHIBITING HEPATITIS C VIRUS WITH DAKLINZA AND AT LEAST ONE ADDITIONAL COMPOUND HAVING ANTI-HCV ACTIVITY

U-1726 REDUCTION IN RISK OF HOSPITALIZATION IN PATIENTS WITH CORONARY HEART DISEASE AND A HISTORY OF PAROXYSMAL OR PERSISTENT AF AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS

U-1727 TOPICAL TREATMENT OF INFLAMMATORY PAPULES AND PUSTULES OF MILD TO MODERATE ROSACEA

U-1728 REDUCTION IN RISK OF HOSPITALIZATION IN PATIENTS WITH STABLE NYHA CLASS III HEART FAILURE AND A HISTORY OF PAROXYSMAL OR PERSISTENT AF AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS

U-1729 REDUCE THE RISK OF RECURRENT DEEP VEIN THROMBOSIS (DVT)

U-1730 REDUCE THE RISK OF RECURRENT PULMONARY EMBOLISM

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1731 TEMPORARY RELIEF OF MINOR ACHES AND PAINS

U-1732 TEMPORARY REDUCTION OF FEVER

U-1733 TREATMENT/PREVENTION OF CARDIOVASCULAR DISEASE

U-1734 USE OF FLIBANSERIN OR A PHARMACEUTICALLY ACCEPTABLE ACID ADDITION SALT THEREOF TO TREAT HYPOACTIVE SEXUAL DESIRE DISORDER (HSDD)

U-1735 METHODS OF TREATING PAIN, INFLAMMATION AND/OR FEVER WITH INTRAVENOUS IBUPROFEN SUCH THAT MEAN ARTERIAL BLOOD PRESSURE DOES NOT INCREASE THE DOSAGE INTERVAL

U-1736 TREATMENT OF THROMBOCYTOPENIA IN ADULT AND PEDIATRIC PATIENTS 1 YEAR AND OLDER WITH CHRONIC IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP)

U-1737 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING ILOPERIDONE TO A PATIENT BY REDUCING THE DOSE IN PATIENTS WHO ARE BEING TREATED WITH FLUOXETINE

U-1738 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) WITH VIBERZI (ELUXADOLINE)

U-1739 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND -THE-CLOCK, LONG-TERM OPIOID TREATMENT, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY

U-1740 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT-NAIVE PATIENTS WITH HIV-1 RNA LESS THAN OR EQUAL TO 100,000 AT THE START OF THERAPY

U-1741 PREVENTION OF DELAYED NAUSEA AND VOMITING ASSOCIATED WITH EMETOGENIC CANCER CHEMOTHERAPY

U-1742 ROLAPITANT IS APPROVED FOR THE PREVENTION OF DELAYED NAUSEA AND VOMITING (I.E., EMESIS) ASSOCIATED WITH EMETOGENIC CANCER CHEMOTHERAPY

U-1743 FOR THE PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH CHEMOTHERAPY

U-1744 PREVENTION OF POST-OPERATIVE NAUSEA AND VOMITING

U-1745 FOR THE TREATMENT OF PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA

U-1746 MONOTHERAPY OR ADJUNCTIVE THERAPY FOR TREATMENT OF PARTIAL-ONSET SEIZURES AND APPROVED IN PATIENTS WITH EPILEPSY

U-1747 FOR CLAIMS 1-3,6-13,16-24 AND 26-32: METHOD OF TREATING ADHD

U-1748 FOR CLAIMS 1-4,6-14,16-24 AND 26-32: METHOD OF TREATING ADHD IN CHILDREN 6 YEARS OF AGE AND OLDER AND ADOLESCENTS

U-1749 ACUTE TREATMENT OF MANIC AND MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER

U-1750 TREATMENT OF SCHIZOPHRENIA AND/OR ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER WITH CARIPRAZINE

U-1751 TREATMENT OF PATIENTS WITH METASTATIC COLORECTAL CANCER WHO HAVE BEEN PREVIOUSLY TREATED WITH FLUOROPYRIMIDINE-, OXALIPLATIN- AND IRINOTECAN-BASED CHEMOTHERAPY, AN ANTI-VEGF BIOLOGICAL THERAPY, AND IF RAS WILD-TYPE, AN ANTI-EGFR THERAPY

U-1752 PROPHYLAXIS OF ORGAN REJECTION

U-1753 TREATMENT OF HCV INFECTION USING DASABUVIR

U-1754 FOR THE TREATMENT OF PULMONARY HYPERTENSION (PAH) IN COMBINATION WITH TADALAFIL

U-1755 FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS

U-1756 METHODS OF TREATING PAIN, INFLAMMATION AND/OR FEVER IN A CRITICALLY ILL PATIENT WITH INTRAVENOUS IBUPROFEN IN NEED THEREOF

U-1757 INHIBITION ON PI3K KINASE

U-1758 METHOD OF TREATING ALLERGIC REACTION VIA INJECTION

U-1759 METHOD OF REVERSING THE ANTICOAGULANT EFFECT OF DABIGATRAN USING IDARUCIZUMAB

U-1760 RISK-REDUCTION OF NSAID GASTRIC ULCER IN PATIENTS REQUIRING CHRONIC NSAID TREATMENT

U-1761 PLAQUE PSORIASIS

U-1762 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT

PATENT AND EXCLUSIVITY TERMS

ADB 106 of 133

PATENT USE

- U-1763 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1764 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1765 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1766 TREATMENT OF HYPERKALEMIA
- U-1767 USE OF CALCIPOTRIENE FOAM FOR THE TOPICAL TREATMENT OF PLAQUE PSORIASIS IN PATIENTS AGED 18 YEARS AND OLDER
- U-1768 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN
- U-1769 TREATMENT OF PAIN BY TRANSMUCOSAL DELIVERY OF BUPRENORPHINE
- U-1770 TREATMENT OF SCHIZOPHRENIA WITH IMPROVEMENT IN NEGATIVE SYMPTOMS AND/OR COGNITIVE DYSFUNCTION OF SCHIZOPHRENIA
- U-1771 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH STERILE WATER FOR INJECTION OR 0.9% SODIUM CHLORIDE INJECTION PRIOR TO ADMINISTRATION
- U-1772 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN
- U-1773 LONG - TERM MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-1774 USE OF A LOTION CONTAINING HALOBETASOL PROPIONATE FOR THE TREATMENT OF CORTICOSTEROID-RESPONSIVE
- U-1775 USE OF A LOTION CONTAINING HALOBETASOL PROPIONATE FOR THE TREATMENT OF CORTICOSTEROID-RESPONSIVE DERMATOSES INCLUDING PSORIASIS
- U-1776 METHOD OF USING COBIMETINIB FOR THE TREATMENT OF MELANOMA
- U-1777 TREATMENT OF PATIENTS WITH METASTATIC EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) T790M MUTATION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC), WHO HAVE PROGRESSED ON OR AFTER EGFR TKI THERAPY
- U-1778 METHOD FOR TREATING MULTIPLE MYELOMA
- U-1779 METHOD FOR TREATING MULTIPLE MYELOMA WITH ONE OR MORE OTHER THERAPEUTIC AGENTS
- U-1780 METHOD FOR TREATING CANCER, INCLUDING MULTIPLE MYELOMA
- U-1781 RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER IN PATIENTS REQUIRING NSAID TREATMENT
- U-1782 FOR HEAD LICE INFESTATIONS
- U-1783 METHOD OF TREATING FREQUENT HEARTBURN BY ADMINISTERING AN ESOMEPRAZOLE MAGNESIUM AS CLAIMED
- U-1784 METHOD OF TREATING FREQUENT HEARTBURN BY ADMINISTERING AN ESOMEPRAZOLE MAGNESIUM TRIHYDRATE AS CLAIMED
- U-1785 METHOD OF TREATING FREQUENT HEARTBURN BY ADMINISTERING AN ESOMEPRAZOLE MAGNESIUM FORMULATION AS CLAIMED
- U-1786 TREATMENT OF PATIENTS WITH RELAPSING FORMS OF MULTIPLE SCLEROSIS WHILE MANAGING THE RISK OF TERIFLUNOMIDE AND ROSUVASTATIN INTERACTION BY LIMITING THE ROSUVASTATIN DOSE TO NO MORE THAN 10MG AND/OR ADMINISTERING ABOUT HALF THE NORMAL DOSE
- U-1787 TREATMENT OF EXOCRINE PANCREATIC INSUFFICIENCY
- U-1788 TREATMENT OF PATIENT HAVING DIABETES MELLITUS VIA ORAL INHALATION OF FDKP MICROPARTICLES COMPRISING INSULIN
- U-1789 METHOD OF ADMINISTERING AN ETHANOL-FREE TAXANE LIQUID NANODISPERSION FORMULATION TO A SUBJECT COMBINING THE FORMULATION WITH AN AQUEOUS MEDIUM TO PROVIDE AN ETHANOL-FREE TAXANE DILUTED SOLUTION
- U-1790 FOR USE IN TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) AND/OR NON-HODGKIN'S LYMPHOMA
- U-1791 EMERGENCY TREATMENT OF ADULT & PEDIATRIC PATIENTS FOLLOWING FLUOROURACIL OR CAPECITABINE OVERDOSE, OR WHO EXHIBIT EARLY-ONSET, SEVERE OR LIFE-THREATENING CARDIAC OR CNS TOXICITY OR UNUSUALLY SEVERE ADVERSE REACTIONS WITHIN 96 HOURS

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-1792 TREATMENT OF OTIC INFECTION OR INFLAMMATION

U-1793 TREATMENT OF PEDIATRIC PATIENTS WITH OTITIS MEDIA WITH EFFUSION UNDERGOING TYMPANOSTOMY TUBE PLACEMENT

U-1794 REVERSAL OF DRUG-INDUCED NEUROMUSCULAR BLOCK

U-1795 REVERSAL OF NEUROMUSCULAR BLOCKAGE INDUCED BY ROCURONIUM BROMIDE OR VECURONIUM BROMIDE

U-1796 TOPICAL TREATMENT OF INFLAMMATORY PAPULES AND PUSTULES OF MILD TO MODERATE ROSACEA

U-1797 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A PHARMACEUTICAL COMPOSITION COMPRISING SELEXIPAG

U-1798 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING SELEXIPAG IN COMBINATION WITH THE ENDOTHELIN RECEPTOR ANTAGONIST MACITENTAN

U-1799 METHOD OF INCREASING GROWTH OF HAIR INCLUDING EYELASHES

U-1800 A METHOD OF TREATING OCULAR PAIN AND/OR ENHANCING OCULAR COMFORT

U-1801 REDUCTION OF SERUM URIC ACID LEVELS

U-1802 TREATMENT OF GOUT

U-1803 TREATMENT OF HYPERURICEMIA

U-1804 ACHIEVING A THERAPEUTIC BENEFIT IN A SUBJECT WITH GOUT

U-1805 USE OF DEXLANSOPRAZOLE IN PATIENTS TAKING CLOPIDOGREL WITHOUT MEANINGFUL CYP2C19 INTERACTIONS

U-1806 COADMINISTERING WITH ALLOPURINOL TO REDUCE SERUM URIC ACID (SUA) BELOW 4 MG/DL; BELOW 6MG/DL IN PATIENTS HAVING URIC ACID DEPOSITS; AND/OR BELOW 6MG/DL WITH SUA INTRADAY CHANGE MORE THAN 50% AND/OR ADVERSE EVENT RATE LESS THAN 15%

U-1807 TREATMENT OF PEDIATRIC PATIENTS 8 TO 17 YEARS OF AGE WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH)

U-1808 USE OF NALTREXONE AND BUPROPION FOR CHRONIC WEIGHT MANAGEMENT FOR TREATING OVERWEIGHT OR OBESITY IN PATIENTS WITH MAJOR DEPRESSIVE DISORDER

U-1809 METHOD OF DRUG DELIVERY VIA THE NASAL CAVITY

U-1810 TREATMENT OF PAIN IN PATIENTS WITH HEPATIC IMPAIRMENT

U-1811 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATIONS AFTER CONFIRMING THE PRESENCE OF BRAF V600E MUTATION

U-1812 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA

U-1813 TREATMENT OF PATIENTS INFECTED WITH HEPATITIS C VIRUS

U-1814 METHOD OF TREATING GLAUCOMA OR ELEVATED INTRAOCULAR PRESSURE

U-1815 TREATMENT OF PARTIAL-ONSET SEIZURES AS ADJUNCTIVE THERAPY IN PATIENTS WITH EPILEPSY AGED 16 YEARS AND OLDER WITH EPILEPSY

U-1816 TREATMENT OF A UREA CYCLE DISORDER

U-1817 PEDIATRIC USE AGES 1 MONTH TO 2 YEARS, GERD AND EROSIIVE ESOPHAGITIS

U-1818 TREATING HR-POS., HER2-NEG. ADVANCED OR METASTATIC BREAST CANCER WITH PALBOCICLIB IN COMBO WITH LETROZOLE AS INITIAL ENDOCRINE BASED THERAPY IN POSTMENOPAUSAL WOMEN, OR FULVESTRANT IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY

U-1819 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE

U-1820 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION BY ADMINISTERING A PHARMACEUTICAL COMPOSITION COMPRISING MACITENTAN AND A POLYSORBATE, WHEREIN THE POLYSORBATE REPRESENTS 0.1 TO 3% OF THE WEIGHT OF SAID PHARMACEUTICAL COMPOSITION

U-1821 METHOD FOR CONTRACEPTION TO A WOMAN COMPRISING ADMINISTERING TO THE WOMAN 30MG OF ULIPRISTAL ACETATE MORE THAN 72 HOURS AND UP TO 120 HOURS AFTER AN UNPROTECTED INTERCOURSE

U-1822 TREATMENT OF SCHIZOPHRENIA OR BIPOLAR DEPRESSION WITH IMPROVEMENT IN ATTENTION FUNCTION IN SCHIZOPHRENIA AND/OR BIPOLAR DISORDER

U-1823 A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT BY COMPENSATING LONG-TERM SENSITIVITY DRIFT OF ELECTROCHEMICAL GAS SENSORS USED IN SYSTEMS FOR

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

DELIVERING THERAPEUTIC NITRIC OXIDE TO A PATIENT

U-1824 A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT BY VERIFYING GAS INFORMATION OF NITRIC OXIDE PRIOR TO DELIVERY TO PATIENT

U-1825 METHOD OF USING VISMODEGIB TO TREAT CANCER IN A MAMMAL

U-1826 TREATMENT OF HR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN COMBINATION WITH PALBOCICLIB IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY

U-1827 A METHOD OF PROVIDING A SUBJECT WITH THERAPEUTICALLY EFFECTIVE AMOUNT OF RACEMIC METHYLPHENIDATE BY ORALLY ADMINISTERING TO SAID SUBJECT A SINGLE METHYLPHENIDATE EXTENDED RELEASE CHEWABLE TABLET ACCORDING TO CLAIM 1

U-1828 INCREASING MEAN ARTERIAL BLOOD PRESSURE IN ADULT PATIENTS WITH HYPOTENSION ASSOCIATED WITH SEPTIC SHOCK

U-1829 EMERGENCY TREATMENT OF ALLERGIC REACTIONS (TYPE I), INCLUDING ANAPHYLAXIS

U-1830 INDUCTION AND MAINTENANCE OF MYDRIASIS DURING INTRAOCULAR SURGERY

U-1831 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A CRYSTALLINE FORM OF SELEXIPAG

U-1832 IMPROVEMENT IN GLYCEMIC CONTROL IN DIABETES MELLITUS PATIENTS BY USE OF A PEN INJECTOR WITH A THREADED DRIVE SLEEVE

U-1833 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION

U-1834 TREATMENT OF POSTOPERATIVE INFLAMMATION AND PREVENTION OF OCULAR PAIN IN PATIENTS UNDERGOING CATARACT SURGERY

U-1835 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WITH 17P DELETION, AS DETECTED BY AN FDA APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY

U-1836 TREATMENT OF HEREDITARY TYROSINEMIA TYPE 1 (HT-1) IN COMBINATION WITH DIETARY RESTRICTION OF TYROSINE AND PHENYLALANINE

U-1837 METHOD FOR TREATING TYPE II DIABETES MELLITUS BY ADMINISTERING SAXAGLIPTIN ALONE OR IN COMBINATION WITH INSULIN, METFORMIN, A THIAZOLIDINEDIONE, GLYBURIDE OR METFORMIN PLUS A SULFONYLUREA

U-1838 METHOD FOR TREATING TYPE II DIABETES MELLITUS BY ADMINISTERING SAXAGLIPTIN IN COMBINATION WITH METFORMIN

U-1839 COMPOSITION AND METHOD FOR PROVIDING A REDUCTION IN SIDE EFFECTS FOR HUMAN PATIENTS IN NEED OF ACETYLCYSTEINE THERAPY

U-1840 TREATMENT OF HCV INFECTION USING PARITAPREVR, OMBITASVIR, RITONAVIR, AND DASABUVIR, WITHOUT RIBAVIRIN

U-1841 USE IN THE LONG-TERM, MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)

U-1842 METHOD OF TREATING EPILEPSY

U-1843 TREATMENT OF PSYCHOSIS

U-1844 TREATMENT OF PARKINSON'S DISEASE PSYCHOSIS

U-1845 TREATMENT OF PSYCHOSIS OR A SYMPTOM THEREOF

U-1846 TREATMENT OF A NEURODEGENERATIVE DISEASE OR A SYMPTOM THEREOF

U-1847 METHOD OF TREATING A BACTERIAL INFECTION

U-1848 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN

U-1849 METHOD OF TREATING PULMONARY HYPERTENSION BY ADMINISTERING TREPROSTINIL OR A SALT THEREOF BY INHALATION USING A DEVICE

U-1850 METHOD OF ADMINISTERING LEVETIRACETAM

U-1851 A DOSING REGIMEN FOR THE TREATMENT OF HYPERCHOLESTEROLEMIA AND HYPERLIPIDEMIA IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA USING AT LEAST THREE STEP-WISE INCREASING DOSES

U-1852 METHOD OF TREATING TYPE 2 DIABETES

U-1853 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH METFORMIN AND, OPTIONALLY, A

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

SULFONYLUREA

- U-1854 TREATMENT OF PRIMARY BILIARY CHOLANGITIS (PBC)
- U-1855 IMPROVEMENT IN GLYCEMIC CONTROL IN DIABETES MELLITUS PATIENTS
- U-1856 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN, IN A PATIENT HOMOZYGOUS FOR THE UGT1A1*28 ALLELE
- U-1857 TO INCREASE BLOOD PRESSURE IN ADULTS WITH VASODILATORY SHOCK (E.G., POST-CARDIOTOMY OR SEPSIS) WHO REMAIN HYPOTENSIVE DESPITE FLUIDS AND CATECHOLAMINES
- U-1858 TREATMENT OF PLAQUE PSORIASIS
- U-1859 TREATMENT OF SCHIZOPHRENIA, ACUTE TREATMENT OF MANIC AND MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER, ADJUNCTIVE TREATMENT OF MAJOR DEPRESSIVE DISORDER, AND TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER
- U-1860 REDUCTION OF THE RATE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION, AND STROKE IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION
- U-1861 USE OF AN INHALER TO ADMINISTER DRY POWDER MEDICAMENT
- U-1862 TREATMENT OF POST-MYOCARDIAL INFARCTION
- U-1863 TREATMENT OF STROKE
- U-1864 TREATMENT OF MYOCARDIAL INFARCTION
- U-1865 TREATMENT OF THROMBOTIC STROKE
- U-1866 TREATMENT OF STABLE AND UNSTABLE ANGINA
- U-1867 METHOD OF INHIBITING PLATELET AGGREGATION
- U-1868 TREATMENT OF ARTERIAL THROMBOTIC COMPLICATIONS SELECTED FROM THE GROUP CONSISTING OF UNSTABLE ANGINA, THROMBOTIC OR EMBOLIC STROKE, TRANSIENT ISCHAEMIC ATTACKS, PERIPHERAL VASCULAR DISEASE AND MYOCARDIAL INFARCTION
- U-1869 TREATMENT OF AN ARTERIAL THROMBOTIC COMPLICATION IN A PATIENT WITH CORONARY ARTERY, CEREBROVASCULAR OR PERIPHERAL VASCULAR DISEASE
- U-1870 ZINGO IS A POWDER INTRADERMAL SYSTEM THAT IS CAPABLE OF DELIVERING FINE DRY POWDERED LIDOCAINE HYDROCHLORIDE MONOHYDRATE FOR LOCAL ANESTHETIC ACTION
- U-1871 TREATMENT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS WITH STAGE 3 OR 4 CHRONIC KIDNEY DISEASE USING CONTROLLED RELEASE, ORAL 25-HYDROXYVITAMIN D
- U-1872 USE OF SUSTAINED RELEASE 25-HYDROXYVITAMIN D IN TREATING PATIENTS HAVING 25-HYDROXYVITAMIN D INSUFFICIENCY OR DEFICIENCY
- U-1873 ADMINISTRATION OF 25-HYDROXYVITAMIN D3 BY CONTROLLED RELEASE
- U-1874 TREATMENT OF FREQUENT HEARTBURN BY ADMINISTERING OMEPRAZOLE ACCORDING TO CLAIMS 1-8
- U-1875 TREATMENT OF FREQUENT HEARTBURN BY ADMINISTERING S-OMEPRAZOLE TRIHYDRATE ACCORDING TO CLAIMS 1-3
- U-1876 METHOD OF ANESTHETIZING AT LEAST A PORTION OF THE MAXILLARY DENTAL ARCH
- U-1877 METHOD OF TREATING PULMONARY HYPERTENSION BY ORALLY ADMINISTERING A FORMULATION OF A PHARMACEUTICALLY ACCEPTABLE SALT OF TREPROSTINIL
- U-1878 FOR OPIOID DEPENDENCE
- U-1879 METHOD OF DIAGNOSING TUMORS USING POSITRON EMISSION TOMOGRAPHY
- U-1880 TREATMENT OF SIGNS AND SYMPTOMS OF DRY EYE DISEASE (DED)
- U-1881 IMPROVEMENT IN GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS BY USE OF A PEN INJECTOR
- U-1882 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS, REDUCTION IN FEVER THROUGH ANALGESIC AND ANTIPYRETIC ACTIVITY
- U-1883 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS (GIST)
- U-1884 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1885 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER

PATENT AND EXCLUSIVITY TERMS

ADB 110 of 133

PATENT USE

- THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1886 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1887 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1888 USE OF CONTROLLED RELEASE 25-HYDROXYVITAMIN D IN TREATING SECONDARY HYPERPARATHYROIDISM IN PATIENTS HAVING CHRONIC KIDNEY DISEASE
- U-1889 TREATMENT OF HCV INFECTION USING DASABUVIR/OMBITASVIR/PARITAPREVIR/RITONAVIR FIXED DOSE COMBINATION
- U-1890 OTC USE: ALLERGY SYMPTOM RELIEVER; TEMPORARILY RELIEVES THESE SYMPTOMS DUE TO HAY FEVER OR OTHER UPPER RESPIRATORY ALLERGIES; NASAL CONGESTION, RUNNY NOSE, SNEEZING, ITCHY NOSE, AND (ITCHY WATER EYES (AGES 12 AND UP))
- U-1891 TREATMENT OR PREVENTION OF NAUSEA AND VOMITING
- U-1892 METHOD OF TREATING LEFT VENTRICULAR DYSFUNCTION
- U-1893 METHOD OF TREATING MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR DISORDER IN PEDIATRIC PATIENTS
- U-1894 COMBINATION TREATMENT WITH A GLITAZONE FOR IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS
- U-1895 METHOD OF TREATING PROSTATE CANCER
- U-1896 SUPPLEMENT FOR VITAMIN B12 DEFICIENCIES
- U-1897 METHOD OF TREATING ACS USING ANGIOPLASTY WITH AGGRASTAT (TIROFIBAN HYDROCHLORIDE)
- U-1898 METHOD OF INHIBITING PLATELET AGGREGATION WITH AGGRASTAT (TIROFIBAN HYDROCHLORIDE)
- U-1899 TREATMENT OF PANCREATIC CANCER THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN
- U-1900 TREATMENT OF THE SIGNS AND SYMPTOMS OF DRY EYE DISEASE (DED)
- U-1901 TREATMENT OF SCHIZOAFFECTIVE DISORDER AS A MONOTHERAPY AND AS AN ADJUNCT TO MOOD STABILIZERS OR ANTIDEPRESSANTS
- U-1902 TREATMENT OR SECONDARY PREVENTION OF CARDIOVASCULAR DISEASE, CARDIOVASCULAR EVENTS, OR CEREBROVASCULAR EVENTS AND RISK-REDUCTION OF ASPIRIN-ASSOCIATED GASTRIC ULCERS
- U-1903 USE OF NALOXONE HYDROCHLORIDE FOR EMERGENCY TREATMENT OF KNOWN OR SUSPECTED OPIOID OVERDOSE, AS MANIFESTED BY RESPIRATORY AND/OR CENTRAL NERVOUS SYSTEM DEPRESSION.
- U-1904 (I) TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY; (II) RESTORING/INCREASING FUNCTIONAL DYSTROPHIN PROTEIN; OR (III) INDUCING SKIPPING; EACH OF (I)-(III) IN PATIENTS HAVING A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- U-1905 METHOD OF TREATING A PATIENT HAVING CYSTIC FIBROSIS, THE PATIENT HAVING A R117H MUTATION IN CFTR, USING N-(5-HYDROXY-2,4-DI-TERT-BUTYL-PHENYL)-4-OXO-1H-QUINOLINE-3-CARBOXAMIDE
- U-1906 METHOD OF TREATING A PATIENT HAVING CYSTIC FIBROSIS, SUCH AS A PATIENT HAVING A G551D MUTATION IN CFTR, USING N-(5-HYDROXY-2,4-DI-TERT-BUTYL-PHENYL)-4-OXO-1H-QUINOLINE-3-CARBOXAMIDE
- U-1907 USE OF A DELIVERY DEVICE TO ADMINISTER A DOSE OF NALOXONE
- U-1908 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING IVACAFTOR AND FORM I LUMACAFTOR
- U-1909 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING IVACAFTOR AND LUMACAFTOR
- U-1910 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING THE DOSAGE UNIT OF CLAIM 1 OF U.S. PATENT NO. 8,716,338
- U-1911 METHOD OF TREATING A PATIENT HAVING CYSTIC FIBROSIS USING IVACAFTOR AND LUMACAFTOR
- U-1912 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING A DOSAGE UNIT AS DEFINED IN CLAIM 1 OF U.S. PATENT NO.

PATENT AND EXCLUSIVITY TERMS

ADB 111 of 133

PATENT USE

9,192,606

- U-1913 TREATMENT OF PEDIATRIC PATIENTS WITH BILATERAL OTITIS MEDIA WITH EFFUSION UNDERGOING TYMPANOSTOMY TUBE PLACEMENT
- U-1914 IN COMBINATION WITH RITUXIMAB, FOR THE TREATMENT OF PATIENTS WITH RELAPSED CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)
- U-1915 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS WITH SEVERE CHRONIC RENAL IMPAIRMENT AND WHO ARE INELIGIBLE FOR METFORMIN THERAPY BY ADMINISTERING LINAGLIPTIN
- U-1916 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH CHEMOTHERAPY (CINV)
- U-1917 TREATMENT OF EXOCRINE PANCREATIC CANCER THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN
- U-1918 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- U-1919 RESTORING AN MRNA READING FRAME TO INDUCE DYSTROPHIN PROTEIN PRODUCTION IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- U-1920 USE OF EXTENDED RELEASE ORAL 25-HYDROXYVITAMIN D3 IN TREATING SECONDARY HYPERPARATHYROIDISM IN ADULT PATIENTS HAVING CHRONIC KIDNEY DISEASE STAGE 3 OR STAGE 4
- U-1921 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE BY PROVIDING AN ABUSE-DETERRENT ORAL CONTROLLED RELEASE COMBINATION DRUG PRODUCT
- U-1922 INTRAVAGINAL PRASTERONE (DEHYDROEPIANDROSTERONE) AT A DAILY DOSE OF 6.5MG FOR THE TREATMENT OF DYSPAREUNIA, A SYMPTOM OF VULVAR AND VAGINAL ATROPHY, DUE TO MENOPAUSE
- U-1923 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS INADEQUATELY CONTROLLED BY BASAL INSULIN OR LIXISENATIDE BY USE OF A PEN INJECTOR WITH A THREADED DRIVE SLEEVE
- U-1924 KYPROLIS IS INDICATED IN COMBINATION WITH LENALIDOMIDE PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY
- U-1925 USE OF AN AUTO INJECTOR TO ADMINISTER NALOXONE HCL
- U-1926 METHOD OF TREATING, REDUCING THE INCIDENCE OF, OR PREVENTING AN ISCHEMIC EVENT IN A PATIENT UNDERGOING PCI BY ADMINISTERING INTRAVENOUSLY 30 UG/KG BOLUS BEFORE PCI AND CONTINUOUS INFUSION OF 4 UG/KG/MIN FOR AT LEAST 2 HOURS OR THE DURATION OF THE PCI
- U-1927 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- U-1928 RUBRACA IS INDICATED AS MONOTHERAPY FOR THE TREATMENT OF PATIENTS WITH DELETERIOUS BRCA MUTATION (GERMLINE AND/OR SOMATIC) ASSOCIATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH TWO OR MORE CHEMOTHERAPIES.
- U-1929 TREATMENT OF DIABETES MELLITUS WITH AN INHALED INSULIN TO IMPROVE GLYCEMIC CONTROL USING A DRY POWDER INHALATION SYSTEM COMPRISING AN INHALER, A CARTRIDGE AND A DRY POWDER MEDICAMENT COMPRISING INSULIN IN A SINGLE INHALATION
- U-1930 METHOD OF AEROSOLIZING/DEAGGLOMERATING AN INSULIN DRY POWDER FOR USE IN TREATING DIABETES MELLITUS VIA ORAL INHALATION USING AN INHALER WITH A CARTRIDGE CONTAINING THE INSULIN DRY POWDER.
- U-1931 PROPHYLAXIS OR TREATMENT OF VENOUS AND ARTERIAL THROMBOTIC DISEASE
- U-1932 METHOD OF TREATING MILD TO MODERATE ATOPIC DERMATITIS.
- U-1933 TREATMENT OF POSTOPERATIVE INFLAMMATION AND REDUCTION OF OCULAR PAIN IN PATIENTS WHO HAVE UNDERGONE CATARACT SURGERY
- U-1934 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH A STRONG CY1A2 INHIBITOR
- U-1935 REDUCTION OF THE RATE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION, AND STROKE IN PATIENTS WITH A HISTORY OF MYOCARDIAL INFARCTION
- U-1936 TREATMENT OF MYOCARDIAL INFARCTION AND STROKE IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION
- U-1937 TREATMENT OF MYOCARDIAL INFARCTION IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

HISTORY OF MYOCARDIAL INFARCTION

- U-1938 TREATMENT OF STROKE IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION
- U-1939 ADMINISTRATION ONCE DAILY WITHIN TWO HOURS AFTER WAKING IN THE MORNING FOR IMPROVEMENT OF GLYCEMIC CONTROL IN A TYPE 2 DIABETES PATIENT
- U-1940 IMPROVEMENT IN THE APPEARANCE OF MODERATE TO SEVERE CONVEXITY OR FULLNESS ASSOCIATED WITH SUBMENTAL FAT IN ADULTS BY MEANS OF REDUCING SUBMENTAL FAT VOLUME AS DESCRIBED IN THE APPROVED LABELING
- U-1941 TREATMENT OF INFANTILE-ONSET SPINAL MUSCULAR ATROPHY
- U-1942 TREATMENT OF SPINAL MUSCULAR ATROPHY BY INCREASING EXON-7 INCLUSION IN SMN2 MRNA
- U-1943 TREATMENT OF SPINAL MUSCULAR ATROPHY
- U-1944 TREATMENT OF SPINAL MUSCULAR ATROPHY BY INHIBITING AN SMN2 PRE-MRNA INTRONIC SPLICING SILENCER SITE
- U-1945 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH METFORMIN.
- U-1946 TREATMENT OF SMALL LYMPHOCYTIC LYMPHOMA
- U-1947 TREATMENT OF MARGINAL ZONE LYMPHOMA
- U-1948 A METHOD FOR TREATING CHRONIC MYELOID LEUKEMIA
- U-1949 FOR USE IN THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)
- U-1950 TREATMENT OF PATIENTS WITH ADVANCED (METASTATIC) NON-SMALL CELL LUNG CANCER WHOSE DISEASE PROGRESSED DURING OR AFTER PLATINUM-BASED CHEMOTHERAPY
- U-1951 TREATMENT OF ONYCHOMYCOSIS OF A TOENAIL
- U-1952 FOR USE IN THE TREATMENT OF PATIENTS WITH INDOLENT B-CELL NON-HODGKIN LYMPHOMA
- U-1953 REDUCE THE RISK OF STROKE IN PATIENTS WITH NONVALVULAR ATRIAL FIBRILLATION WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1954 TREATMENT OF DEEP VEIN THROMBOSIS WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1955 TREATMENT OF PULMONARY EMBOLISM WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1956 FOLLOWING INITIAL 6 MONTHS TREATMENT FOR DEEP VEIN THROMBOSIS (DVT) AND/OR PULMONARY EMBOLISM (PE), REDUCTION IN THE RISK OF RECURRENCE OF DVT AND OF PE WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1957 PROPHYLAXIS OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM IN PATIENTS UNDERGOING KNEE OR HIP REPLACEMENT SURGERY, WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1958 FOR THE TREATMENT OF GENOTYPE 1, 2, 3 OR 4 CHRONIC HEPATITIS C VIRUS (HCV) INFECTION AS A COMPONENT OF A COMBINATION ANTIVIRAL TREATMENT REGIMEN WITH RIBAVIRIN
- U-1959 TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN ADULTS WITH 1% OXYMETAZOLINE HYDROCHLORIDE CREAM
- U-1960 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF SCHIZOPHRENIA IN ADULTS
- U-1961 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: ACUTE MONOTHERAPY OF MANIC OR MIXED EPISODES (AGES 10 TO ADULT)
- U-1962 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: MAINTENANCE MONOTHERAPY IN ADULTS
- U-1963 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: AS ADJUNCTIVE TREATMENT TO LITHIUM OR VALPROATE IN ADULTS
- U-1964 ELEVATION OF INTRACELLULAR CGMP RESULTING IN INCREASED INTESTINAL FLUID AND ACCELERATED TRANSIT
- U-1965 FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) IN COMBINATION WITH TADALAFIL, WHEREIN THE WEIGHT RATIO OF AMBRISENTAN TO TADALAFIL IS ABOUT 1:2 TO ABOUT 1:3

PATENT AND EXCLUSIVITY TERMS

ADB 113 of 133

PATENT USE

U-1966 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: ACUTE MONOTHERAPY OF MANIC OR MIXED EPISODES IN PEDIATRIC PATIENTS AGE 10-17

U-1967 METHOD OF TREATING TYPE 2 DIABETES IN PATIENTS WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE THERAPY WITH ONE OR MORE CONVENTIONAL ANTIHYPERGLYCEMIC AGENTS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH METFORMIN

U-1968 METHOD OF TREATING TYPE 2 DIABETES IN PATIENTS WHO HAVE NOT BEEN PREVIOUSLY TREATED WITH AN ANTIHYPERGLYCEMIC AGENT BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH METFORMIN

U-1969 TOPICAL TREATMENT OF ONYCHOMYCOSIS OF THE TOENAIL(S) DUE TO TRICHOPHYTON RUBRUM AND TRICHOPHYTON MENTAGROPHYTES

U-1970 TREATMENT OF ONYCHOMYCOSIS OF A TOENAIL CAUSED BY TRICHOPHYTON RUBRUM OR TRICHOPHYTON MENTAGROPHYTES

U-1971 FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA

U-1972 FOR THE TREATMENT OF PATIENTS WITH INDOLENT B-CELL NON-HODGKIN LYMPHOMA

U-1973 METHOD OF TREATING CYSTIC FIBROSIS USING N-(5-HYDROXY-2,4-DITERT-BUTYL-PHENYL)-4-OXO-1H-QUINOLINE-3-CARBOXAMIDE AND 3-(6-(1-2,2-DIFLUOROBENZO[D][1,3]DIOXOL-5-YL) CYCLOPROPANECARBOXAMIDO)-3-METHYLPYRIDIN-2-YL)BENZOIC ACID

U-1974 TREATMENT OF HALLUCINATIONS AND DELUSIONS ASSOCIATED WITH PARKINSON'S DISEASE PSYCHOSIS

U-1975 METHOD OF INCREASING EYELASH GROWTH WITH BIMATOPROST

U-1976 METHOD FOR TREATING TYPE 2 DIABETES MELLITUS (T2DM) IN PATIENTS WHO HAVE INADEQUATE CONTROL WITH DAPAGLIFLOZIN

U-1977 METHOD FOR TREATING TYPE 2 DIABETES MELLITUS (T2DM) IN PATIENTS WHO ARE ALREADY TREATED WITH DAPAGLIFLOZIN AND SAXAGLIPTIN

U-1978 TREATMENT OF ADVANCED PROSTATE CANCER WITH A REDUCED LIKELIHOOD OF CAUSING A GONADOTROPHIN RELEASING HORMONE AGONIST SIDE-EFFECT

U-1979 THE TREATMENT OF CARCINOID SYNDROME DIARRHEA IN COMBINATION WITH SOMATOSTATIN ANALOG (SSA) THERAPY IN ADULTS INADEQUATELY CONTROLLED BY SSA THERAPY

U-1980 A METHOD OF TREATING NOCTURIA DUE TO NOCTURNAL POLYURIA IN ADULTS

U-1981 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER-2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER

U-1982 USE OF REVLIMID (LENALIDOMIDE) FOR TREATMENT OF PATIENTS WITH TRANSFUSION-DEPENDENT ANEMIA DUE TO LOW-OR INTERMEDIATE-1-RISK MYELOYDYSPLASTIC SYNDROMES ASSOCIATED WITH A DELETION 5Q ABNORMALITY WITH OR WITHOUT ADDITIONAL CYTOGENETIC ABNORMALITIES

U-1983 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHOSE DISEASE HAS RELAPSED OR PROGRESSED AFTER TWO PRIOR THERAPIES, ONE OF WHICH INCLUDED BORTEZOMIB

U-1984 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA, IN COMBINATION WITH DEXAMETHASONE

U-1985 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA, AS MAINTENANCE FOLLOWING AUTOLOGOUS HEMATOPOIETIC STEM CELL TRANSPLANTATION (AUTO-HSCT)

U-1986 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA, IN COMBINATION WITH DEXAMETHASONE, WHEREIN THOSE PATIENTS HAVE NOT RECEIVED PREVIOUS TREATMENT FOR MULTIPLE MYELOMA

U-1987 METHOD OF CONTROLLING GLYCEMIA IN DIABETICS BY ADMINISTERING AN INITIAL DOSE OF INSULIN-FDKP WITH A MEAL; DETERMINING BLOOD GLUCOSE LEVEL 1-2 HRS AFTER AND ADMINISTERING A SUPPLEMENTAL DOSE OF INSULIN-FDKP IF POSTPRANDIAL GLUCOSE LEVEL IS >140 MG/DL

U-1988 METHOD TO TREAT INFANTILE HEMANGIOMA

U-1989 INTRAVITREAL TREATMENT OF MACULAR EDEMA FOLLOWING BRANCH RETINAL VEIN OCCLUSION (BRVO) OR CENTRAL RETINAL VEIN OCCLUSION (CRVO)

U-1990 INTRAVITREAL TREATMENT OF DIABETIC MACULAR EDEMA

U-1991 REDUCTION OF MORTALITY IN ACUTE MYOCARDIAL INFARCTION

PATENT AND EXCLUSIVITY TERMS

ADB 114 of 133

PATENT USE

U-1992 USE OF TROKENDI XR FOR PROPHYLACTIC TREATMENT OF MIGRAINE

U-1993 ADJUNCTIVE TREATMENT TO LEVODOPA/CARBIDOPA IN PATIENTS WITH PARKINSON'S DISEASE EXPERIENCING "OFF" EPISODES

U-1994 REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) IN ADULTS

U-1995 TREATMENT OF TARDIVE DYSKINESIA

U-1996 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

U-1997 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH METFORMIN AND/OR A PPAR-GAMMA AGONIST AND/OR SULFONYLUREA AND/OR INSULIN

U-1998 TREATING HR-POS., HER2-NEG. ADVANCED OR METASTATIC BREAST CANCER WITH PALBOCICLIB IN COMBO WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE BASED THERAPY IN POSTMENOPAUSAL WOMEN OR FULVESTRANT IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY

U-1999 CHRONIC IDIOPATHIC CONSTIPATION

U-2000 MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS

U-2001 USE FOR THE TREATMENT OF ASTHMA IN PATIENTS 6 YEARS OF AGE AND OLDER

U-2002 USE FOR MAINTENANCE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE

U-2003 A METHOD OF POSITIONING AN INTRAUTERINE SYSTEM BY HOLDING AN INSERTER HANDLE WITH ONE HAND, ADVANCING THE INSERTER THROUGH THE CERVIX AND INTO THE UTERUS, AND RETRACTING A SLIDER ON THE HANDLE TO RELEASE THE INTRAUTERINE SYSTEM

U-2004 REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE IN ADULTS WITH PARTIAL SEIZURES WITH COMPLEX SYMPTOMATOLOGY

U-2005 REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE IN ADULTS WITH GENERALIZED TONIC-CLONIC SEIZURES

U-2006 REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE IN ADULTS WITH MIXED SEIZURE PATTERNS THAT INCLUDE PARTIAL SEIZURES WITH COMPLEX SYMPTOMATOLOGY, GENERALIZED TONIC-CLONIC SEIZURES, OR OTHER PARTIAL OR GENERALIZED SEIZURES

U-2007 TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) WHO ARE FLT3 MUTATION-POSITIVE, IN COMBINATION WITH STANDARD CYTARABINE AND DAUNORUBICIN INDUCTION AND CYTARABINE CONSOLIDATION CHEMOTHERAPY

U-2008 TREATMENT OF ADULT PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM), SYSTEMIC MASTOCYTOSIS WITH ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN), OR MAST CELL LEUKEMIA (MCL)

U-2009 METHOD OF TREATING POSTMENOPAUSAL WOMEN WITH OSTEOPOROSIS AT HIGH RISK FOR FRACTURE.

U-2010 ACUTE TREATMENT OF MIGRAINE BY DELIVERING A POWDERED SUBSTANCE COMPRISING SUMATRIPTAN VIA A BREATH-POWERED DELIVERY DEVICE

U-2011 TREATMENT OF MIGRAINE VIA DELIVERY OF SUMATRIPTAN VIA THE NASAL CAVITY

U-2012 A METHOD FOR TREATING OVARIAN CANCER BY ADMINISTERING RUCAPARIB, WHEREIN THE CANCER IS ASSOCIATED WITH A DELETERIOUS BRCA MUTATION

U-2013 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS)

U-2014 A METHOD OF TREATING SECONDARY HYPERPARATHYROIDISM (SHPT)

U-2015 SODIUM THIOSULFATE INJECTION IS INDICATED FOR SEQUENTIAL USE WITH SODIUM NITRITE FOR THE TREATMENT OF ACUTE CYANIDE POISONING

U-2016 TREATMENT FOR ONYCHOMYCOSIS THAT IS TINEA UNGUIUM

U-2017 TREATMENT OF OPIOID DEPENDENCE

U-2018 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS IN A CRITICALLY ILL PATIENT WITH INTRAVENOUS IBUPROFEN IN NEED THEREOF

U-2019 METHOD OF DELIVERING TO A PATIENT WITH DIABETES MELLITUS IN A SINGLE INHALATION, GREATER THAN 75% OF A DRY POWDER DOSE COMPRISING INSULIN AND FUMARYL DIKETOPIPERAZINE USING A HIGH RESISTANCE TO FLOW DRY POWDER INHALER.

U-2020 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-2021 METHOD OF ADMINISTERING LEVETIRACETAM UNDER FASTED CONDITIONS

U-2022 METHOD OF ADMINISTERING LEVETIRACETAM UNDER FED CONDITIONS

U-2023 A METHOD OF INCREASING THE BIOAVAILABILITY OF GUAIFENESIN IN A SOLUTION CONTAINING 54% TO 66% BY WEIGHT OF PROPYLENE GLYCOL AND GLYCEROL, WHEREIN THE METHOD INCREASES THE CMAX BY AT LEAST 1.5 AND/OR INCREASES THE AUC (0-INF) BY AT LEAST 1.4

U-2024 METHOD FOR TRANSDERMALLY DELIVERING A DRUG TO A USER IN NEED THEREOF

U-2025 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER

U-2026 TAFINLAR(R) IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST.

U-2027 TAFINLAR(R) IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST.

U-2028 TREATMENT OF ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS CAUSED BY DESIGNATED SUSCEPTIBLE BACTERIA IN ADULTS

U-2029 PREVENTING CONDITION CHARACTERIZED BY UNDESIRED THROMBOSIS

U-2030 PROPHYLAXIS OF VENOUS THROMBOSIS

U-2031 TAFINLAR IS INDICATED AS A SINGLE AGENT FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST

U-2032 TAFINLAR IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST.

U-2033 MEKINIST IS INDICATED, AS A SINGLE AGENT OR IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS

U-2034 INHIBITING COAGULATION

U-2035 PROPHYLAXIS OF VENOUS THROMBOEMBOLISM

U-2036 A METHOD OF TREATING PULMONARY HYPERTENSION COMPRISING PARENTERALLY ADMINISTERING A FORMULATION COMPRISING A) 0.1 TO 5% W/V OF TREPROSTINIL OR A PHARMACEUTICALLY ACCEPTABLE SALT THEREOF AND B) A CITRATE BUFFER

U-2037 MEKINIST IS INDICATED, AS A SINGLE AGENT OR IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST

U-2038 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST

U-2039 TREATMENT OF ADULT PATIENTS WITH CHRONIC HCV INFECTION WHO HAVE GENOTYPE 1, 2, 3, 4, 5, OR 6 INFECTION AND HAVE PREVIOUSLY BEEN TREATED WITH AN HCV REGIMEN CONTAINING AN NS5A INHIBITOR

U-2040 TREATMENT OF ADULT PATIENTS WITH CHRONIC HCV INFECTION WHO HAVE GENOTYPE 1A OR 3 INFECTION AND HAVE PREVIOUSLY BEEN TREATED WITH AN HCV REGIMEN CONTAINING SOFOSBUVIR WITHOUT AN NS5A INHIBITOR

U-2041 TREATMENT OF PARTIAL-ONSET SEIZURES

U-2042 DISCONTINUING ADMINISTRATION OF FLUVOXAMINE TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE AND THEN ADMINISTERING PIRFENIDONE

U-2043 EXTENDED ADJUVANT TREATMENT OF ADULT PATIENTS WITH EARLY STAGE HER2-OVEREXPRESSED/AMPLIFIED BREAST CANCER, TO FOLLOW ADJUVANT TRASTUZUMAB BASE THERAPY

U-2044 DOSE REDUCTION OF PIRFENIDONE BY ABOUT ONE HALF DURING CONCURRENT ADMINISTRATION OF CIPROFLOXACIN AT A DOSE OF 750 MG TWICE DAILY (1500 MG/DAY) TO REDUCE DRUG INTERACTIONS IN TREATMENT OF A FIBROTIC, INFLAMMATORY, OR AUTOIMMUNE DISORDER

U-2045 ADMINISTRATION OF PIRFENIDONE AND AVOIDING CONCURRENT ADMINISTRATION OF CIPROFLOXACIN AT A DOSE OF 750 MG TO REDUCE DRUG INTERACTIONS IN TREATMENT OF A FIBROTIC, INFLAMMATORY, OR AUTOIMMUNE DISORDER

U-2046 ADMINISTERING PIRFENIDONE WHILE AVOIDING CONCOMITANT USE OF A CYP1A2 INHIBITOR THAT IS A MODERATE TO STRONG INHIBITOR OF BOTH CYP1A2 AND ANOTHER CYP ENZYME

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

SELECTED FROM CYP2C9, CYP2C19, AND CYP2D6

- U-2047 ADMINISTERING PIRFENIDONE CONCURRENTLY WITH FLUVOXAMINE, THE PIRFENIDONE AT A DOSE OF ABOUT 801 MG/DAY TO REDUCE DRUG INTERACTIONS WITH FLUVOXAMINE
- U-2048 ADMINISTERING PIRFENIDONE WHILE AVOIDING CO-ADMINISTRATION OF A STRONG CYP1A2 INHIBITOR TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE
- U-2049 DISCONTINUING ADMINISTRATION OF A STRONG CYP1A2 INHIBITOR TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE AND THEN ADMINISTERING PIRFENIDONE
- U-2050 ADMINISTERING PIRFENIDONE WHILE AVOIDING CO-ADMINISTRATION OF FLUVOXAMINE TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE
- U-2051 DISCONTINUING SMOKING TO AVOID REDUCED PIRFENIDONE EFFICACY AND THEN ADMINISTERING PIRFENIDONE
- U-2052 DISCONTINUING ADMINISTRATION OF A STRONG CYP1A2 INDUCER TO AVOID REDUCED PIRFENIDONE EFFICACY AND THEN ADMINISTERING PIRFENIDONE
- U-2053 ADMINISTERING PIRFENIDONE WHILE AVOIDING CONCOMITANT ADMINISTRATION OF A STRONG INDUCER OF CYP1A2, INCLUDING CIGARETTE SMOKE, TO AVOID REDUCED PIRFENIDONE EFFICACY
- U-2054 ADMINISTERING PIRFENIDONE WHILE AVOIDING CONCOMITANT ADMINISTRATION OF A STRONG INDUCER OF CYP1A2 TO AVOID REDUCED PIRFENIDONE EFFICACY
- U-2055 DOSING OF AT LEAST 1600 MG/DAY FOLLOWING GRADE 2 LIVER ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2056 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-1600 MG/DAY, FOLLOWING BY ADMINISTERING AT LEAST 1600 MG/DAY IN TREATMENT OF IPF
- U-2057 DOSING 2403 MG/DAY PIRFENIDONE FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2058 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400MG/DAY DOSE, FOLLOWED BY ADMINISTERING 2403MG/DAY IN TREATMENT OF IPF
- U-2059 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, FOLLOWED BY FULL DAILY DOSE IN TREATMENT OF IPF
- U-2060 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, THEN AT LEAST 1600MG/DAY IN TREATMENT OF IPF
- U-2061 DOSING OF AT LEAST 1600 MG/DAY FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2062 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-1600 MG/DAY DOSE, FOLLOWED BY ADMINISTERING AT LEAST 1600 MG/DAY DOSE IN TREATMENT OF IPF
- U-2063 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, FOLLOWED BY ADMINISTERING AT LEAST 1600 MG/DAY IN TREATMENT OF IPF
- U-2064 DOSING AT LEAST 1602 MG/DAY FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION
- U-2065 FULL DAILY DOSING FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION
- U-2066 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400 MG/DAY DOSE, FOLLOWED BY FULL DAILY DOSE
- U-2067 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE, FOLLOWED BY ADMINISTERING AT LEAST 1602 MG/DAY
- U-2068 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, FOLLOWED BY FULL DAILY DOSE

PATENT AND EXCLUSIVITY TERMS

ADB 117 of 133

PATENT USE

- U-2069 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING A SUB-1600 MG/DAY DOSE, FOLLOWED BY ADMINISTERING AT LEAST 1602 MG/DAY
- U-2070 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, THEN SUB-1600 MG/DAY, THEN AT LEAST 1602 MG/DAY
- U-2071 FULL DAILY DOSING FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2072 FULL DAILY DOSING FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2073 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, FOLLOWED BY ADMINISTERING FULL DAILY DOSE IN TREATMENT OF IPF
- U-2074 DOSING 1602 MG/DAY PIRFENIDONE FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2075 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS FOLLOWED BY ADMINISTERING 1602 MG/DAY IN TREATMENT OF IPF
- U-2076 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING 801 MG/DAY FOLLOWED BY ADMINISTERING 1602 MG/DAY IN TREATMENT OF IPF
- U-2077 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400 MG/DAY DOSE THEN FULL DAY DAILY DOSE IN TREATMENT OF IPF
- U-2078 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN LIMITS, THEN SUB-2400MG/DAY DOSE, THEN FULL DAILY DOSE IN TREATMENT OF IPF
- U-2079 PIRFENIDONE DOSE ESCALATION REGIMEN FOR TREATMENT OF FIBROSIS AS 801 MG/DAY FOR DAYS 1-7 OF THE REGIMEN, 1602 MG/DAY FOR DAYS 8-14 OF THE REGIMEN, AND 2403 MG/DAY FOR AT LEAST DAY 15 OF THE REGIMEN
- U-2080 PIRFENIDONE DOSE ESCALATION REGIMEN FOR TREATMENT OF IPF AS 801 MG/DAY FOR DAYS 1-7 OF THE REGIMEN, 1602 MG/DAY FOR DAYS 8-14 OF THE REGIMEN, AND 2403 MG/DAY FOR AT LEAST DAY 15 OF THE REGIMEN
- U-2081 DISCONTINUING USE OF A CYP1A2 INHIBITOR THAT IS A MODERATE TO STRONG INHIBITOR OF BOTH CYP1A2 AND ANOTHER CYP ENZYME SELECTED FROM CYP2C9, CYP2C19, AND CYP2D6 AND THEN ADMINISTERING PIRFENIDONE
- U-2082 MODIFYING PIRFENIDONE ADMINISTRATION FROM A DOSE OF ABOUT 2400 MG/DAY DOWNWARD BY ABOUT 1600 MG/DAY WHILE CO-ADMINISTERING FLUVOXAMINE TO REDUCE DRUG INTERACTIONS WITH FLUVOXAMINE
- U-2083 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, FOLLOWED BY 801 MG/DAY, DOSE, THEN 1602 MG/DAY IN TREATMENT OF IPF
- U-2084 TREATMENT OF SEVERE CHRONIC PAIN VIA INTRATHECAL INFUSION OF ZICONOTIDE IN PATIENTS ALSO RECEIVING MORPHINE
- U-2085 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH RIFAMPIN
- U-2086 A METHOD FOR ADMINISTERING ESTRADIOL COMPRISING A MONOLITHIC TRANSDERMAL DRUG DELIVERY SYSTEM CONSISTING OF (I) A BACKING LAYER AND (II) A SINGLE ADHESIVE POLYMER MATRIX LAYER AS CLAIMED IN US PATENT NO. 9730900
- U-2087 TREATMENT OF RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) WITH AN ISOCITRATE DEHYDROGENASE-2 (IDH2) MUTATION
- U-2088 TREATMENT OF PARTIAL-ONSET SEIZURES WITH OR WITHOUT SECONDARILY GENERALIZED SEIZURES IN PATIENTS WITH EPILEPSY
- U-2089 TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES AS ADJUNCTIVE THERAPY IN PATIENTS WITH EPILEPSY

PATENT AND EXCLUSIVITY TERMS

ADB 118 of 133

PATENT USE

- U-2090 FOR THE TREATMENT OF ADULTS WITH NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC)
- U-2091 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN, IN A PATIENT NOT HOMOZYGOUS FOR THE UGT1A1*28 ALLELE
- U-2092 METHOD FOR CONFIRMING DOSE DELIVERY
- U-2093 TREATMENT OF TYPE II SPINAL MUSCULAR ATROPHY
- U-2094 TREATMENT OF TYPE III SPINAL MUSCULAR ATROPHY
- U-2095 MITOSOL IS AN ANTIMETABOLITE INDICATED AS AN ADJUNCT TO AB EXTERNO GLAUCOMA SURGERY. IT IS INTENDED FOR TOPICAL APPLICATION TO THE SITE OF GLAUCOMA FILTRATION SURGERY
- U-2096 SOTYLIZE IS INDICATED FOR THE MAINTENANCE OF NORMAL SINUS RHYTHM [DELAY IN TIME TO RECURRENCE OF ATRIAL FIBRILLATION/ATRIAL FLUTTER (AFIB/AFL)] IN PATIENTS WITH SYMPTOMATIC AFIB/AFL WHO ARE CURRENTLY IN SINUS RHYTHM
- U-2097 TREATMENT OF DMD IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- U-2098 INCREASING PRODUCTION OF FUNCTIONAL DYSTROPHIN PROTEIN IN DMD PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- U-2099 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING BRONCHITIS AND/OR EMPHYSEMA
- U-2100 INDICATED FOR THE ONCE-DAILY TREATMENT OF ASTHMA IN PATIENTS 18 YEARS AND OLDER
- U-2101 MAINTENANCE TREATMENT OF RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER, WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- U-2102 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH THREE OR MORE PRIOR LINES OF CHEMOTHERAPY BASED ON AN FDA-APPROVED COMPANION DIAGNOSTIC FOR LYNPARZA
- U-2103 MAINTENANCE TREATMENT OF BRCA-MUTATED RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER, WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- U-2104 TREATMENT OF HYPERURICEMIA ASSOCIATED WITH GOUT IN PATIENTS WHO HAVE NOT ACHIEVED TARGET SERUM URIC ACID LEVELS WITH A MEDICALLY APPROPRIATE DAILY DOSE OF ALLOPURINOL ALONE
- U-2105 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING IMMEDIATE RELEASE LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- U-2106 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- U-2107 TREATMENT OF LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, DIFFERENTIATED THYROID CARCINOMA REFRACTORY TO RADIOACTIVE IODINE TREATMENT
- U-2108 TREATMENT OF HORMONE RECEPTOR POSITIVE HER2-NEGATIVE ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN NOT PREVIOUSLY TREATED WITH ENDOCRINE THERAPY
- U-2109 CAROSPIR IS INDICATED FOR TREATMENT OF NYHA CLASS III-IV HEART FAILURE AND REDUCED EJECTION FRACTION TO INCREASE SURVIVAL, MANAGE EDEMA, AND TO REDUCE THE NEED FOR HOSPITALIZATION FOR HEART FAILURE
- U-2110 METHOD FOR CHRONIC WEIGHT MANAGEMENT IN PATIENTS WITH MODERATE RENAL IMPAIRMENT WHO ARE OBESE, OR OVERWEIGHT AND HAVE AT LEAST ONE WEIGHT RELATED COMORBID CONDITION
- U-2111 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1-5 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 1-5
- U-2112 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 6 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 6
- U-2113 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN

PATENT AND EXCLUSIVITY TERMS

ADB 119 of 133

PATENT USE

- SECRETAGOGUE AS RECITED IN CLAIM 7 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 7
- U-2114 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 9 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 9
- U-2115 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 10 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 10
- U-2116 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 12 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 12
- U-2117 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 14-15 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 14-15
- U-2118 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 16-18 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 16-18
- U-2119 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 19 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 19
- U-2120 TREATMENT OF PATIENTS 18 YEARS OF AGE AND OLDER WITH COMPLICATED URINARY TRACT INFECTIONS CAUSED BY SUSCEPTIBLE MICROORGANISMS
- U-2121 TREATMENT OF PARTIAL-ONSET SEIZURES IN A PATIENT SUFFERING FROM OR SUSCEPTIBLE TO ABSENCE SEIZURES
- U-2122 USE FOR REDUCING EXACERBATIONS OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE
- U-2123 TREATMENT OF PARTIAL-ONSET SEIZURES IN PATIENTS WITH EPILEPSY WHO HAVE BEEN PREVIOUSLY TREATED WITH OXCARBAZEPINE
- U-2124 TREATMENT OF ADULT PATIENTS WITH RELAPSED FOLLICULAR LYMPHOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- U-2125 THE TREATMENT OF AN INFLAMMATORY DISORDER OF THE RESPIRATORY TRACT BY ONCE-PER-DAY ADMINISTRATION OF A PHARMACEUTICAL FORMULATION COMPRISING FLUTICASONE FUROATE AND A LONG-ACTING BETA2 ADRENORECEPTOR AGONIST
- U-2126 USE OF FLUTICASONE FUROATE FOR THE TREATMENT OF AN INFLAMMATORY OR ALLERGIC CONDITIONS, INCLUDING CHRONIC OBSTRUCTIVE PULMONARY DISEASE
- U-2127 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA
- U-2128 METHOD OF INHIBITING THE BINDING OF ACETYLCHOLINE TO AN ACETYLCHOLINE RECEPTOR IN THE RESPIRATORY TRACT OF A HUMAN, COMPRISING CONTACTING THE RECEPTOR WITH AN EFFECTIVE AMOUNT OF UMECLIDINIUM, VIA INHALATION
- U-2129 METHOD OF INHIBITING THE BINDING OF ACETYLCHOLINE TO AN ACETYLCHOLINE RECEPTOR IN THE RESPIRATORY TRACT OF A HUMAN, COMPRISING CONTACTING THE RECEPTOR WITH AN EFFECTIVE AMOUNT OF UMECLIDINIUM, VIA TOPICAL APPLICATION
- U-2130 TREATMENT OF PARTIAL ONSET SEIZURES IN PATIENTS WITH EPILEPSY AGED 16 YEARS AND OLDER WITH EPILEPSY
- U-2131 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION, WITH COMPARABLE EFFICACY, AND A REDUCTION IN SPECIFIED ADVERSE EVENTS, COMPARED TO BRIMONIDINE 0.2% TID
- U-2132 IN COMBINATION WITH FULVESTRANT FOR THE TREATMENT OF WOMEN WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY
- U-2133 METHOD OF DELIVERING FLUTICASONE PROPIONATE TO A NASAL AIRWAY
- U-2134 THE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE BY ONCE-PER-DAY ADMINISTRATION OF A PHARMACEUTICAL FORMULATION COMPRISING FLUTICASONE FUROATE

PATENT AND EXCLUSIVITY TERMS

ADB 120 of 133

PATENT USE

AND A LONG-ACTING BETA2 ADRENORECEPTOR

- U-2135 AS MONOTHERAPY FOR THE TREATMENT OF ADULT PATIENTS WITH HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY AND PRIOR CHEMOTHERAPY IN THE METASTATIC SETTING
- U-2136 TREATMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY
- U-2137 TREATMENT OF POSTHERPETIC NEURALGIA
- U-2138 TOPICAL TREATMENT OF ACTINIC KERATOSIS OF THE FACE OR SCALP USING MORE THAN ONE TREATMENT COURSE OF INGENOL MEBUTATE
- U-2139 TREATMENT OF TYPE 2 DIABETES MELLITUS IN COMBINATION WITH EXENATIDE
- U-2140 METHOD OF TREATING PARTIAL ONSET SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- U-2141 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6
- U-2142 REDUCTION IN THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS (DVT) AND/OR PULMONARY EMBOLISM (PE) IN PATIENTS AT CONTINUED RISK FOR RECURRENT DVT AND/OR AFTER COMPLETION OF INITIAL TREATMENT LASTING AT LEAST 6 MONTHS
- U-2143 AFTER COMPLETION OF INITIAL TREATMENT LASTING AT LEAST 6 MONTHS, TO REDUCE THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS AND/OR PULMONARY EMBOLISM IN CERTAIN PATIENTS WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST 5 CONSECUTIVE DAYS
- U-2144 REDUCTION OF INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- U-2145 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-2146 IMPROVEMENT IN GLYCEMIC CONTROL IN DIABETES MELLITUS PATIENTS BY USE OF A PEN INJECTOR WITH A ROTATING DRIVE SLEEVE
- U-2147 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY ORALLY ADMINISTERING 20MG OF TASIMELTEON ONCE DAILY BEFORE BEDTIME
- U-2148 A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT BY MEASURING AND DISPLAYING AN INDICATION OF THE CALCULATED DELIVERY CONCENTRATION OF NITIRIC OXIDE AS COMPARED TO THE DESIRED DELIVERY CONCENTRATION OF NITRIC OXIDE
- U-2149 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY ADMINISTERING TASIMELTEON
- U-2150 TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE
- U-2151 METHOD OF TREATING PAIN OR INFLAMMATION WITH AN INJECTABLE CONTROLLED OR SUSTAINED RELEASE FORMULATION OF TRIAMCINOLONE ACETONIDE
- U-2152 TREATMENT OF PAIN ASSOCIATED WITH IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) WITH VIBERZI (ELUXADOLINE)
- U-2153 REDUCING FASTING PLASMA GLUCOSE IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2154 REDUCING FASTING PLASMA GLUCOSE IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2155 REDUCING BODY WEIGHT IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2156 REDUCING HBA1C IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2157 TREATING TYPE 2 DIABETES MELLITUS BY STIMULATING INSULIN RELEASE
- U-2158 DECREASING GASTRIC MOTILITY OR DELAYING GASTRIC EMPTYING BY USING A SUSTAINED-RELEASE COMPOSITION
- U-2159 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA/SMALL LYMPHOCYTIC LYMPHOMA
- U-2160 MANAGEMENT OF OSTEOARTHRITIS PAIN BY ADMINISTERING 5 MG OF MELOXICAM
- U-2161 TREATMENT OF NAUSEA AND VOMITING, INCLUDING THE PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF HIGHLY OR MODERATELY EMETOGENIC CANCER CHEMOTHERAPY
- U-2162 FOR CLEANSING THE LARGE INTESTINE AS A PREPARATION FOR COLONOSCOPY
- U-2163 TREATMENT OF HR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER-2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN COMBINATION WITH PALBOCICLIB OR ABEMACICLIB IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY

PATENT AND EXCLUSIVITY TERMS

ADB 121 of 133

PATENT USE

U-2164 ZELBORAF IS INDICATED FOR THE TREATMENT OF PATIENTS WITH ERDHEIM-CHESTER DISEASE WITH BRAF V600 MUTATION

U-2165 MANAGEMENT OF OSTEOARTHRITIS PAIN BY ADMINISTERING 10 MG OF MELOXICAM

U-2166 TREATMENT OF MAJOR DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I DISORDER

U-2167 METHOD OF USING A TABLET EMBEDDED WITH A SENSOR THAT COMMUNICATES INFORMATION VIA A SIGNAL THROUGH THE BODY OF A PATIENT TO A RECEIVER

U-2168 METHOD OF USING A LOGIC CIRCUIT TO STABILIZE BATTERY VOLTAGE SUPPLIED TO A SENSOR EMBEDDED WITH A TABLET AND THAT COMMUNICATES INFORMATION VIA A SIGNAL THROUGH THE BODY OF A PATIENT TO A RECEIVER

U-2169 METHOD OF USING A RECEIVER TO IDENTIFY A SIGNAL FROM A TABLET EMBEDDED WITH A SENSOR THAT COMMUNICATES INFORMATION THROUGH THE BODY OF A PATIENT

U-2170 METHOD OF USING A RECEIVER TO RECEIVE A SIGNAL FROM A TABLET EMBEDDED WITH A SENSOR THAT COMMUNICATES INFORMATION THROUGH THE BODY OF A PATIENT

U-2171 ADJUVANT TREATMENT OF ADULT PATIENTS AT HIGH RISK OF RECURRENT RCC FOLLOWING NEPHRECTOMY

U-2172 METHOD TO TREAT SEVERE ALLERGIC EMERGENCIES IN PATIENTS WEIGHING 7.5 TO 15 KG (16.5 TO 33 LBS)

U-2173 TREATING OPIOID DEPENDENCE BY ADMINISTERING BUPRENORPHINE

U-2174 TREATING OPIOID DEPENDENCY BY ADMINISTERING BUPRENORPHINE ONCE PER MONTH

U-2175 TREATING OPIOID DEPENDENCY BY ADMINISTERING BUPRENORPHINE ONCE MONTHLY

U-2176 TREATING OPIOID ADDICTION BY ADMINISTERING BUPRENORPHINE

U-2177 TREATING OPIOID ADDICTION BY SUBCUTANEOUS INJECTION OF BUPRENORPHINE

U-2178 TREATING OPIOID ADDICTION BY ADMINISTERING BUPRENORPHINE COMPOSITION WITH 28 DAY DOSE DURATION

U-2179 IN SITU FORMATION OF SOLID BUPRENORPHINE COMPOSITION

U-2180 TREATING ADDICTION WITH 100 MG OR 300 MG DOSE OF BUPRENORPHINE

U-2181 TREATING OPIOID DEPENDENCY BY SUBCUTANEOUSLY ADMINISTERING BUPRENORPHINE

U-2182 IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS

U-2183 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 13

U-2184 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1, 13, AND 14

U-2185 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 15 AND 27

U-2186 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 15, 27, AND 28

U-2187 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 29 AND 39

U-2188 ADJUNCT TO DIET AND EXECRISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 29, 39, AND 40

U-2189 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 41 AND 52

U-2190 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 41, 52, AND 53

U-2191 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 54 AND 64

U-2192 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 54, 64, AND 65
- U-2193 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 66 AND 75
- U-2194 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 66, 75, AND 76
- U-2195 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 77 AND 87
- U-2196 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 77, 87, AND 88
- U-2197 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 89 AND 99
- U-2198 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 89, 99, AND 100
- U-2199 TREATMENT OF SCHIZOPHRENIA WITH IMPROVEMENT IN ATTENTION FUNCTION IN SCHIZOPHRENIA
- U-2200 COMBINATION TREATMENT WITH INSULIN GLARGINE WITH OR WITHOUT METFORMIN FOR IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS
- U-2201 TREATMENT OF BIPOLAR DEPRESSION WITH IMPROVEMENT IN ATTENTION FUNCTION IN BIPOLAR DISORDER
- U-2202 OZEMPIC IS INDICATED AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2203 A METHOD OF PROVIDING A SUBJECT WITH A THERAPEUTICALLY EFFECTIVE AMOUNT OF RACEMIC METHYLPHENIDATE BY ORALLY ADMINISTERING TO SAID SUBJECT A SINGLE METHYLPHENIDATE EXTENDED RELEASE CHEWABLE TABLET AS CLAIMED
- U-2204 TREATING PATIENTS WITH ACUTE PROMYELOCYTIC LEUKEMIA (APL) WHO ARE REFRACTORY TO, OR HAVE RELAPSED FROM, RETINOID AND ANTHRACYCLINE CHEMOTHERAPY, AND WHOSE APL IS CHARACTERIZED BY THE PRESENCE OF THE T(15;17) TRANSLOCATION OR PML/RAR-ALPHA GENE EXPRESSION
- U-2205 TREATMENT OF SEBORRHEIC KERATOSES THAT ARE RAISED
- U-2206 TREATING OPIOID DEPENDENCY BY ADMINISTERING BUPRENORPHINE
- U-2207 TREATING ADDICTION BY SUBCUTANEOUS INJECTION OF BUPRENORPHINE
- U-2208 TREATING ADDICTION BY ONCE PER MONTH ADMINISTRATION OF BUPRENORPHINE
- U-2209 TREATING OPIOID ADDICTION BY ADMINISTERING BUPRENORPHINE ONCE PER MONTH
- U-2210 TREATING OPIOID ADDICTION BY 100 MG OR 300 MG DOSE BUPRENORPHINE
- U-2211 TREATING OPIOID ADDICTION BY ADMINISTRATION OF BUPRENORPHINE
- U-2212 REDUCING FASTING PLASMA GLUCOSE IN A HUMAN IN NEED THEREOF IN COMBINATION WITH A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2213 REDUCING HBA1C IN A HUMAN IN NEED THEREOF IN COMBINATION WITH A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2214 AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES
- U-2215 ERTUGLIFLOZIN IN COMBINATION WITH SITAGLIPTIN AND IN FURTHER COMBINATION WITH METFORMIN AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2216 ERTUGLIFLOZIN AND SITAGLIPTIN IN COMBINATION AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2217 TREATING HIGH OUTPUT SHOCK WITH ANGIOTENSIN II BY INCREASING MEAN ARTERIAL PRESSURE IN PATIENTS TREATED WITH CATECHOLAMINES AND REDUCING CATECHOLAMINE USE
- U-2218 MAINTAINING MEAN ARTERIAL PRESSURE OF ABOUT 65 MMHG OR HIGHER WITH ANGIOTENSIN II IN SHOCK PATIENTS TREATED WITH CATECHOLAMINES AND REDUCING CATECHOLAMINE USE

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-2219 TREATMENT OF CHRONIC SMALL LYMPHOCYTIC LEUKEMIA

U-2220 A METHOD FOR THE DIAGNOSIS OF ADULT GROWTH HORMONE DEFICIENCY BY MEASURING THE LEVEL OF GROWTH HORMONE AFTER ORAL ADMINISTRATION OF MACIMORELIN

U-2221 TREATING REFRACTORY HYPOTENSION WITH ABOUT 20 NG/KG/MIN ANGIOTENSIN II IN A PATIENT RECEIVING VASOPRESSOR

U-2222 RELIEVES REDNESS OF THE EYE DUE TO MINOR EYE IRRITATIONS

U-2223 METHOD OF TREATING ANGINA PECTORIS

U-2224 TREATMENT OF DYSKINESIA AND INCREASING ON TIME WITHOUT TROUBLESOME DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS

U-2225 METHOD OF ADMINISTERING A LOCAL ANESTHETIC TO THE MUCOUS MEMBRANES IN PATIENTS WITH HEPATIC IMPAIRMENT

U-2226 METHOD OF ADMINISTERING A LOCAL ANESTHETIC TO THE MUCOUS MEMBRANES IN PATIENTS WITH RENAL IMPAIRMENT

U-2227 METHOD OF ADMINISTERING A LOCAL ANESTHETIC TO THE MUCOUS MEMBRANES IN GERIATRIC PATIENTS

U-2228 TREATMENT OF SMALL LYMPHOCYTIC LEUKEMIA

U-2229 IN COMBINATION WITH TRETINOIN, TREATING ADULTS AND PEDIATRIC PATIENTS 1 YEAR AND OLDER WITH NEWLY-DIAGNOSED LOW-RISK ACUTE PROMYELOCYTIC LEUKEMIA (APL) CHARACTERIZED BY THE PRESENCE OF THE T(15;17) TRANSLOCATION OR PML/RAR-A GENE EXPRESSION

U-2230 IRRITABLE BOWEL SYNDROME WITH CONSTIPATION

U-2231 TREATING REFRACTORY HYPOTENSION WITH ABOUT 5 NG/KG/MIN TO ABOUT 20 NG/KG/MIN ANGIOTENSIN II IN A PATIENT RECEIVING VASOPRESSOR

U-2232 TREATMENT OF PSORIATIC ARTHRITIS USING A DOSAGE TITRATION SCHEDULE

U-2233 TREATMENT OF PSORIATIC ARTHRITIS WITH APREMILAST USING A DOSAGE TITRATION SCHEDULE AND A SECOND ACTIVE AGENT

U-2234 USE OF IVACAFTOR FOR TREATING CYSTIC FIBROSIS IN A PATIENT WITH A MILD TO MODERATE CF PHENOTYPE WITH AT LEAST ONE MUTATION IN THE CFTR GENE THAT IS RESPONSIVE TO IVACAFTOR BASED ON CLINICAL AND/OR IN VITRO ASSAY DATA

U-2235 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC HIGH-RISK CASTRATION-SENSITIVE PROSTATE CANCER

U-2236 REDUCING THE RISK OF PRETERM BIRTH IN WOMEN WITH A SINGLETON PREGNANCY WHO HAVE A HISTORY OF SINGLETON SPONTANEOUS PRETERM BIRTH

U-2237 TREATMENT OF NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NM-CRPC)

U-2238 METHOD OF IMPROVING GLYCEMIC CONTROL IN PATIENTS WITH DIABETES MELLITUS BY ADMINISTERING A MIXTURE OF INSULIN DEGLUDEC AND INSULIN ASPART DURING OR AROUND THE TIME OF THE LARGEST MEAL OF THE DAY

U-2239 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION, WITH A REDUCTION IN SPECIFIED ADVERSE EVENTS, COMPARED TO BRIMONIDINE 0.2% TID

U-2240 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION, WITH COMPARABLE EFFICACY TO BRIMONIDINE 0.2% TID

U-2241 TREATMENT OF SMALL LYMPHOCYTIC LYMPHOMA WITH 17P DELETION

U-2242 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA WITH 17P DELETION

U-2243 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA/SMALL LYMPHOCYTIC LYMPHOMA WITH 17P DELETION

U-2244 A METHOD OF TREATING BACTERIAL INFECTIONS IN HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) PATIENTS COMPRISING ADMINISTERING A BACTERICIDALLY EFFECTIVE AMOUNT OF AVIBACTAM SODIUM

U-2245 A METHOD OF TREATING A BACTERIAL INFECTION IN HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA AND VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) PATIENTS COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AVIBACTAM SODIUM

U-2246 TEZACAFTOR AND IVACAFTOR FOR THE TREATMENT OF CYSTIC FIBROSIS IN PATIENTS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVING AT LEAST ONE CFTR GENE MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR BASED ON IN VITRO DATA AND/OR CLINICAL EVIDENCE

PATENT AND EXCLUSIVITY TERMS

ADB 124 of 133

PATENT USE

- U-2247 TEZACAFTOR AND IVACAFTOR FOR THE TREATMENT OF PATIENTS WITH A MILD TO MODERATE CLINICAL PHENOTYPE OF CYSTIC FIBROSIS HAVING AT LEAST ONE CFTR GENE MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR BASED ON IN VITRO DATA AND/OR CLINICAL EVIDENCE
- U-2248 TEZACAFTOR AND IVACAFTOR FOR THE TREATMENT OF CYSTIC FIBROSIS IN PATIENTS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HETEROZYGOUS FOR THE F508DEL MUTATION AND A SECOND MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR
- U-2249 MANAGEMENT OF ACUTE PAIN SEVERE ENOUGH TO REQUIRE AN OPIOID ANALGESIC AND FOR WHICH ALTERNATIVE TREATMENTS ARE INADEQUATE
- U-2250 DETECTION OF CARCINOMA IN THE BLADDER BY PHOTODYNAMIC CYSTOSCOPY
- U-2251 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-2252 THE TREATMENT OF ACUTE OTITIS EXTERNA IN PATIENTS 6 MONTHS OF AGE AND OLDER DUE TO PSEUDOMONAS AERUGINOSA AND STAPHYLOCOCCUS AUREUS
- U-2253 PROPHYLACTIC TREATMENT OF NAUSEA AND VOMITING, INCLUDING PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING ASSOCIATED CHEMOTHERAPY
- U-2254 USE OF POMALIDOMIDE WITH DEXAMETHASONE FOR PATIENTS WITH MULTIPLE MYELOMA AFTER AT LEAST TWO PRIOR THERAPIES INCLUDING LENALIDOMIDE AND A PROTEASOME INHIBITOR AND DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETING THE LAST THERAPY
- U-2255 TREATING SECONDARY HYPERPARATHYROIDISM IN CHRONIC KIDNEY DISEASE WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE LEVEL AND THE SUSTAINED RELEASE IS OVER AT LEAST 10 HOURS
- U-2256 TREATING SECONDARY HYPERPARATHYROIDISM IN CHRONIC KIDNEY DISEASE WITH SUSTAINED RELEASE 25-HYDROXYVITAMIN D TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE LEVEL AND CMAX IS REDUCED COMPARED TO BOLUS IV INJECTION AND IMMEDIATE-RELEASE, ORAL DOSING
- U-2257 TREATING SHPT IN CKD WITH SUSTAINED RELEASE CALCIFEDIOL TO REDUCE SERUM PARATHYROID HORMONE LEVEL AND CHANGE IN SERUM CONCENTRATION OF CALCIFEDIOL IN DOSE INTERVAL IS REDUCED COMPARED TO BOLUS IV INJECTION AND IMMEDIATE-RELEASE, ORAL DOSING
- U-2258 TREATING SECONDARY HYPERPARATHYROIDISM IN CKD WITH SUSTAINED RELEASE CALCIFEDIOL TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE LEVEL AND CMAX24HR/C24HR IS REDUCED COMPARED TO BOLUS IV INJECTION AND IMMEDIATE-RELEASE, ORAL DOSING
- U-2259 TREATING SECONDARY HYPERPARATHYROIDISM IN CKD WITH SUSTAINED RELEASE CALCIFEDIOL TO REDUCE THE PATIENT'S SERUM PARATHYROID HORMONE LEVEL AND TMAX IS INCREASED COMPARED TO BOLUS IV INJECTION AND IMMEDIATE-RELEASE, ORAL DOSING
- U-2260 METHOD OF REDUCING THE RISK OF PERIPROCEDURAL MYOCARDIAL INFARCTION, AND STENT THROMBOSIS IN A PATIENT UNDERGOING PCI BY ADMINISTERING INTRAVENOUSLY 30 UG/KG BOLUS BEFORE PCI AND THEN A CONTINUOUS INFUSION
- U-2261 MODIFIED DOSING REGIMEN FOR THE MANAGEMENT OF MILD TO MODERATE PAIN OR MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS
- U-2262 MODIFIED DOSING REGIMEN FOR THE REDUCTION OF FEVER
- U-2263 MODIFIED DOSING REGIMEN FOR THE MANAGEMENT OF MODERATE TO SEVERE PAIN WITH ADJUNCTIVE OPIOID ANALGESICS
- U-2264 METHODS OF TREATING PAIN, INFLAMMATION, FEVER, PATENT DUCTUS ARTERIOSIS WITH AQUEOUS COMPOSITION
- U-2265 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH HEC AND MEC IN ADULT AND PEDIATRIC PATIENTS
- U-2266 METHODS OF MAKING AQUEOUS COMPOSITION AND TREATING PAIN, INFLAMMATION, FEVER, PATENT DUCTUS ARTERIOSIS WITH AQUEOUS COMPOSITION
- U-2267 METHOD FOR RELIEVING THE PAIN ASSOCIATED WITH POST-HERPETIC NEURALGIA
- U-2268 DISCONTINUING A STRONG CYP1A2 INDUCER TO AVOID REDUCED PIRFENIDONE EFFICACY AND THEN ADMINISTERING PIRFENIDONE
- U-2269 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400MG/DAY DOSE THEN FULL DAILY DOSE IN TREATMENT OF IPF

PATENT AND EXCLUSIVITY TERMS

ADB 125 of 133

PATENT USE

- U-2270 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, THEN SUB-2400MG/DAY DOSE, THEN FULL DAILY DOSE IN TREATMENT OF IPF
- U-2271 THERAPEUTIC TREATMENT OF PATIENTS WITH CASTRATION-RESISTANT PROSTATE CANCER, SYMPTOMATIC BONE METASTASES AND NO KNOWN VISCERAL METASTATIC DISEASE
- U-2272 TREATMENT OF NASAL POLYPS IN PATIENTS ≥ 18 YEARS OF AGE WHO HAVE HAD ETHMOID SINUS SURGERY USING A CORTICOSTEROID-ELUTING (MOMETASONE FUROATE) IMPLANT
- U-2273 A METHOD FOR TREATING EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER, WHEREIN THE CANCER IS ASSOCIATED WITH A DELETERIOUS BRCA MUTATION
- U-2274 MAINTAINING SERUM 25-HYDROXYVITAMIN D AT A LEVEL OF AT LEAST 30 NG/ML WITH ORAL, SUSTAINED RELEASE 25-HYDROXYVITAMIN D
- U-2275 TREATING CYSTIC FIBROSIS PATIENTS AGES 12 AND OLDER, WHO ARE HOMOZYGOUS FOR F508DEL OR HAVE AT LEAST 1 CFTR GENE MUTATION RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS ($<30\%$ CRYSTALLINE) IVACAFTOR
- U-2276 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT AGE 6 OR OLDER HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS (LESS THAN ABOUT 30% CRYSTALLINE) IVACAFTOR
- U-2277 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS INADEQUATELY CONTROLLED BY LIXISENATIDE
- U-2278 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS INADEQUATELY CONTROLLED BY LIXISENATIDE IN COMBINATION WITH METFORMIN
- U-2279 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS INADEQUATELY CONTROLLED BY LIXISENATIDE IN COMBINATION WITH METFORMIN AND A SECOND ORAL ANTIDIABETIC DRUG
- U-2280 ADJUNCTIVE TREATMENT OF PATIENTS WITH TSC-ASSOCIATED PARTIAL-ONSET SEIZURES
- U-2281 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 1 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 1
- U-2282 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 2 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 2
- U-2283 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 3-7 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 3-7
- U-2284 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 8 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 8
- U-2285 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 11 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 11
- U-2286 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 14 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 14
- U-2287 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 16-19 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 16-19
- U-2288 TREATMENT OF TYPE 2 DIABETES MELLITUS WITH EXENATIDE AS AN ADD-ON TO BASIL INSULIN OR BASAL INSULIN PLUS METFORMIN THERAPY
- U-2289 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21L858R MUTATIONS

PATENT AND EXCLUSIVITY TERMS

ADB 126 of 133

PATENT USE

- U-2290 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT (45 ML/MIN/1.73 M2<=EGFR<60 ML/MIN/1.73 M2) BY ONCE DAILY ADMINISTRATION OF 10 MG OR 25 MG OF EMPAGLIFLOZIN
- U-2291 REDUCTION OF THROMBOTIC CARDIOVASCULAR EVENTS IN PATIENTS WITH A HISTORY OF MYOCARDIAL INFARCTION (MI) OR WITH PERIPHERAL ARTERIAL DISEASE (PAD)
- U-2292 METHOD OF REDUCING THE RISK OF CARDIOVASCULAR DEATH IN ADULT PATIENTS WITH TYPE 2 DIABETES MELLITUS AND CARDIOVASCULAR DISEASE BY ONCE DAILY ADMINISTRATION OF 10 MG OR 25 MG OF EMPAGLIFLOZIN
- U-2293 USE IN COMBINATION WITH DEXAMETHASONE IN ADULTS FOR THE PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF CANCER CHEMOTHERAPY, INCLUDING, BUT NOT LIMITED TO, HIGHLY EMETOGENIC CHEMOTHERAPY
- U-2294 TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC IMMUNE THROMBOCYTOPENIA (ITP) WHO HAVE HAD AN INSUFFICIENT RESPONSE TO A PREVIOUS TREATMENT
- U-2295 TREATMENT OF PARTIAL-ONSET SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- U-2296 TAFINLAR IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- U-2297 IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES PATIENTS BY ADMINISTERING A STARTING DOSE OF 10 MCG FOR 14 DAYS AND INCREASING TO A MAINTENANCE DOSE OF 20 MCG ON DAY 15
- U-2298 TAFINLAR IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ANAPLASTIC THYROID CANCER (ATC) WITH BRAF V600E MUTATION AND WITH NO SATISFACTORY LOCOREGIONAL TREATMENT OPTIONS
- U-2299 TAFINLAR IS INDICATED AS A SINGLE AGENT FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATION
- U-2300 USE IN COMBINATION WITH THE MUSCARINIC ANTAGONIST SOLIFENACIN SUCCINATE FOR THE TREATMENT OF OVERACTIVE BLADDER (OAB) WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY
- U-2301 USE IN COMBINATION WITH DEXAMETHASONE IN ADULTS FOR THE PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF HIGHLY EMETOGENIC CANCER CHEMOTHERAPY
- U-2302 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST, AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- U-2303 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE ADJUVANT TREATMENT OF PATIENTS WITH MELANOMA WITH BRAF V600E OR V600K MUTATIONS AND INVOLVEMENT OF LYMPH NODE(S), FOLLOWING COMPLETE RESECTION
- U-2304 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION
- U-2305 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC ANAPLASTIC THYROID CANCER (ATC) WITH BRAF V600E MUTATION AND WITH NO SATISFACTORY LOCOREGIONAL TREATMENT OPTIONS
- U-2306 ONCE DAILY TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN ADULTS WITH 1% OXYMETAZOLINE HYDROCHLORIDE CREAM
- U-2307 TREATMENT OF AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE
- U-2308 TREATMENT OF ADULT PATIENTS WITH SHORT BOWEL SYNDROME WHO ARE DEPENDENT ON PARENTERAL SUPPORT
- U-2309 USE IN THE TREATMENT OF MAJOR DEPRESSIVE DISORDER TO IMPROVE PROCESSING SPEED, AN ASPECT OF COGNITIVE FUNCTION
- U-2310 FOR CLEANSING OF THE COLON IN PREPARATION FOR COLONOSCOPY IN ADULTS
- U-2311 TREATMENT OF HYPERURICEMIA ASSOCIATED WITH GOUT IN PATIENTS WHO HAVE NOT ACHIEVED TARGET SERUM URIC ACID LEVELS WITH A XANTHINE OXIDASE INHIBITOR ALONE
- U-2312 TREATMENT OF HYPERKALEMIA IN ADULTS
- U-2313 METHOD OF REDUCING THE RISK OF CARDIOVASCULAR DEATH, NON-FATAL MYOCARDIAL INFARCTION, AND/OR NON-FATAL STROKE IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- ESTABLISHED CARDIOVASCULAR DISEASE BY ADMINISTERING LIRAGLUTIDE
- U-2314 TREATMENT OF THROMBOCYTOPENIA IN AN ADULT PATIENT WITH CHRONIC LIVER DISEASE WHO IS SCHEDULED TO UNDERGO A PROCEDURE USING DOPTELET
- U-2315 TREATMENT OF MULTIPLE SCLEROSIS IN THE PEDIATRIC PATIENT POPULATION WITH 0.25 MG FINGOLIMOD
- U-2316 TREATMENT OF DYSPAREUNIA
- U-2317 TREATMENT OF A SYMPTOM OF VULVAR AND VAGINAL ATROPHY
- U-2318 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HAVE AT LEAST ONE CFTR GENE MUTATION THAT IS RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND IVACAFTOR
- U-2319 KYPROLIS IS INDICATED IN COMBINATION WITH DEXAMETHASONE OR WITH LENALIDOMIDE PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY
- U-2320 KYPROLIS IS INDICATED AS A SINGLE AGENT FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE OR MORE LINES OF THERAPY
- U-2321 A METHOD OF APPLYING TRYPAN BLUE ONTO AN OUTER SURFACE OF THE ANTERIOR LENS CAPSULE TO FACILITATE REMOVAL OF THE LENS SUBSTANCE
- U-2322 TREATMENT OF ADULT PATIENTS WITH MODERATELY TO SEVERELY ACTIVE ULCERATIVE COLITIS (UC)
- U-2323 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL), WITH OR WITHOUT 17P DELETION, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-2324 FOR SECONDARY PREVENTION OF CARDIOVASCULAR AND CEREBROVASCULAR EVENTS IN PATIENTS AT RISK OF DEVELOPING ASPIRIN-ASSOCIATED GASTRIC ULCERS
- U-2325 EMERGENCY TREATMENT OF ALLERGIC REACTIONS (TYPE 1), INCLUDING ANAPHYLAXIS; A METHOD OF TREATING ALLERGIC REACTION, ANAPHYLAXIS, ANAPHYLACTIC SHOCK, OR COMBINATION THEREOF BY AN INJECTION OF AT LEAST ONE DOSAGE OF THE INJECTABLE LIQUID PHARMACEUTICAL
- U-2326 TREATMENT OF NOCTURIA DUE TO NOCTURNAL POLYURIA IN ADULTS
- U-2327 TREATMENT OF NOCTURIA DUE TO NOCTURNAL POLYURIA IN ADULTS, COMPRISING MONITORING A PATIENT'S SERUM SODIUM CONCENTRATION
- U-2328 METHOD OF USING PLAZOMICIN TO TREAT BACTERIAL INFECTIONS
- U-2329 METHOD OF ADMINISTERING A LOCAL ANESTHETIC PRIOR TO PERFORMING A DIAGNOSTIC OR SURGICAL PROCEDURE ON A SUBJECT WITH HEPATIC OR RENAL IMPAIRMENT
- U-2330 METHOD OF TREATING MELANOMA
- U-2331 INDICATED IN COMBINATION WITH ENCORAFENIB FOR THE TREATMENT OF MELANOMA
- U-2332 INDICATED IN COMBINATION WITH ENCORAFENIB FOR THE TREATMENT OF MELANOMA MEDIATED BY A B-RAF PROTEIN KINASE
- U-2333 INDICATED IN COMBINATION WITH ENCORAFENIB FOR THE TREATMENT OF MELANOMA WITH A BRAF MUTATION
- U-2334 TREATMENT OF MELANOMA WITH A BRAF MUTATION
- U-2335 TREATMENT OF MELANOMA
- U-2336 TREATMENT OF MELANOMA MEDIATED BY A B-RAF PROTEIN KINASE
- U-2337 INDICATED IN COMBINATION WITH BINIMETINIB FOR THE TREATMENT OF MELANOMA WITH A BRAF MUTATION
- U-2338 MAINTAINING MEAN ARTERIAL PRESSURE OF ABOUT 65 MMHG OR ABOVE WITH ABOUT 1 NG/KG/MIN TO ABOUT 40 NG/KG/MIN ANGIOTENSIN II IN HYPOTENSIVE PATIENTS TREATED WITH VASOPRESSIN OR A VASOPRESSIN ANALOGUE AND REDUCING VASOPRESSIN OR VASOPRESSIN ANALOGUE USE
- U-2339 USE OF A PHARMACEUTICAL COMPOSITION COMPRISING LINAGLIPTIN, METFORMIN AND A BASIC AMINO ACID TO TREAT TYPE 2 DIABETES MELLITUS
- U-2340 TREATMENT OF POSTOPERATIVE INFLAMMATION
- U-2341 METHOD OF RECONSTITUTING A LYOPHILIZED LIPOSOMAL COMPOSITION FOR ADMINISTERING CYTARABINE AND DAUNORUBICIN TO TREAT ADULTS WITH NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- MRC)
- U-2342 METHOD OF ADMINISTERING A RECONSTITUTED LIPOSOMAL COMPOSITION CONTAINING CYTARABINE AND DAUNORUBICIN TO TREAT ADULTS WITH NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC)
- U-2343 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGED 12 YEARS AND OLDER, WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION OR HETEROZYGOUS FOR F508DEL AND A SECOND CFTR MUTATION PREDICTED TO BE RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH TEZACAFTOR AND IVACAFTOR
- U-2344 TREATMENT OF THROMBOCYTOPENIA IN ADULT PATIENTS WITH CHRONIC LIVER DISEASE WHO ARE SCHEDULED TO UNDERGO A PROCEDURE
- U-2345 TREATMENT OF PATIENTS WITH CASTRATION-RESISTANT PROSTATE CANCER (CRPC)
- U-2346 TREATMENT OF HUMAN SMALLPOX DISEASE CAUSED BY VARIOLA VIRUS IN ADULTS AND PEDIATRIC PATIENTS WEIGHING AT LEAST 13 KG
- U-2347 TREATMENT OF TYPE 2 DIABETES MELLITUS IN A PATIENT WITH RENAL IMPAIRMENT AND FOR WHOM METFORMIN THERAPY IS INAPPROPRIATE BY ADMINISTERING LINAGLIPTIN WITHOUT DOSE ADJUSTMENT
- U-2348 A METHOD FOR PREVENTION OF PREGNANCY
- U-2349 FOR ONCE-DAILY MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN PATIENTS AGED 5 YEARS AND OLDER
- U-2350 A METHOD OF TREATING A CANCER CHARACTERIZED BY AN IDH1 MUTATION WHERE THE CANCER IS ACUTE MYELOGENOUS LEUKEMIA (AML)
- U-2351 TREATMENT OF ACUTE MYELOID LEUKEMIA (AML) WITH AN IDH1 MUTATION
- U-2352 TREATMENT OF HIV-1 INFECTION IN ADULTS WHO HAVE NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY OR ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS
- U-2353 TX OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PK ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENATES IN ADULTS WHO HAVE NO PRIOR ANTIRETROVIRAL TX HISTORY OR ARE VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS
- U-2354 COMBINATION WITH OTHER ANTIRETROVIRALS (ATV) FOR TREATMENT OF HIV-1 IN ATV TREATMENT-EXPERIENCED PATIENTS 2 YEARS AND OLDER WITH EVIDENCE OF VIRAL REPLICATION AND HIV-1 STRAINS RESISTANT TO NON-NUCLEOSIDE REVERSE TRANSCRIPTASE INHIBITOR AND OTHER ATV
- U-2355 IN COMBINATION WITH AN AROMATASE INHIBITOR FOR THE TREATMENT OF PRE/PERIMENOPAUSAL OR POSTMENOPAUSAL WOMEN WITH HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER, AS INITIAL ENDOCRINE-BASED THERAPY
- U-2356 IN COMBINATION WITH FULVESTRANT FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER, AS INITIAL ENDOCRINE BASED THERAPY OR FOLLOWING DISEASE PROGRESSION ON ENDOCRINE THERAPY
- U-2357 METHOD OF TREATING ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)
- U-2358 TREATMENT OF PATIENTS WITH HORMONE RECEPTOR (HR)-NEGATIVE BREAST CANCER WITH DELETERIOUS OR SUSPECTED DELETERIOUS GBRCAM, HER2-NEGATIVE METASTATIC BREAST CANCER, WHO HAVE BEEN TREATED WITH CHEMOTHERAPY IN NEOADJUVANT, ADJUVANT, OR METASTATIC SETTING
- U-2359 TREATMENT OF PATIENTS WITH HORMONE RECEPTOR (HR)-POSITIVE BREAST CANCER WHO SHOULD HAVE BEEN TREATED WITH PRIOR ENDOCRINE THERAPY OR BE CONSIDERED INAPPROPRIATE FOR ENDOCRINE THERAPY
- U-2360 MANAGEMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH ENDOMETRIOSIS
- U-2361 METHOD OF ADMINISTERING A GRANULATE FORMULATION OF 5-METHYL-1-PHENYL-2-(1H)-PYRIDONE AS RECITED IN CLAIM 1, TO TREAT IDIOPATHIC PULMONARY FIBROSIS
- U-2362 TREATMENT OF HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6
- U-2363 ADMINISTRATION OF RISPERIDONE
- U-2364 TREATMENT OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PHARMACOKINETIC ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENASE IN ADULTS WHO HAVE NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY
- U-2365 TREATMENT OF HIV-1 INFECTION USING A COMPOSITION CONTAINING A PHARMACOKINETIC ENHANCER THAT INIBITS CYTOCHROME P450 MONOOXYGENASE IN ADULTS WHO ARE

PATENT AND EXCLUSIVITY TERMS

ADB 129 of 133

PATENT USE

- VIROLOGICALLY SUPPRESSED ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS
- U-2366 TREATMENT OF LIVER DISEASE THROUGH NUTRITION FOR PATIENTS UNDER THE AGE OF 12
- U-2367 USE FOR PATIENTS WITH PARENTERAL NUTRITION ASSOCIATED CHOLESTASIS OR PARENTERAL NUTRITION ASSOCIATED LIVER DISEASE
- U-2368 TOPICAL TREATMENT OF ACNE VULGARIS IN PATIENTS 9 YEARS OF AGE AND OLDER
- U-2369 FOR THE TREATMENT OF GENOTYPE 1, 4, 5 OR 6 CHRONIC HEPATITIS C VIRUS (HCV) INFECTION
- U-2370 FOR TREATMENT-NAIVE GENOTYPE 1 PATIENTS WITH CHRONIC HEPATITIS C VIRUS (HCV) INFECTION FOR A DURATION OF 8-WEEKS
- U-2371 THE TREATMENT OF FABRY PATIENTS
- U-2372 A METHOD OF REDUCING LEFT VENTRICULAR MASS INDEX (LVMI) IN A FABRY PATIENT BY ADMINISTERING MIGALASTAT
- U-2373 A METHOD OF REDUCING PODOCYTE GLOBOTRIAOSYL CERAMIDE (GL-3) IN A FABRY PATIENT BY ADMINISTERING MIGALASTAT
- U-2374 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR AND IVACAFTOR
- U-2375 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR FORM I AND IVACAFTOR
- U-2376 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING LUMACAFTOR AND A SOLID COMPOSITION COMPRISING AMORPHOUS AND LESS THAN ABOUT 30% CRYSTALLINE IVACAFTOR
- U-2377 USE OF VITAL DYE FOR FACILITATING SURGICAL PROCEDURES FOR VITREO-RETINAL SURGERY
- U-2378 TREATMENT OF POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS
- U-2379 USE IN IDENTIFICATION OF INTRAOCULAR MEMBRANES TO FACILITATE REMOVAL DURING OPHTHALMIC SURGERY
- U-2380 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTIONS IN PATIENTS 18 YEARS OF AGE AND OLDER
- U-2381 TREATMENT IN COMBINATION WITH A GNRH AGONIST OF NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NM-CRPC)
- U-2382 TREATMENT IN COMBINATION WITH A GNRH AGONIST OF HIGH RISK NON-METASTATIC, CASTRATION-RESISTANT PROSTATE CANCER (NM-CRPC)
- U-2383 METHOD OF CONTROLLING GLYCEMIA IN A DIABETIC PATIENT WITH DELAYED OR PROLONGED FOOD ABSORPTION BY ADMINISTERING 50 TO 75% OF A PREDETERMINED DOSE OF INSULIN-FDKP AT MEALTIME, AND ADMINISTERING REMAINDER OF DOSE 30-120 MINUTES AFTER BEGINNING OF MEAL
- U-2384 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 10
- U-2385 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1,10 AND 11
- U-2386 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 12 AND 19
- U-2387 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 12, 19 AND 20
- U-2388 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 21 AND 28
- U-2389 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 21, 28, AND 29
- U-2390 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 30 AND 41

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

- U-2391 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 30, 41, AND 42
- U-2392 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 43 AND 50
- U-2393 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 43, 50 AND 51
- U-2394 FOR USE IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN ADULT PATIENTS WITH NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY
- U-2395 FOR THE TREATMENT OF HIV-1 INFECTION IN ADULT PATIENTS WITH NO PRIOR ANTIRETROVIRAL TREATMENT HISTORY
- U-2396 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE DOSAGE UNIT COMPRISING LUMACAFTOR AS RECITED IN CLAIM 1 OF US PATENT 8716338 AND IVACAFTOR
- U-2397 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 2-5 YEARS WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE DOSAGE UNIT COMPRISING LUMACAFTOR AND IVACAFTOR AS RECITED IN CLAIM 1 OF US PATENT 9192606
- U-2398 TOPICAL TREATMENT OF PRIMARY AXILLARY HYPERHIDROSIS IN ADULTS AND PEDIATRIC PATIENTS 9 YEARS OF AGE AND OLDER
- U-2399 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 12 YEARS AND OLDER, WITH A F508DEL OR G551D CFTR GENE MUTATION AND A A455E, 2789+5G->A, OR 3849+10KBC->T MUTATION, COMPRISING CONCURRENT COADMINISTRATION OF THE COMPOSITIONS OF CLAIM 1 OF U.S. PATENT 10058546
- U-2400 REDUCING ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- U-2401 A METHOD OF TREATING AMYOTROPHIC LATERAL SCLEROSIS IN A PATIENT IN NEED OF SUCH TREATMENT, SAID METHOD COMPRISING ADMINISTERING TO SAID PATIENT AN EFFECTIVE AMOUNT OF A SUSPENSION ACCORDING TO CLAIM 1
- U-2402 TREATMENT OF SCHIZOPHRENIA BY RAPID AND CONTINUOUS INTRAMUSCULAR INJECTION
- U-2403 TREATMENT OF PSORIASIS USING A DOSAGE TITRATION SCHEDULE
- U-2404 METHOD OF DELIVERING SUMATRIPTAN TO A NASAL CAVITY
- U-2405 A METHOD FOR TREATING A BACTERIAL INFECTION IN INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS PATIENTS 9 YEARS OF AGE AND OLDER COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF SARECYCLINE HYDROCHLORIDE
- U-2406 A METHOD FOR TREATING A PATIENT 9 YEARS OF AGE AND OLDER SUFFERING FROM AN INFLAMMATORY SKIN DISORDER OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF SARECYCLINE HYDROCHLORIDE
- U-2407 A METHOD FOR TREATING ACNE IN INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS PATIENTS 9 YEARS OF AGE AND OLDER COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF SARECYCLINE HYDROCHLORIDE CRYSTALLINE SALT
- U-2408 A METHOD FOR TREATING A BACTERIAL INFECTION IN INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS PATIENTS 9 YEARS OF AGE AND OLDER COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF SARECYCLINE HYDROCHLORIDE CRYSTALLINE SALT
- U-2409 A METHOD FOR TREATING ACNE IN INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS PATIENTS 9 YEARS OF AGE AND OLDER COMPRISING ADMINISTERING SARECYCLINE HYDROCHLORIDE IN 60 MG, 100 MG OR 150 MG EQUIVALENT DOSES
- U-2410 TREATMENT OF ADULT PATIENTS FOR WHOM TREATMENT WITH BOTH AMLODIPINE FOR HYPERTENSION AND CELECOXIB FOR OSTEOARTHRITIS ARE APPROPRIATE
- U-2411 TREATMENT OF CYSTIC FIBROSIS IN A PATIENT AGE 12 YEARS OR OLDER WHO IS HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE USING THE TABLET COMPRISING LUMACAFTOR AS RECITED IN CLAIM 1, 19, OR 21 OF U.S. PATENT NO. 10,076,513 AND IVACAFTOR
- U-2412 FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) AND/OR SMALL LYMPHOCYTIC LEUKEMIA (SLL)
- U-2413 FOR THE TREATMENT OF PATIENTS WITH FOLLICULAR LYMPHOMA (FL)
- U-2414 TREATING MYCOBACTERIUM AVIUM COMPLEX (MAC) LUNG DISEASE IN ADULTS AS PART OF A

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

COMBINATION DRUG REGIMEN

- U-2415 TREATING MYCOBACTERIUM AVIUM COMPLEX (MAC) LUNG DISEASE IN ADULTS AS PART OF A COMBINATION ANTIBACTERIAL DRUG REGIMEN
- U-2416 TREATING MYCOBACTERIUM AVIUM COMPLEX (MAC) LUNG DISEASE IN ADULTS WITH CYSTIC FIBROSIS AS PART OF A COMBINATION DRUG REGIMEN
- U-2417 TREATING MYCOBACTERIUM AVIUM COMPLEX (MAC) LUNG DISEASE IN NON-CYSTIC FIBROSIS ADULTS AS PART OF A COMBINATION ANTIBACTERIAL DRUG REGIMEN
- U-2418 METHOD OF ADMINISTERING TESTOSTERONE ENANTHATE SUBCUTANEOUSLY
- U-2419 METHOD OF OPERATING AN INJECTION DEVICE
- U-2420 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGES 12 AND OLDER, WHO HAVE TWO COPIES OF THE F508DEL MUTATION OR ONE F508DEL MUTATION AND A CFTR MUTATION PREDICTED TO BE RESPONSIVE TO TEZACAFTOR/IVACAFTOR, WITH THE COMPOSITION OF CLAIM 1 OF US 10,081,621
- U-2421 USE IN COMBINATION WITH CLOBAZAM FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH DRAVET SYNDROME
- U-2422 USE IN COMBINATION WITH CLOBAZAM FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH LENNOX GASTAUT SYNDROME WHO HAVE BEEN PREVIOUSLY TREATED WITH CLOBAZAM
- U-2423 USE IN COMBINATION WITH CLOBAZAM FOR THE TREATMENT OF SEIZURES IN PATIENTS WITH DRAVET SYNDROME WHO HAVE BEEN PREVIOUSLY TREATED WITH CLOBAZAM
- U-2424 USE IN COMBINATION WITH CLOBAZAM FOR TREATMENT OF SEIZURES IN PATIENTS WITH LENNOX GASTAUT SYNDROME
- U-2425 USE FOR THE TREATMENT OF CONVULSIVE SEIZURES IN PATIENTS WITH DRAVET SYNDROME
- U-2426 USE FOR THE TREATMENT OF CONVULSIVE SEIZURES IN PATIENTS WITH LENNOX GASTAUT SYNDROME
- U-2427 USE FOR THE TREATMENT OF DROP SEIZURES IN PATIENTS WITH DRAVET SYNDROME
- U-2428 TREATMENT OF PARTIAL-ONSET SEIZURES WITH OR WITHOUT SECONDARILY GENERALIZED SEIZURES IN PATIENTS WITH EPILEPSY 4 YEARS OF AGE AND OLDER
- U-2429 TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES AS ADJUNCTIVE THERAPY IN PATIENTS WITH EPILEPSY 12 YEARS OF AGE AND OLDER
- U-2430 TREATMENT OF POLYNEUROPATHY OF HEREDITARY TRANSTHYRETIN AMYLOIDOSIS
- U-2431 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS
- U-2432 LONG-TERM, MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-2433 METHOD OF TREATING A BIOLOGICAL RHYTHM DISORDER, SUCH AS INSOMNIA
- U-2434 USE IN COMBINATION WITH LENALIDOMIDE AND DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-2435 REDUCTION OF RISK OF MAJOR CARDIOVASCULAR EVENTS (CV DEATH, MI, AND STROKE) IN CHRONIC CAD OR PAD
- U-2436 USE IN THE TREATMENT OF MAJOR DEPRESSIVE DISORDER TO IMPROVE TREATMENT EMERGENT SEXUAL DYSFUNCTION (TESD) INDUCED BY PRIOR SEROTONIN REUPTAKE INHIBITOR TREATMENT
- U-2437 TREATMENT OF ADULT PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BREAST CANCER SUSCEPTIBILITY GENE (BRCA)-MUTATED (GBRCAM) HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE LOCALLY ADVANCED OR METASTATIC BREAST CANCER
- U-2438 CARDIOVASCULAR OUTCOMES TRIAL OF LIRAGLUTIDE 1.8 MG IN PATIENTS WITH TYPE 2 DIABETES AND CARDIOVASCULAR DISEASE
- U-2439 TREATMENT OF MENOPAUSE SYMPTOMS, INCLUDING VASOMOTOR SYMPTOMS
- U-2440 FOR THE MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-2441 REDUCTION OF RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS IN THE TREATMENT OF TYPE 2 DIABETES MELLITUS PATIENTS
- U-2442 USE FOR THE TREATMENT OF ATONIC SEIZURES IN PATIENTS WITH LENNOX-GASTAUT SYNDROME
- U-2443 USE FOR THE TREATMENT OF ATONIC SEIZURES IN PATIENTS WITH DRAVET SYNDROME

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

| | |
|--------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| U-2444 | TREATMENT OF SUBJECTS HAVING BACTERIAL SKIN OR SKIN STRUCTURE INFECTION |
| U-2445 | TREATMENT IN COMBINATION WITH AZACITIDINE OR DECITABINE OR LOW-DOSE CYTARABINE OF NEWLY-DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) IN ADULTS WHO ARE AGE 75 YEARS OR OLDER, OR WHO HAVE COMORBIDITIES THAT PRECLUDE USE OF INTENSIVE INDUCTION CHEMOTHERAPY |
| U-2446 | TREATMENT OF ADULT PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL), WITH OR WITHOUT 17P DELETION, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY |
| U-2447 | TREATMENT OF SEVERE HYPERTRIGLYCERIDEMIA (500 MG/DL) IN ADULT PATIENTS AS AN ADJUNCT TO DIET |
| U-2448 | TREATMENT OF TRAVELERS' DIARRHEA CAUSED BY NON-INVASIVE STRAINS OF ESCHERICHIA COLI IN ADULTS |
| U-2449 | TREATMENT OF BACTERIAL SKIN AND SKIN STRUCTURE INFECTION |
| U-2450 | POSITRON EMISSION TOMOGRAPHY DIAGNOSTIC AGENT IN ADULTS WITH SUSPECTED PROSTATE CANCER RECURRENCE BASED ON ELEVATED BLOOD PROSTATE SPECIFIC ANTIGEN LEVELS FOLLOWING PRIOR TREATMENT |
| U-2451 | TREATMENT OF THROMBOCYTOPENIA IN ADULT AND PEDIATRIC PATIENTS 1 YEAR AND OLDER WITH CHRONIC IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP) |
| U-2452 | COMBINATION WITH IMMUNOSUPPRESSIVE THERAPY FOR FIRST-LINE TREATMENT OF ADULT AND PEDIATRIC PATIENTS 2 YEARS AND OLDER WITH SEVERE APLASTIC ANEMIA |
| U-2453 | TREATMENT OF FUNGAL INFECTIONS, INCLUDING BLASTOMYCOSIS, HISTOPLASMOSIS, AND ASPERGILLOSIS |
| U-2454 | USE FOR THE TREATMENT OF DROP SEIZURES IN PATIENTS WITH LENNOX-GASTAUT SYNDROME |
| U-2455 | USE IN COMBINATION WITH CLOBAZAM FOR TREATMENT OF DROP SEIZURES IN PATIENTS WITH LENNOX GASTAUT SYNDROME |
| U-2456 | TREATMENT OF ACUTE MYELOID LEUKEMIA (AML) |
| U-2457 | REINITIATION OF SCHIZOPHRENIA TREATMENT FOLLOWING A MISSED DOSE MORE THAN 9 MONTHS AGO |
| U-2458 | REINITIATION OF SCHIZOPHRENIA TREATMENT FOLLOWING A MISSED DOSE 4-9 MONTHS AGO |
| U-2459 | TREATMENT OF DYSKINESIA AND DECREASING OFF TIME IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS |
| U-2460 | VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF CORONARY ARTERY BYPASS GRAFT IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY |
| U-2461 | VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF CARDIOVASCULAR BYPASS GRAFT AND VASCULATURE IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY |
| U-2462 | VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF VESSEL WITH ARTERIOVENOUS MALFORMATION IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY |
| U-2463 | VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION IN SURGICAL FLAPS IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY |
| U-2464 | VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF TRANSPLANTED ORGAN OR ATTACHED VESSEL IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY |
| U-2465 | VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF VESSEL GRAFT IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY |
| U-2466 | VISUALIZATION OF VESSELS, BLOOD FLOW AND TISSUE PERFUSION OF DONOR ORGAN OR ATTACHED VESSEL IN VASCULAR, GASTROINTESTINAL, ORGAN TRANSPLANT, AND PLASTIC, MICRO- AND RECONSTRUCTIVE, INCLUDING MINIMALLY INVASIVE, SURGERY |
| U-2467 | VISUALIZATION OF EXTRAHEPATIC BILIARY DUCT ATTACHED TO DONOR ORGAN IN PATIENTS 12 YEARS AND OLDER |
| U-2468 | VISUALIZATION OF EXTRAHEPATIC BILIARY DUCT ATTACHED TO TRANSPLANTED ORGAN IN PATIENTS 12 YEARS AND OLDER |
| U-2469 | METHOD OF TREATING CANCEROUS SOLID TUMORS |

PATENT AND EXCLUSIVITY TERMS**PATENT USE**

U-2470 METHOD OF TREATING SOLID TUMORS THAT EXHIBIT AN NTRK GENE FUSION

U-2471 METHOD OF TREATING SOLID TUMORS THAT EXHIBIT AN NTRK FUSION GENE IN A PEDIATRIC PATIENT

U-2472 METHOD OF TREATING NEUROBLASTOMA, GLIOMA, THYROID, AND BREAST CANCER SOLID TUMORS THAT EXHIBIT AN NTRK GENE FUSION

U-2473 METHOD OF TREATING CMN, IFS, HGG, DIPGS, PTC, SOFT TISSUE SARCOMA, AND SPINDLE CELL SARCOMA SOLID TUMORS EXHIBITING AN NTRK GENE FUSION IN A PEDIATRIC PATIENT WITH AN ORAL SOLUTION

U-2474 METHOD OF TREATING SOLID TUMORS THAT EXHIBIT AN NTRK GENE FUSION AFTER SURGICAL RESECTION

U-2475 METHOD OF TREATING SOLID TUMORS THAT EXHIBIT AN NTRK GENE FUSION IN A PEDIATRIC PATIENT

U-2476 USE OF A DELIVERY DEVICE TO DELIVER A DOSE OF NALOXONE

U-2477 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH CYP1A2 STRONG INHIBITORS

U-2478 METHOD FOR THE INDUCTION OF LOCAL ANESTHESIA PRIOR TO PERFORMING A PROCEDURE ON, THROUGH, OR ADJACENT TO THE MUCOUS MEMBRANES

U-2479 METHOD OF ADMINISTERING A LOCAL ANESTHETIC TO THE MUCOUS MEMBRANES